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GRADUATE COLLEGE

SUBSURFACE AND OUTCROP ORGANIC GEOCHEMISTRY OF THE EAGLE
FORD SHALE (CENOMANIAN-CONIACIAN) IN WEST, SOUTHWEST,
CENTRAL, AND EAST TEXAS

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CENTRAL, AND EAST TEXAS

A DISSERTATION APPROVED FOR THE
CONOCOPHILLIPS SCHOOL OF GEOLOGY AND GEOPHYSICS

BY

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To Alessandro and Oswaldo

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Table of Contents

Acknowledgements	iv
Table of Contents	vii
List of Tables	xi
List of Figures.....	xiii
Abstract.....	xix
CHAPTER I.....	1
1. INTRODUCTION	1
1.1 Geologic Framework	4
1.1.1 Regional Geology	4
1.1.2 West and South Texas Eagle Ford Shale Geology.....	9
1.1.3 Central Texas Eagle Ford Shale Geology	14
1.1.4 East Texas Eagle Ford Shale Geology	19
1.1.5 Paleogeography and Climate	22
1.2 Objectives	26
1.3 Methodology	27
1.3.1 Study area and sample locations	27
1.3.2 Experimental	29
1.3.2.1 Preliminary rock sample treatment	32
1.3.2.2 Total Organic Carbon (TOC) and Rock Eval Pyrolysis	32
1.3.2.3 Petrographic analysis	32
1.3.2.4 Extraction and Fractionation	33
1.3.2.5 Molecular sieving.....	34
1.3.2.6 Gas Chromatography (GC)	35

1.3.2.7	Gas Chromatography-Mass Spectrometry (GC-MS)	35
1.3.2.8	Quantitative Biomarker Analysis	36
1.3.2.9	Gas Chromatography Isotope-Ratio Mass Spectrometry (GC-IRMS)	41
CHAPTER II		42
2.	RESULTS AND DISCUSSION.....	42
2.1	Geochemical Screening	42
2.1.1	Organic Richness	43
2.1.2	Organic Matter Type.....	49
2.1.3	Thermal Maturity from Rock Eval and Vitrinite Reflectance Analysis.....	56
2.1.4	Gas Chromatography	61
2.1.4.1	n-Alkanes distribution.....	61
2.1.4.2	Pristane and Phytane	67
2.2	Biomarker and geochemical parameters for evaluation of organic matter source	71
2.2.1	Steranes.....	71
2.2.1.1	Regular steranes	71
2.2.1.2	Diasteranes (rearranged steranes).....	77
2.2.1.3	Pregnanes	79
2.2.2	Terpanes.....	80
2.2.2.1	Tricyclic terpanes	81
2.2.2.2	Tetracyclic terpanes	86
2.2.2.3	Hopanes.....	88
2.2.3	Aryl Isoprenoids.....	96
2.2.4	Monoaromatic Steroids.....	101
2.3	Biomarkers and geochemical parameters for evaluation of thermal maturity	104
2.3.1	Steranes.....	104

2.3.2	Terpanes	107
2.3.2.1	Hopanes.....	107
2.3.2.2	Moretanes.....	108
2.3.2.3	Ts and Tm	109
2.3.3	Monoaromatic and Triaromatic Steroids	111
2.3.4	Phenanthrenes	114
2.4	Isotope Analysis.....	117
3.	CHAPTER III.....	122
3.1	Regional and local organic geochemistry of the Eagle Ford Shale	122
4.	CONCLUSIONS	129
5.	REFERENCES	131
6.	APPENDIX	146
A.	Abbreviations and formulas used for calculation of geochemical biomarker ratios	146
B.	Total Organic Carbon (TOC) and Rock Eval (RE) parameters for the Eagle Ford Shale samples.....	150
C.	Geochemical logs of TOC, HI, Pr/Ph and AIR for the Eagle Ford Shale bitumens analyzed	160
D.	Gas chromatograms of the saturate fractions for the Eagle Ford Shale samples analyzed in this study (Pr = pristane; Ph = phytane, n-C ₂₅ = C ₂₅ normal alkane)	167
E.	Geochemical ratios of <i>n</i> -alkanes and isoprenoids for the saturate fractions of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined).....	180
F.	Geochemical logs of biomarker ratios of steranes for the Eagle Ford Shale bitumens analyzed	187

G. Geochemical logs of biomarker ratios of terpanes for the Eagle Ford Shale bitumens analyzed	194
H. Geochemical ratios of steranes for the branched and cyclic fractions (B&C) of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined).....	201
I. Geochemical ratios of terpanes for the branched and cyclic fractions (B&C) of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined).....	207
J. Aryl isoprenoids ratio (AIR) for the branched and cyclic fractions (B&C) of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined).....	214
K. Geochemical ratios for aromatic biomarkers of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined)	218
L. Quantitative biomarker analysis results for steranes (Concentrations are expressed as μg biomarkers/g TOC; ND = not determined).....	224
M. Quantitative biomarker analysis results for terpanes (Concentrations are expressed as μg biomarkers/g TOC; ND = not determined).....	242
N. Quantitative biomarker analysis results for aryl isoprenoids (Concentrations are expressed as μg biomarkers/g TOC; ND = not determined).....	258
O. Quantitative biomarker analysis results for monoaromatic steroids (Concentrations are expressed as μg biomarkers/g TOC; ND = not determined).....	267
P. Quantitative biomarker analysis results for phenanthrenes and triaromatic steroids (Concentrations are expressed as μg biomarkers/g TOC; ND = not determined).....	276
Q. ^{13}C values for <i>n</i> -alkanes of Eagle Ford Shale bitumens and oils analyzed in this study. ^{13}C values are expressed in per mil (‰) relative to the Vienna Pee Dee belemnite (VPDB) standard.....	285

List of Tables

Table 1. List of samples for organic geochemical analyses	30
Table 2. List of ions selected for GC-MS analysis of branched and cyclic and aromatic fractions from Eagle Ford Shale samples	36
Table 3. Parameters, terms, and definitions derived from Rock Eval pyrolysis analyses (modified from Peters, 1986 and Jarvie et al., 2007)	43
Table 4. Average TOC values for the Eagle Ford Shale outcrop and subsurface rock samples analyzed in this study	44
Table 5. Summary table of <i>n</i> -alkanes analyses from the saturate fraction of Eagle Ford Shale bitumens and oils (ND = not determined)	64
Table 6. Identification of steranes in the partial m/z 217.3 fragmentogram of the B&C fractions	75
Table 7. Identification of terpanes in the partial m/z 191.3 fragmentogram of the B&C fractions	85
Table 8. Identification of monoaromatic steroids (MAS) in the partial m/z 253.3 fragmentogram of the B&C fractions	103
Table 9. Average steranes isomerization ratios for the Eagle Ford Shale bitumens and liquids (ND = not determined; *not averaged)	107
Table 10. Average C ₃₁ 22S/(22S+22R) hopane ratios for Eagle Ford Shale bitumens and oils (*not averaged)	108
Table 11. Average C ₃₀ Moretane/Hopane ratios for Eagle Ford Shale bitumens and oils (*not averaged)	109

Table 12. Identification of triaromatic steroids (TAS) in the partial m/z 231.3 fragmentogram of the aromatic fractions	114
Table 13. Identification of phenanthrenes in the partial mass chromatograms of m/z 178.3, 192.3, 206.3 of the aromatic fractions.....	116
Table 14. Calculated vitrinite reflectance (R_c) for Eagle Ford Shale bitumens and oils from MPI-1 values.....	116
Table 15. Average of bulk and <i>n</i> -alkanes geochemical parameters for the Eagle Ford Shale members across West, Southwest, Central and East Texas (ND = not determined)	127
Table 16. Average biomarker ratios for the Eagle Ford Shale members across West, Southwest, Central and East Texas (ND = not determined).....	128

List of Figures

Figure 1. Composite map of the Eagle Ford Shale in Southwest Texas, showing production, depth, thickness, and thermal maturity of this gas-shale play (EIA, 2010) ..	3
Figure 2. Texas map showing the main structural and physiographic features influencing the Gulf of Mexico Coastal Plain (modified from Donovan et al., 2013a)...	6
Figure 3. Generalized chronostratigraphic chart for the Cretaceous of South Texas (modified from Donovan et al., 2012).....	7
Figure 4. Generalized A) Paleogeography of the Comanche Platform; B) Cross section X-X' illustrating inherited seafloor paleobathymetry during Late Cenomanian time (from Gardner et al., 2013).....	8
Figure 5. Sequence stratigraphic and facies subdivision proposed by Donovan et al. (2014) for the Eagle Ford Group at the Lozier Canyon outcrop based on petrophysical and geochemical data (from Donovan et al., 2014).....	11
Figure 6. Sequence stratigraphic framework for the Eagle Ford Shale Group at the Fasken “A” #1 well in Webb County, South Texas based on petrophysical and geochemical data (modified from Donovan et al., 2013a)	13
Figure 7. Core description, facies analysis, and gamma ray logs for the ACC #1 core from Fairbanks (2012; GR = gamma ray, CGR = computed gamma ray).....	17
Figure 8. Stratigraphic cross-section showing facies variation and units continuity along the San Marcos Arch (modified from Fairbanks, 2012).....	18
Figure 9. Generalized SW-NE schematic strike cross section illustrating the relationships among lithostratigraphic units across the Eagle Ford Shale Play (modified from Hentz and Ruppel, 2010)	20

Figure 10. SW-NE regional strike cross section of the East Texas Basin and northeast flank of the San Marcos Arch (modified from Hentz and Ruppel, 2010)	21
Figure 11. Map showing the extent of Western Interior Seaway, locations of the Foreland Basin, and the Eastern Stable Platform during the Late Cretaceous (~85 Ma) (modified from Robinson and Kirschbaum, 1995 and Blakey, 2011)	25
Figure 12. Texas map showing location of the samples analyzed in this study	28
Figure 13. Schematic workflow used for laboratory analyses of the Eagle Ford Shale rock, whole oil and condensate samples (modified from Miceli Romero and Philp, 2012).....	31
Figure 14. TOC, HI, and Pr/Ph geochemical logs for the Eagle Ford Shale outcrop and subsurface samples analyzed in this study (Pr = pristane; Ph = phytane)	45
Figure 15. Modified Van Krevelen diagram for the Eagle Ford Shale samples. HI values for EFFA samples are unreliable due to their low remaining hydrocarbon potential	51
Figure 16. Rock Eval Remaining Hydrocarbon Potential (S ₂) vs. TOC plot for determination of kerogen type and maturity of Eagle Ford Shale samples (plot template modified from GeoMark Research Ltd.)	53
Figure 17. S ₁ vs. TOC plot for the Eagle Ford Shale samples showing possible hydrocarbon contamination of the EFMK samples (plot template modified from GeoMark Research Ltd.)	54
Figure 18. T _{max} vs. Production Index (PI) plot for determination of thermal maturity and sample contamination of Eagle Ford Shale samples (plot template modified from GeoMark Research Ltd.)	55

Figure 19. T_{max} vs. HI plot showing maturity and kerogen type of the Eagle Ford Shale samples. HI values for EFFA samples are unreliable due to their low remaining hydrocarbon potential 59

Figure 20. Photomicrographs from the EFLH-9 Eagle Ford Shale sample (Photomicrographs courtesy of Brian Cardott, 2013) 60

Figure 21. Gas chromatograms of the saturate fractions from bitumen extracts of the EFLC outcrop (Pr = pristane; Ph = phytane, $n-C_{25} = C_{25}$ normal alkane) 65

Figure 22. Isoprenoids plot of Pristane/ $n-C_{17}$ versus Phytane/ $n-C_{18}$ showing redox conditions, maturity, and depositional environments for samples of the Eagle Ford Shale ($n-C_{17} = C_{17}$ normal alkane; $n-C_{18} = C_{18}$ normal alkane) 70

Figure 23. Partial fragmentograms of the m/z 217.3 ion showing distribution of steranes in the B&C fractions of the EFLC samples. Peak identification is presented in Table 6 73

Figure 24. Ternary diagram of C_{27} , C_{28} , and C_{29} regular steranes for the Eagle Ford Shale samples 76

Figure 25. Geochemical logs of steranes ratios for the EFLC samples. Formulas for calculation of ratios are in Appendix A. Geochemical logs from other locations analyzed are in Appendix F. Numerical values of biomarker ratios are in Appendix H 78

Figure 26. Partial fragmentograms of the m/z 191.3 ion showing distribution of terpanes in the B&C fractions of the EFLC samples. Peak identification is presented in Table 7 83

Figure 27. Geochemical logs of biomarker ratios of terpanes for the EFLC samples. Formulas for calculation of ratios are in Appendix A. Geochemical logs from other

locations analyzed are in Appendix G. Numerical values of biomarker ratios are in Appendix I.....	87
Figure 28. Plot of C_{22}/C_{21} versus C_{24}/C_{23} tricyclic terpanes shows source rock depositional environments for Eagle Ford Shale bitumens and oils (Dotted lines are used as a guide and do not represent fixed fields on the diagram. Plot modified from Peters et al., 2005)	88
Figure 29. Plot of C_{26}/C_{25} versus $C_{31}R/C_{30}$ Hopane shows that the Eagle Ford Shale bitumens and oils are mainly of marine origin (Dotted lines are used as a guide and do not represent fixed fields on the diagram. Plot modified from Peters et al., 2005).....	90
Figure 30. Plot of 30-Nor/ C_{30} hopane versus $C_{35}S/C_{34}S$ homohopanes suggest most of the Eagle Ford Shale rock samples are of marine origin (Dotted lines are used as a guide and do not represent fixed fields on the diagram. Plot modified from Peters et al. 2005)	91
Figure 31. Homohopanes distribution for bitumens and oils from the Eagle Ford Shale	94
Figure 32. Partial fragmentograms of the m/z 133.1 ion showing the aryl isoprenoids distribution in the B&C fractions of the EFLC samples. Number of carbon atoms for the 2,3,6-trimethyl substituted aryl isoprenoids is indicated above each peak ($C_{24}D$ = internal standard)	98
Figure 33. Geochemical logs of biomarker ratios of aryl isoprenoids for Eagle Ford Shale samples at Lozier Canyon. Formulas for calculation of ratios are in Appendix A. Geochemical logs from other locations analyzed are in Appendix C. Numerical values of AIR are in Appendix J.....	99

Figure 34. AIR versus Pr/Ph plot for the Eagle Ford Shale samples (Dotted line is used as a guide and does not represent fixed fields on the diagram. Plot template from Schwark and Frimmel, 2004) 100

Figure 35. Partial fragmentograms of the m/z 253.3 ion showing distribution of the monoaromatic steroids (MAS) in the B&C fractions of the EFLC samples. Peak identification is presented in Table 8..... 102

Figure 36. Ternary diagram of C₂₇, C₂₈, and C₂₉ monoaromatic steroids for Eagle Ford Shale bitumens and oils (Plot modified from Moldowan et al., 1985)..... 104

Figure 37. Plot of C₂₉ / (+) steranes versus C₂₉ 20S/(20S+20R) steranes showing variations in thermal maturity for the Eagle Ford Shale source rocks and liquids (Gray area represents end points of isomerization reactions. Plot modified from Peters, 1999) 106

Figure 38. Plot of diasteranes/(diasteranes+regular) C₂₇ Steranes versus Ts/(Ts+Tm) ratios shows differences in thermal maturity, source, and redox potential for Eagle Ford Shale samples (Plot modified from Moldowan et al., 1994)..... 111

Figure 39. Partial fragmentograms of the m/z 231.3 ion showing distribution of the triaromatic steroids (TAS) in the aromatic fractions of sample EFLC-1. Peak identification is presented in Table 12..... 113

Figure 40. Summed mass chromatograms of m/z 106.3, 192.3, 206.3 ions showing distribution of phenanthrenes compounds in the aromatic fractions of sample EFLC-1. Peak identification is presented in Table 13..... 115

Figure 41. *n*-Alkane isotope profile from the EFLC extracts (immature)..... 118

Figure 42. *n*-Alkane isotope profile from the EFMK extracts (oil window)..... 119

Figure 43. <i>n</i> -Alkane isotope profile from the EFFA extracts (peak oil window)	119
Figure 44. <i>n</i> -Alkane isotope profile from the EFAC extracts (immature)	120
Figure 45. <i>n</i> -Alkane isotope profile from the EFLH extracts (oil window).....	120
Figure 46. <i>n</i> -Alkane isotope profile from oils and condensate samples (condensate/wet gas window).....	121

Abstract

A comprehensive organic geochemical analysis was performed to a suite of outcrops and core samples from the Eagle Ford Shale Group from West, Southwest, Central and East Texas with the aim of determining variations in organic matter source, thermal maturity and depositional environments. Oils and condensates produced from Eagle Ford Shale and San Miguel reservoirs were analyzed to determine the origin of these liquids. A total of 180 samples were subjected to total organic carbon (TOC) and Rock Eval analysis for geochemical screening, and one sample from each location was analyzed for vitrinite reflectance (%R_o). Rock samples with good source rock potential (TOC>1.0% wt.) were selected for biomarker and isotope analyses. These analyses were carried out by means of gas chromatography (GC), gas chromatography-mass spectrometry (GC-MS), and gas chromatography-isotope ratio mass spectrometry (GC-IRMS).

TOC and Rock Eval parameters show that the Eagle Ford Shale has excellent source rock potential and is dominated by Type II kerogen indicating a marine origin. Distributions of regular steranes, hopanes and monoaromatic steroids (MAS) point towards a marine carbonate depositional environment. Aryl isoprenoids suggest the occurrence of intermittent photic zone anoxia (PZA). In addition *n*-alkanes, steranes distribution, and the tentative identification of gammacerane suggest deposition under hypersaline conditions in West and East Texas. Biomarker parameters indicate that in East Texas the Eagle Ford Shale was partly sourced by terrigenous organic matter, reflecting the influence of the Harris Delta. Thermal maturity parameters indicate that the Eagle Ford Shale is immature to marginally mature in West and Central Texas, and

show a progressive increase in maturity increase towards the southeast following the regional dip. In East Texas, the Eagle Ford Shale is in the main stage of oil generation.

Geochemical logs show a minimal range of vertical variation within the Eagle Ford Shale Group. The Lower Eagle Ford Shale Formation has the highest TOC and hydrogen index (HI) values and in particular, the Lozier Canyon Member is the most organic-rich. Pristane and phytane (Pr/Ph) and biomarker ratios suggest the establishment of stronger anoxic conditions during deposition of the Lower Eagle Ford Shale Formation compared to the Upper Eagle Ford Shale, where the latter may have received an additional siliciclastic and terrigenous organic matter input. In East Texas, Pr/Ph ratios indicate deposition under oxic-suboxic conditions for the entire Eagle Ford Shale interval analyzed. Isotope data indicates a marine organic matter source for the Eagle Ford Shale Group, but ^{13}C values do not show significant organic facies, depositional environment, or thermal maturity changes.

Geochemical data were correlated with an already established sequence stratigraphic framework for the Eagle Ford Shale Group in order to identify relationships between organic geochemical trends and stratigraphic sequences. In this study, these observations reached a general interpretation and it is recommended they be further refined by high-resolution organic geochemistry studies.

CHAPTER I

1. INTRODUCTION

The Eagle Ford Shale (Cenomanian-Coniacian) is one of the most actively explored and produced shale-gas plays in the United States. This mixed siliciclastic/carbonate unit has proved to be the source rock for oil and gas in conventional reservoirs across Texas (Liro et al., 1994). The Eagle Ford Shale Play trends northeast for about 400 miles (640 Km) from the Mexican border into East Texas (Figure 1). It is approximately 50 miles wide (80 Km) and has a variable thickness from 660 ft (~200 m) in the Maverick Basin, to approximately 12 to 17 ft (~4 to 21m) in the San Marcos Arch, to about 450 ft (~150 m) in the center of the East Texas Basin (Hentz and Ruppel, 2010; RRC, 2013). In northwest Texas, the Eagle Ford Shale produces oil from about 5,000 feet (~1,500 m) deep. In the southwest, the Eagle Ford Shale produces mainly dry gas from about 12,000 feet deep (~3,600 m; Figure 1; EIA, 2010; Durham, 2013; RRC, 2013). Up-to-date there are about 22 active fields within the Eagle Ford trend. Gas, condensate, and oil production from the Eagle Ford Shale has been rapidly increasing since its development as an unconventional play in 2008. As of August 2013, hydrocarbon production has reached over 1 million barrels of oil equivalent per day (boe/d; RRC, 2013).

The Upper Cretaceous Eagle Ford Shale has worldwide age equivalent units, which comprise an important fraction of the hydrocarbon source rocks in the world. Deposition of organic rich rocks during the Cretaceous period was influenced by variations in different factors such as tectonism, volcanism, atmospheric and ocean chemistry, climate, sea-level, and sediment supply (Dean and Arthur, 1998). The Eagle

Ford Shale was deposited during a major transgressive episode on a carbonate platform south of the Western Interior Seaway during the Late Cretaceous. During this time, North America experienced warm climatic conditions, which when combined with the series of events leading to onset of the Oceanic Anoxic Event 2 (OAE2), significantly contributed on making the Eagle Ford Shale an organic-rich source rock (Robinson and Kirschbaum, 1995).

Several studies have demonstrated the stratigraphic and geochemical heterogeneities present in shales and shale-gas systems, previously thought to have no significant vertical or lateral variations. The understanding of these changes is essential to improve the efficiency of hydrocarbon prediction and production in unconventional reservoirs. Recent studies of the Eagle Ford Shale from outcrop and subsurface data have added significant value to the understanding of this play (Adams and Carr, 2010; Donovan and Staerker, 2010; Hentz and Ruppel, 2010; Harbor, 2011; Fairbanks, 2012; Donovan et al., 2012, 2013a, 2013b). However, few studies have reported detailed organic geochemical data and interpretations beyond Rock Eval pyrolysis and organic carbon content (Robison, 1997; Liro et al., 1994; Edman and Pitman, 2010; Lewan et al., 2012). The present research provides a comprehensive regional organic geochemical analysis of the Eagle Ford Shale including GC, GC-MS, GC-IRMS in addition to the traditional Rock Eval and TOC screening data, emphasizing the uses and applications of biomarker and isotope analysis in shale-gas studies. In order to achieve this, outcrop and core samples from the Eagle Ford Shale as well as oil and condensate samples produced from the Eagle Ford Shale and the San Miguel Formations were analyzed using several organic geochemistry techniques. The main objective of this research was

to geochemically characterize the Eagle Ford Shale and to determine if the liquid samples available were generated from this source rock. This study will improve our understanding of the lateral and vertical variability of the Eagle Ford Shale in outcrop and subsurface, and the value of organic geochemistry for evaluating unconventional shale-gas plays as an aid in sequence stratigraphic studies.

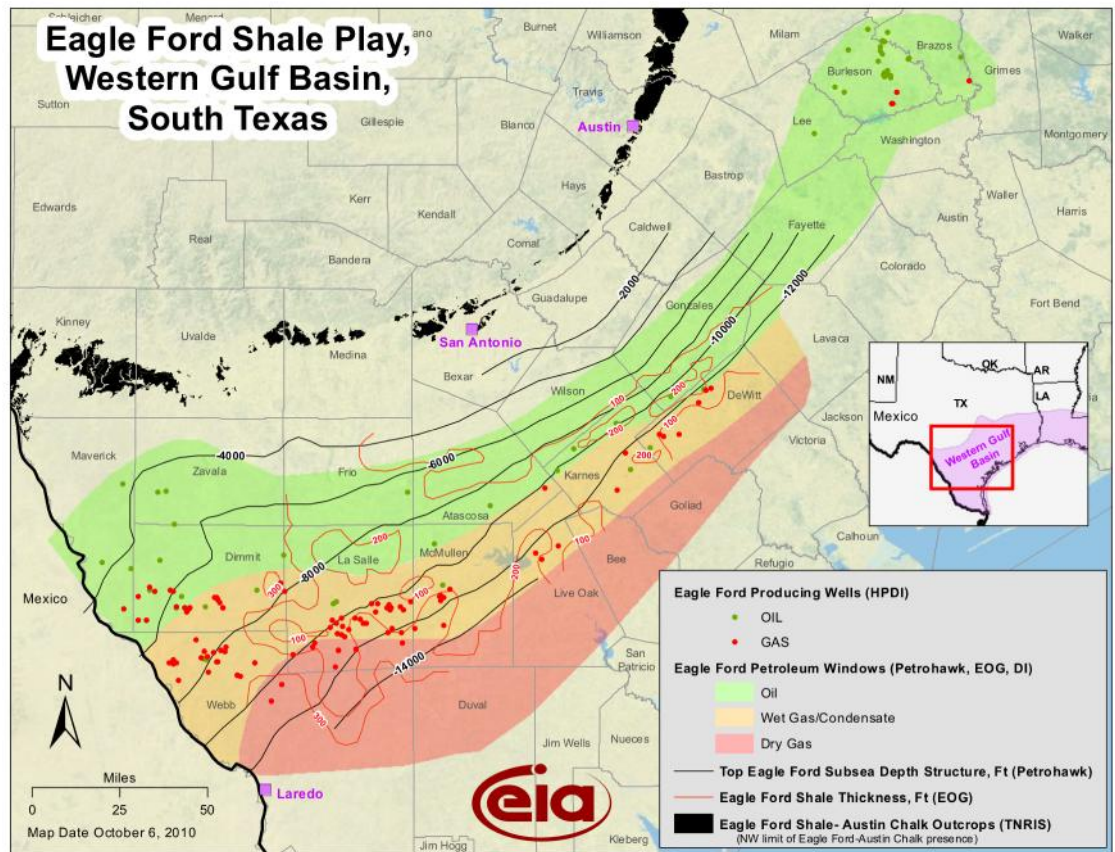


Figure 1. Composite map of the Eagle Ford Shale in Southwest Texas, showing production, depth, thickness, and thermal maturity of this gas-shale play (EIA, 2010)

1.1 Geologic Framework

1.1.1 Regional Geology

The Eagle Ford Shale was deposited during the Late Cretaceous in the Gulf of Mexico Coastal Plain. In Texas, the Gulf of Mexico Coastal Plain is delineated by important structural features that have influenced sediment deposition over geologic time. From west to east these structures correspond to the Maverick Basin of the Rio Grande Embayment, San Marcos Arch, East Texas Basin, and Sabine Uplift (Hentz and Ruppel, 2010; Donovan et al., 2012; Figure 2). A chronostratigraphic chart for South Texas (Figure 3) illustrates the major formations deposited during Cretaceous time. Hill (1887) defined two major successions for the Cretaceous, named Comanche and Gulfian Series. The Sligo, Pearsall, Glen Rose, Edwards, Georgetown, Del Rio and Buda formations comprise the carbonate-dominated Comanche Series, whereas the Eagle Ford, Austin, Anacacho, San Miguel, Olmos, and Escondido formations correspond to the siliciclastic-dominated Gulfian Series (Figure 3; Hill, 1887; Donovan et al., 2012).

The Comanche Platform was developed during the Albian and Early Cenomanian time across Central Texas. This carbonate platform was rimmed by reef build-ups corresponding to the Stuart City Reef Trend (Edwards, Georgetown, Del Rio and Buda formations), and it was centered in the San Marcos Arch. In South Texas, the South Texas Submarine Plateau formed (Gardner et al., 2013; Donovan et al., 2012). Throughout the Middle to Late Cenomanian, differential subsidence generated the Maverick and East Texas Basins (Figure 4a). During this time, a major marine transgression started the accumulation of organic-rich carbonate mudstones

corresponding to the Eagle Ford Shale Group (Hentz and Ruppel, 2010; Donovan et al., 2012). Donovan et al. (2012) presented an idealized cross section of the Comanche Platform to describe the inherited seafloor paleobathymetry at the beginning of Eagle Ford Shale deposition (Figure 4b). These authors proposed shallow water depths of 100 to 200 ft (~30 to 60 m) in most of the Comanche Platform, and water depths of 400 to 600 ft (~120 to 180 m) for the deeper portions of the seaway including the South Texas Submarine Plateau, Rio Grande Embayment, and the Texas Basin.

The stratigraphic characteristics of the Eagle Ford Shale Group vary across the region, since deposition of these units was influenced by several structural and physiographic features. The following sections will describe in detail the stratigraphic variations of the Eagle Ford Shale from Southwest, Central, and East Texas, based on recent studies published on this shale-gas play.

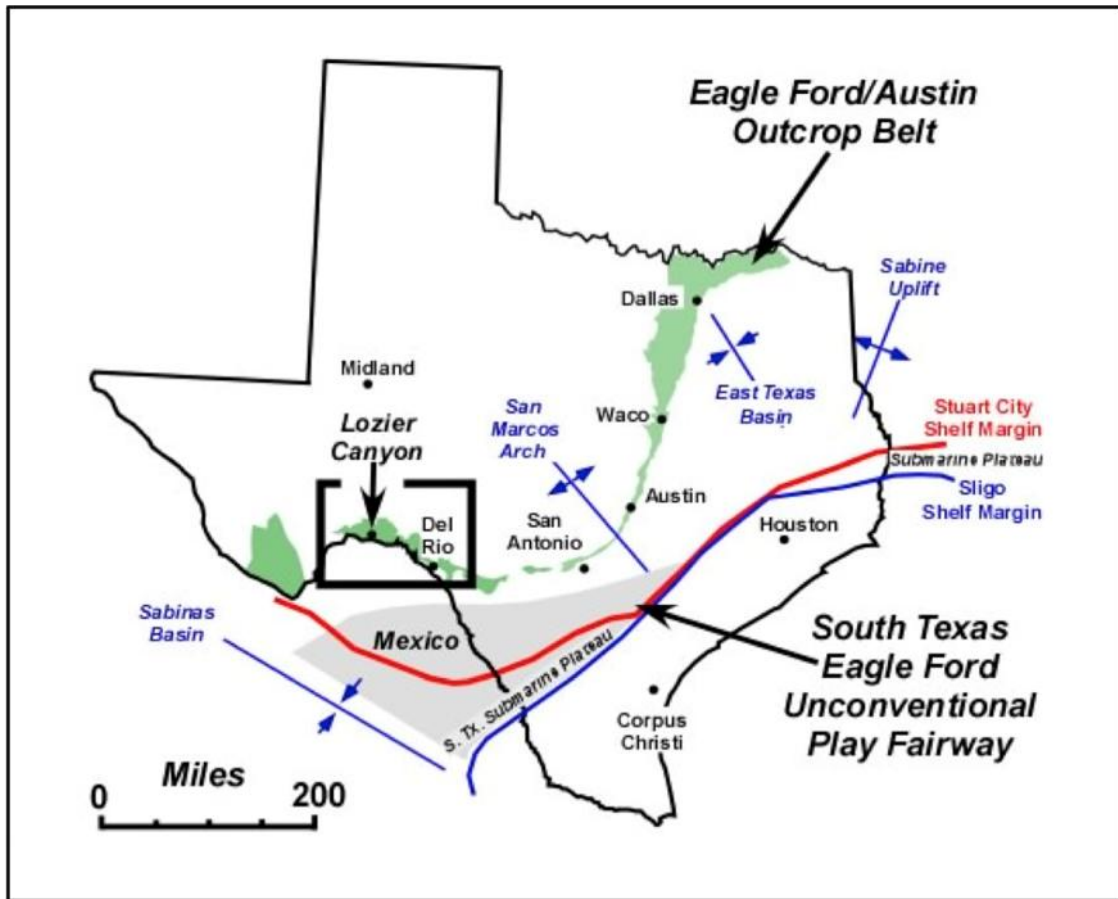


Figure 2. Texas map showing the main structural and physiographic features influencing the Gulf of Mexico Coastal Plain (modified from Donovan et al., 2013a)

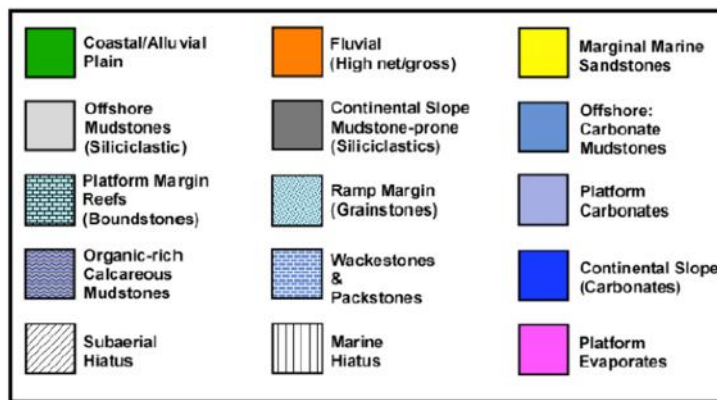
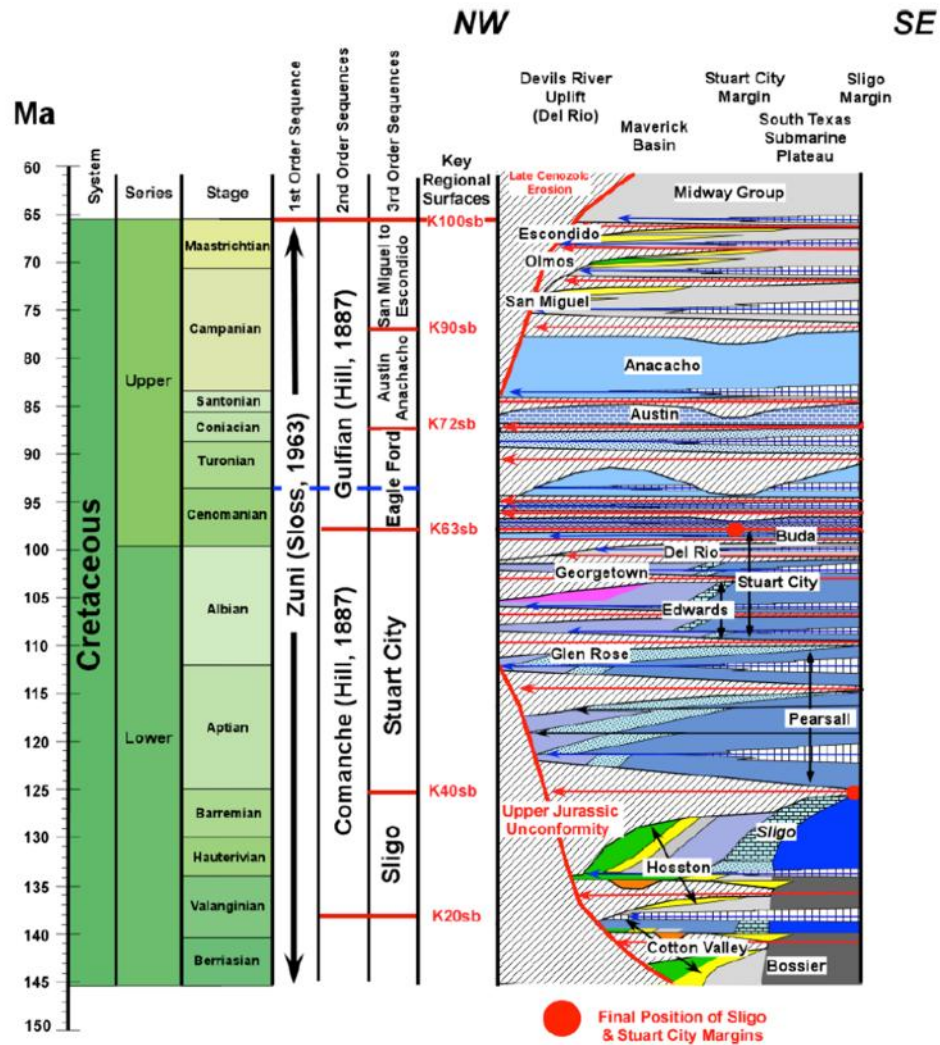


Figure 3. Generalized chronostratigraphic chart for the Cretaceous of South Texas (modified from Donovan et al., 2012)

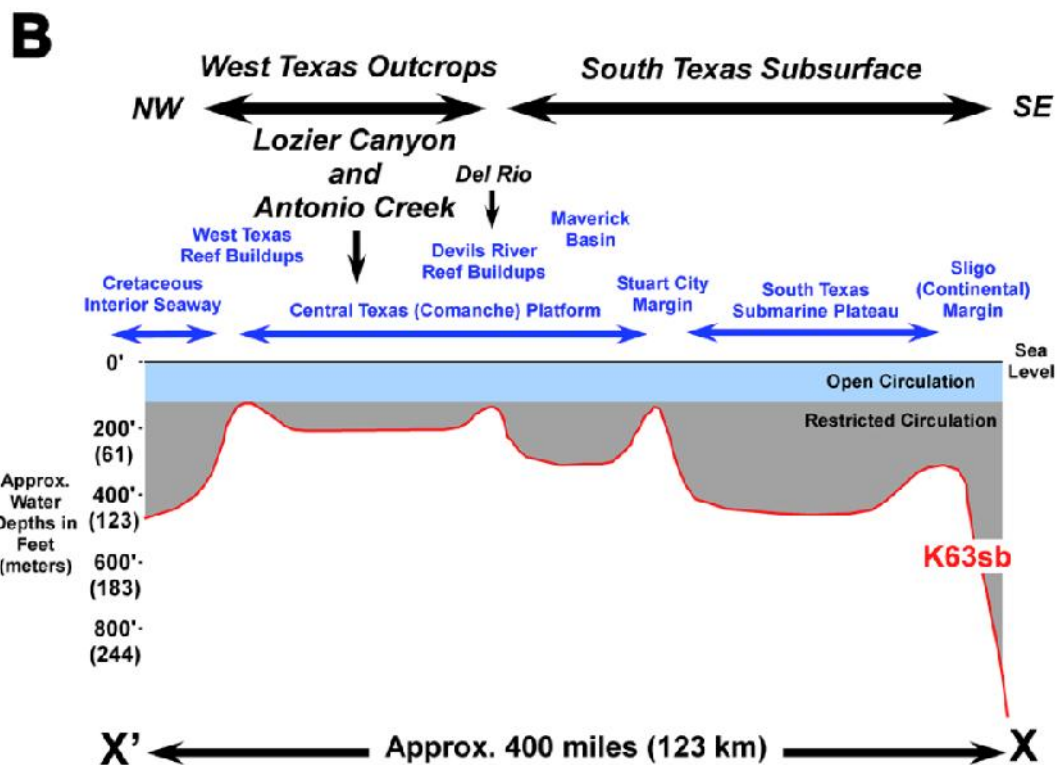
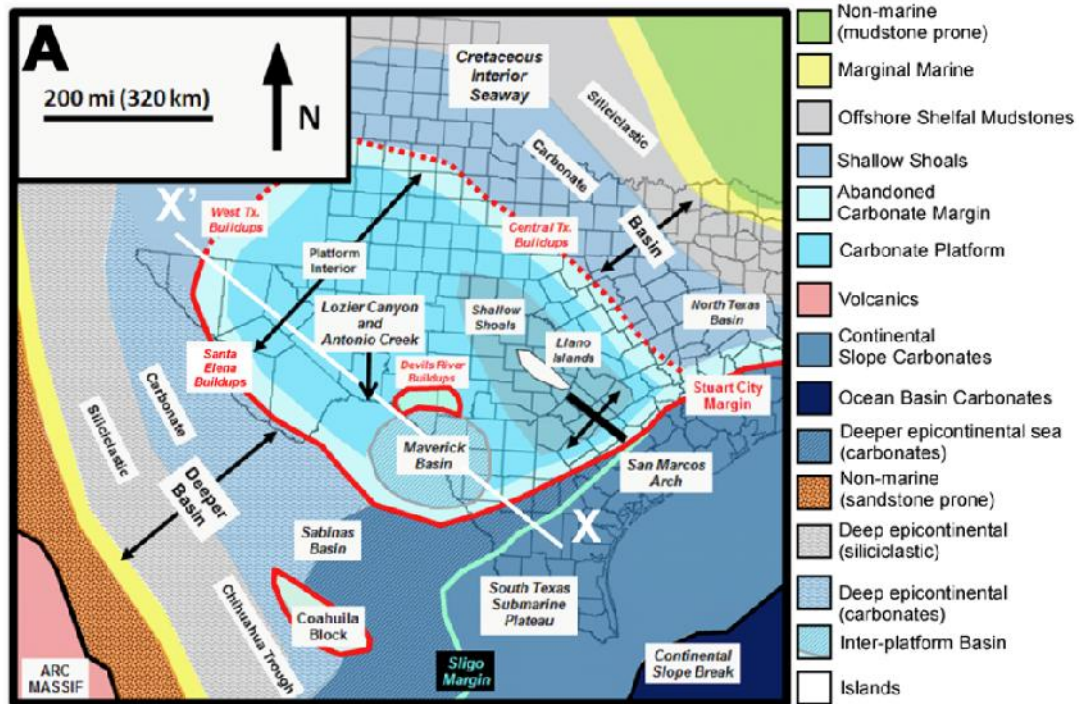


Figure 4. Generalized A) Paleogeography of the Comanche Platform; B) Cross section X-X' illustrating inherited seafloor paleobathymetry during Late Cenomanian time (from Gardner et al., 2013)

1.1.2 West and South Texas Eagle Ford Shale Geology

Donovan and Staerker (2010) summarized the different subdivisions that have been proposed to describe Eagle Ford Shale in West Texas, and raised attention to dual and arbitrary terminology that has been used to describe this interval west and east of the Devils River. Because of the different, and sometimes confusing nomenclature published on the Eagle Ford Shale, Donovan et al. (2013a) encouraged the use of the name Eagle Ford and not Boquillas for the strata located between the Buda and Austin Chalk formations in Texas, and proposed a new facies subdivision to standardize the Eagle Ford Group stratigraphic terminology (Donovan et al., 2013a; 2013b).

The Eagle Ford Shale outcrop belt trends in a northeast direction from El Paso in West Texas towards San Antonio, where the outcrops follow the edge of the East Texas Basin northward to the Oklahoma state line and eastward into Arkansas (Liro et al., 1994). In Terrell County, West Texas, the Lozier Canyon outcrop is a northeast-facing cutbank approximately 2000 ft long (~610 m) and 250 ft high (~76 m), which provides an excellent panoramic view of the complete Eagle Ford Group, the overlain Austin Chalk and the underlying Buda Limestone (Donovan and Staerker, 2010; Donovan et al., 2012; Donovan et al., 2013a and b). Donovan et al. (2012) referred to the Eagle Ford succession at Lozier Canyon as the Eagle Ford Group. These authors proposed a vertical facies succession for this unit based on their field observations and previous studies from other investigators. This succession, from base to top comprises the following five facies (Donovan et al., 2012; 2013a and b):

Facies A: Light gray hummocky cross-stratified limestones (grainstones) separated by thin calcareous mudstone beds.

Facies B: Black organic-rich calcareous mudstones with scattered limestones (packstone/grainstone) interbeds.

Facies C: Medium gray thick-bedded limestones (packstones) with mudstone interbeds.

Facies D: Pale yellow-ochre, echinoid-bearing marls and nodular limestones.

Facies E: Yellow ochre, thin-bedded limestones (grainstones) interbedded with calcareous mudstones.

Donovan et al. (2012; 2013a and b) further subdivided these five facies into 16 sub-units, which helped them define four unconformity-bounded depositional sequences (Figure 5). The Eagle Ford Group is divided into Lower and Upper formations by a regional unconformity at the contact between facies B and C. The Lower Eagle Ford Formation contains two depositional sequences, the Lozier Canyon and the Antonio Creek members, separated at the contact between sub-facies B2 and B3. Likewise, the Upper Eagle Ford Formation also contains two depositional sequences; the Scott Ranch and Langtry members divided at the contact between facies C and D.

At the Lozier Canyon outcrop, the Lozier Canyon Member (Figure 5) consists of interbedded grainstones and mudstones (subunits A1-A4) of approximately 18 to 20 ft (~6 m) overlain by about 30 ft (~9 m) of organic-rich mudstones and mudstones with hummocky-stratified grainstones (subunits B1 and B2). The Antonio Creek Member is characterized by bentonite-rich, dark gray mudstones, which show a marked decrease in organic content (Donovan et al., 2013b). The Scott Ranch Member is composed of interbedded light gray limestones and medium gray mudstones. Its basal section contains organic-poor, clay-rich mudstones and limestones (subunit C1), while the

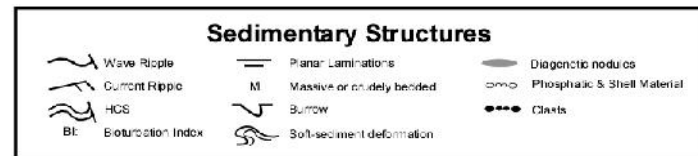
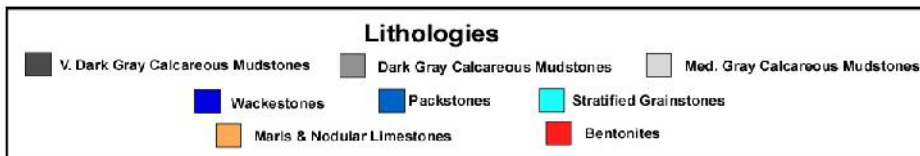
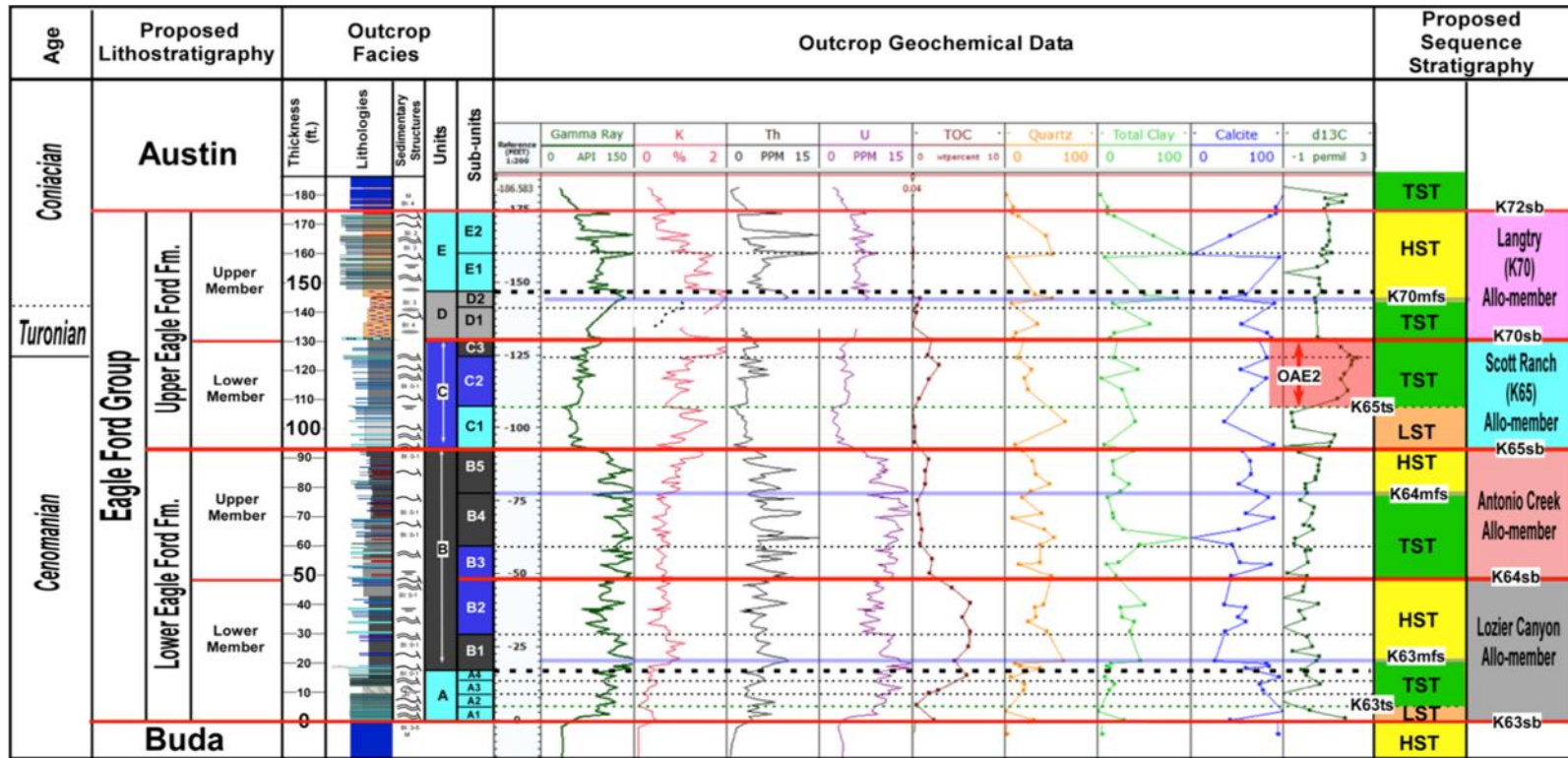


Figure 5. Sequence stratigraphic and facies subdivision proposed by Donovan et al. (2014) for the Eagle Ford Group at the Lozier Canyon outcrop based on petrophysical and geochemical data (from Donovan et al., 2014)

upper section (subunit C2) is characterized by high organic content, and a ^{13}C excursion that corresponds to the Oceanic Anoxic Event 2 (OAE2) and the Cenomanian-Turonian boundary. The top of this member contains a clay-rich, low organic content interval (subunit C3) marked by the end of the ^{13}C excursion (Donovan et al., 2012; 2013b). From base to top, the Langtry Member consists of pebble-size clasts, highly bioturbated yellowish-ochre echinoid-rich packstones (subunit D1), and wackestones (subunit D2) overlain by interbedded carbonate mudstones and hummocky-bedded packstones (subunit E). In addition, this member is also bentonite-rich, containing 2 to 6-inches-thick (5 to 15 cm) bentonite beds (Donovan et al., 2013b).

Based on geophysical and geochemical logs, Donovan et al. (2012; 2013a) were able to identify the same depositional sequences described at the Lozier Canyon outcrop in the South Texas subsurface (Figure 6). At the Fasken “A” #1 well (Webb County, Texas) the Eagle Ford Group is about 300 (~91 m) thick. However, in this area the Lozier Canyon and Scott Ranch members have a larger stratigraphic thickness compared to the equivalent units at the Lozier Canyon location. Moreover, the Lozier Canyon Member displays the highest organic content and corresponds to the main unconventional reservoir zone. Donovan et al. (2012; 2013a) identified a “marker beds” interval towards the base of the Upper Eagle Ford Formation characterized by high-clay, low-carbonate content, low gamma ray and resistivity, and a distinct density and neutron logs separation. Both, the Antonio Creek and Langtry members show a distinctive high gamma ray signature as a result of high bentonite content. In addition, a ^{13}C excursion within the Scott Ranch Member could be observed at the occurrence of the OAE2 and the Cenomanian-Turonian boundary.

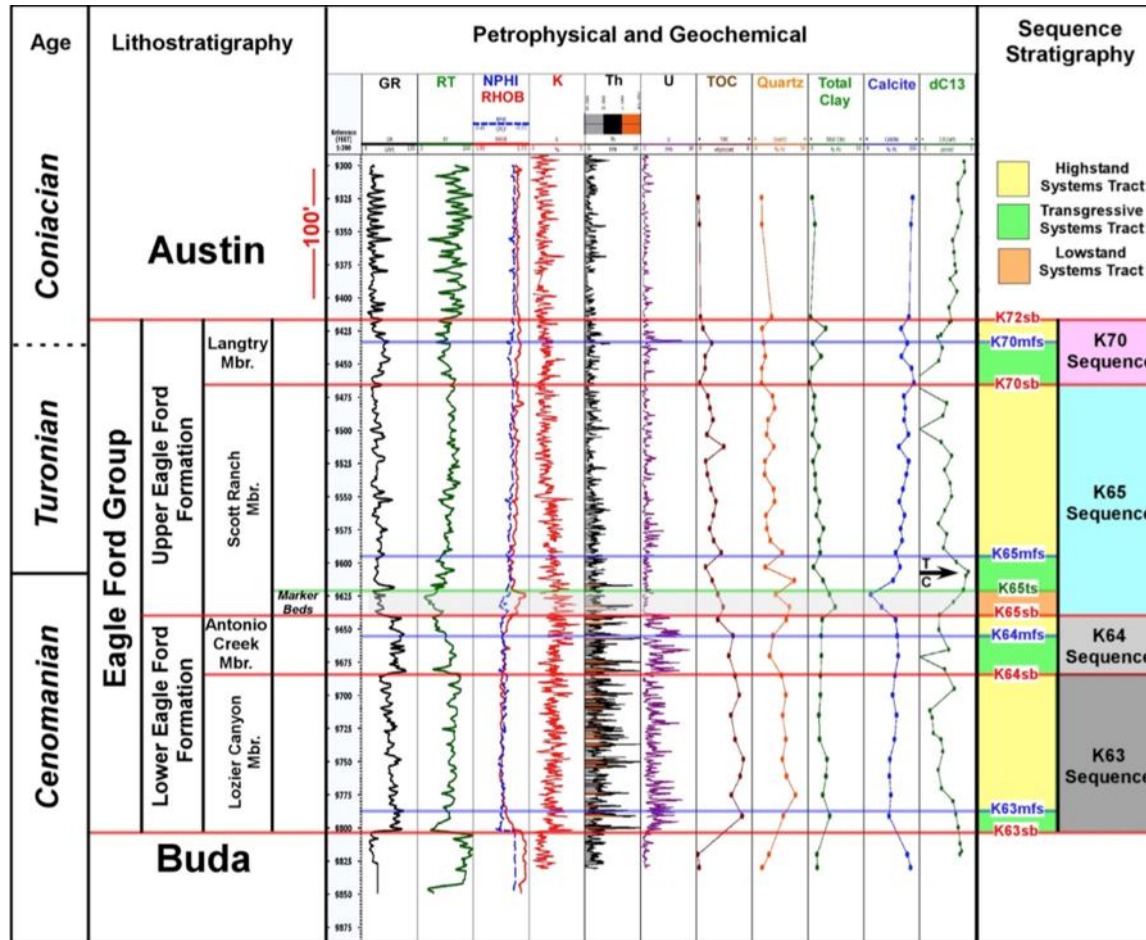


Figure 6. Sequence stratigraphic framework for the Eagle Ford Shale Group at the Fasken “A” #1 well in Webb County, South Texas based on petrophysical and geochemical data (modified from Donovan et al., 2013a)

1.1.3 Central Texas Eagle Ford Shale Geology

Fairbanks (2012) performed a facies analysis of the Eagle Ford Shale in Central Texas including samples from the Bouldin Creek outcrop, and the ACC#1 core. Harbor (2011) did a similar study in the Central Texas subsurface using samples from the W. Brechtel #1, and C.J. Hendershot #1, among other wells. The ACC#1 core is considered as a type section core for the Eagle Ford Group in the Austin area (Figure 7; Fairbanks, 2012). In this core, Fairbanks (2012) performed a detailed description of the Eagle Ford Shale and described the contacts with the underlying Buda Limestone and the overlying Austin Chalk. The author subdivided the Eagle Ford Group into four different units similar to studies previously done in the same area. From base to top these intervals correspond to the Pepper Shale, Lake Waco Formation (Waller and Bouldin Members), and South Bosque Formation (Figure 7). The facies described by Fairbanks (2012) in this area include massive argillaceous mudrocks; massive argillaceous foraminiferal mudrocks; laminated argillaceous foraminiferal mudrocks; laminated foraminiferal wackestone; cross-laminated foraminiferal packstones and grainstones massive bentonitic claystones; and nodular foraminiferal packstones and grainstones.

According to Harbor (2011) and Fairbanks (2012), the contact between the Buda and the Eagle Ford is sharp, marking a lithologic change from a light gray, highly-bioturbated, massive skeletal wackestone/packstone, which characterizes the Buda Limestone, to a dark gray, massive argillaceous mudstone that corresponds to the Eagle Ford Shale. The Pepper Shale of the Eagle Ford Group is a relatively thin (4 to 6 ft; ~1 to 2 m), recessive, dark gray, argillaceous claystone (massive argillaceous mudrock facies) interval with moderate TOC (ave. 3.2%; Fairbanks, 2012). This unit is overlain

by the Waller Member, which is mainly composed of massive and laminated argillaceous foraminiferal mudrock. The Waller Member has high TOC (ave. 3.7%) and higher carbonate content compared to the Pepper Shale and also contains smaller amounts of massive argillaceous mudrock, cross-laminated foraminiferal packstone/grainstone and nodular foraminiferal packstone/grainstone. In outcrop it is poorly exposed and its thickness in core is about 10 ft (~3 m). A sharp contact separates the Waller Member from the overlying Bouldin Member. According to Fairbanks (2012) the Bouldin Member is about 10 to 12 ft (~3 m) thick, contains the lowest clay and TOC content (ave. 2.1%), and is mainly composed of interbedded calcite-rich limestones and mudrock (laminated foraminiferal wackestone and cross-laminated and nodular foraminiferal packstones/grainstones facies). In this interval Fairbanks (2012) identified significant amounts of bentonitic claystones and enrichment in Mo, U, Mn, and V/Cr, which the author described as indication of maximum basin restriction during deposition of this member. Overlying the Bouldin Member is the South Bosque Formation. This unit is about 16 ft (~5 m) thick and is composed of the massive and laminated argillaceous foraminiferal mudrock facies, occasional amounts of bentonitic claystones, and moderate organic content (ave. 2.5%; Fairbanks, 2012). Fairbanks (2012) describes a gradational contact of about 3 ft (~91 cm) as the transition towards the overlying Austin Chalk Formation. The author describes this formation as a light gray, heavily bioturbated mudstone/wackestone.

Fairbanks (2012) assigns the Pepper Shale, Lake Waco Formation (Waller and Bouldin Members), and the basal part of the South Bosque Formation as the Lower Eagle Ford, and the rest of the South Bosque Formation as the Upper Eagle Ford Shale.

Harbor (2011) identified an additional upper member in the Eagle Ford interval south of the Austin area. This interval named “Harbor 2011” by Fairbanks (2012) is composed of disrupted bedded foraminiferal packstones, massive inoceramid packstones, and laminated foraminiferal peloidal packstones (Figure 8; Harbor, 2011). In general, all the Eagle Ford members described by Fairbanks (2012) show a southward thickening along the San Marcos Arch axis, possibly resulting from an increase in accommodation due to subsidence. This author observed an overall facies change to more fine-grained, low energy facies, with exception of the South Bosque Formation, which does not show a significant facies nor thickness variations (Fairbanks, 2012).

The Bouldin Creek outcrop is located on the northeastern flank of the San Marcos Arch and is the type locality for the Bouldin Member (Fairbanks, 2012). In this outcrop, separated for about 10 miles southwest from the ACC #1 only the intervals corresponding to the upper section of the Waller Member, the Bouldin Member, and the lower section of the South Bosque Formation are present (Fairbanks, 2012). A northeast-southwest cross-section from the Austin area across the San Marcos Arch (Figure 8; Fairbanks, 2012), which includes the ACC #1, W. Brechtel #1 and C.J. Hendershot #1 wells analyzed in this study show that the Pepper Shale is present across the region; however, the Waller Member pinches out towards the southwest being absent in the W. Brechtel #1 location (Fairbanks, 2012). On the contrary, the Bouldin Member thickens in the same direction.

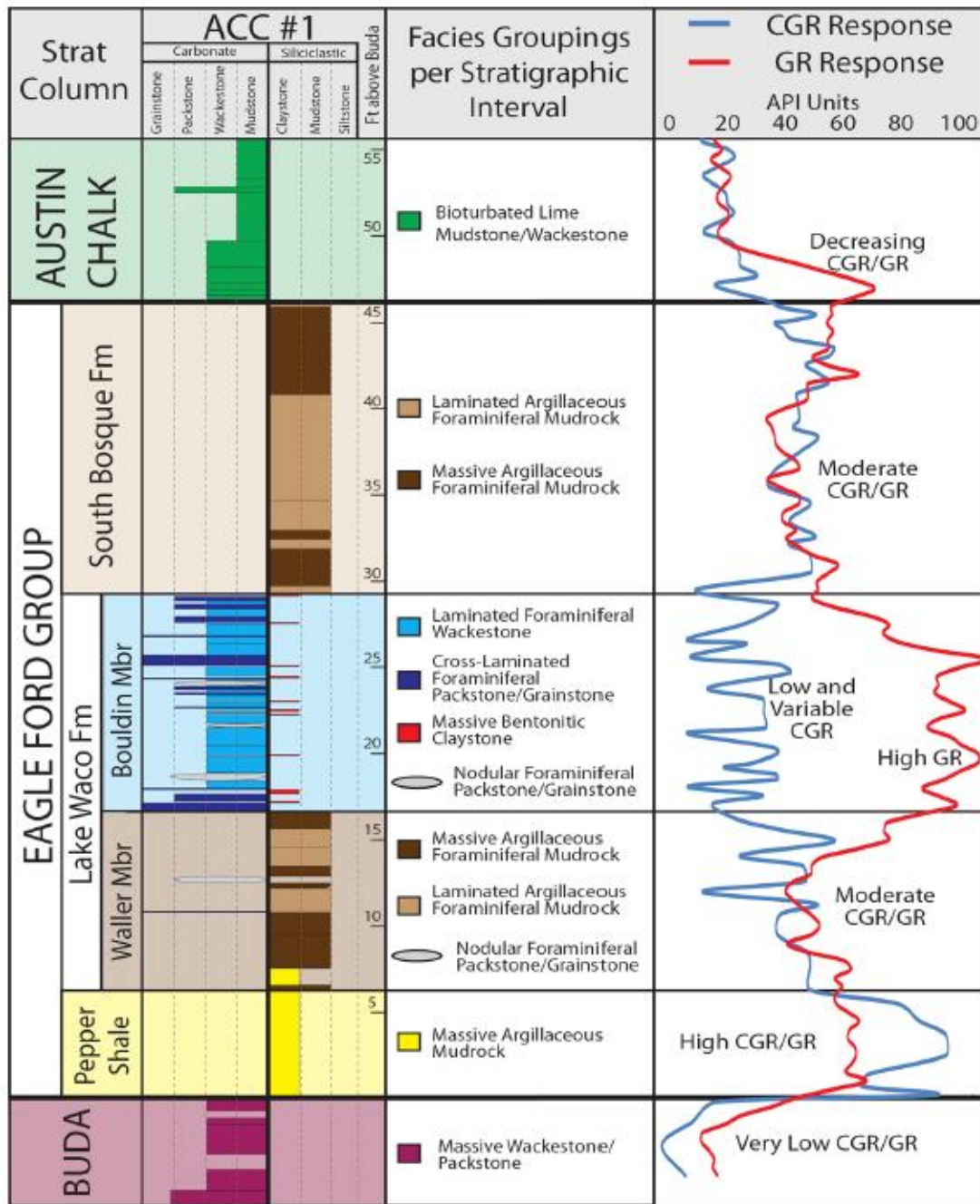


Figure 7. Core description, facies analysis, and gamma ray logs for the ACC #1 core from Fairbanks (2012; GR = gamma ray, CGR = computed gamma ray)

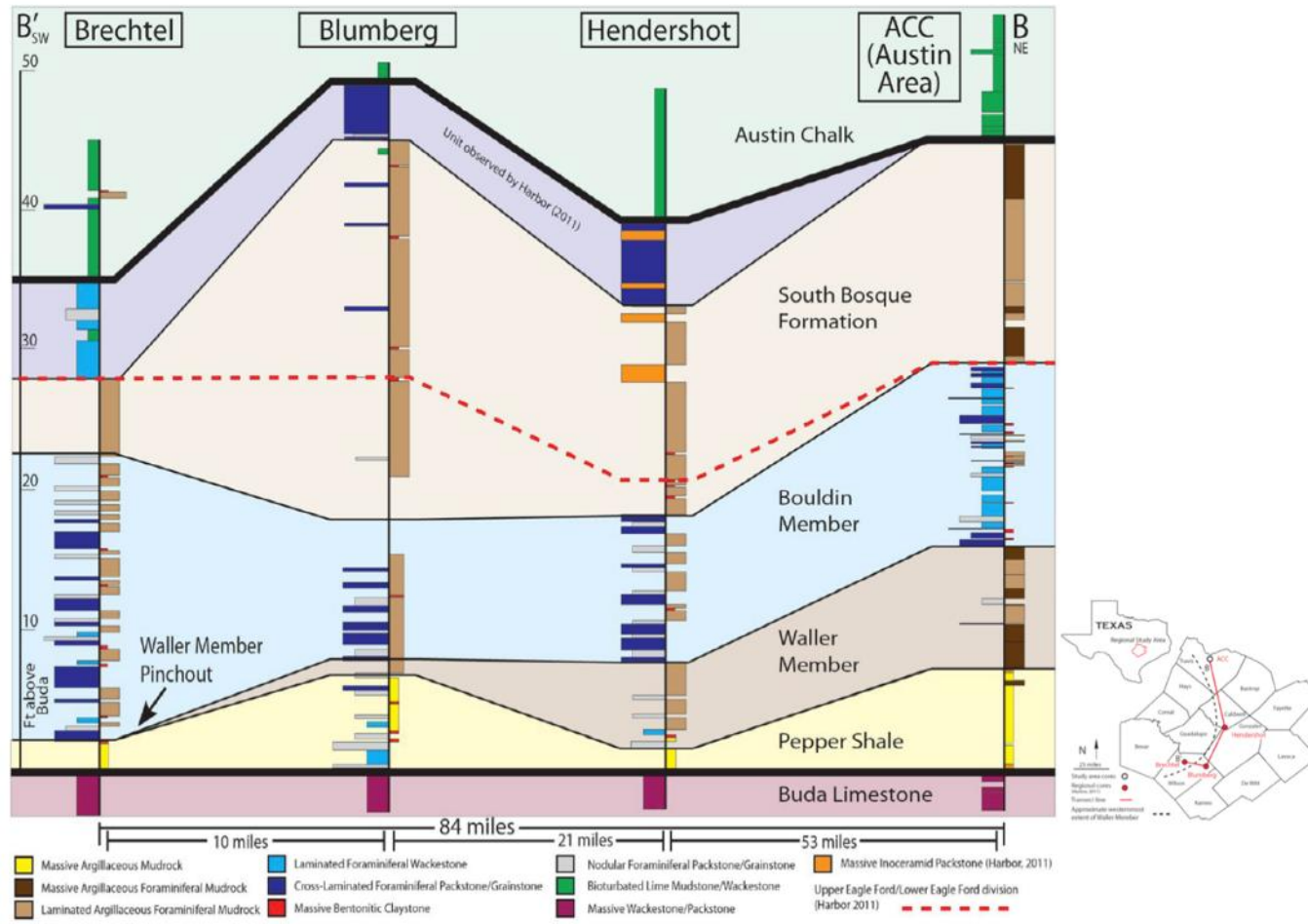


Figure 8. Stratigraphic cross-section showing facies variation and units continuity along the San Marcos Arch (modified from Fairbanks, 2012)

1.1.4 East Texas Eagle Ford Shale Geology

The Eagle Ford Shale thins towards the northwest from the Maverick Basin to the San Marcos Arch and then thickens in the same direction towards the East Texas Basin (Figure 9; Hentz and Ruppel, 2010). In East Texas, the interval between the Buda Limestone and the Austin Chalk includes from base to top the Maness Shale, Pepper Shale, and the Eagle Ford Shale. In the East Texas Basin, the siliciclastic succession of the Woodbine Group occurs between the Eagle Ford Shale and the mudrocks of the Maness Shale. The Woodbine sands pinchout southwest of the basin, and grade into the mudrocks of the Pepper Shale to the north of the San Marcos Arch (Figures 9 and 10; Hentz and Ruppel, 2010). In this area, the Eagle Ford Shale mainly consists of faintly laminated calcareous shales; organic-rich calcareous shales; laminated dolomitic shales; laminated marls, and occasional thin, reworked ash beds. These facies are interpreted as delta to prodelta deposits (Arthur and Sageman, 2004; Jennings and Antia, 2013). To the southwest, the Eagle Ford Shale contains sandstones in its uppermost part, which are referred as to the Sub-Clarksville Sand (Surlles, 1987). Underlying the Eagle Ford Shale, the Maness Shale is composed of silty, slightly calcareous or dolomitic, massive, and thinly laminated shales, which are described as prodelta deposits (Arthur and Sageman, 2004; Hentz and Ruppel, 2010; Jennings and Antia, 2013).

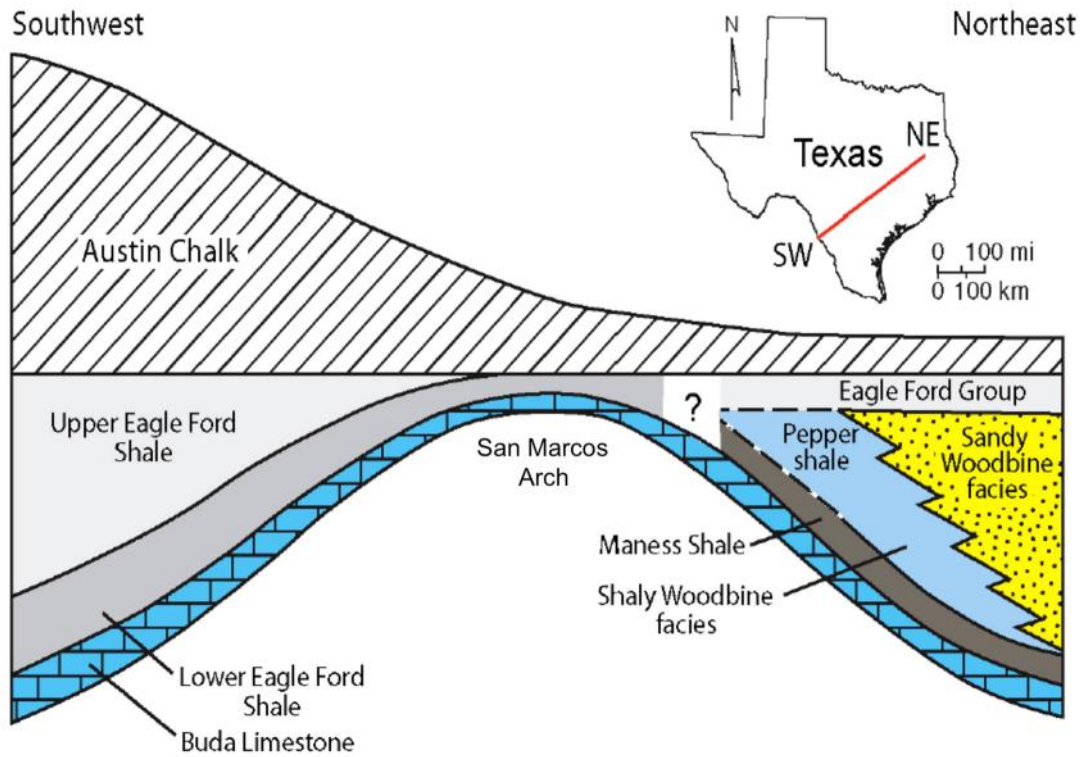


Figure 9. Generalized SW-NE schematic strike cross section illustrating the relationships among lithostratigraphic units across the Eagle Ford Shale Play (modified from Hentz and Ruppel, 2010)

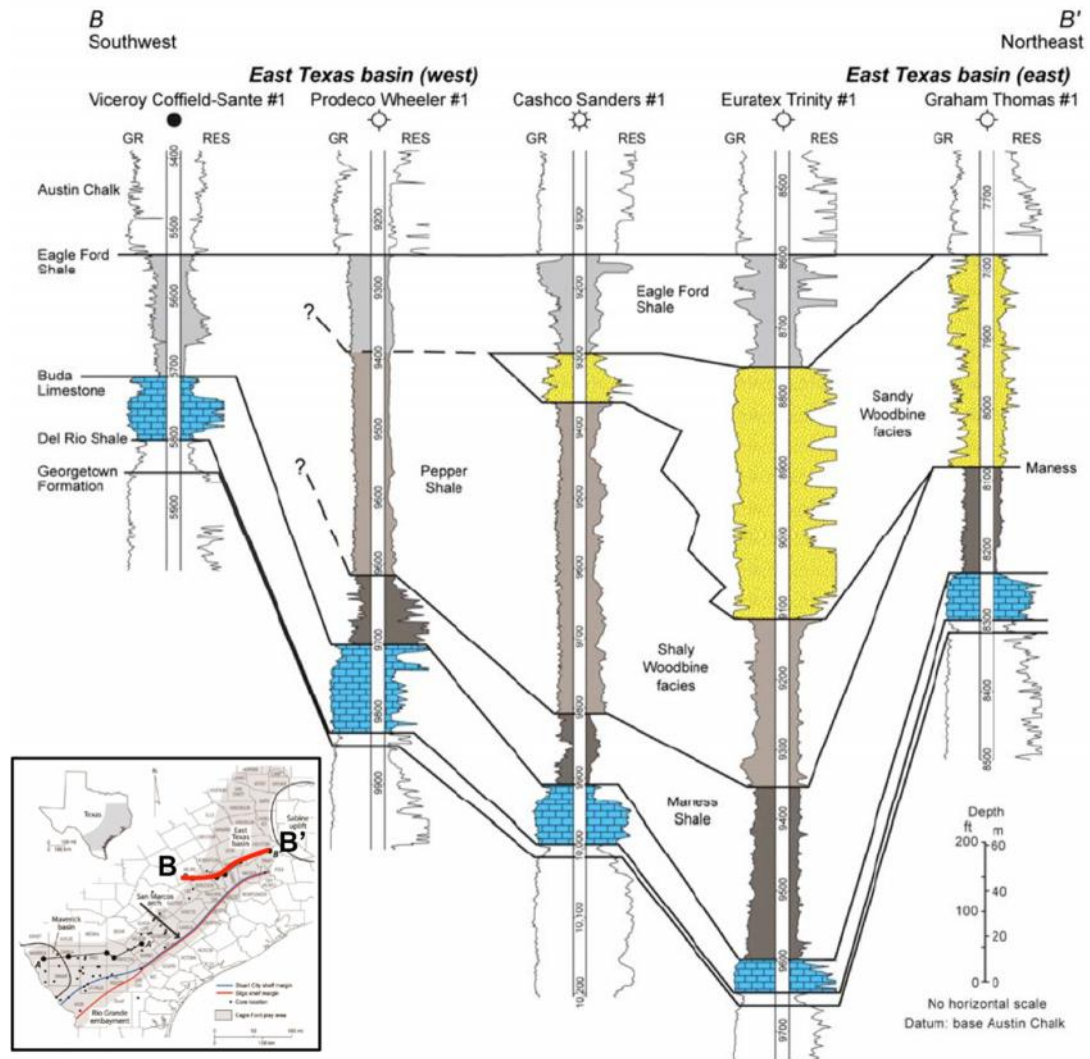


Figure 10. SW-NE regional strike cross section of the East Texas Basin and northeast flank of the San Marcos Arch (modified from Hentz and Ruppel, 2010)

1.1.5 Paleogeography and Climate

Deposition of organic rich rocks during the Cretaceous period was influenced by variations in different factors such as tectonism, volcanism, atmospheric and ocean chemistry, climate, sea-level, and sediment supply (Dean and Arthur, 1998). The impact of these phenomena produced important global eustatic sea level changes, which affected sedimentation and the extent of shelves and epicontinental seas both regionally and globally (Robinson and Kirschbaum, 1995; Dean and Arthur, 1998). In terms of plate tectonics, during the Late Cretaceous the South Atlantic Ocean started to open. North America was connected to Europe, and Australia and Antarctica were still joined. India and Madagascar began to separate, sending India on its collision course with the Eurasian plate (Scotese, 2001).

Globally, the Cretaceous climate was characterized by warm and equable temperatures, low to moderate precipitation, and a lack of ice caps, unlike today. These conditions have been attributed to a possible major “greenhouse” effect, where great amounts of CO₂ were released into the atmosphere due to an increase in the Earth’s volcanic activity (Robinson and Kirschbaum, 1995; Dean and Arthur, 1998).

From the middle Cretaceous, the interplay of these factors led to the development of “Oceanic Anoxic Events” (OAEs), in which mid- and deep-water oceanic masses across the Earth became highly oxygen depleted. OAEs reflect regional to global scale oceanographic and climatic changes due to a major perturbation in the ocean-atmosphere-terrestrial carbon cycle (Dean and Arthur, 1998). The Cenomanian-Turonian (C-T) boundary is marked by one of these events known as Oceanic Anoxic Event 2 (OAE2), characterized by a global sea level rise, an increase in burial of

organic carbon and consequent drop in atmospheric CO₂, and biotic turnover (Corbett et al., 2011). Before the beginning of the OAE2, CO₂ concentration (pCO₂) was about 4 to 18 times higher than present which contributed to the development of a greenhouse Earth (Arthur et al., 1988; Barclay et al., 2006). Increased nutrient levels resulting from higher weathering rates and upwelling produced a global increase in organic matter primary productivity, which in turn produced an important increase in organic carbon burial rates (Arthur et al., 1988; Fortwengler et al., 2003; Barclay et al., 2006; Sageman et al., 2008; Sinninghe Damsté et al., 2009). The burial of ¹³C-depleted organic carbon generated a rapid positive excursion (¹³C enrichment) in the stable carbon isotope composition of carbonate (¹³C_{carb}, 1.5-2‰), organic carbon (¹³C_{org}; 3.5-4‰), and phytoplankton, which can be determined at or near the C-T boundary in the sedimentary record worldwide (Arthur et al., 1988; Fortwengler et al., 2003; Sinninghe Damsté et al., 2009).

In North America, rock deposition during the Upper Cretaceous was highly influenced by the relationship between foreland basins, eustatic sea level variations, and climate. The paleogeographic location of North America during this time places the continent in a temperate to subtropical climate zone (Robinson and Kirschbaum, 1995). At the beginning of Cenomanian time, a transgression of the seas from the north and south into the North American craton began, creating the Western Interior Seaway (WIS; Figure 11) (Robinson and Kirschbaum, 1995). The WIS connected the Circum-Boreal sea with the proto-Gulf of Mexico, and during maximum transgressions it constituted a large north-south arm of the Tethys Ocean extending from the Gulf Coast to the Arctic region (Robinson and Kirschbaum, 1995; Dean and Arthur, 1998, 1998b).

WIS expansion during this time was mainly influenced by an increase in seafloor spreading rates and ice cap melting, while sediment accumulation was driven by Cordilleran tectonics, epeirogenic flexure, oceanographic variations, and climate change (Leckie et al., 1998; Scott et al., 2010). During this time, marine shale deposition was occurring in the central part of the seaway, along with extensive platform and basin carbonate buildup in southern Texas. By early Turonian the WIS had reached its maximum transgression, and extensive, highly calcareous muds were produced and deposited (e.g. upper Greenhorn Limestone, Eagle Ford Shale). It is thought that during this time climate was predominantly warm and an influx of terrigenous organic matter was minimal. The presence of volcanic ash as bentonite beds in some sequences from early to middle Turonian suggest that during this time moderate to intense volcanism was occurring (Robinson and Kirschbaum, 1995).

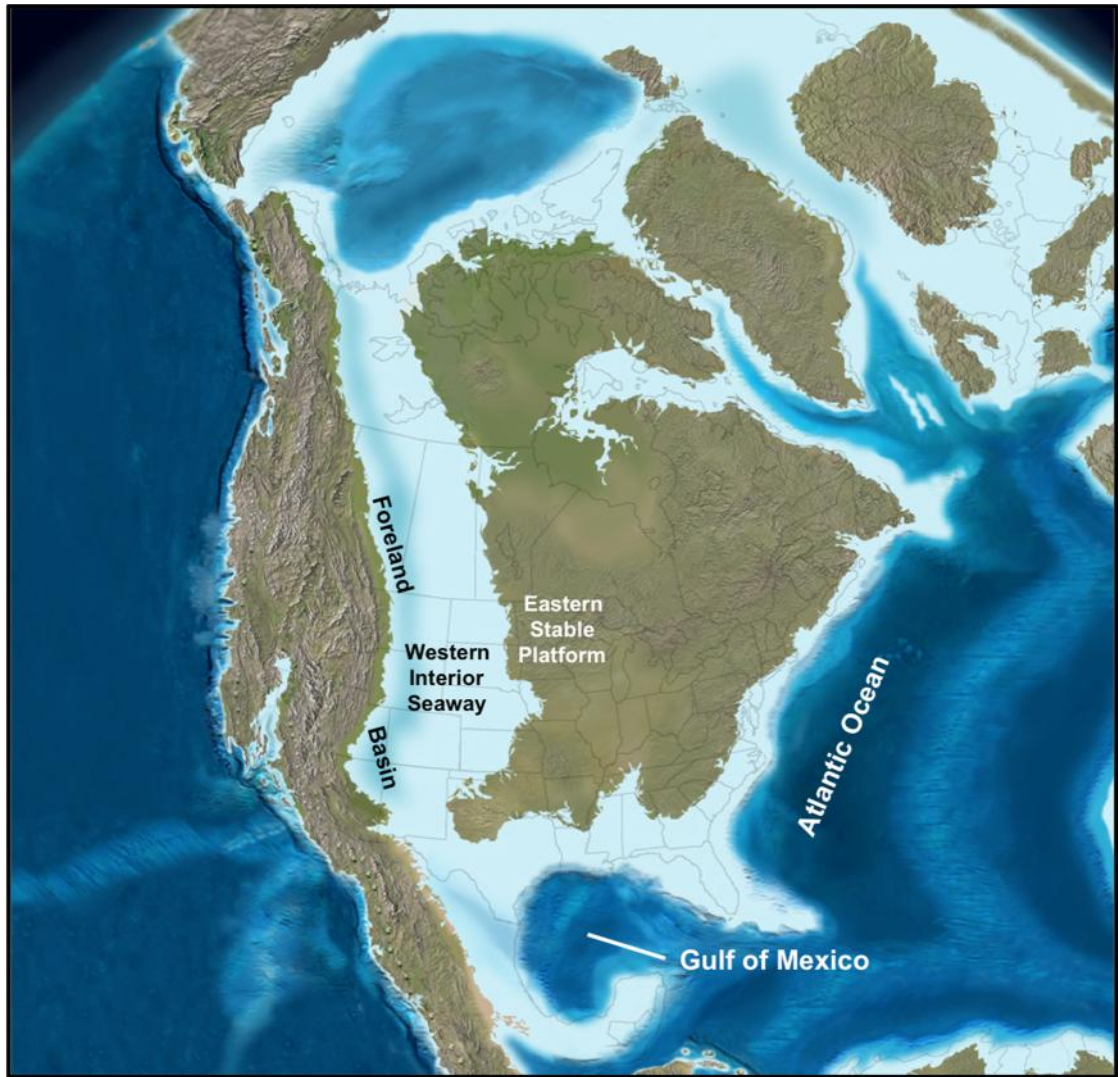


Figure 11. Map showing the extent of Western Interior Seaway, locations of the Foreland Basin, and the Eastern Stable Platform during the Late Cretaceous (~85 Ma) (modified from Robinson and Kirschbaum, 1995 and Blakey, 2011)

1.2 Objectives

The main objective of the present study is to geochemically characterize the Eagle Ford Shale in West, Southwest, Central and East Texas and demonstrate the usefulness of organic geochemistry as a tool for complementing and refining sequence stratigraphic frameworks, especially in unconventional shale gas plays. For this, an assessment of possible variations in different organic geochemical parameters in the Eagle Ford Shale from outcrop to subsurface in a local and regional context is intended. In order to accomplish these goals, the following specific objectives were defined:

- Determination of the organic richness and type of organic matter present in the Eagle Ford Shale.
- Evaluation of the source rock generation potential for hydrocarbons and thermal maturity.
- Assessment of variations in organic matter source and depositional environment conditions.
- Integration and correlation of the organic geochemistry data obtained from this research with available sequence stratigraphy data for the study area.
- Characterization of oils and condensates produced from Upper Cretaceous reservoirs and correlation with Eagle Ford Shale geochemical parameters in order to increase the understanding of the petroleum systems in the study area.

1.3 Methodology

1.3.1 Study area and sample locations

A total of 178 samples were collected for this study from outcrops and wells located throughout West, South-central, and East Texas (Figure 12, Table 1). In West Texas, the BP-leased Lozier Canyon outcrop provides an excellent exposure to evaluate organic facies variations that may be present within the Eagle Ford Shale. Results from a multidisciplinary study of this outcrop have been published in several field guidebooks and in Donovan and Staerker (2010) and Donovan et al. (2012, 2013a; 2013b). In South Texas, core samples obtained from the Newfield F. McKnight #526-1H and Swift Fasken “A” #1H wells were used to understand organic geochemical variations in the subsurface and in areas where the Eagle Ford Shale is highly mature. The West Bouldin Creek outcrop in Austin, Texas includes a thinner section of the Eagle Ford Shale and corresponds to the type locality of the Bouldin Member (Figure 12). Close to this area, core samples from the ACC#1 well, the type section for the Eagle Ford Group in Central Texas were also collected for comparison with the Bouldin Creek outcrop. Further south, two additional cores were sampled, the C.J. Hendershot #1 and W. Brechtel #1, which have been described by Harbor (2011). Finally, in East Texas, a core obtained from the Lily Hoppess #1 well containing sections corresponding to the Lower Eagle Ford and Maness Shale was used to geochemically characterize what is thought to be the most prolific interval for hydrocarbons in the Eagle Ford Shale and to evaluate regional variations that may be present between west, central, and east Texas. Additionally, oil and condensate samples from five wells

(Figure 12) located in southwest and central Texas were analyzed for and source rock/oil correlations.

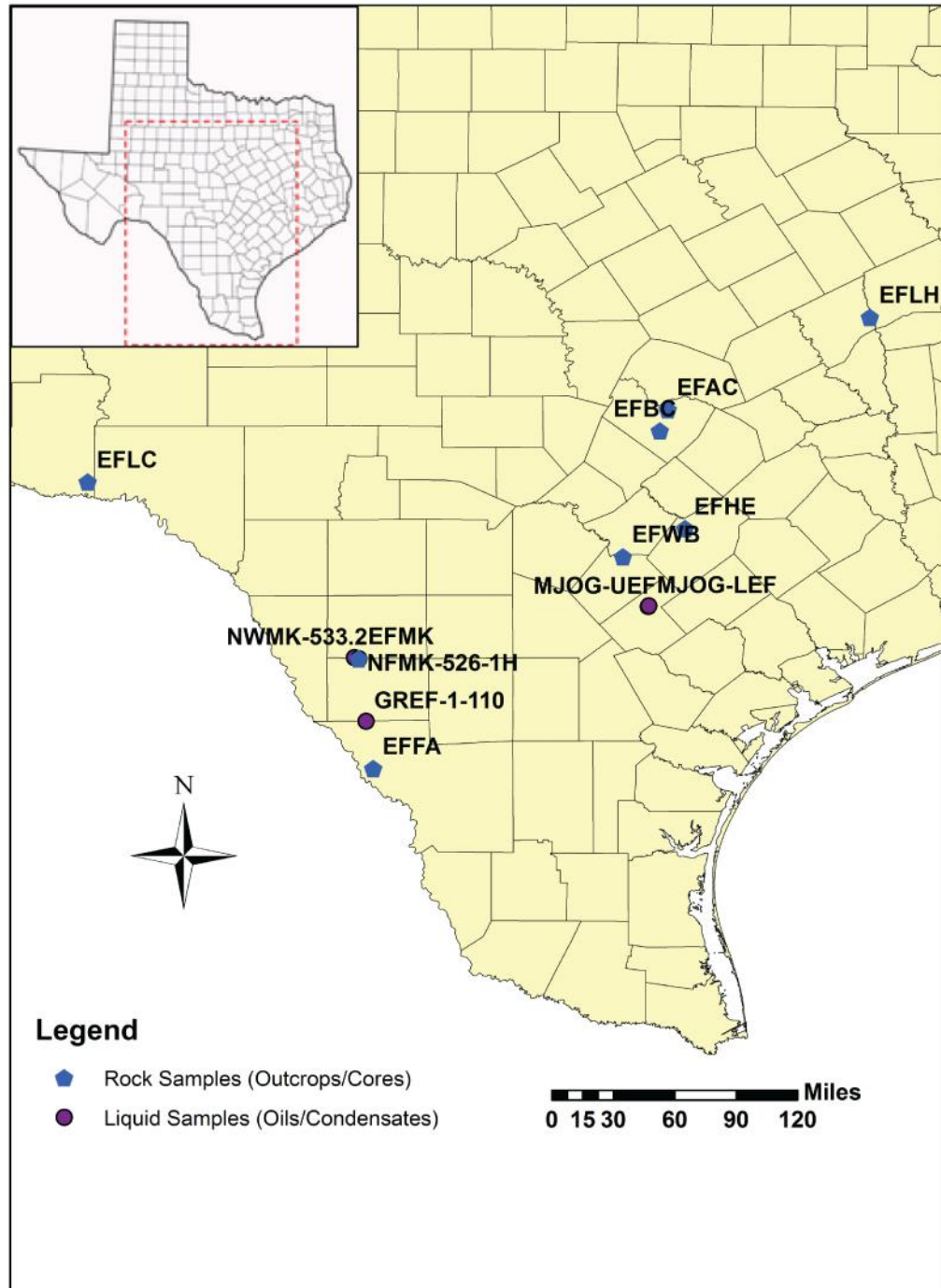


Figure 12. Texas map showing location of the samples analyzed in this study

1.3.2 Experimental

Total Organic Carbon (TOC) determination and Rock Eval pyrolysis were performed on all rock samples available for this study (Table 1). A TOC value of 1% was established as a cutoff for selecting samples for bitumen extraction. Crushed rock extracts were separated into maltenes and asphaltenes by addition of excess *n*-pentane. The maltene fraction was further fractionated into saturates, aromatics, and NSO compounds by high performance liquid chromatography (HPLC). Screening analyses of saturates and aromatics were performed by gas chromatography (GC). Selected samples were analyzed by gas chromatography-mass spectrometry (GCMS) for biomarker analyses, and gas chromatography isotope-ratio mass spectrometry (GC-IRMS) for compound-specific isotope analysis. Oils and condensates were analyzed using GC and GCMS. Rock samples from Lozier Canyon, Bouldin Creek, Fasken A #1-H, and Lily Hoppess #1 were chosen for vitrinite reflectance measurements. Rock Eval pyrolysis and TOC analyses were performed at GeoMark Research, Inc. in Humble, Texas and Core Laboratories in Houston, Texas. The remaining geochemical analyses were performed at the University of Oklahoma Organic Geochemistry Laboratories located in Norman, Oklahoma. A summary of the laboratory workflow used in this project is depicted in Figure 13.

Table 1. List of samples for organic geochemical analyses

Well/Outcrop name	Abbreviation	Sample type	# of samples	Location
Lozier Canyon	EFLC	Outcrop	43	Terrell Co.
Newfield Ferguson McKnight #526-1H	EFMK	Core	16	Dimmit Co.
Swift Fasken "A" #1-H	EFFA	Core	40	Webb Co.
Bouldin Creek	EFBC	Outcrop	34	Travis Co.
ACC#1	EFAC	Core	10	Travis Co.
W. Brechtel #1	EFWB	Core	9	Wilson Co.
C.J. Hendershot #1	EFHE	Core	10	Caldwell Co
Encana Lily Hoppess #1	EFLH	Core	11	Robertson Co.
Newfield W. McKnight #533.2	NWMK-533.2	Oil	1	Zavala Co.
Newfield F. McKnight #526-1H	NFMK-526-1H	Oil	1	Dimmit Co.
Murphy Oil Murphy JOG #1-H	MJOG-UEF	Oil	1	Karnes Co.
Murphy Oil Murphy JOG #E 1-H	MJOG-LEF	Oil	1	Karnes Co.
Lewis Gates Ranch #1-110	GRAF-1-110	Condensate	1	Dimmit Co.

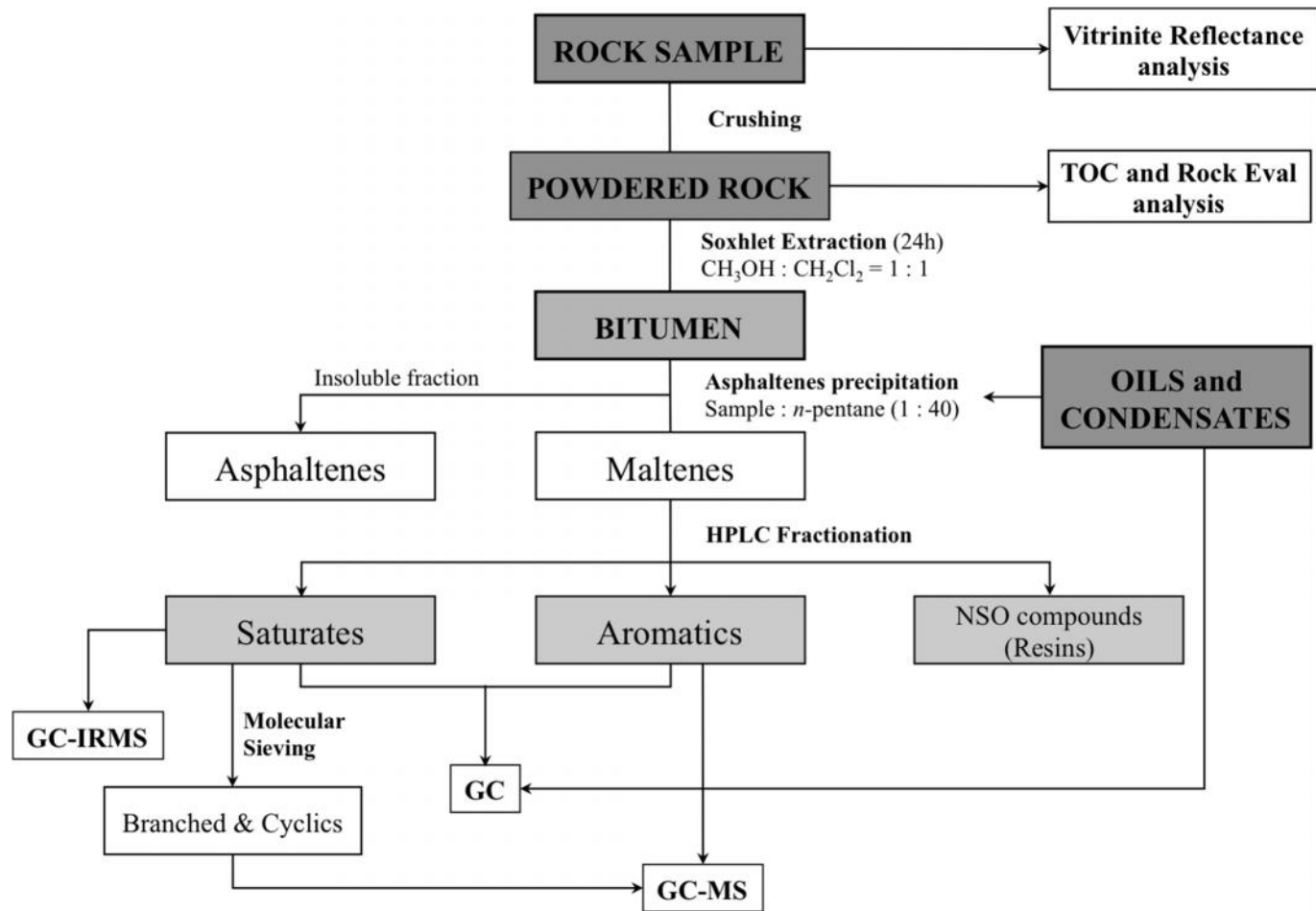


Figure 13. Schematic workflow used for laboratory analyses of the Eagle Ford Shale rock, whole oil and condensate samples (modified from Miceli Romero and Philp, 2012)

1.3.2.1 Preliminary rock sample treatment

Individual rock samples were washed with hot water, distilled water and a 1:1 mixture of dichloromethane (CH_2Cl_2) and methanol (CH_3OH) in order to remove any possible contaminants (e.g. drilling mud, plastic wrap, and handling). After the samples were completely dried, they were crushed using a combination of a small shatterbox and an alumina mortar and pestle until a fine powder was obtained for screening analysis (TOC and Rock Eval) and soxhlet extraction.

1.3.2.2 Total Organic Carbon (TOC) and Rock Eval Pyrolysis

Total Organic Carbon (TOC) analysis and Rock Eval pyrolysis were performed on 173 rock samples at GeoMark Research, Inc. in Humble, Texas and Core Laboratories in Houston, Texas. Approximately 1-2 grams of crushed rock were used for determination of TOC and Rock Eval parameters. From the results of these analyses samples were chosen for further bitumen extraction and maltenes fractionation.

1.3.2.3 Petrographic analysis

Measured vitrinite reflectance (Measured % R_o) values were obtained from pellets prepared at the Oklahoma Geological Survey Organic Petrography Laboratories in Norman, Oklahoma and measured at the University of Oklahoma Organic Geochemistry Laboratories under the guidance of Mr. Brian Cardott. Preparation of dispersed organic pellets involved grinding about 10 grams of whole rock samples to 18 mesh and placing them in plastic ring forms. A mixture of epoxide epoxy resin and hardener (5:1) was centrifuged for 3 minutes to remove air bubbles from it and then poured into the ring forms until half full. Each crushed rock sample was sprinkled into

its designated ring form and mixed with the epoxy until all the grains were covered with the epoxy mixture and evenly spread across the bottom of the ring form. Additional bubble-free epoxy was poured into the ring forms up to the rim. The pellets were left to harden at room temperature for about 12 hours.

Dispersed organic pellets were polished using a Buehler Ecomet III Grinding and Polishing Apparatus in order to remove any scratches and obtain a relief-free surface for visual analysis. Pellets were polished with 320, 400, and 600 grit paper for 3 minutes using tap water as lubricant. Each sample was subsequently rinsed off using distilled water and placed in an ultrasonic bath with distilled water and few drops of Kodak Photo-Flo 200 solution for 1 to 2 minutes. Sample pellets were further polished using a Buehler Texmet polishing cloth with Wendt Dunnington 0.3 μ m alumina slurry and distilled water as lubricant for 4 minutes. Pellets were then washed with distilled water and placed in the ultrasonic bath for 2 minutes. After this step, the pellets were rinsed with distilled water and blow-dried using filtered air. Finally, the pellets were polished with Wendt Dunnington 0.05 μ m alumina slurry following the same workflow indicated above, and placed in a desiccator to dry.

Mean random vitrinite reflectance measurements were performed using a Leitz Ortholux 2 Microscope under oil immersion and reflected white light following the methodology from ASTM (2011).

1.3.2.4 Extraction and Fractionation

A soxhlet extraction device was pre-extracted for 24 hours using a 1:1 mixture of dichloromethane (CH₂Cl₂) and methanol (CH₃OH) in order to remove contaminants. Selected samples (50g approximately) were extracted with the same solvent mixture for

24 hours. Excess solvent from the extract was evaporated using a rotary evaporator and the bitumen transferred to a glass centrifuge tube. The bitumen was separated into maltenes and asphaltenes by adding an excess (40:1) of *n*-pentane (C₅H₁₂). The centrifuge tubes were placed in a freezer overnight to ensure complete asphaltene precipitation.

Maltene fractions were transferred to 250mL round bottom flasks to evaporate the solvent excess and then placed in glass vials. This fraction was then diluted in a ratio of 20mg sample per 50uL *n*-hexane (C₆H₁₄) for compound class fractionation by HPLC using an HPLC Agilent 1260 Infinity. Saturates, aromatics and NSO (nitrogen, sulfur, and oxygen) compounds (or resins) were separated using different flow rates and solvent mixtures of *n*-hexane (C₆H₁₄) and dichloromethane, dichloromethane and ethyl acetate (C₄H₈O₂), and ethyl acetate and methanol respectively.

Saturate and aromatic fractions were diluted using 1mL of dichloromethane per 3mg of sample for GC and GC-MS analyses. The same dilution rate was used on the saturates for GC-IRMS analyses.

1.3.2.5 *Molecular sieving*

n-Alkanes were isolated from the saturate fraction using molecular sieves following the methodology described by West et al. (1990). A glass wool-plugged Pasteur pipette was packed with approximately 2g of silicalite powder that was previously activated by heating overnight at 500°C and washed with three bed volumes of *n*-pentane in order to remove impurities. The sample was added to the column and allowed to stand for 2 minutes. Then three additional bed volumes of *n*-pentane were used to elute the sample. The *n*-alkanes were retained in the sieve while the branched

and cyclic (B&C) compounds eluted from the column. This fraction was also diluted with 1mL of dichloromethane per 3mg of sample for biomarker analyses.

1.3.2.6 Gas Chromatography (GC)

GC analyses were performed using an Agilent 6890 series gas chromatograph equipped with a 30m x 0.32mm Agilent/J&W Scientific HP-5 fused silica capillary column coated with a 0.25 μ m liquid film and a flame ionization detector (FID). The volume injected in the GC instrument was 1 μ L per sample. The temperature program started with an initial temperature of 40°C with 10 minutes hold time. The temperature was increased to 300°C at a rate of 4°C per minute followed by an isothermal period of 24 minutes for a total run of 99 minutes. The injector and flame ionization detector (FID) temperatures were set at 300°C and 310°C respectively. Samples were analyzed in splitless mode using helium (He) as the carrier gas at a flow rate of 1.4mL per minute. *n*-alkanes and isoprenoids were identified in each chromatogram using relative retention times.

1.3.2.7 Gas Chromatography-Mass Spectrometry (GC-MS)

Biomarker analyses of the branched and cyclic alkanes (B&C) and aromatic fractions were carried out using an Agilent 7890 gas chromatograph coupled with a 5975 mass selective detector (MSD) using single ion monitoring (SIM, Table 2). The GC used a 60m x 0.25mm Agilent/J&W Scientific DB-5 fused silica capillary column coated with a 0.25 μ m film thickness. The injected volume of branched and cyclic and aromatic fractions was 1 μ L per run. The GC program started at 40°C with 1.5 minutes hold time and was later increased to 300°C at a rate of 4°C per minute and then held

constant for 34 minutes for a total run time of 100.5 minutes. Samples were run in splitless mode and Helium was used as the carrier gas with a flow rate of 1.4mL per minute. Biomarkers were identified from fragmentograms corresponding to each ion using relative retention times and by comparison with published data.

Table 2. List of ions selected for GC-MS analysis of branched and cyclic and aromatic fractions from Eagle Ford Shale samples

Fraction	m/z	Biomarker
Branched & Cyclics	66.0	Deuterated <i>n</i> -Tetracosane (internal standard)
	191.3	Tri/tetra/pentacyclic Hopanes
	217.3	Regular steranes, Diasteranes, Pregnanes
	133.1	Aryl Isoprenoids
	253.3	Monoaromatic Steroids
Aromatics	188.2	Deuterated Phenanthrene (internal standard)
	178.3	Phenanthrene
	192.3	Methylphenanthrenes
	206.3	Dimethylphenanthrenes
	231.3	Triaromatic Steranes

1.3.2.8 Quantitative Biomarker Analysis

Quantitation of biomarkers present in the branched and cyclics, and aromatic fractions of source rock extracts, condensates, and oils was done by adding a known aliquot of internal standard (ISTD) to each sample before GCMS analysis. The branched and cyclics, and aromatic samples were quantitatively spiked with deuterated *n*-tetracosane (C₂₄D₅₀) and deuterated phenanthrene (C₁₄D₁₀) respectively. The resulting solution was injected into the GCMS. The relative concentration of biomarkers in ppm or µg/g SAT (saturates) or ARO (aromatics) reported here was calculated from peak areas of each biomarker compared to that of the ISTD using a response factor.

In particular, the response factor of the ISTD (RF_{ISTD}) for each sample was calculated using the peak area of the ISTD (A_{ISTD}) as follows:

$$RF_{ISTD} = \frac{C_{ISTD\ inj}}{A_{ISTD}} \quad [1]$$

The concentration of the ISTD in the solution prepared for GCMS injection ($C_{ISTD\ inj}$) was calculated using the formula:

$$C_{ISTD\ inj} = \frac{C_{ISTD} \times V_{ISTD}}{V_{B\&C\ or\ ARO} + V_{ISTD}} \quad [2]$$

where,

C_{ISTD} = Concentration of ISTD in the internal standard solution

V_{ISTD} = Total volume of internal standard added to the B&C or Aromatics solution

$V_{B\&C\ or\ ARO}$ = Total volume of B&C or Aromatics solution prepared for injection

In our experiments, $V_{B\&C\ or\ ARO} \gg V_{ISTD}$ (e.g. $V_{B\&C\ or\ ARO} = 100\mu\text{L}$ and $V_{ISTD} = 2\mu\text{L}$), therefore equation [2] can be approximately calculated as:

$$C_{ISTD\ inj} \approx \frac{C_{ISTD} \times V_{ISTD}}{V_{B\&C\ or\ ARO}} \quad [3]$$

The concentration of the biomarker (C_a) in the solution prepared for GCMS injection was calculated using the formula:

$$C_a = RF_{ISTD} \times A_{bio}$$

$$C_a \approx \frac{C_{ISTD} \times \left[\frac{V_{ISTD}}{V_{B\&C\ or\ ARO}} \right]}{A_{ISTD}} \times A_{bio} \quad [4]$$

where,

A_{bio} = Peak area of the biomarker

The concentration of the solution prepared for GCMS injection (C_b) was calculated as follows:

$$C_b = \frac{C_{B\&C \text{ or } ARO} \times V_{B\&C \text{ or } ARO}}{V_{B\&C \text{ or } ARO} + V_{ISTD}} \quad [5]$$

and $C_b = C_{B\&C}$

where,

$C_{B\&C \text{ or } ARO}$ = Concentration of the B&C or Aromatics solution

Hence, the concentration of biomarkers in the saturate or aromatic fraction ($C_{bio X}$) is:

$$C_{bio X} = \frac{C_a}{C_b} \times F_X \times (1 \times 10^6) \quad [6]$$

or

$$C_{bio X} \approx \frac{C_{ISTD} \times V_{ISTD}}{A_{ISTD} \times V_{B\&C \text{ or } ARO}} \times \frac{1}{C_{B\&C \text{ or } ARO}} \times A_{bio} \times F_X \times 1 \times 10^6 \quad [7]$$

where,

Subscript X = Saturate or aromatic fraction

If X = Saturate, $F_{Sat} = \frac{m_{B\&C}}{m_{Sat}}$

IF X = Aromatic $F_{Aro} = 1$

$m_{B\&C}$ = Amount of B&C obtained from molecular sieving

m_{Sat} = Amount of saturate fraction used for molecular sieving

1×10^6 = Conversion factor to μg biomarker/g Saturates or Aromatics

In case of source rocks extracts, it is recommended to express the concentrations in μg biomarker/g TOC, whereas crude oil concentrations should be reported in μg biomarker/g whole oil. Therefore, after obtaining the biomarker concentrations relative to the saturate or aromatic fractions, another factor needs to be added to the calculations in order to normalize the biomarker concentrations relative to TOC and whole oil, respectively.

For source rock extracts:

$$C_{bio\ X} = \frac{\mu\text{g biomarker}}{\text{g Saturates}} \times \frac{m_{Sat}}{m_{TOC}}$$

and

$$\frac{m_{Sat}}{m_{TOC}} = \frac{\frac{m_{Sat}}{m_{Maltenes}} \times \frac{m_{Maltenes}}{m_{Extract}} \times \frac{m_{Extract}}{m_{Rock}}}{\frac{m_{TOC}}{m_{Rock}}}$$

where

$\frac{m_{Sat}}{m_{TOC}}$ = Amount of saturate fraction, m_{Sat} (g) within one gram (1g) of total

organic carbon (TOC)

$\frac{m_{Sat}}{m_{Maltenes}}$ = Amount of saturate fraction, m_{Sat} (g) recovered from the maltenes

fraction, $m_{Maltenes}$ (g) by HPLC

$\frac{m_{Maltenes}}{m_{Extract}}$ = Amount of maltenes, $m_{Maltenes}$ (g) recovered from the source rock

extract, $m_{Extract}$ (g) by n -C₅ precipitation

$\frac{m_{Extract}}{m_{Rock}}$ = Amount of extract, $m_{Extract}$ (g) recovered from the source rock, m_{Rock}

(g) by Soxhlet extraction

$\frac{n_{TOC}}{m_{Rock}}$ = Amount of TOC m_{TOC} (g) determined in the source rock m_{Rock} (g) by

LECO TOC

TOC is generally reported as weight percent (% wt.), which corresponds to the amount of TOC (g) in 100g of rock.

For oil samples:

$$C_{bio X} = \frac{\mu g \text{ biomarker}}{g \text{ Saturates}} \times \frac{m_{Sat}}{m_{WO}}$$

and

$$\frac{m_{Sat}}{m_{WO}} = \frac{m_{Sat}}{m_{Maltenes}} \times \frac{m_{Maltenes}}{m_{WO}}$$

$\frac{n_{Sat}}{n_{WO}}$ or g Sat/g oil is the number of grams saturate fraction recovered from one gram oil.

$\frac{m_{Sat}}{m_{Maltenes}}$ is the amount of saturate fraction m_{Sat} (g) recovered from $m_{Maltene}$ (g) maltene by liquid chromatography.

$\frac{n_{Maltenes}}{n_{WO}}$ is the amount of Maltene $m_{Maltene}$ (g) recovered from m_{oil} (g) oil by n -C₅ precipitation.

Regarding the biomarker concentrations in the aromatic fraction, the term m_{Sat} was substituted for the term m_{Aro} , which corresponds to the amount of aromatic fraction recovered from the maltenes fraction by HPLC.

1.3.2.9 Gas Chromatography Isotope-Ratio Mass Spectrometry (GC-IRMS)

Individual compounds of the saturate fractions were analyzed for compound specific isotopes using an Agilent 7890 gas chromatograph with a Isolink and Carbon combustion reactor interfaced to an Conflo IV and a Thermo MAT 253 isotope-ratio mass spectrometer. The GC used a 60m x 0.25mm x 0.25 μ m film Agilent/J&W Scientific DB-1MS capillary column. The injection volume of sample was 1 μ L. The temperature program was initially set at 40°C and held isothermal for 1.5 minutes. Then it was increased at a rate of 4°C per minute to 300°C and held constant for 24 minutes for a total run time of 90 minutes. Samples were analyzed in splitless mode injection using He as the carrier gas. This method pulses CO₂ standard according to the time events set up. The samples are run with deuterated *n*-nonane (C₉D₂₀), *n*-decane (C₁₀D₂₂), *n*-hexadecane (C₁₆D₃₄), *n*-nonadecane (C₁₉D₄₀), *n*-tetracosane (C₂₄D₅₀), and *n*-dotriacontane (C₃₂D₆₆) as external standards.

CHAPTER II

2. RESULTS AND DISCUSSION

Data presented in the following chapters uses the Formations, Members, and Facies subdivision proposed by Donovan et al. (2012, 2013a and b, 2014) for the Eagle Ford Group at the Lozier Canyon outcrop (EFLC), Fasken “A” #1 core (EFFA), Bouldin Creek outcrop (EFBC), and ACC #1 core (EFAC). Harbor (2011) and Fairbanks (2012) also defined a facies subdivision for the Bouldin Creek outcrop (EFBC) and ACC #1 (EFAC) cores. These data overlap allowed the assignment of the same nomenclature from Donovan et al. (2012, 2013a and b) to these cores and to extend it to the W. Brechtel #1 (EFWB) and C.J. Hendershot #1 (EFHE) cores. This nomenclature was adopted to standardize the different geologic units and facilitate the interpretations and correlations of geochemical data. In the case of the Ferguson McKnight #526-1H (EFMK) and Lily Hoppess #1 (EFLH) wells, the available data was not sufficient to assign Donovan et al.’s (2012, 2013a and b) terminology and the original Formation assignment will be used.

2.1 Geochemical Screening

Outcrop and subsurface rock samples analyzed in this study were initially screened using Rock Eval pyrolysis and TOC. From these data, sets of subsamples were selected for biomarker and isotope analyses. Table 3 shows the definitions of Rock Eval pyrolysis terms used in the discussion and presented in subsequent Figures and Tables. Appendix B shows detailed Rock Eval pyrolysis and TOC data obtained from the outcrop and core samples analyzed.

Table 3. Parameters, terms, and definitions derived from Rock Eval pyrolysis analyses (modified from Peters, 1986 and Jarvie et al., 2007)

TOC	Total Organic Carbon	wt. %
S₁	Free volatile hydrocarbons thermally flushed from a rock sample at 300°C (free oil content)	mg HC/g rock
S₂	Products that crack during standard Rock Eval pyrolysis temperatures (remaining potential)	mg HC/g rock
S₃	Organic carbon dioxide released from rock samples	mg CO ₂ /g rock
T_{max}	Temperature at peak evolution of S ₂ hydrocarbons	°C
HI	Hydrogen Index = $S_2 \times 100/TOC$	mg HC/g TOC
OI	Oxygen Index = $S_3 \times 100/TOC$	mg CO ₂ /g TOC
S₁/TOC	Normalized Oil Content = $S_1 \times 100/TOC$	
S₂/S₃	Describes type of hydrocarbons generated	Values from 0.00 to >5.00
PI	Production Index = $S_1/(S_1+S_2)$ (or transformation ratio)	Values from 0.00 to 1.00

2.1.1 Organic Richness

Throughout Texas, the Eagle Ford Shale proves to be an excellent potential source rock based on its TOC, HI, and thermal maturity. Table 4 shows average TOC for the outcrop and subsurface rock samples analyzed in this study. In general, the Eagle Ford Shale shows good to excellent source rock potential based on the guidelines of Peters and Cassa (1994). Several EFLC and EFBC samples may have higher “pre-weathering” TOC and HI values, since some of these outcrop samples were affected by weathering, which tends to decrease the amount of organic carbon. In particular, the upper section from the Lozier Canyon outcrop is highly weathered, and when

recalculating TOC for the EFLC samples including only Facies A through C, the average TOC values increase to 2.44% (Figure 14).

Table 4. Average TOC values for the Eagle Ford Shale outcrop and subsurface rock samples analyzed in this study

Well name	# of samples	Average TOC (wt.%)
EFLC	40	1.8
EFMK	26	2.1
EFFA	40	4.0
EFBC	34	3.0
EFAC	10	3.8
EFWB	9	4.6
EFHE	10	4.7
EFLH	22	4.2

Figure 14 shows the TOC geochemical logs for the outcrops and wells analyzed. The logs show a fair variability in terms of organic richness, which may be a result of variations in organic facies. In general, the outcrops and wells analyzed contain greater amounts of organic carbon in the Lower Eagle Ford Shale, particularly in the Lozier Canyon Member. In the case of the EFFA samples, it is possible that the TOC values have been affected by the high maturity of the Eagle Ford Shale in this area.

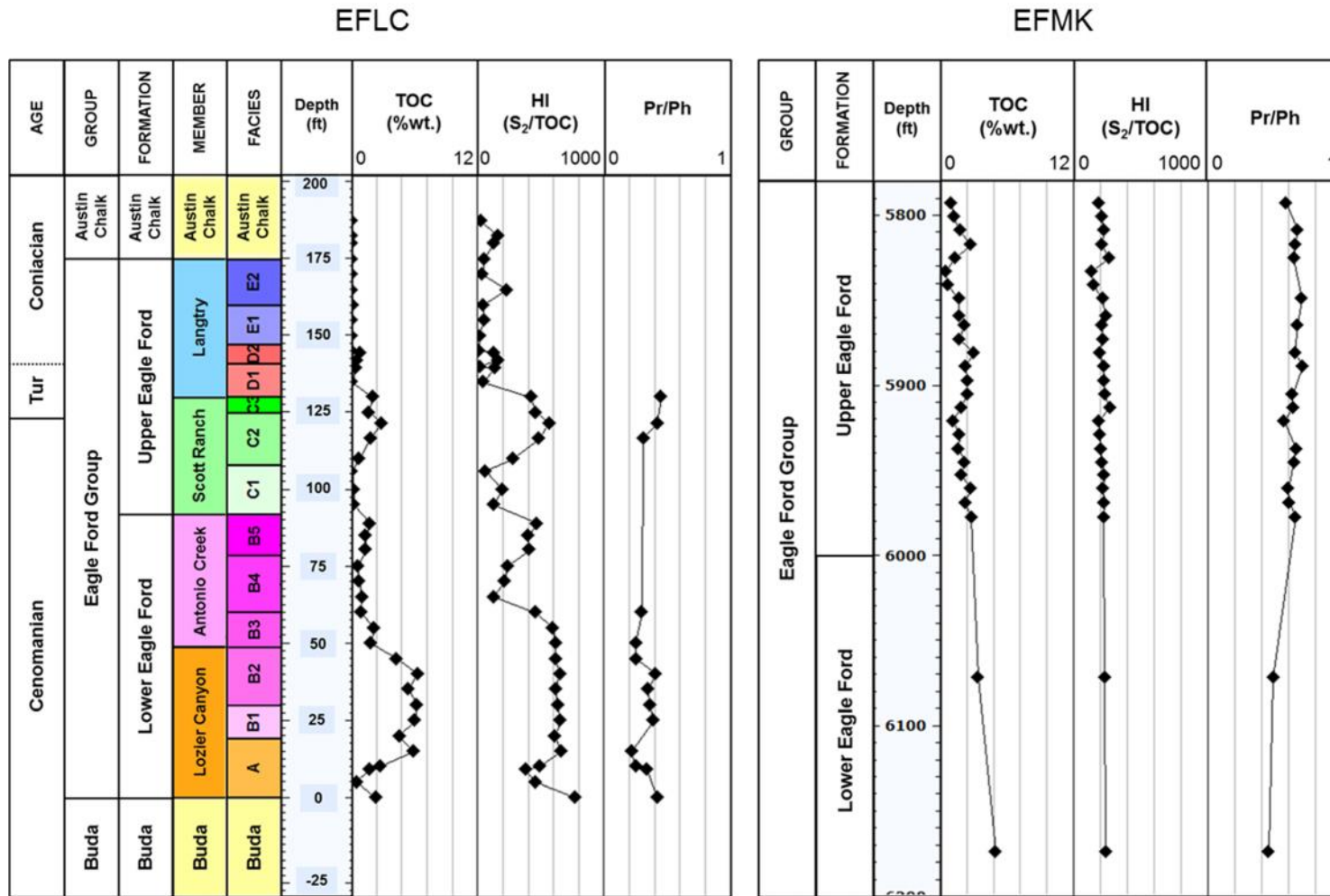


Figure 14. TOC, HI, and Pr/Ph geochemical logs for the Eagle Ford Shale outcrop and subsurface samples analyzed in this study (Pr = pristane; Ph = phytane)

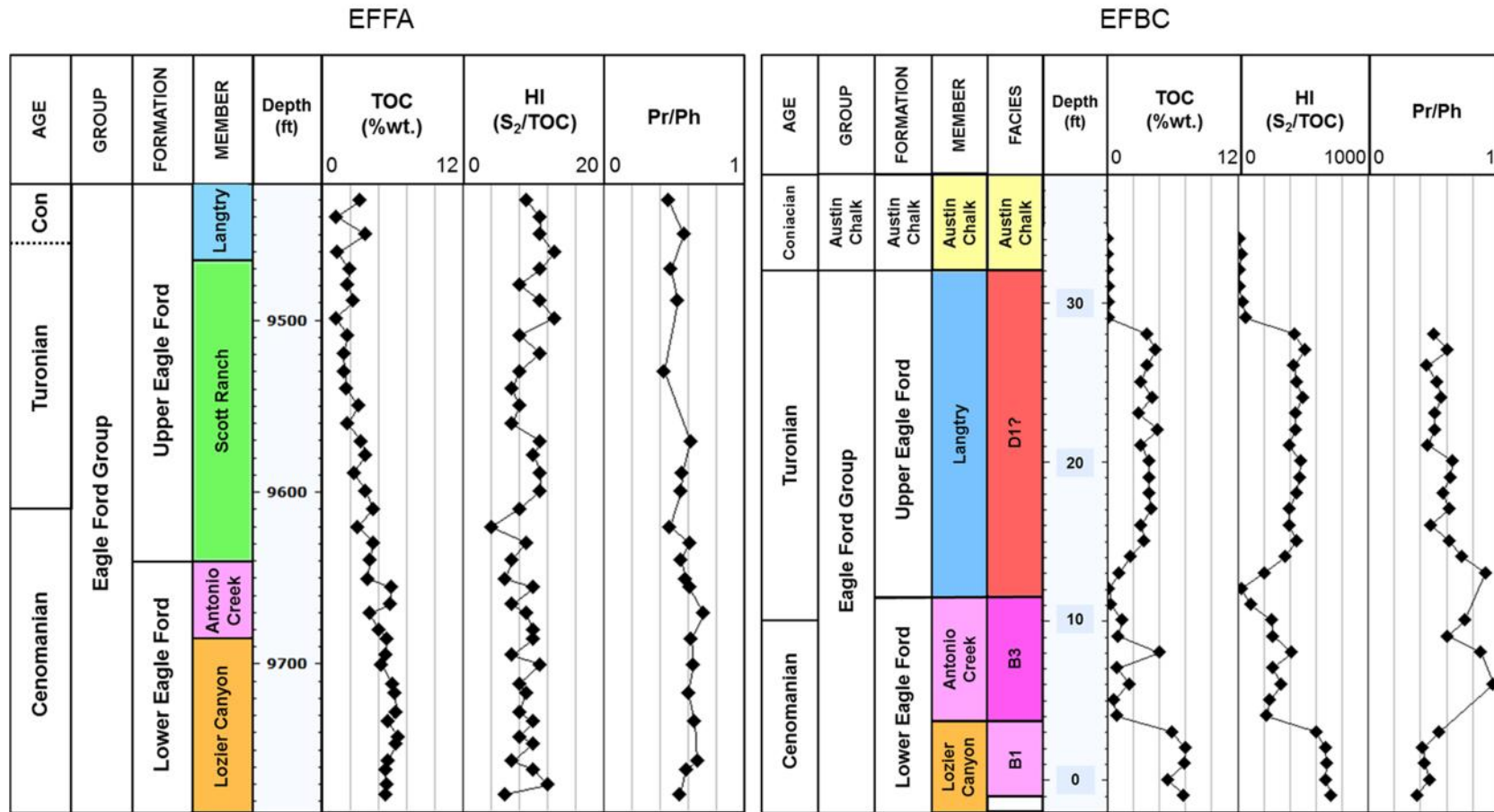


Figure 14. TOC, HI, and Pr/Ph geochemical logs for the Eagle Ford Shale outcrop and subsurface samples analyzed in this study (Pr = pristane; Ph = phytane; cont.)

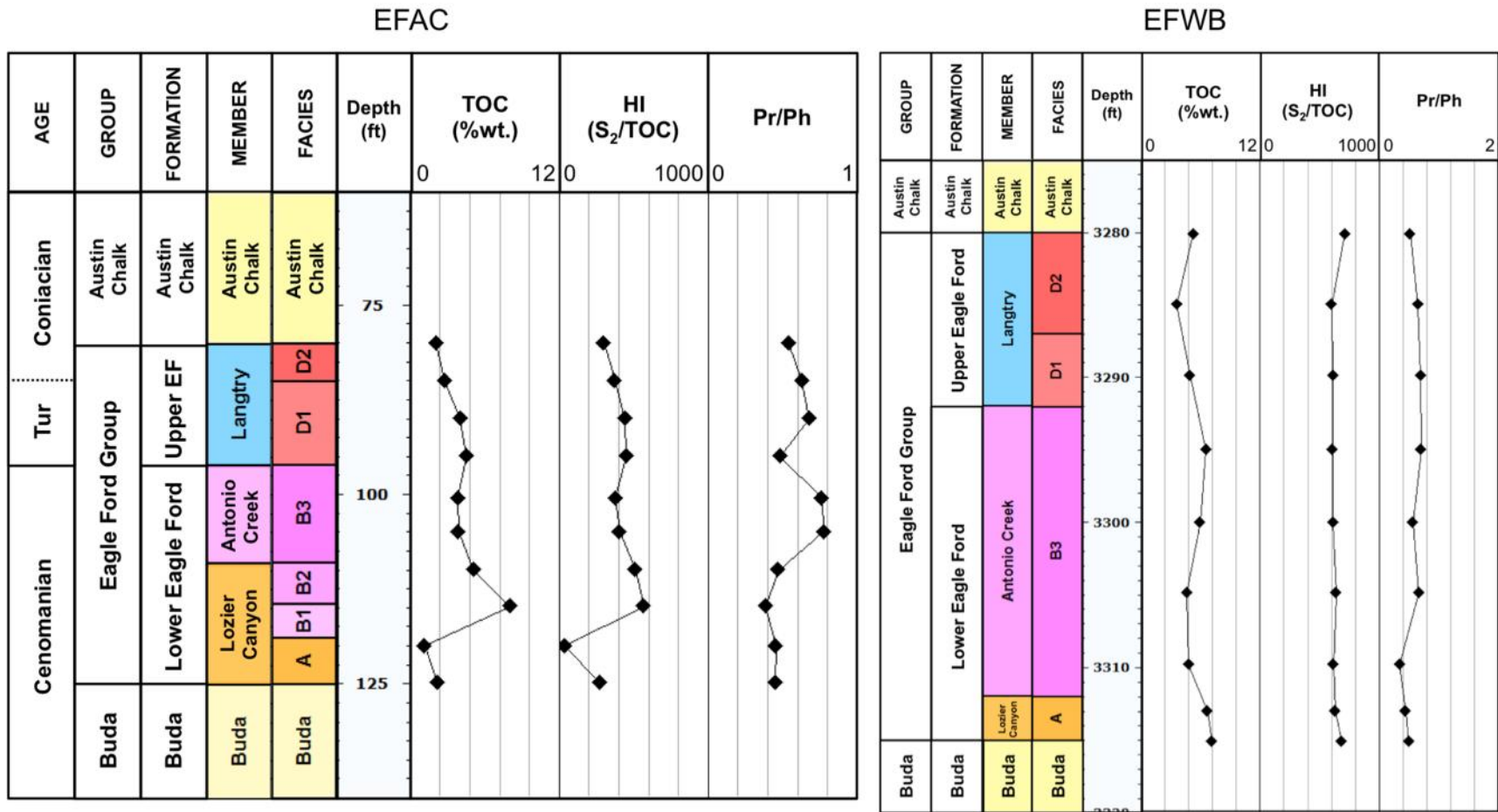


Figure 14. TOC, HI, and Pr/Ph geochemical logs for the Eagle Ford Shale outcrop and subsurface samples analyzed in this study (Pr = pristane; Ph = phytane; cont.)

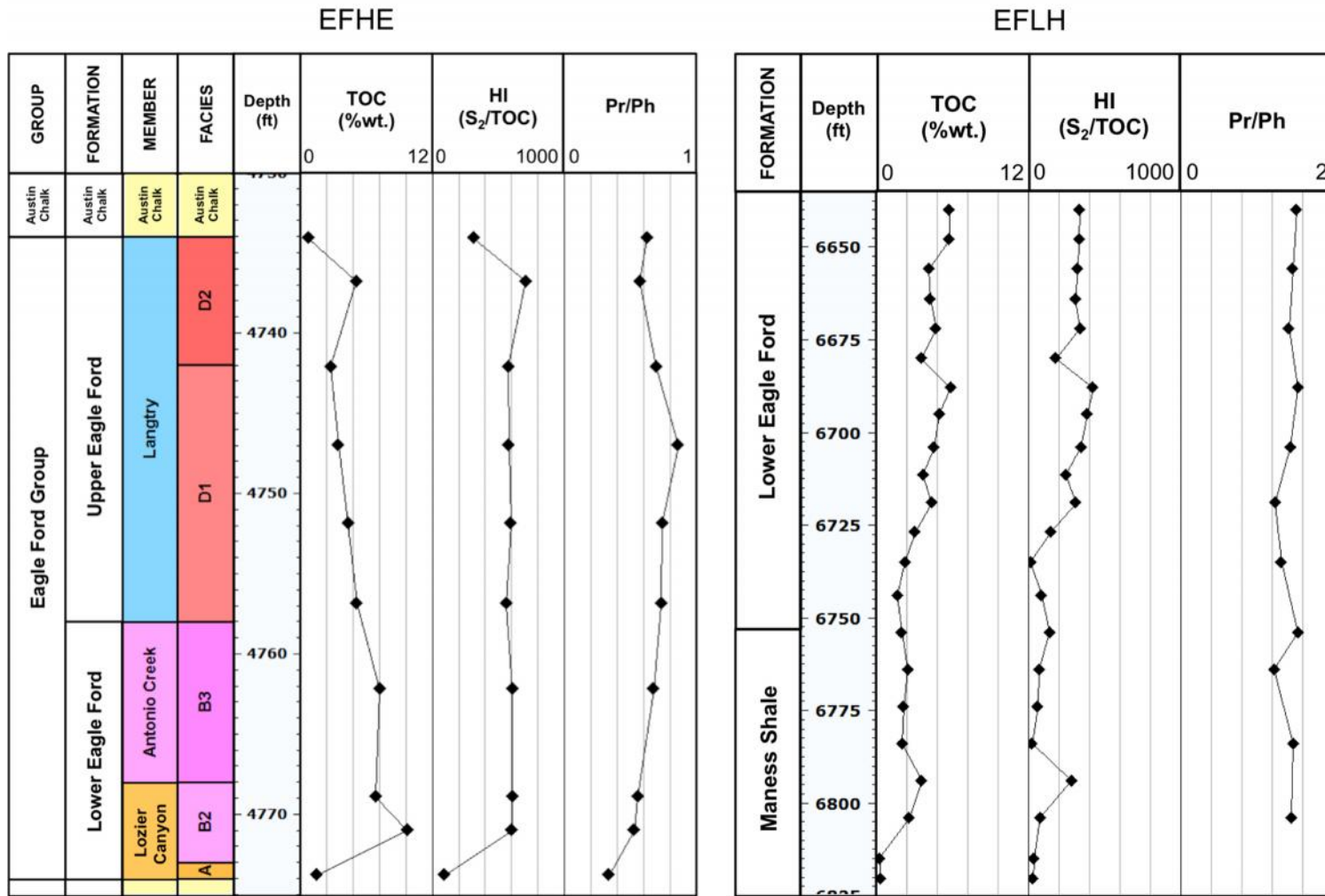


Figure 14. TOC, HI, and Pr/Ph geochemical logs for the Eagle Ford Shale outcrop and subsurface samples analyzed in this study (Pr = pristane; Ph = phytane; cont.)

2.1.2 Organic Matter Type

Rock Eval parameters were used for an initial evaluation of the kerogen type in the Eagle Ford Shale. A plot of Hydrogen Index (HI) versus Oxygen Index (OI), also known as a Van Krevelen-type diagram (Figure 15; Tissot et al., 1974; Tissot and Welte, 1984; Hunt, 1979) shows great variations of HI between the samples. Rock samples from EFAC, EFWB, EFHE, EFBC and some from EFLC show a predominance of Type II kerogen, which is mainly of marine origin. In addition, the high HI values for these samples may be a result of their low maturity. The few EFLC samples plotted towards the Type III field of the Van Krevelen-type diagram correspond to samples from the upper section of the outcrop, which are affected by weathering. In this case, their low HI values may be a result of organic matter alteration and not due to a terrigenous source. Very low HI values from the EFFA samples are the result of its high thermal maturity. Consequently, these samples show a Type III, gas-prone kerogen signature, which in this case is not related to organic matter source. In general, the EFLH samples are dominated by Type II-III kerogen. Interestingly, the samples with HI values lower than 200 correspond to Maness Shale samples. Deposition of Maness Shale sediments in the East Texas region was influenced by the Dexter fluvial system, which distributed siliciclastic sediments sourced from the Ouachita Highlands south into Texas and Louisiana (Adams and Carr, 2010). This sedimentary process may have possibly produced dilution of the organic matter due to a clastic input into the Maness Shale sediments. From the current sample distribution on the Van Krevelen-type diagram, one would be cautious to establish a definite organic matter type. Nevertheless, source rocks have mixes of different types of kerogen due to

variations in environmental conditions throughout the depositional history of the rock, and in some cases, due to their maturation history. Maturity influences the amounts and composition of kerogen and bitumen in the source rock, which in turn affects the HI and OI values calculated to construct this diagram.

Geochemical logs of HI (Figure 14) show a similar trend to that of TOC, indicating that in general, the Eagle Ford Shale has good oil and gas potential, particularly in the interval corresponding to the Lozier Canyon Member.

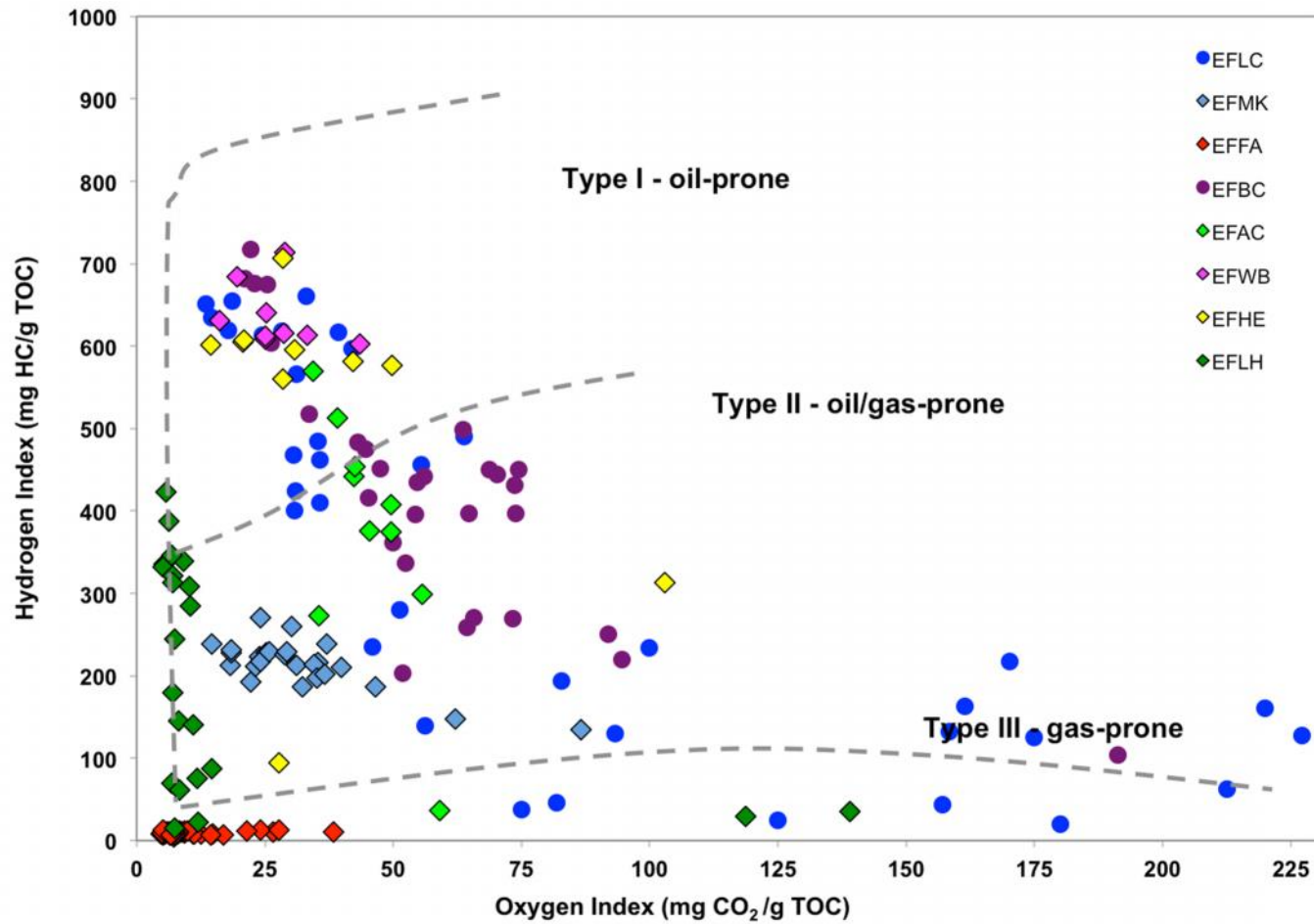


Figure 15. Modified Van Krevelen diagram for the Eagle Ford Shale samples. HI values for EFFA samples are unreliable due to their low remaining hydrocarbon potential

An alternative approach to assess the kerogen quality in potential source rock samples, especially in those rocks with high carbonate content is by using a diagram of S_2 versus TOC (Cornford et al., 1998; Figure 16). This diagram confirms the predominance of Type-II oil-prone kerogen for most of the Eagle Ford Shale samples and shows the low remaining hydrocarbon potential of the EFFA samples as a result of their high maturity. Initially, it was thought that the mix of Type II-III kerogen that characterizes the EFMK samples was related to an organic matter mix or variations in thermal maturity. However, when diagrams of S_1 versus TOC (Figure 17) and T_{max} versus PI (Figure 18) were generated, it was observed that these samples might be contaminated or stained with non-indigenous hydrocarbons or migrated oil. According to Hunt (1979), sidewall cores, cores, and cuttings can be contaminated by drilling mud and handling during the drilling process, affecting the source rock extracts. Therefore, these samples needed to be carefully evaluated on subsequent analyses.

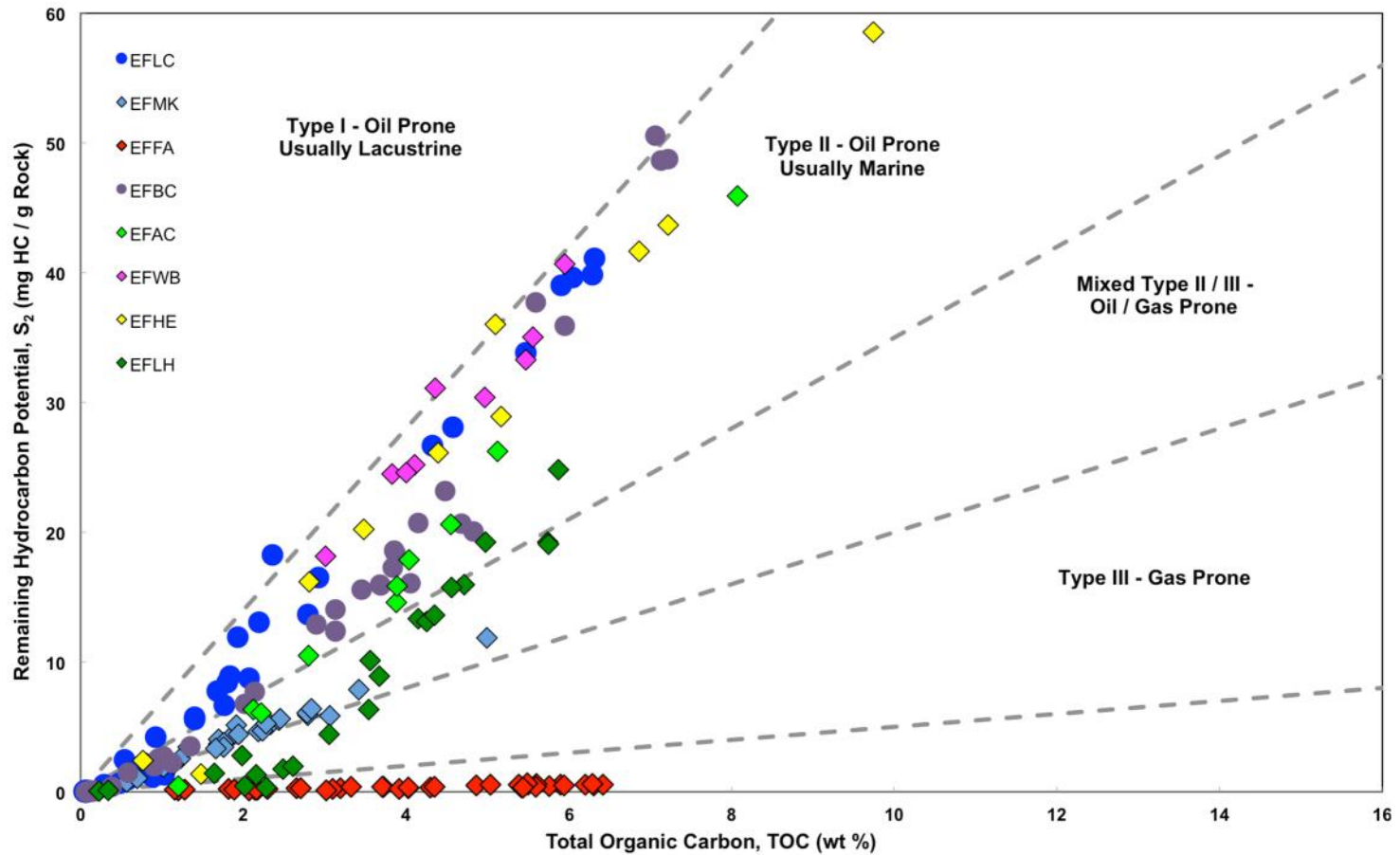


Figure 16. Rock Eval Remaining Hydrocarbon Potential (S_2) vs. TOC plot for determination of kerogen type and maturity of Eagle Ford Shale samples (plot template modified from GeoMark Research Ltd.)

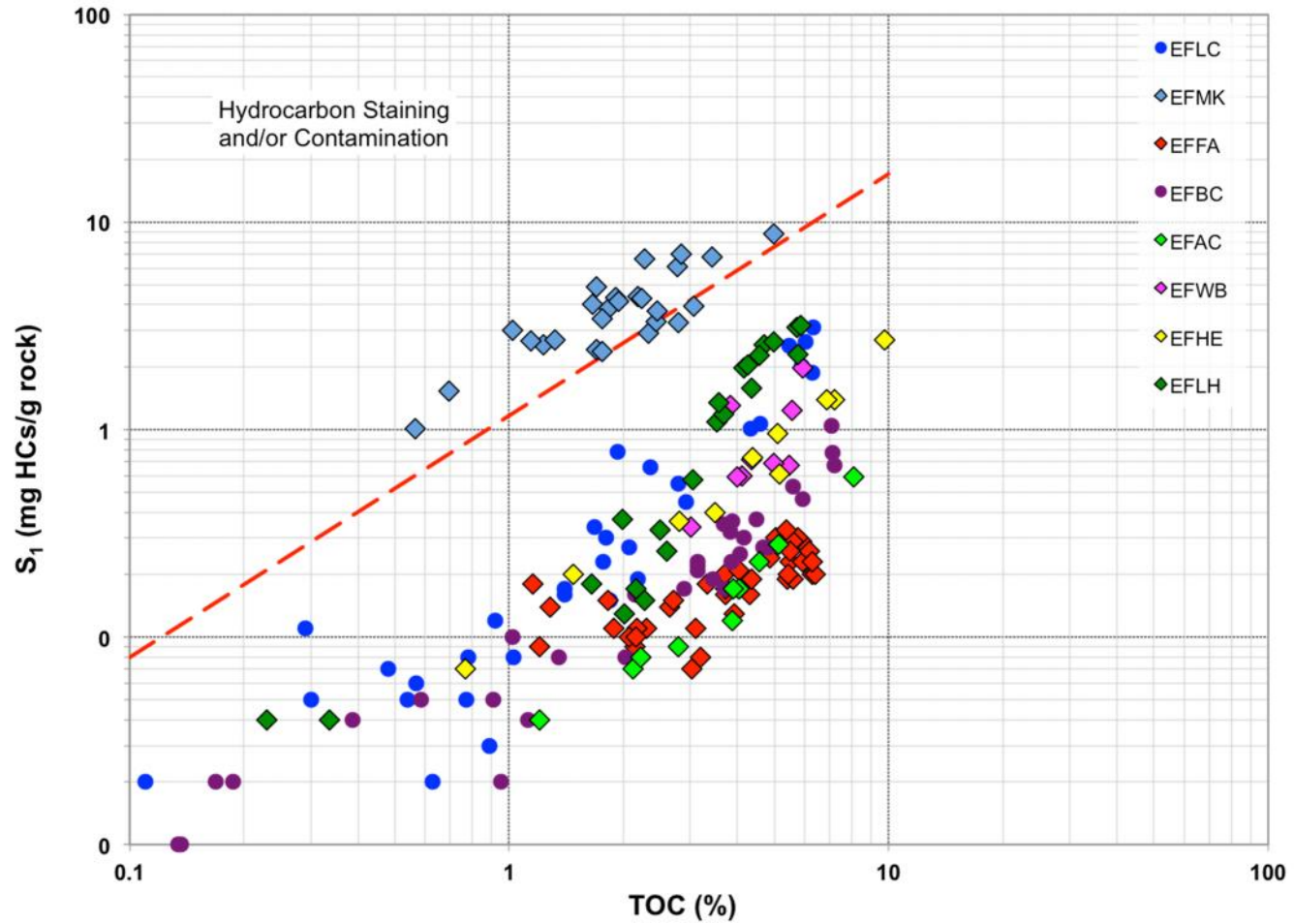


Figure 17. S_1 vs. TOC plot for the Eagle Ford Shale samples showing possible hydrocarbon contamination of the EFMK samples (plot template modified from GeoMark Research Ltd.)

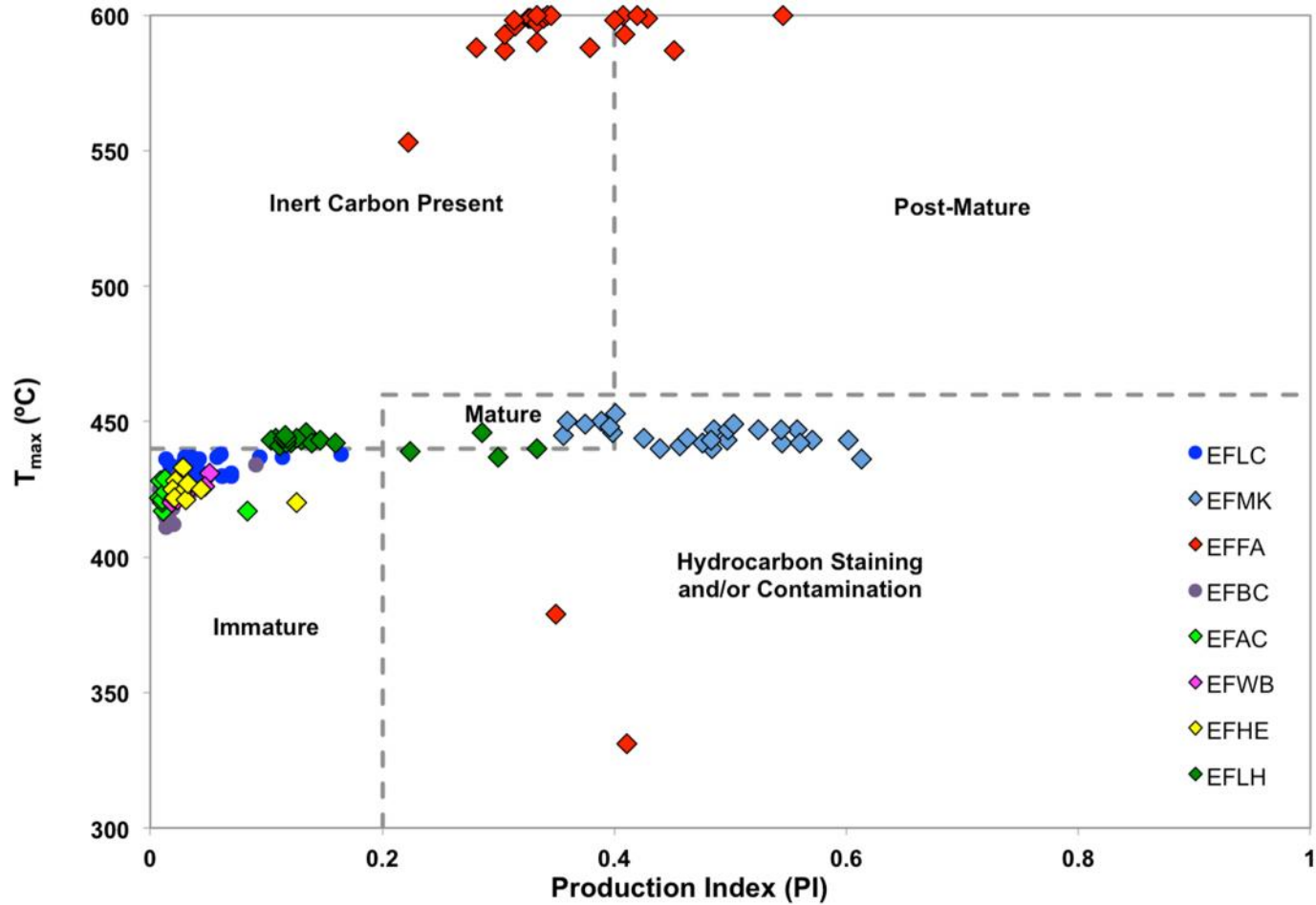


Figure 18. T_{max} vs. Production Index (PI) plot for determination of thermal maturity and sample contamination of Eagle Ford Shale samples (plot template modified from GeoMark Research Ltd.)

2.1.3 Thermal Maturity from Rock Eval and Vitrinite Reflectance Analysis

Based on Rock Eval T_{\max} values (Figures 18 and 19), most of the Eagle Ford Shale samples analyzed from East and West Texas are thermally immature to marginally mature (early oil window; $T_{\max} < 445^{\circ}\text{C}$). The EFFA samples have very low S_2 values (Appendix B) indicating the low remaining hydrocarbon potential for these rocks. As a result, HI and T_{\max} parameters, which are calculated from S_2 are unreliable for these samples. The T_{\max} versus PI plot (Figure 18) shows that the EFFA samples may contain dead or inert carbon and some samples from the EFMK core may be contaminated with hydrocarbons as previously mentioned.

Vitrinite reflectance ($\%R_o$) measurements were attempted on four Eagle Ford Shale samples from four different locations: EFLH-9, EFLC-10, EFBC-8, and EFFA-3. Overall, vitrinite identification of these samples was challenging. In the Eagle Ford Shale samples selected for R_o analysis the vitrinite is scarce. This fact was rather expected since vitrinite is mainly derived from land-plant material (ASTM, 2011). Because the Eagle Ford Shale was deposited in a carbonate environment there is very low (if any) terrigenous organic matter contribution. Most of the vitrinite fragments identified were isolated and pitted. Isolated grains are generally unreliable for measuring vitrinite reflectance, since these may be unrelated to the sample and correspond to contamination acquired during sample preparation. Pitted vitrinite usually shows lower reflectance values. In addition, some vitrinite clasts were smaller than the measuring spot of the microscope used for the analyses, which also lead to unreliable measurements. In the East Texas EFLH-9 sample, ten (10) vitrinite clasts were identified and measured, ranging from 0.91 to 1.22 $\%R_o$ with a mean random R_o

measurement of 1.04%, indicating that in this area the Eagle Ford Shale is in the oil window. Figure 20 shows a photomicrograph of a vitrinite clast (1.06% R_o), bitumen (0.62% R_o), and lignite contamination (0.29% R_o) from sample EFLH-9 (Brian Cardott, 2013, personal communication). This figure illustrates the variability of macerals and bitumen content observed in the Eagle Ford Shale samples that could potentially lead to errors in the identification and measurement of vitrinite reflectance. The amount of vitrinite observed in this sample compared to the Central and West Texas samples might indicate an input of terrigenous organic matter for the East Texas area. Rock pellets EFLC-10 and EFBC-8 corresponding to outcrop samples were too weathered for obtaining a statistically representative sample, and sample EFFA-3 was too mature to observe any type of maceral. However, Liro et al. (1994) were able to measure vitrinite reflectance in samples from the Bouldin Creek outcrop (samples EFBC in this study), obtaining values of 0.45% R_o , which indicates that in this area the Eagle Ford Shale is thermally immature. Slatt et al. (2012) characterized samples from the Comstock West outcrop (US Highway 90, Val Verde County, Texas), and determined that in this location the Eagle Ford Shale is immature, with T_{max} values between 423 and 429°C and a mean random vitrinite reflectance value of 0.53%. The interval analyzed in their study corresponds to Facies B of Donovan and Staerker (2010) and Donovan et al. (2012; 2013a and b) at the Lozier Canyon site. Since the Lozier Canyon outcrop is located about 40 miles northwest from the Comstock West outcrop, it is inferred that the thermal maturity at Lozier Canyon would be similar. Edman (2012) analyzed the thermal maturity of several Eagle Ford Shale rocks, bitumens, and oils from the First Shot oil field (Wilson, Gonzales, and Karnes Counties, Texas) using a combination of

vitrite reflectance measurements and various maturity geochemical parameters. The author determined that the thermal maturity of the Eagle Ford Shale at the oil field location ranged from early to late oil window (0.50 to 1.21% R_o ; Edman, 2012). The First Shot oil field is located about 20 miles south from the EFWB and EFHE wells analyzed in this study; hence, the maturity in the Austin area should be similar to that observed by Edman (2012).

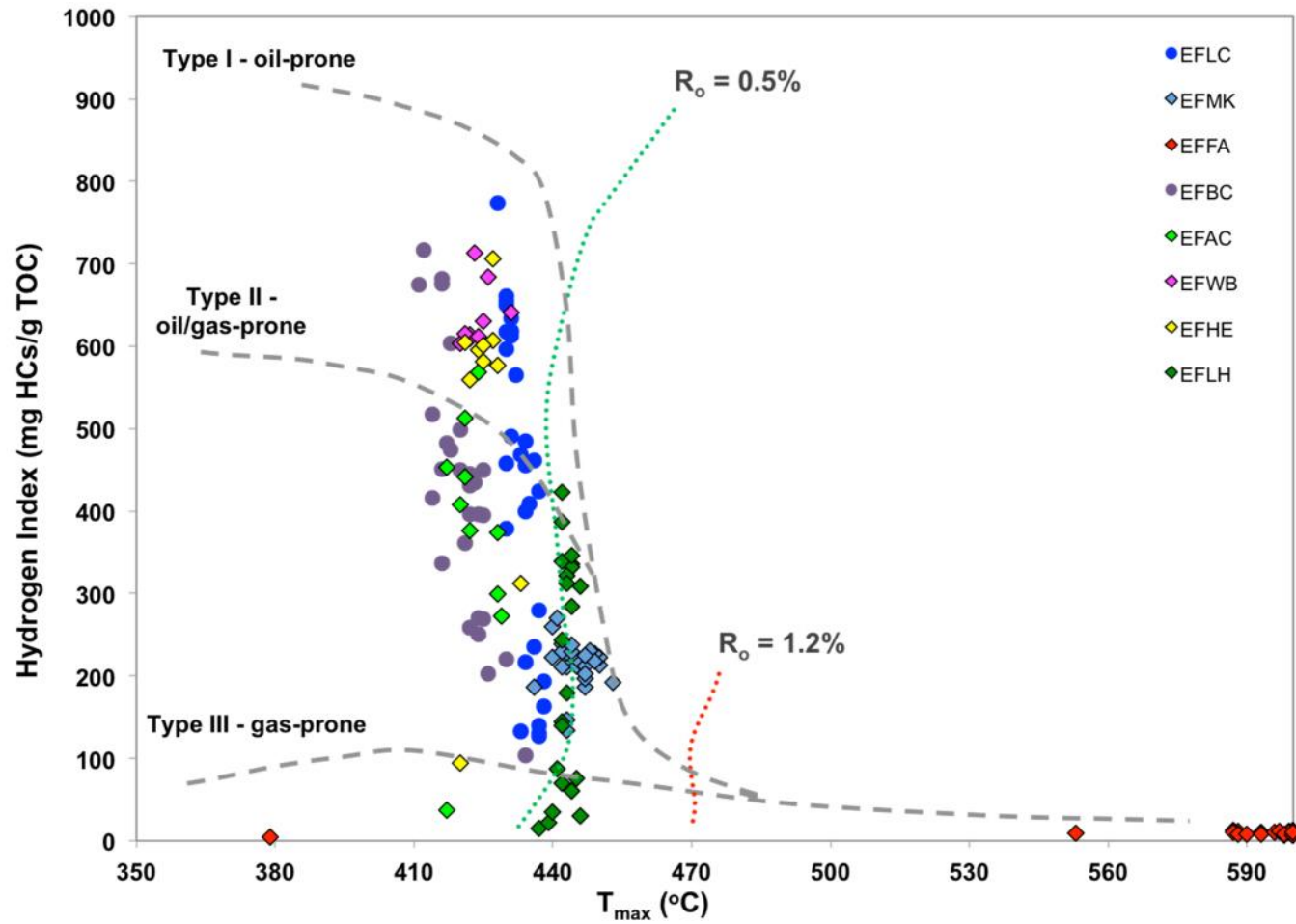


Figure 19. T_{max} vs. HI plot showing maturity and kerogen type of the Eagle Ford Shale samples. HI values for EFFA samples are unreliable due to their low remaining hydrocarbon potential

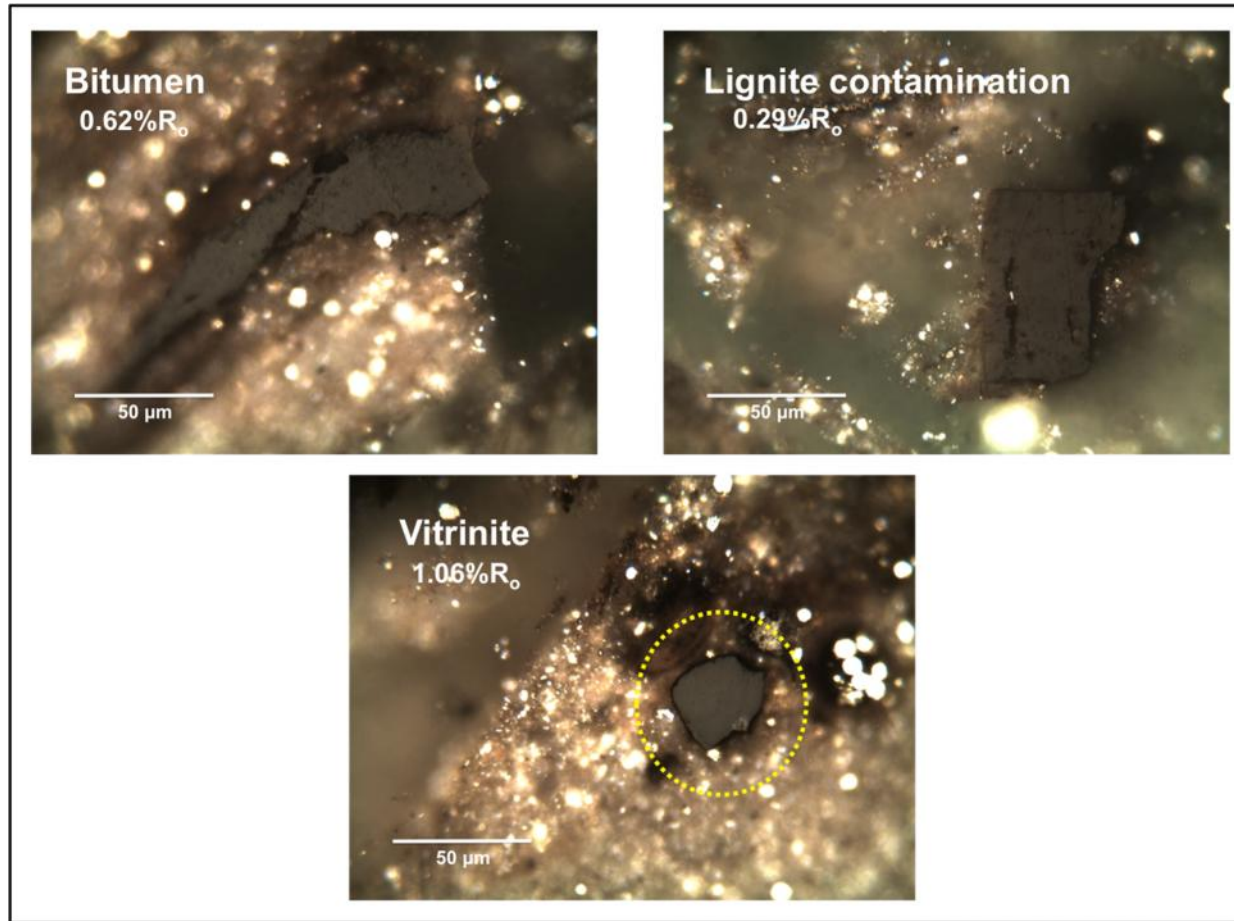


Figure 20. Photomicrographs from the EFLH-9 Eagle Ford Shale sample (Photomicrographs courtesy of Brian Cardott, 2013)

2.1.4 Gas Chromatography

2.1.4.1 *n*-Alkanes distribution

Gas chromatograms from the saturate fraction of bitumen extracted from Eagle Ford Shale samples show minimal compositional variations between the different members (Figure 21). Detailed results of these analyses including formulas, selected chromatograms, and ratios calculated are presented in Appendices A, C, and D respectively. Table 5 summarizes some of the geochemical parameters calculated from the *n*-alkanes fingerprints for each sample set. The terrigenous to aquatic ratio (TAR), is a parameter calculated to evaluate the type of organic matter input that compares the high-odd-carbon-numbered *n*-alkanes (terrigenous organic matter) with the low-odd-carbon-numbered *n*-alkanes (marine organic matter; Peters et al., 2005). The carbon preference index (CPI; Bray and Evans, 1961) compares the odd-carbon-numbered *n*-alkanes ($n\text{-C}_{25}\text{-}n\text{-C}_{33}$) against the even-carbon-numbered *n*-alkanes ($n\text{-C}_{24}\text{-}n\text{-C}_{34}$). In general, *n*-alkanes originated from marine organic matter do not show carbon-number preference, whereas *n*-alkanes derived from terrigenous organic matter tend to show odd-carbon-number predominance. Low maturity hypersaline and carbonate source rocks usually show low CPI values due to a predominance of even-numbered *n*-alkanes (Peters et al., 2005). In some cases, CPI can be used to roughly assess thermal maturity. According to Peters et al. (2005), oils and bitumens with CPI values significantly above 1.0 (odd predominance) are indicative of low thermal maturity. The TAR and CPI ratios are therefore influenced by source and thermal maturity (Peters et al., 2005), and additional geochemical parameters need to be correlated with these ratios when characterizing oils and source rocks.

In case of the Eagle Ford Shale samples analyzed, the majority shows a unimodal distribution towards the low-carbon-number members, and a slight preference of even-carbon-numbered *n*-alkanes (Figure 21). The bitumen samples show a maximum around *n*-C₁₇, whereas the oil and condensate samples have *n*-alkanes maxima around *n*-C₁₄ and *n*-C₁₅. The discrepancy between *n*-alkanes maxima between extracts and oils is mainly due to the loss of low carbon number *n*-alkanes during source rock extraction. Based on their *n*-alkane distributions and low TAR and CPI ratios (~0.2 and ~1.0 respectively), it is inferred that these samples are genetically related. These observations point towards a marine origin and the existence of reducing (anoxic) conditions during source rock deposition. In some samples, peak identification beyond *n*-C₂₆ was problematic due to the low intensity of the *n*-alkanes and coelution with steranes and hopanes. This case was particularly true for EFBC, EFAC, EFWB, EFHE, and EFFA samples, for which the TAR ratio could not be calculated. The *n*-alkanes distribution for the EFFA samples is also affected by its high thermal maturity, as indicated by its high CPI values (>1.2). Relatively high CPI and TAR values (>0.25 and >1.0 respectively) for the EFLH samples could be a result of high thermal maturity, although it may also suggest a contribution of terrigenous organic matter. A number of chromatograms from EFLC and EFBC show low *n*-alkanes intensities, which could be related to weathering. According to Hunt (1979), outcrops are commonly exposed to complex weathering processes that significantly decrease the amount of hydrocarbon and organic carbon content in the rocks. Therefore, the chromatographic signature can show similar characteristics to those of biodegraded oil (Philp et al., 1992). The degree of weathering that may affect outcrops is dependent on its porosity, permeability,

fracturing, microbiological activity, and climate (Hunt, 1979). However, Hunt (1979) also points out that the presence of pyrite may indicate low weathering degree, and in turn, a small decrease in the hydrocarbon and organic carbon content. Both, the Lozier Canyon and Bouldin Creek outcrops contain dispersed and framboidal pyrite (Aris Pramudito 2014, personal communication). Consequently, it is considered here that the data derived from these samples can be used with confidence.

Table 5. Summary table of *n*-alkanes analyses from the saturate fraction of Eagle Ford Shale bitumens and oils (ND = not determined)

Samples	Range	n-alkane distribution	n-alkane maximum	Pr/Ph range	TAR range	CPI range
EFLC	<i>n</i> -C ₁₄ - <i>n</i> -C ₂₉	unimodal	<i>n</i> -C ₂₂	0.22-0.45	0.20-3.70	0.41-1.48
EFMK	<i>n</i> -C ₁₃ - <i>n</i> -C ₃₆	unimodal	<i>n</i> -C ₁₆	0.91-1.42	0.10-0.18	0.95-1.01
EFFA	<i>n</i> -C ₁₃ - <i>n</i> -C ₂₇	unimodal	<i>n</i> -C ₁₇	0.86-1.42	ND	1.21-1.69
EFBC	<i>n</i> -C ₁₅ - <i>n</i> -C ₂₅	unimodal	<i>n</i> -C ₁₇	0.75-1.92	ND	0.56-0.61
EFAC	<i>n</i> -C ₁₄ - <i>n</i> -C ₂₆	bimodal	<i>n</i> -C ₁₇ and <i>n</i> -C ₂₂	0.77-1.55	ND	0.80-1.29
EFWB	<i>n</i> -C ₁₄ - <i>n</i> -C ₂₆	bimodal	<i>n</i> -C ₁₇ and <i>n</i> -C ₂₂	0.35-0.71	ND	0.61-0.83
EFHE	<i>n</i> -C ₁₄ - <i>n</i> -C ₂₆	unimodal	<i>n</i> -C ₁₇	0.67-1.72	ND	0.49-0.67
EFLH	<i>n</i> -C ₁₃ - <i>n</i> -C ₃₇	unimodal	<i>n</i> -C ₁₇	1.23-1.54	0.25-0.58	1.05-1.13
NWMK-533.2	<i>n</i> -C ₁₁ - <i>n</i> -C ₃₅	unimodal	<i>n</i> -C ₁₅	1.07	0.23	0.99
NFMK-526-1H	<i>n</i> -C ₁₁ - <i>n</i> -C ₃₅	unimodal	<i>n</i> -C ₁₅	0.80	0.21	0.98
MJOG-UEF	<i>n</i> -C ₁₁ - <i>n</i> -C ₃₆	unimodal	<i>n</i> -C ₁₄	0.80	0.23	0.97
MJOG-LEF	<i>n</i> -C ₁₁ - <i>n</i> -C ₃₆	unimodal	<i>n</i> -C ₁₄	1.14	0.21	1.00
GREF-1-110	<i>n</i> -C ₁₁ - <i>n</i> -C ₃₆	unimodal	<i>n</i> -C ₁₃	1.25	0.18	1.04

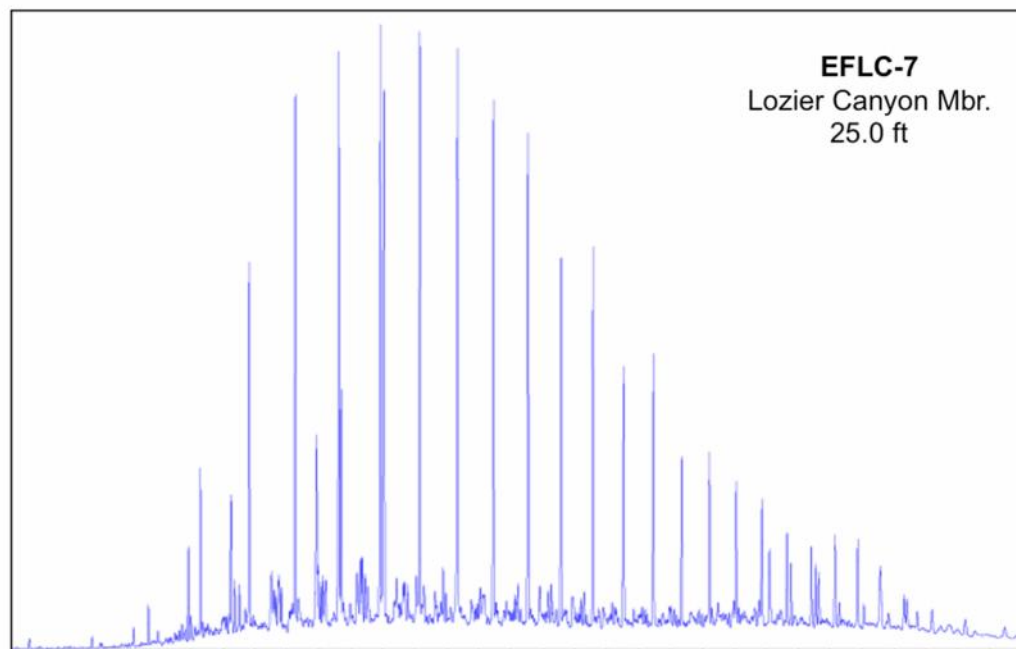
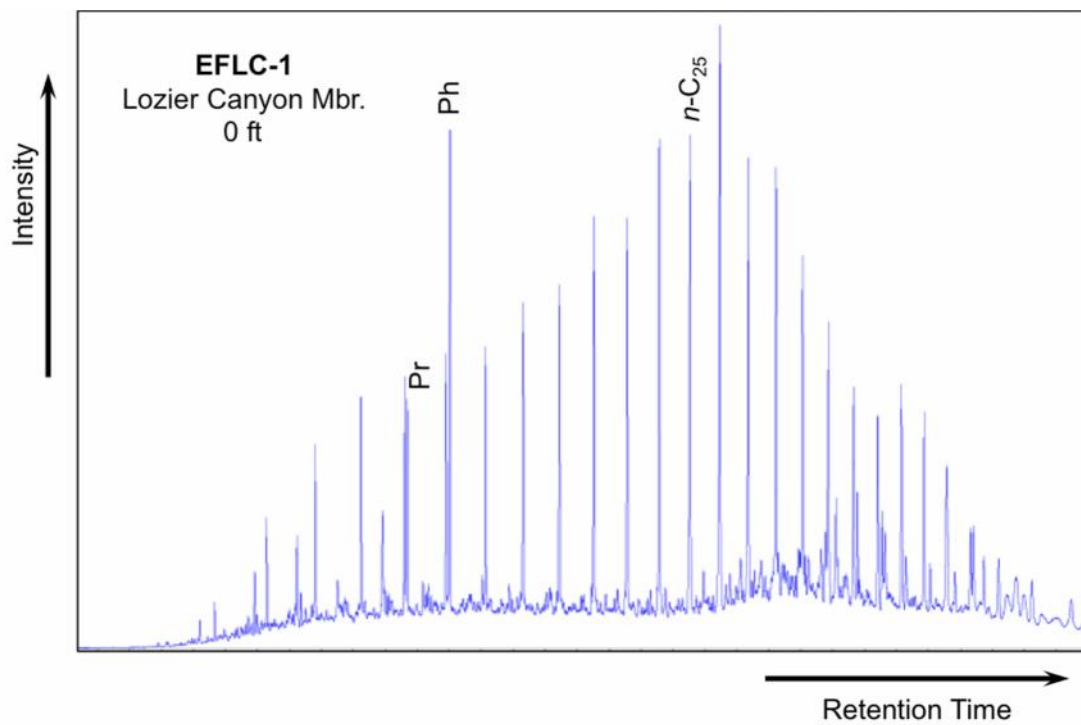


Figure 21. Gas chromatograms of the saturate fractions from bitumen extracts of the EFLC outcrop (Pr = pristane; Ph = phytane, $n-C_{25}$ = C_{25} normal alkane)

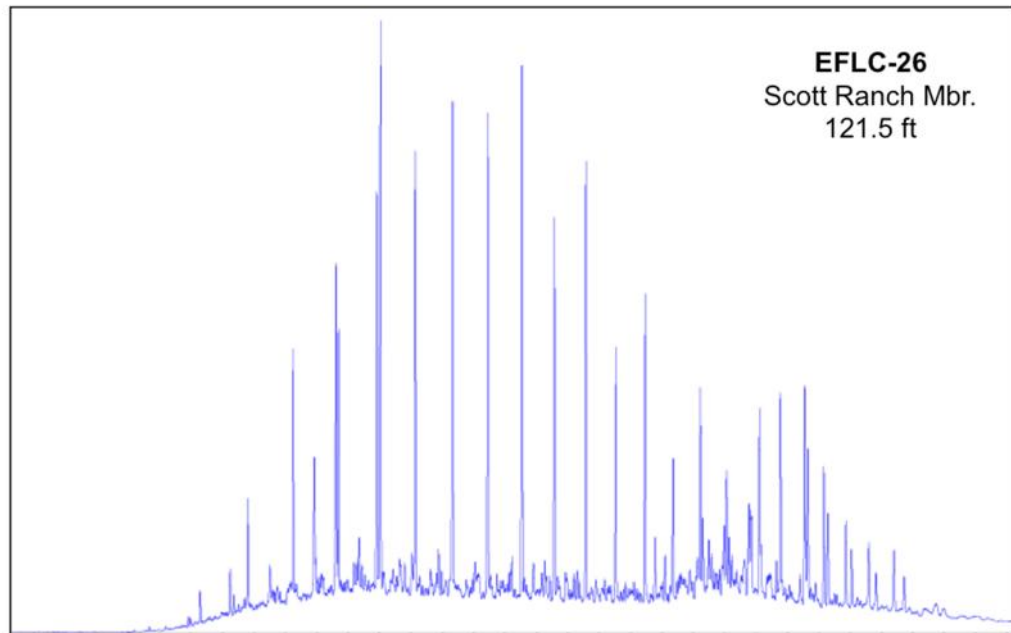
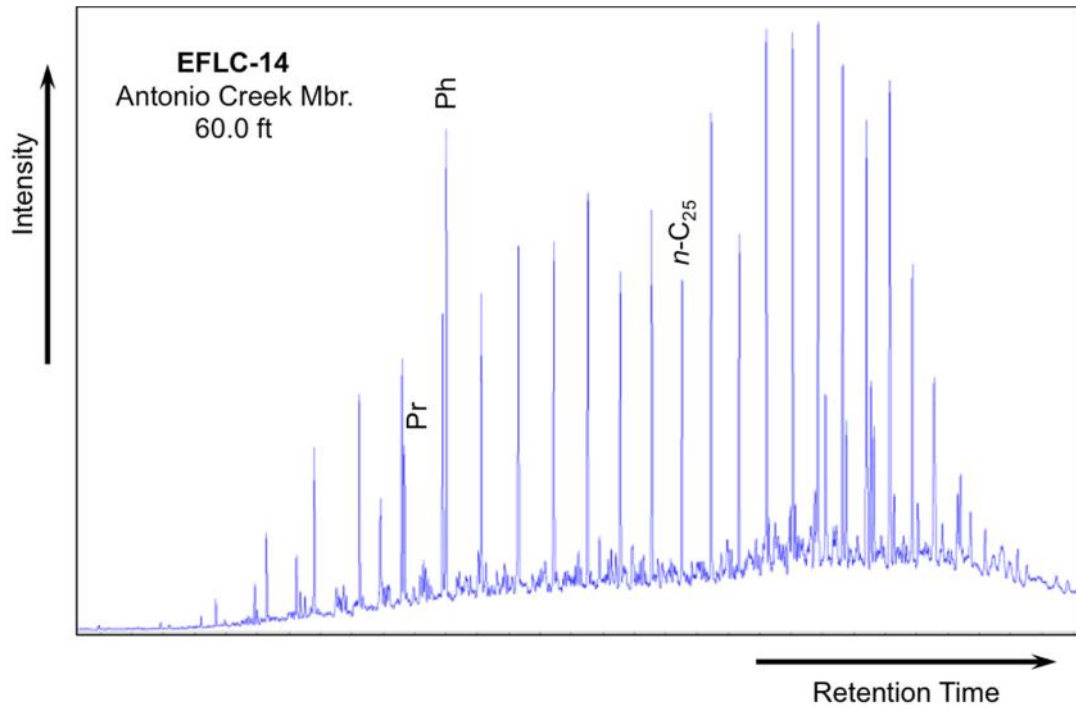


Figure 21. Gas chromatograms of the saturate fractions from bitumen extracts of the EFLC outcrop (Pr = pristane; Ph = phytane, *n*-C₂₅ = C₂₅ normal alkane; cont.)

2.1.4.2 *Pristane and Phytane*

Pristane (Pr) and phytane (Ph) were determined in the *n*-alkanes fingerprints from the saturate fractions of the bitumens and oils (Figure 21, Appendix D). These acyclic isoprenoids have been widely used for evaluating redox conditions during source rock deposition, since their origin is dependent on oxygen availability (Didyk et al., 1978; Waples, 1985; Peters et al., 2005). Pr and Ph are primarily derived from the phytyl side chain of the chlorophyll-a during diagenesis. However, other sources, such as unsaturated isoprenoids in zooplankton and higher animals, tocopherols, and archaea, have been proposed as precursors of Pr and Ph (Blumer et al., 1963; Blumer and Thomas, 1965; Goosens et al., 1984; Rowland, 1990; Bechtel et al., 2007), and Pr can also be derived from isoprenoids bound to kerogen during catagenesis (Larter et al., 1979).

Several chemical pathways generate Pr and Ph from the phytyl chain of chlorophyll. Under reducing conditions, phytol undergoes hydrogenation and reduction to generate phytane. Conversely, under oxic conditions, oxidation and decarboxylation of the phytol chain produces pristane (Tissot and Welte, 1984; Peters et al., 2005). Consequently, the Pr/Ph ratio assists in evaluating the redox potential of the depositional environment of the source rock (Didyk et al., 1978; Shanmugam, 1985; Peters et al., 2005). Pr/Ph ratios lower than 1 are associated with anoxic conditions, whereas Pr/Ph greater than 1 suggest oxic conditions during source rock deposition. In addition, for rocks and oils within the oil window Peters et al. (2005) pointed out that Pr/Ph values greater than 3 are related to terrigenous organic matter input under oxic-

suboxic conditions, while Pr/Ph values lower than 0.8 indicate anoxic, hypersaline or carbonate depositional environments (ten Haven et al., 1987; Peters et al., 2005).

Overall, the majority of the bitumen and oil samples show low Pr/Ph ratios indicating the presence of reducing conditions (anoxia) during source rock deposition (Table 5; Appendix E). Geochemical logs of Pr/Ph ratios for outcrop and core samples show variations in redox potential that inversely correlates with TOC and HI values (Figure 14). In general, Pr/Ph ratios of samples from EFLC, EFBC, EFAC, EFWB, EFHE, EFMK, and EFFA, suggest that the Lozier Canyon Member experienced stronger reducing conditions compared to the upper members of the Eagle Ford Group. This may have contributed to organic matter preservation, which resulted in its high TOC values. Additionally, a progressive increase in Pr/Ph ratios towards the upper members is observed, suggesting an increase in oxygen content in the water column during deposition of these units. Similarly, Pr/Ph ratios for the EFMK core suggest occurrence of stronger reducing conditions during deposition of the Lower Eagle Ford (Figure 14). On the other hand, the Lower Eagle Ford and underlying Maness Shale analyzed in the East Texas EFLH core do not indicate marked changes in redox conditions during deposition.

Another set of parameters useful for evaluating variations in redox conditions, organic matter source, maturity, and alteration in source rock extracts and oils involves the relationship between Pr and Ph and *n*-alkanes C₁₇, and *n*-C₁₈ (Shanmugam, 1985). Figure 22 shows a crossplot of Pr/*n*-C₁₇ versus Ph/*n*-C₁₈ for the bitumens and oils analyzed in this study. This diagram indicates that these samples are mainly composed of marine organic matter. However, few samples corresponding to EFLH, EFHE, and

EFBC also plot in the mixed organic matter field of the diagram, suggesting a marine and terrigenous organic matter input. Liro et al. (1994) found a similar result between Pr/*n*-C₁₇ and Ph/*n*-C₁₈ for samples in the upper section of the Bouldin Creek outcrop (EFBC samples in the present study). These authors attributed the variations in organic matter source to a change to more proximal environmental conditions during deposition of the younger section of the Eagle Ford Shale at this locality. The EFBC samples plotted in the mixed organic matter field of the diagram (Figure 22) correspond to samples from the Langtry Member, suggesting a similar interpretation to that of Liro et al. (1994) for the Upper Eagle Ford Shale at the Austin area. Rock samples from EFFA, EFMK, EFLH, and oils show the highest maturity level according to the lower proportion of isoprenoids relative to *n*-alkanes.

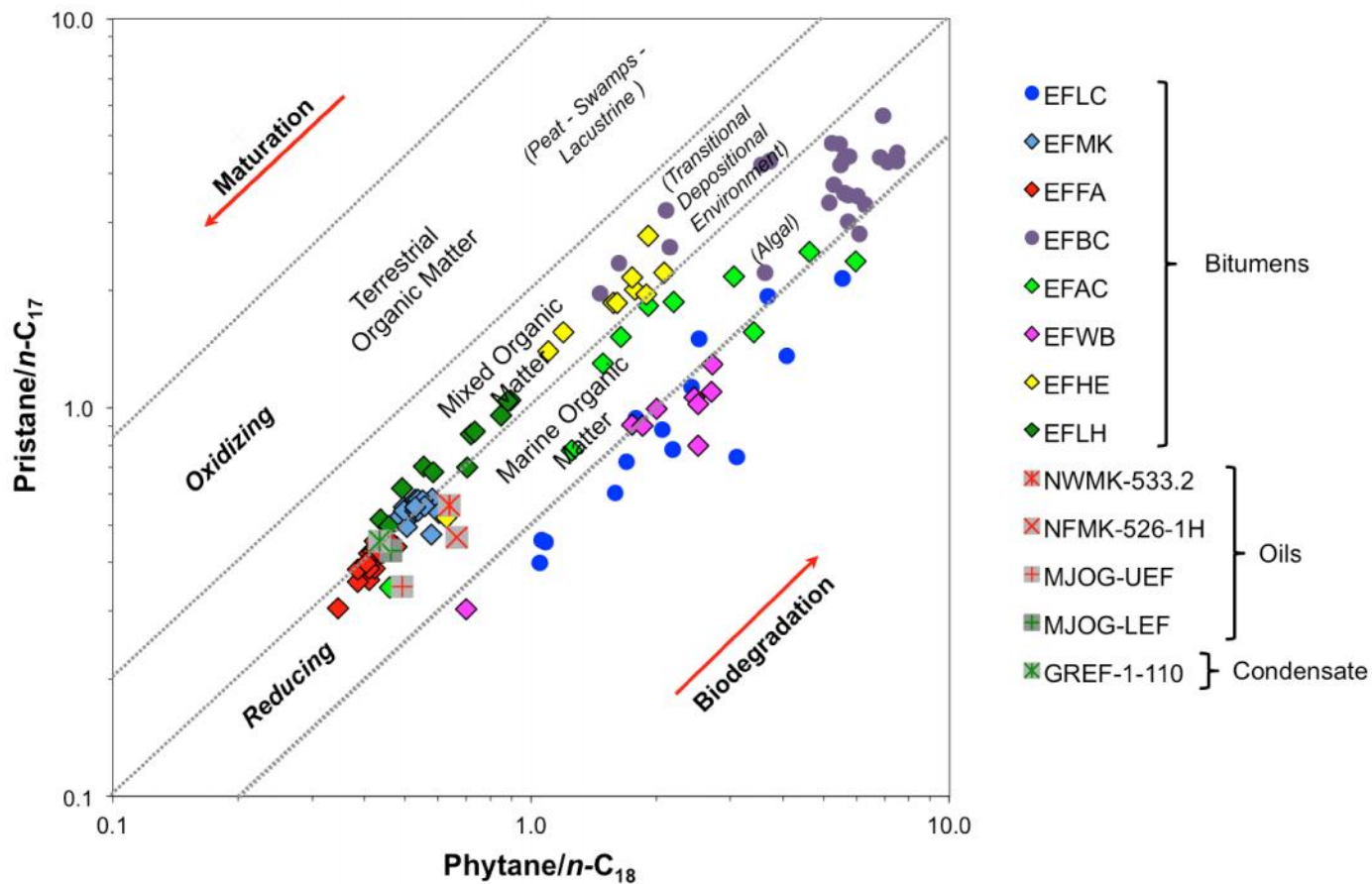


Figure 22. Isoprenoids plot of Pristane/ n -C₁₇ versus Phytane/ n -C₁₈ showing redox conditions, maturity, and depositional environments for samples of the Eagle Ford Shale (n -C₁₇ = C₁₇ normal alkane; n -C₁₈ = C₁₈ normal alkane)

2.2 Biomarker and geochemical parameters for evaluation of organic matter source

2.2.1 Steranes

This group of biomarkers is commonly used for evaluation of organic matter source, maturity, and correlation studies. Steroids are derived from sterols, which are compounds present in eukaryotic organisms, mainly algae and higher plants (Tissot and Welte, 1984; Peters et al., 2005). Distributions of steranes, diasteranes, and pregnanes were determined in the B&C fractions of bitumens and oils by SIM/GC-MS through analysis of the m/z 217.3 ion. Fragmentograms of steranes distributions for the EFLC samples are presented in Figure 23. Peak identification is presented in Table 6. Formulas for calculation of geochemical ratios are in Appendix A. Numerical values of the geochemical ratios calculated are in Appendix H. Biomarkers quantitation results are in Appendix L.

2.2.1.1 Regular steranes

Most of the Eagle Ford Shale analyzed samples show a higher proportion of C₂₇ steranes compared to that of the C₂₈ and C₂₉ homologs, suggesting a marine organic matter source for these samples. The distribution of regular steranes for samples EFLC, EFMK, and EFLH also shows a higher proportion of the C₂₇, C₂₈, and C₂₉ isomers compared to their counterparts (Figure 23). This feature has been attributed to hypersaline environments (ten Haven et al., 1988), suggesting that deposition of the Eagle Ford Shale in West and East Texas was restricted enough to allow development of water-density stratification and hypersaline conditions.

Samples EFBC and EFAC feature a steranes distribution typical of immature source rocks (Figure 23). These samples are characterized by a predominance of the 20R steranes (biological configuration) and higher amounts of the C₂₈ relative to the C₂₉ steranes, which are consistent with Waples and Machihara (1991) descriptions of steranes distribution of low maturity oils and extracts.

A ternary diagram of the distribution of C₂₇, C₂₈ and C₂₉ regular steranes (Figure 24) was used to assess possible variations in depositional environment of the Eagle Ford Shale samples following a similar approach to that of Huang and Meinschein (1979) and Moldowan et al. (1985). C₂₇ steranes (cholestane) originate from precursors found in plankton and marine invertebrates. C₂₈ steranes (ergostane) are also derived from similar precursors, although they can also be generated from terrigenous organisms. (Huang and Meinschein, 1979; Moldowan et al., 1985). Similarly, C₂₉ steranes (stigmastane) are derived from terrigenous sources and marine algae (Volkman, 1986). The regular steranes ternary plot for the Eagle Ford Shale samples (Figure 24) shows that the analyzed bitumens and oils are grouped together in a single area of the plot, indicating a common genetic relationship. Moreover, these samples cluster in the “open marine” and “marine shale/carbonate” fields of the Huang and Meinschein (1979) and Moldowan et al. (1985) ternary plots, respectively.

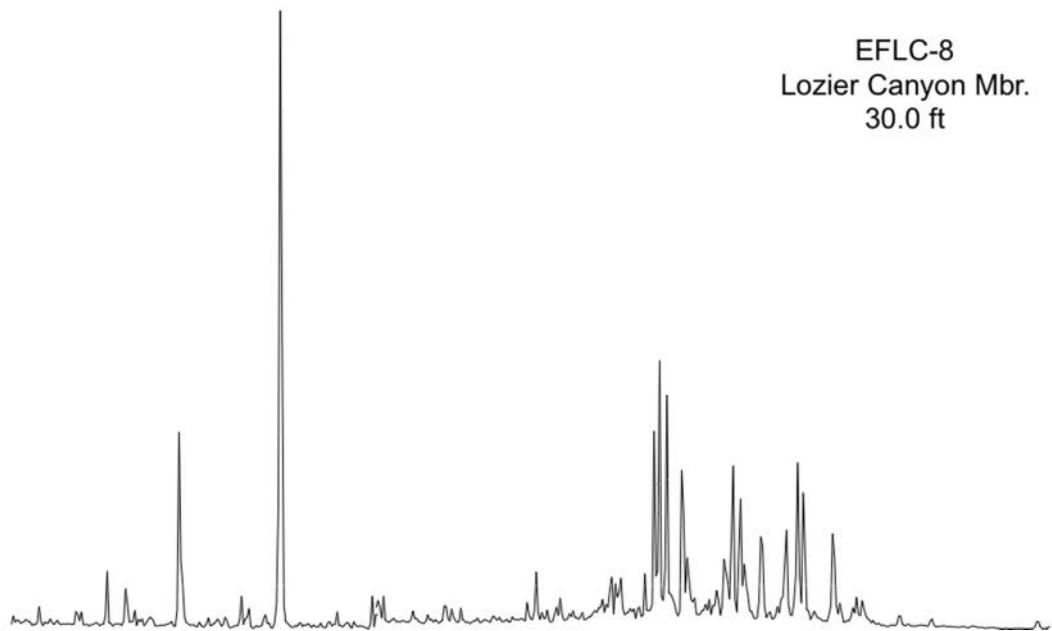
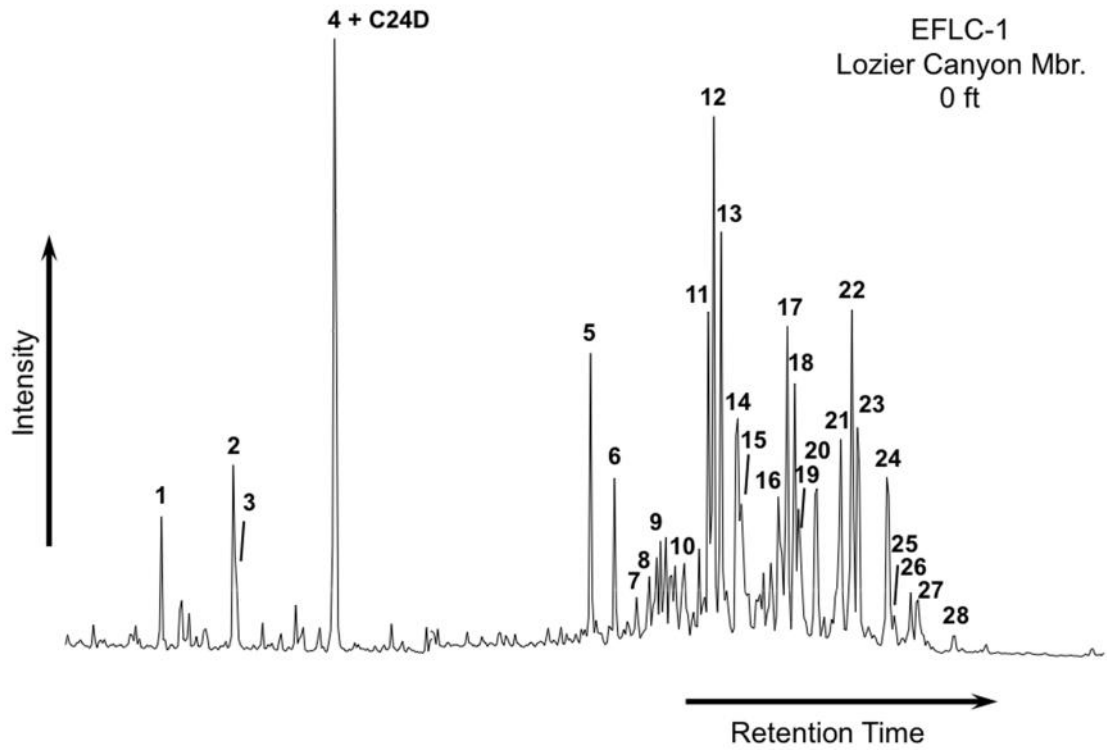


Figure 23. Partial fragmentograms of the m/z 217.3 ion showing distribution of steranes in the B&C fractions of the EFLC samples. Peak identification is presented in Table 6

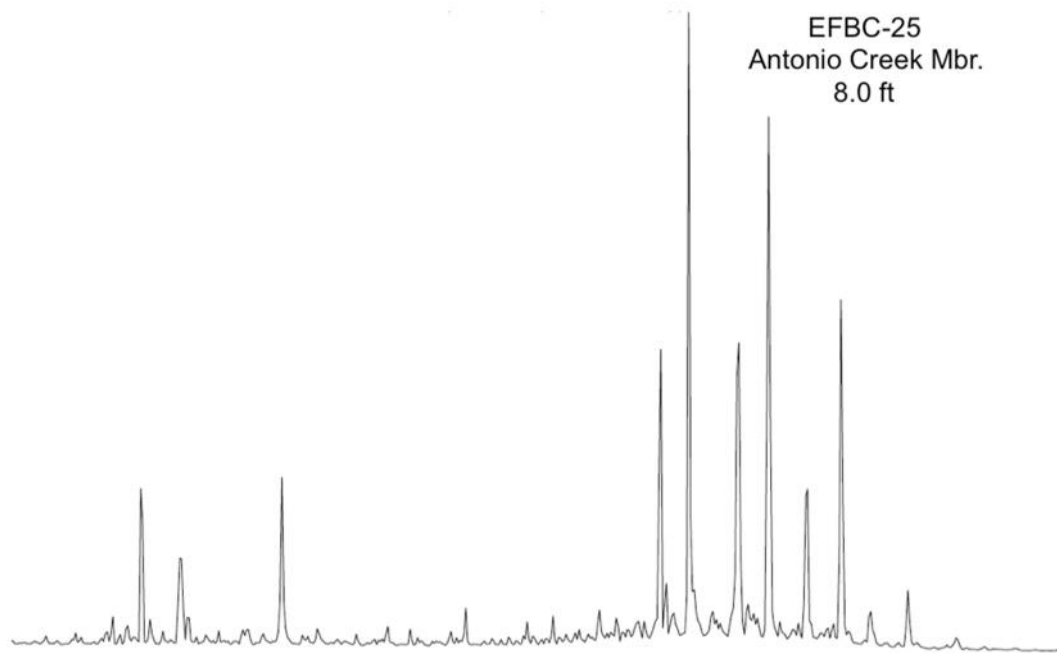
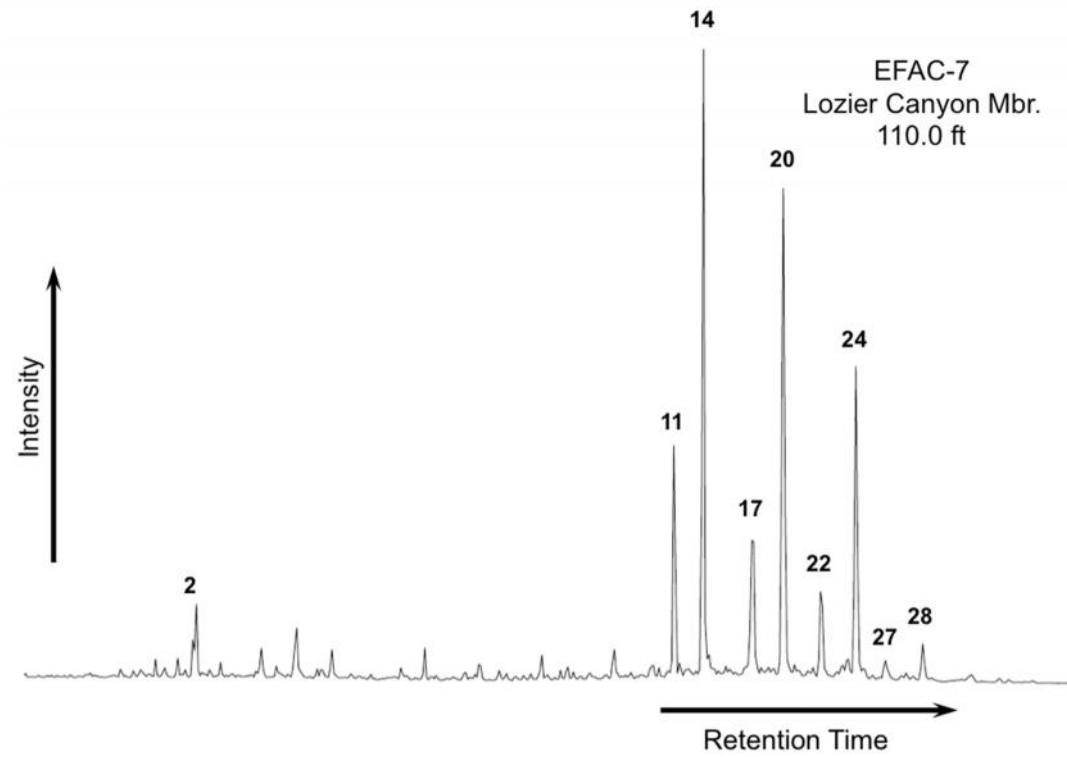


Figure 23. Partial fragmentograms of the m/z 217.3 ion showing distribution of steranes in the B&C fractions of the EFBC and EFAC samples. Peak identification is presented in Table 6 (cont.)

Table 6. Identification of steranes in the partial m/z 217.3 fragmentogram of the B&C fractions

Peak #	Compound
C24D	Deuterated <i>n</i> -tetracosane (ISTD)
1	C ₂₁ Diapregnane
2	C ₂₁ 14 (H),17 (H)-Pregnane
3	C ₂₂ Diahomopregnane
4	C ₂₂ 14 (H),17 (H)-Homopregnane
5	C ₂₇ 13 (H),17 (H)-Diacholestane (20S)
6	C ₂₇ 13 (H),17 (H)-Diacholestane (20R)
7	C ₂₇ 13 (H),17 (H)-Diacholestane (20S)
8	C ₂₇ 13 (H),17 (H)-Diacholestane (20R)
9	C ₂₈ 24-Methyl-13 (H),17 (H)-Diacholestane (20S)
10	C ₂₈ 24-Methyl-13 (H),17 (H)-Diacholestane (20R)
11	C ₂₈ 24-Methyl-13 (H),17 (H)-Diacholestane (20S) + C ₂₇ 14 (H),17 (H)-Cholestane (20S)
12	C ₂₉ 24-Ethyl-13 (H),17 (H)-Diacholestane (20S) + C ₂₇ 14 (H),17 (H)-Cholestane (20R)
13	C ₂₇ 14 (H),17 (H)-Cholestane (20S) + C ₂₈ 24-Methyl-13 (H),17 (H)-Diacholestane (20R)
14	C ₂₇ 14 (H),17 (H)-Cholestane (20R)
15	C ₂₉ 24-Ethyl-13 (H),17 (H)-Diacholestane (20R)
16	C ₂₉ 24-Ethyl-13 (H),17 (H)-Diacholestane (20S)
17	C ₂₈ 24-Methyl-14 (H),17 (H)-Cholestane (20S)
18	C ₂₈ 24-Methyl-14 (H),17 (H)-Cholestane (20R) + C ₂₉ 24-Ethyl-13 (H),17 (H)-Diacholestane (20R)
19	C ₂₈ 24-Methyl-14 (H),17 (H)-Cholestane (20S)
20	C ₂₈ 24-Methyl-14 (H),17 (H)-Cholestane (20R)
21	C ₂₉ 24-Ethyl-14 (H),17 (H)-Cholestane (20S)
22	C ₂₉ 24-Ethyl-14 (H),17 (H)-Cholestane (20R)
23	C ₂₉ 24-Ethyl-14 (H),17 (H)-Cholestane (20S)
24	C ₂₉ 24-Ethyl-14 (H),17 (H)-Cholestane (20R)
25	C ₃₀ 24-Propyl-14 (H),17 (H) -Cholestane (20S)
26	C ₃₀ 24-Propyl-14 (H),17 (H) -Cholestane (20R)
27	C ₃₀ 24-Propyl-14 (H),17 (H) -Cholestane (20S)
28	C ₃₀ 24-Propyl-14 (H),17 (H) -Cholestane (20R)

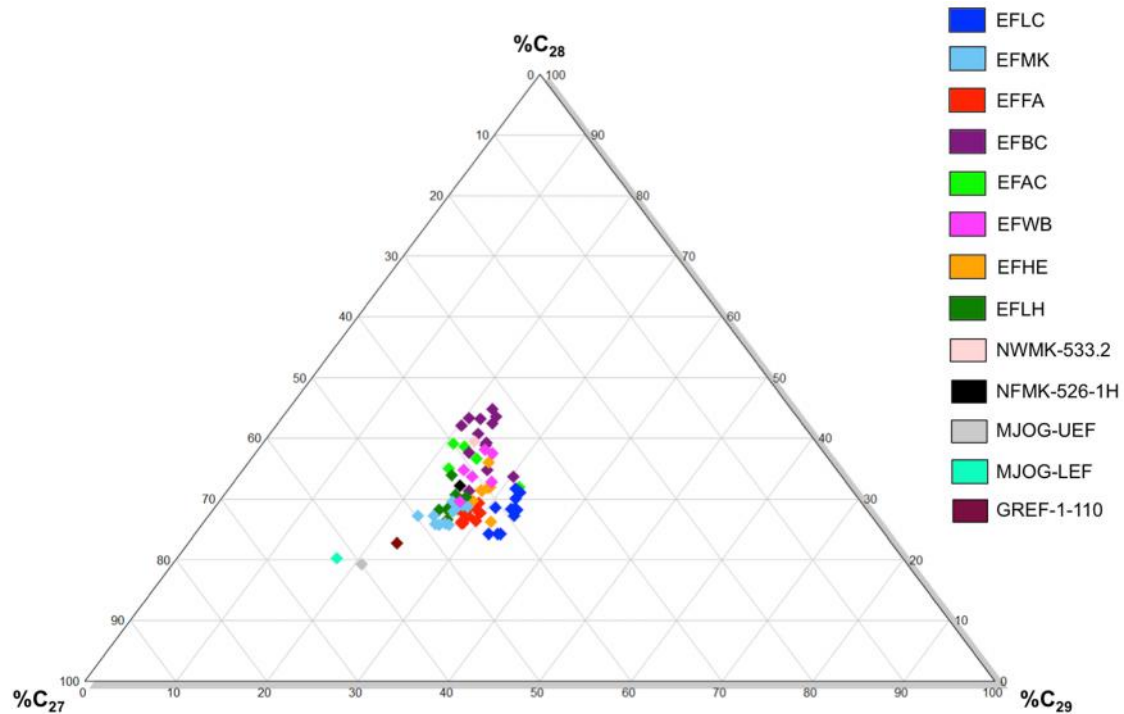


Figure 24. Ternary diagram of C₂₇, C₂₈, and C₂₉ regular steranes for the Eagle Ford Shale samples

Another group of steranes that are used for determining organic matter type are the C₃₀ 4-desmethylsteranes (also named 24-*n*-propylcholestanes or C₃₀ steranes). These compounds are derived from Chrysophyte, a marine algae of the order *Sarcinochrysidales*, which makes them specific biomarkers for marine organic matter (Moldowan et al., 1985; Peters et al., 2005). C₃₀ steranes were detected in all the samples analyzed. The steranes index, which combines the proportions of the C₃₀ sterane versus the C₂₇ to C₃₀ steranes (Appendix H) can be used to evaluate marine organic input (Peters et al., 2005). Geochemical logs of this ratio for the bitumen samples show small variations in source input within the different Eagle Ford members (Figure 25, Appendix F), suggesting a greater marine input in the lower Lozier Canyon Member and Antonio Creek Member.

2.2.1.2 *Diasteranes (rearranged steranes)*

Diasteranes are generated by the reduction of diasterenes, which are derived from sterols during diagenesis. This conversion reaction is catalyzed by acidic sites on clays, and promoted by low pH and oxic conditions (Moldowan et al., 1986; Peters, et al., 2005). Therefore, these biomarkers can be used as lithology and redox conditions indicators. The C_{27} diasteranes/ C_{27} steranes ratio can help differentiate between carbonate and siliciclastic source rocks in unaltered samples of similar thermal maturity (Mello et al., 1988; Peters et al., 2005). Low C_{27} diasteranes/ C_{27} steranes ratios are associated with anoxic carbonate sediments; whereas high ratios usually indicate clay-rich source rocks. Nevertheless, this evidence needs to be correlated with other geochemical parameters, since significant amounts of diasteranes have also been found in some clay-poor carbonate environments (Clark and Philp, 1989; Peters et al., 2005).

Geochemical logs of C_{27} diasteranes/ C_{27} steranes for the Eagle Ford Shale samples show lower values of this ratio in the Lozier Canyon Member, suggesting that this interval has lower clay content relative to the other members (Figure 25, Appendix F). These observations are supported by the descriptions of Donovan et al. (2013b) for the different Eagle Ford Shale members at Lozier Canyon, which characterize the Scott Ranch, Antonio Creek and Langtry members as clay-rich and containing significant amounts of bentonite beds. It is possible that these clays promoted steranes rearrangement to diasteranes. Diasteranes in samples EFAC and EFBC were present in very low concentrations or not detected at all possibly due to their low maturity.

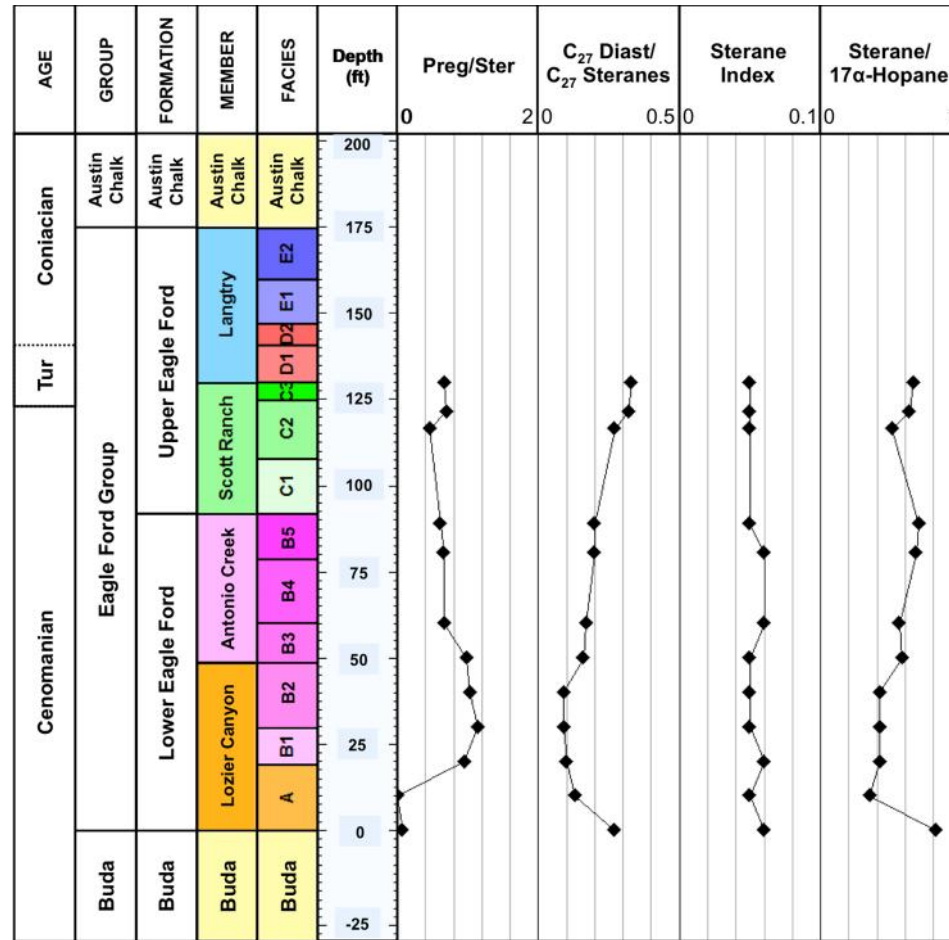


Figure 25. Geochemical logs of steranes ratios for the EFLC samples. Formulas for calculation of ratios are in Appendix A. Geochemical logs from other locations analyzed are in Appendix F. Numerical values of biomarker ratios are in Appendix H

2.2.1.3 *Pregnanes*

The origin of pregnane (C₂₀ sterane) and homopregnane (C₂₁ sterane) is not well understood (ten Haven et al., 1985, 1988). However, these compounds have been determined in high concentrations in bitumen extracted from source rocks associated with hypersaline depositional conditions. Wang (1993) analyzed extracts from the Springer Formation and found that pregnanes were more resistant to weathering compared to regular steranes and diasteranes. Pregnanes are also produced from degradation of the higher-carbon-numbered steranes at high maturity levels (R. Paul Philp, personal communication, 2014).

Geochemical logs of the pregnanes/steranes ratio for the Eagle Ford Shale samples show higher values in the Lower member, indicating the possibility of hypersaline, anoxic conditions present in the water column during deposition of this unit (Figure 25, Appendix F). In contrast, overall concentrations of pregnanes are much higher than those of dia- and regular steranes in samples from the EFMK and EFFA wells (Appendix L). These high maturity samples do not show any evidence of alteration, suggesting an enrichment of pregnanes relative to regular and rearranged steranes. Zhang et al. (2014) found that high pregnane/sterane and low 30-Norhopane/C₃₀ hopane in oil and bitumens from the Tazhong Uplift suggested enrichment of pregnane due to thermal maturation. A similar relation between these two geochemical parameters was observed for the EFMK, EFFA, and EFLH samples (Appendices H and I).

2.2.2 Terpanes

Terpanes constitute a group of compounds derived from prokaryotic organisms (eubacteria and blue-green algae), and plants. (Ourisson et al., 1982; Tissot and Welte, 1984; Peters et al. 2005). Terpene compounds include acyclic, bicyclic, tricyclic, tetracyclic, and pentacyclic homologous series (Peters et al., 2005). Tri-, tetra-, and pentacyclic terpanes were identified in the Eagle Ford Shale bitumens and oils by analysis of their B&C fractions through SIM/GC-MS of the m/z 191.3 ion. Fragmentograms of these compounds are presented in Figure 26. Peak identifications are in Table 7. Formulas for calculation of geochemical ratios are in Appendix A. Biomarkers quantitation results are in Appendix M. Geochemical ratios of terpanes (Appendix I) and their relationships with other biomarker groups helped assessing variations in organic matter, redox conditions, and thermal maturity for the Eagle Ford Shale bitumens and oils.

The regular steranes/17 α -Hopanes ratio is frequently used to evaluate eukaryote (plankton and/or benthic algae) versus prokaryote (bacteria) organic matter input in sediments. High values of this ratio are indicative of marine depositional environments, whereas low, near zero values are associated with terrigenous and/or reworked organic matter (Peters et al., 2005). In general, high steranes/17 α -Hopanes ratios are identified throughout the Eagle Ford Shale Group, indicating a greater marine organic matter input during deposition of these units (Figure 25, Appendix I). In the EFLH well, steranes/17 α -hopanes does not show much variation except for a decrease in this ratio near the boundary between the Maness and Eagle Ford Shale. This change might be

related to variations in depositional conditions that favored an increase in prokaryotic input at the end of Maness Shale deposition.

2.2.2.1 *Tricyclic terpanes*

Tricyclic terpanes (cheilanthanes) are thought to be derived from algae and bacterial precursors (Ourisson et al., 1982; Aquino Neto et al., 1981; Peters et al., 2005). Volkman et al. (1989) found significant amounts of tricyclic terpanes in bitumen from rocks containing *Tasmanites*, suggesting an origin from these organisms. Philp and Gilbert (1986) determined that tricyclic terpanes are abundant in marine oils. These compounds are highly resistant to thermal maturation and biodegradation compared to the hopanes. For that reason, tricyclic terpanes are commonly used in source rock characterization, evaluation of thermal maturity, biodegradation, and in oil-source rock correlations (Zumberge, 1987; Peters et al., 2005). Ratios of tricyclics/17 -hopanes and C₂₃ tricyclic/C₃₀ hopane help determining organic matter source by comparing bacterial and/or algal input versus prokaryotic contribution in source rock extracts and oils (Peters et al., 2005). Mello et al. (1988) observed high amounts of tricyclic terpanes associated with lacustrine saline and marine carbonate environments. Overall, the analyzed samples show higher tricyclics/17 -hopanes and C₂₃ tricyclic/C₃₀ hopane ratios in Lower Eagle Ford Shale (Figure 27, Appendix I), indicating a greater marine input during deposition of this member, as previously suggested.

Tricyclic terpane ratios are also helpful to differentiate depositional environments and identify source rock lithology (Peters et al., 2005). Particularly, the C₂₂/C₂₁ and the C₂₄/C₂₃ tricyclic terpane ratios assist in the identification of bitumens and oils derived from carbonate rocks. Extracts and oils derived from these source rocks

typically show high C_{22}/C_{21} and low C_{24}/C_{23} tricyclic terpane ratios, while marine source rocks show low C_{22}/C_{21} (0.20-0.50) and high C_{24}/C_{23} (0.50-1.00) tricyclic terpanes (Peters et al., 2005). Figure 28 shows a plot of C_{22}/C_{21} versus C_{24}/C_{23} tricyclic terpane ratios for bitumens and oils samples from the Eagle Ford Shale. From this graph it is suggested that most of these samples are derived from marine shales, with exception of few EFLC bitumens, which point to be derived from carbonate source rocks. Another geochemical parameter useful for determining source type is the C_{26}/C_{25} tricyclic terpane ratio (Figure 29), which helps distinguishing marine (0.50-1.30) from lacustrine source rocks (>1.0 ; Peters et al., 2005). Combination of this ratio with the $C_{31}R/C_{30}$ hopane ratio also assists in determining source rock type. Figure 29 shows the relation between these two parameters and indicate that the Eagle Ford Shale samples analyzed have a marine origin.

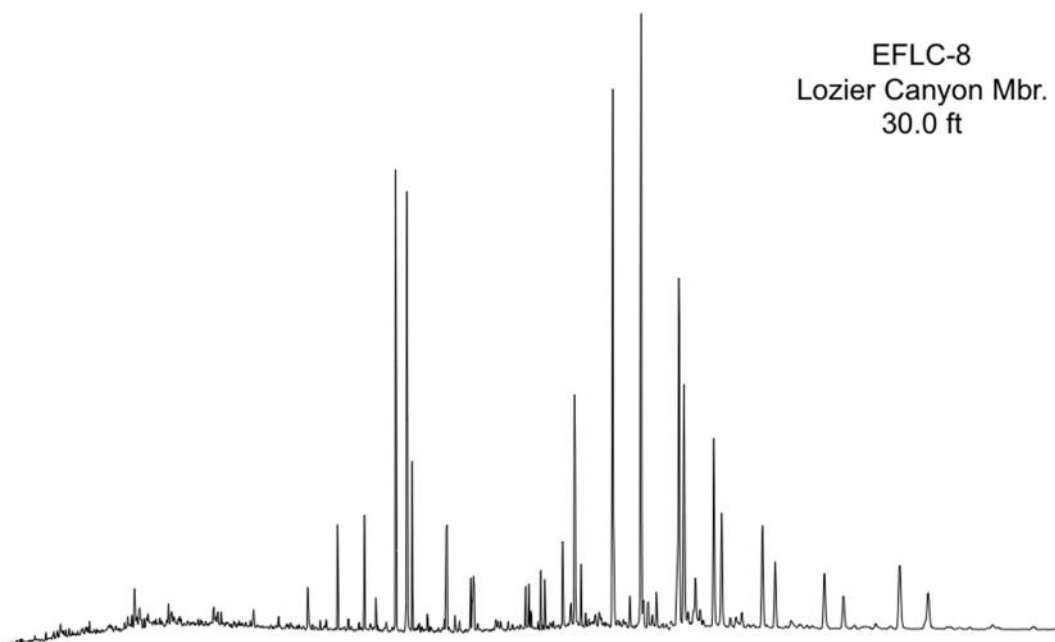
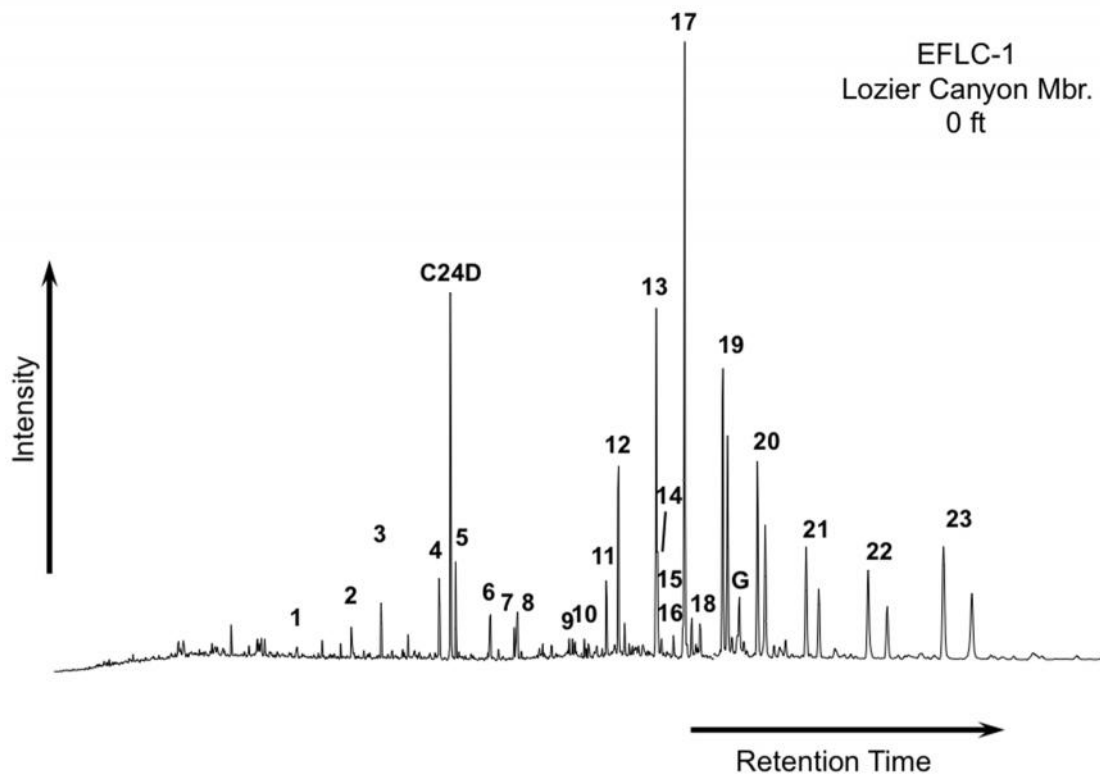


Figure 26. Partial fragmentograms of the m/z 191.3 ion showing distribution of terpanes in the B&C fractions of the EFLC samples. Peak identification is presented in Table 7

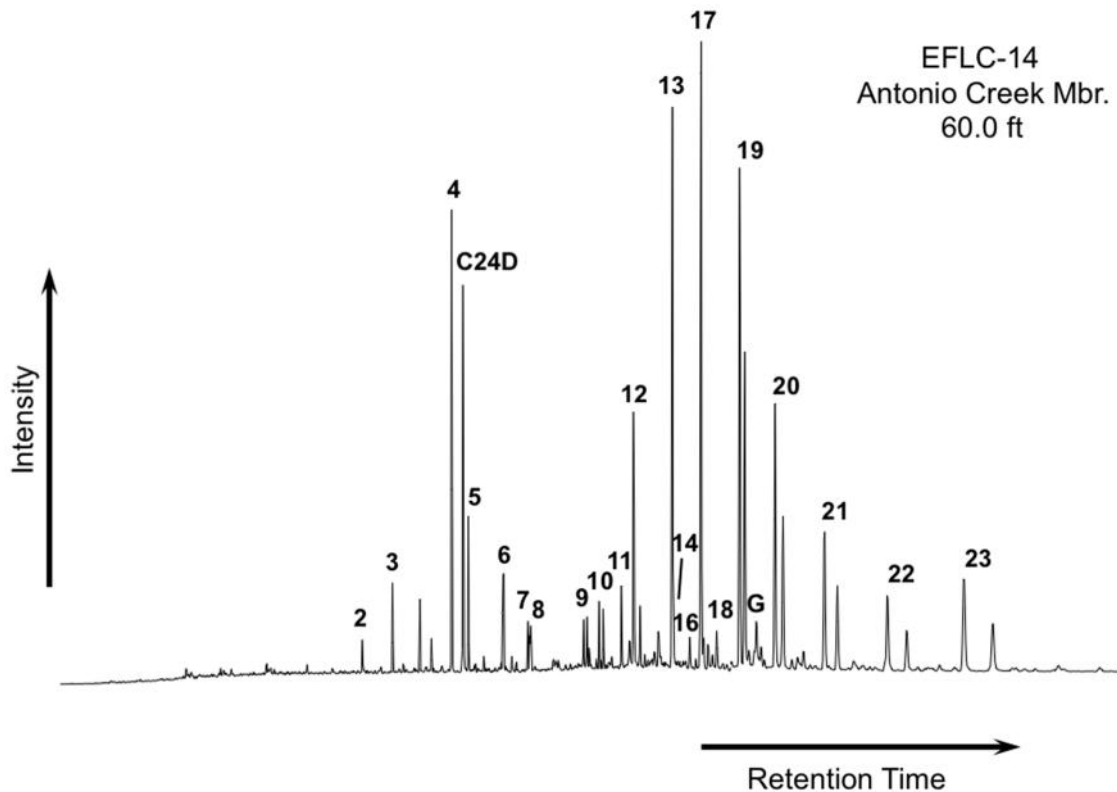


Figure 26. Partial fragmentograms of the m/z 191.3 ion showing distribution of terpanes in the B&C fractions of the EFLC samples. Peak identification is presented in Table 7 (cont.)

Table 7. Identification of terpanes in the partial m/z 191.3 fragmentogram of the B&C fractions

Peak #	Compound
C24D	Deuterated <i>n</i> -tetracosane (ISTD)
1	C₂₀ Tricyclic terpane (Cheilanthane)
2	C₂₁ Tricyclic terpane (Cheilanthane)
3	C₂₂ Tricyclic terpane (Cheilanthane)
4	C₂₃ Tricyclic terpane (Cheilanthane)
5	C₂₄ Tricyclic terpane (Cheilanthane)
6	C₂₅ Tricyclic terpanes (Cheilanthanes 22S and 22R)
7	C₂₄ Tetracyclic terpane
8	C₂₆ Tricyclic terpanes (Cheilanthanes 22S and 22R)
9	C₂₈ Tricyclic terpanes (Cheilanthanes 22S and 22R)
10	C₂₉ Tricyclic terpanes (Cheilanthanes 22S and 22R)
11	C₂₇ 18 (H)-22,29,30-Trisnorneohopane (Ts)
12	C₂₇ 17 (H)-22,29,30-Trisnorhopane (Tm)
13	C₂₉ 17 (H),21 (H)-30-Norhopane
14	C₂₉Ts 18 (H)-30-Norneohopane
15	C₃₀D 15 -methyl-17 (H)-27-Norhopane (Diahopane)
16	C₂₉ 17 (H),21 (H)-30-Normoretane
17	C₃₀ 17 (H),21 (H)-Hopane
18	C₃₀ 17 (H),21 (H)-Moretane
19	C₃₁ 17 (H),21 (H)-Homohopanes (22S & 22R)
21	C₃₂ 17 (H),21 (H)-Bishomohopane (22S & 22R)
22	C₃₃ 17 (H),21 (H)-Trishomohopane (22S & 22R)
23	C₃₄ 17 (H),21 (H)-Tetrakishomohopane (22S & 22R)
24	C₃₅ 17 (H),21 (H)-Pentakishomohopane (22S & 22R)
G	Gammacerane

2.2.2.2 *Tetracyclic terpanes*

Tetracyclic terpanes can be synthesized from bacteria or derived from hopanoid precursors. Similarly to the tricyclic terpanes, these compounds are more resistant to thermal maturity and biodegradation (Peters et al., 2005). Tetracyclic terpanes range from C₂₄ to C₂₇, where the C₂₄ homolog has a widespread occurrence (Aquino Neto et al., 1981; Peters et al., 2005). The C₂₄ tetracyclic terpane is frequently used as an indicator of carbonate and evaporite depositional environments (Palacas et al., 1984; Clark and Philp, 1989; Peters et al., 2005). The C₂₄ tetracyclic terpane was identified in all the analyzed bitumens and oils, and the C₂₄ tetracyclic terpane/C₃₀ hopane ratio show small variations within the different Eagle Ford Shale members (Figure 27, Appendix D). Because of the carbonate platform depositional environment of the Eagle Ford Shale it is unlikely that the presence of the C₂₄ tetracyclic terpane be related to a significant higher plant input. It is possible that these variations are more dependent on lithology than in organic facies variations. Rangel et al. (2000) observed significant amounts of C₂₄ tetracyclic terpane in the siliciclastic facies of La Luna Formation. A correlation between C₂₄ tetracyclic terpane/ C₃₀ hopane ratio and C₂₇ diasteranes/C₂₇ steranes was observed in the Antonio Creek and the upper interval of the Scott Ranch Member, suggesting that in this case, the presence of C₂₄ tetracyclic terpane may be dependent on clastic input.

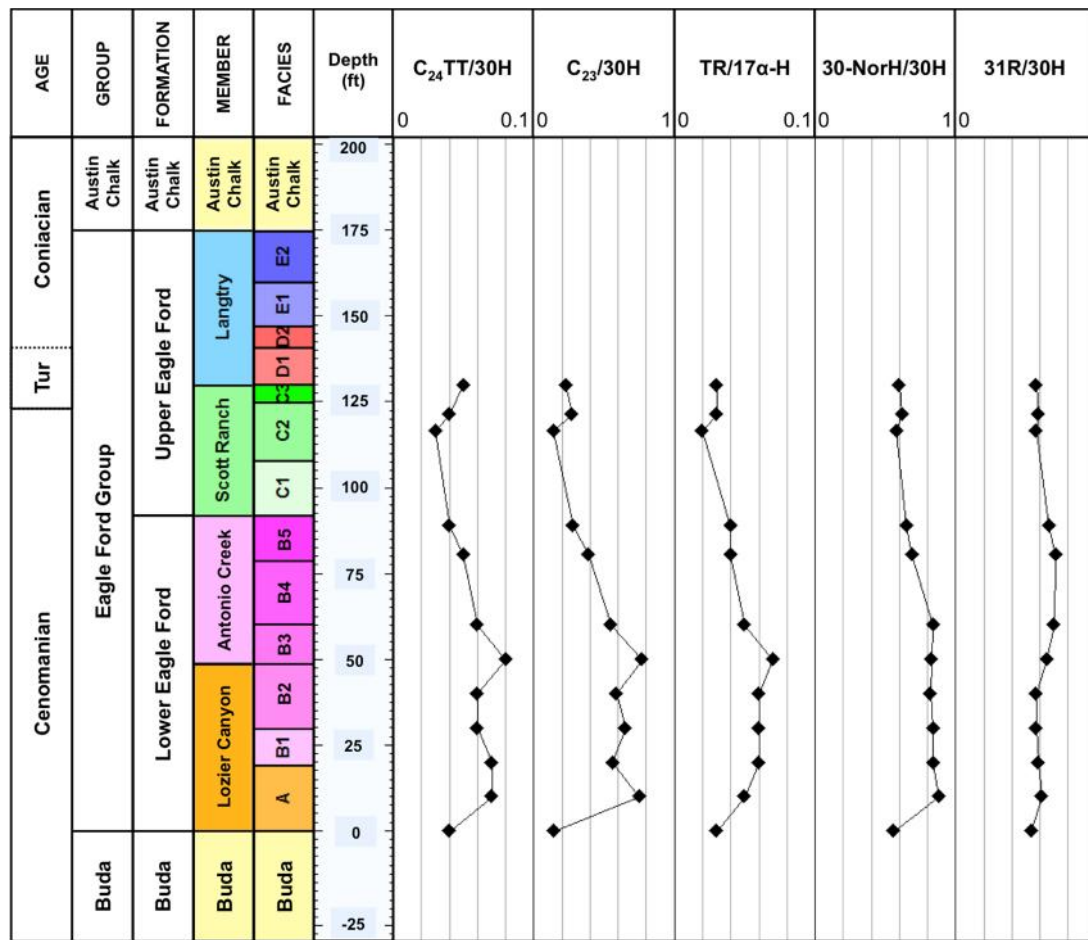


Figure 27. Geochemical logs of biomarker ratios of terpanes for the EFLC samples. Formulas for calculation of ratios are in Appendix A. Geochemical logs from other locations analyzed are in Appendix G. Numerical values of biomarker ratios are in Appendix I

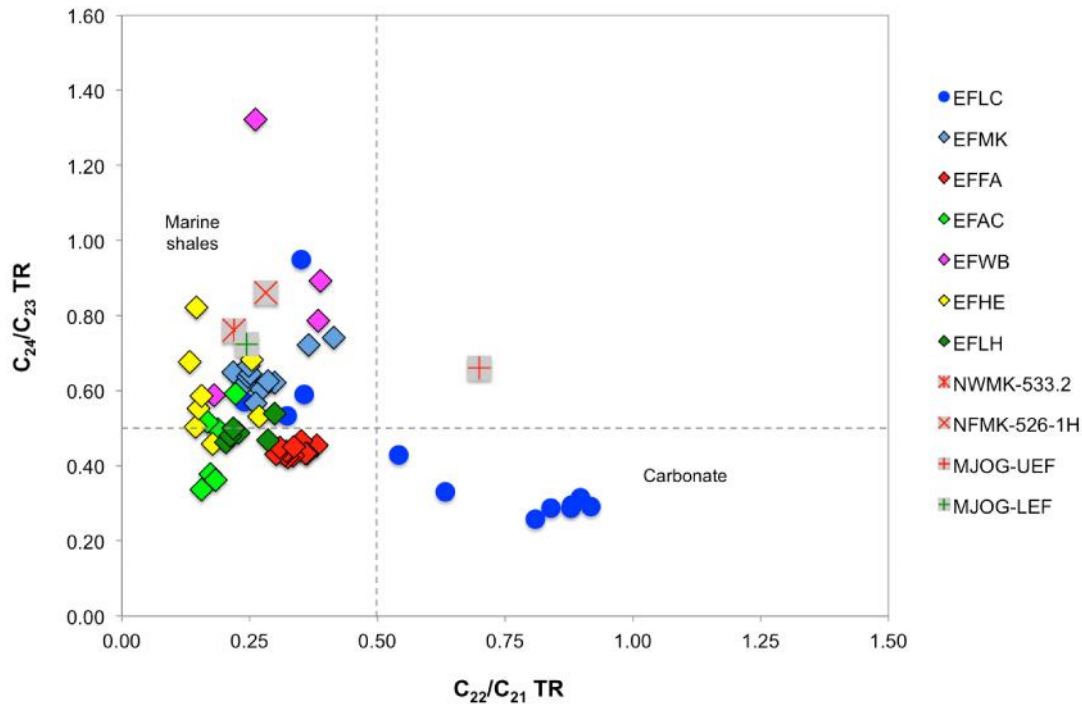


Figure 28. Plot of C_{22}/C_{21} versus C_{24}/C_{23} tricyclic terpanes shows source rock depositional environments for Eagle Ford Shale bitumens and oils (Dotted lines are used as a guide and do not represent fixed fields on the diagram. Plot modified from Peters et al., 2005)

2.2.2.3 Hopanes

Hopanes or pentacyclic terpanes are biomarkers derived from hopanoid compounds present in cell membranes of prokaryotes (bacteria and cyanobacteria). Because of their origin, hopanes are ubiquitous in bitumens and oils (Peters and Moldowan, 1991). The C_{29} 17 (H)-norhopane (30-Norhopane) and the C_{30} 17 (H)-hopane are usually the major peaks in the m/z 191 fragmentograms and these biomarkers may be used as environmental indicators (Waples and Machihara, 1991; Peters et al., 2005). The 30-NorH/30H hopane ratio increases with increasing thermal maturity due to the higher stability of the C_{29} hopane compared to the C_{30} homolog (Peters et al., 2005). Most of the Eagle Ford Shale samples analyzed have 30-NorH/30H

values lower than 1.0; however, higher concentrations of 30-Norhopane were observed in the Lozier Canyon Member, probably due to higher amounts of carbonates in this interval (Figure 27, Appendix I). Immature samples EFBC and EFAC have 30-NorH/30H ratios greater than 1.0, which point towards a marine carbonate origin for the Eagle Ford Shale. The higher 30-NorH/30H ratios in the EFFA samples and oils are possibly the result of their high thermal maturity. When compared to the $C_{35}S/C_{34}S$ hopanes ratio, the 30-NorH/30H can help identify source facies of oils and extracts (Figure 30). According to Peters et al. (2005), the majority of oils and bitumens generated from marine carbonates have high $C_{35}S/C_{34}S$ and 30-NorH/30H ratios (>0.8 and >0.6 respectively). This relationship is observed in samples EFLC, EFMK, and EFHE (Figure 30). On the contrary, samples from EFWB, EFLH, and the NWMK-533.2 oil suggest to be derived from a siliciclastic source. In addition, the low values of the $C_{35}S/C_{34}S$ ratio for the EFLH samples suggest dysoxic to oxic source rock deposition. The relationship between $C_{35}S/C_{34}S$ as redox indicators will be explained in the next section. Geochemical logs of the 30-NorH/30H for the bitumens (Figure 27, Appendix G) show higher values of this ratio in the Lozier Canyon Member, possibly suggesting a higher carbonate input during deposition of this interval. In addition, it is inferred that the 30-NorH/30H values greater than 1.0 for the EFFA bitumens are not indicative of source type, but the result of the high thermal maturity of these samples. Correlation of this parameter with other geochemical ratios, such as tricyclics/17-hopanes, C_{23} tricyclic/ C_{30} hopane, and C_{24} tetracyclic terpane/ C_{30} hopane ratios point towards a marine source rock for these Eagle Ford Shale bitumens and oils.

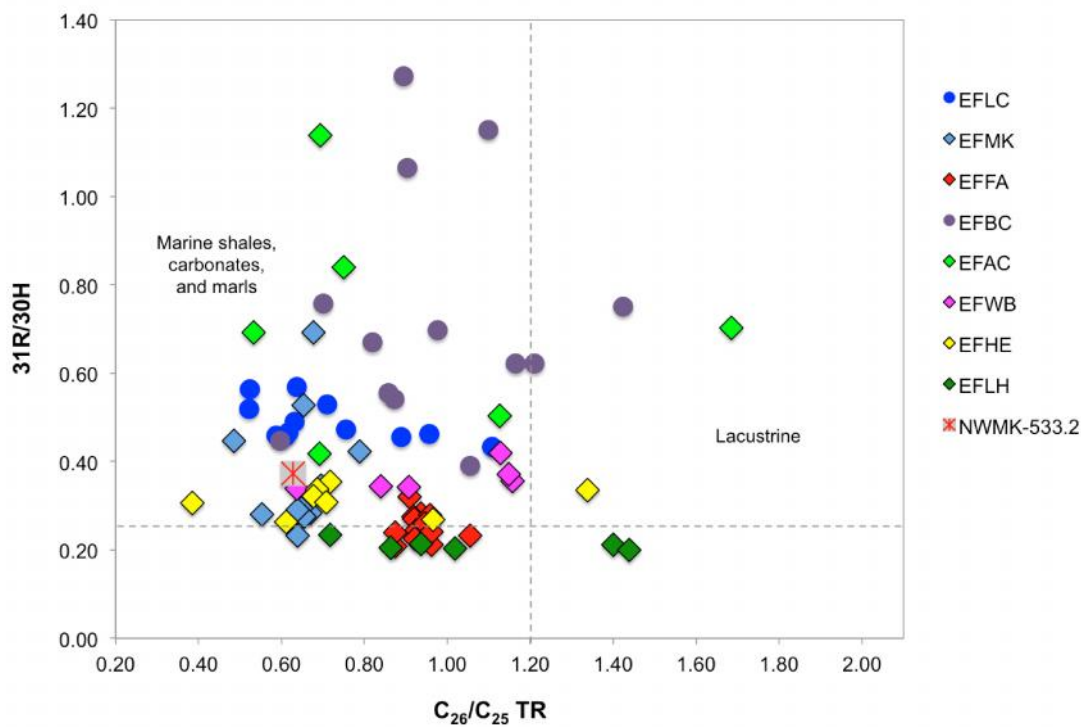


Figure 29. Plot of C_{26}/C_{25} versus $C_{31}R/C_{30}$ Hopane shows that the Eagle Ford Shale bitumens and oils are mainly of marine origin (Dotted lines are used as a guide and do not represent fixed fields on the diagram. Plot modified from Peters et al., 2005)

The $C_{31} 22R/C_{30}$ hopane ratio (31R/30H) is applied to differentiate source rocks of marine versus lacustrine origin. $C_{31} 22R/C_{30}$ hopane ratios greater than 0.25 are associated with marine shale, carbonate, and marl source rocks (Peters et al., 2005). Geochemical logs of this ratio for the Eagle Ford Shale extracts (Figure 27, Appendix G) do not show much variation between the different members, and the majority of samples have values of 31R/30H indicative of a marine origin (Figure 29).

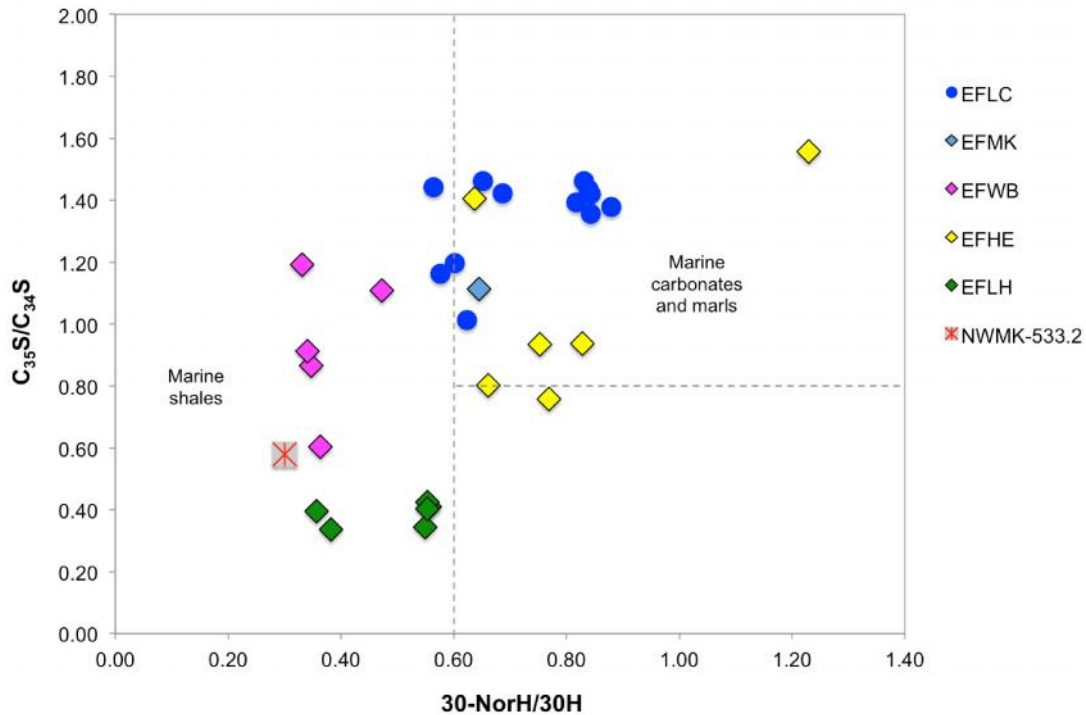


Figure 30. Plot of 30-Nor/C₃₀ hopane versus C₃₅S/C₃₄S homohopanes suggest most of the Eagle Ford Shale rock samples are of marine origin (Dotted lines are used as a guide and do not represent fixed fields on the diagram. Plot modified from Peters et al. 2005)

The relative distribution of the extended hopanes or homohopanes (C₃₁-C₃₅) has been widely used to assess redox potential during marine source rock deposition. These compounds originated from bacteriohopanetetrol and other C₃₅ hopanoids (Peters and Moldowan, 1991; Waples and Machihara, 1991). Particularly, higher proportions of the C₃₅ homohopane with respect to its C₃₄ homolog are associated with anoxic marine depositional environments due to preferential preservation of the C₃₅ homolog (Peters and Moldowan, 1991; Moldowan et al., 1992; Peters et al., 2005). The C₃₅S/C₃₄S ratio is one of the geochemical parameters used for evaluation of redox variations at the time of source rock deposition. Values of this ratio greater than 0.8 are frequently related to

marine carbonate source rocks (Figure 30; Peters et al., 2005). Moreover, the homohopane index ($C_{35}/C_{31-C_{35}}$) is commonly calculated in extracts and oils to evaluate redox conditions, identifying variations in organic facies in the same source rock, and for oils and source rocks correlation studies (Clark and Philp, 1989; Peters and Moldowan, 1991; Peters et al., 2005).

Plots of homohopane distributions for the Eagle Ford Shale bitumens and oils analyzed (Figure 31) indicate that in general, these source rocks were deposited under anoxic conditions, which promoted preservation of the C_{35} homohopane. Samples from the East Texas EFLH core, and the NWMK-533.2 oil do not show this predominance of the C_{35} homohopane, suggesting source rock deposition under suboxic-oxic conditions. C_{34} and C_{35} homohopanes were not detected in most of the EFMK and EFFA extracts, and the rest of the analyzed oils. This could be attributed to the high maturity of these samples, since it has been observed that thermal maturation can increase the proportion of the lower molecular weight homohopanes (Moldowan et al., 1992). Another observation is the difference in homohopane distributions within each Eagle Ford Shale member for all outcrops and cores analyzed, which could indicate organic facies variations between the different units (Figure 31). For example, samples EFLH-15 and -20 from the EFLH core, which correspond to the Maness Shale, show different homohopane distributions from samples EFLH-1 to -13, which are from the Lower Eagle Ford Shale. In the case of the EFLC outcrop, samples EFLC-25, -26, and -28 corresponding to the Scott Ranch Member (lower Upper Eagle Ford Shale) do not show a strong predominance of C_{35} over C_{34} homohopanes, possibly indicating deposition of this member under suboxic-oxic conditions. Variations in homohopane distributions in

samples from the EFHE core, suggest organic facies changes between samples EFHE-1 and -2 (Langtry Member); EFHE-3 to -7 (Langtry and Antonio Creek members); and EFHE-9 and -10 (Lozier Canyon Member).

Gammacerane was tentatively identified in all the Eagle Ford Shale samples analyzed with exception of the MJOG oils, where terpanes in general show very low concentrations (Figure 26). This compound is a C₃₀ triterpane originated during diagenesis by reduction of tetrahymanol, whose precursor corresponds to *Tetrahymena*, a fresh water ciliated protozoan (ten Haven et al., 1989; Peters et al., 2005). Gammacerane is a specific biomarker for water-column stratification and hypersaline conditions in marine and non-marine environments (Sinninghe Damsté et al., 1995; Peters et al., 2005).

The occurrence of gammacerane in the Eagle Ford Shale samples suggests that regional hypersaline conditions and water stratification may have developed during deposition of these sediments. This observation positively correlated with the low Pr/Ph ratios and the predominance of the steranes over the steranes, both parameters also related to hypersalinity (ten Haven et al., 1988). Nevertheless, the presence of gammacerane in these samples needs to be further confirmed by fullscan GCMS analysis in order to analyze variations in the concentration of this biomarker in the Eagle Ford Shale.

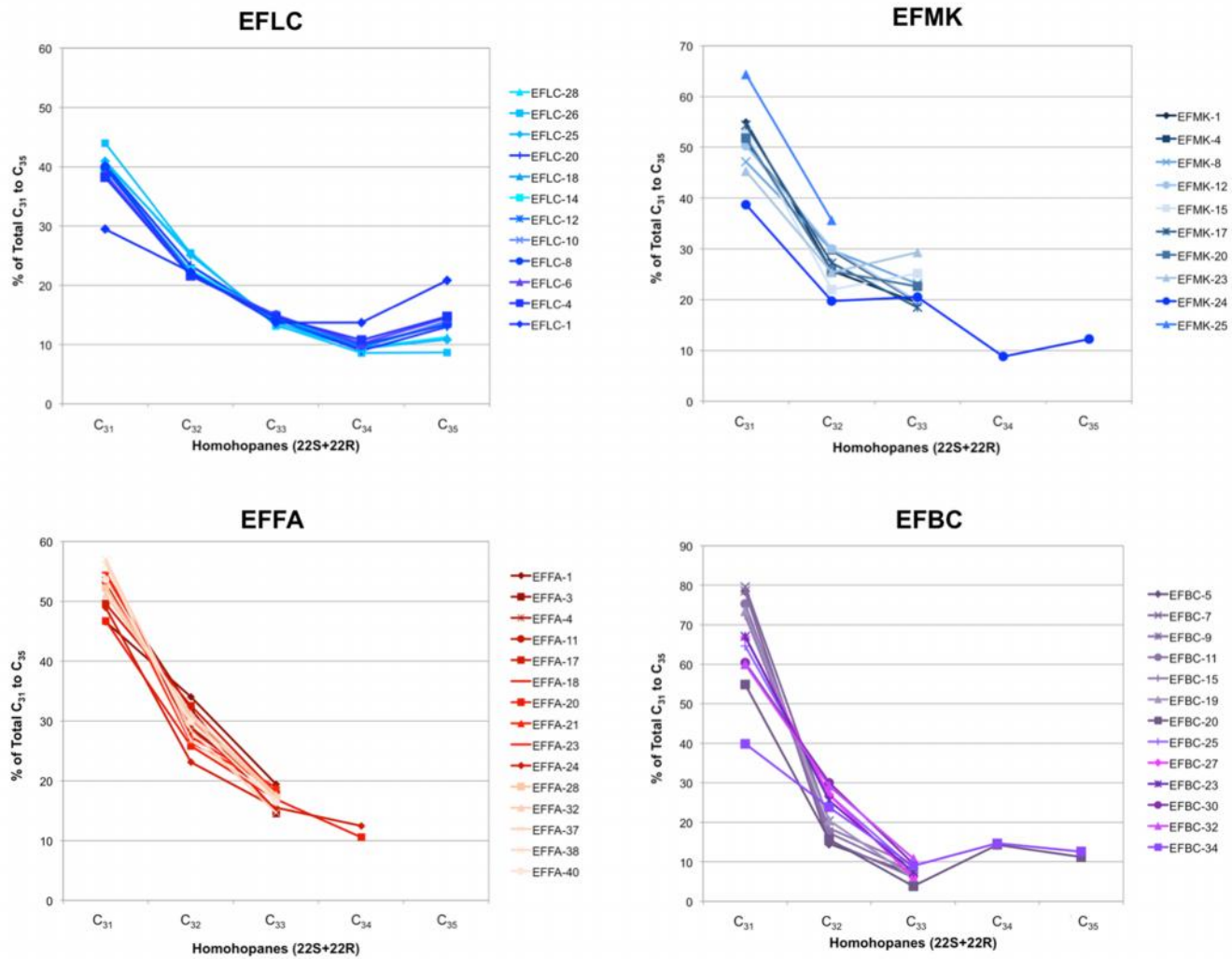


Figure 31. Homohopanes distribution for bitumens and oils from the Eagle Ford Shale

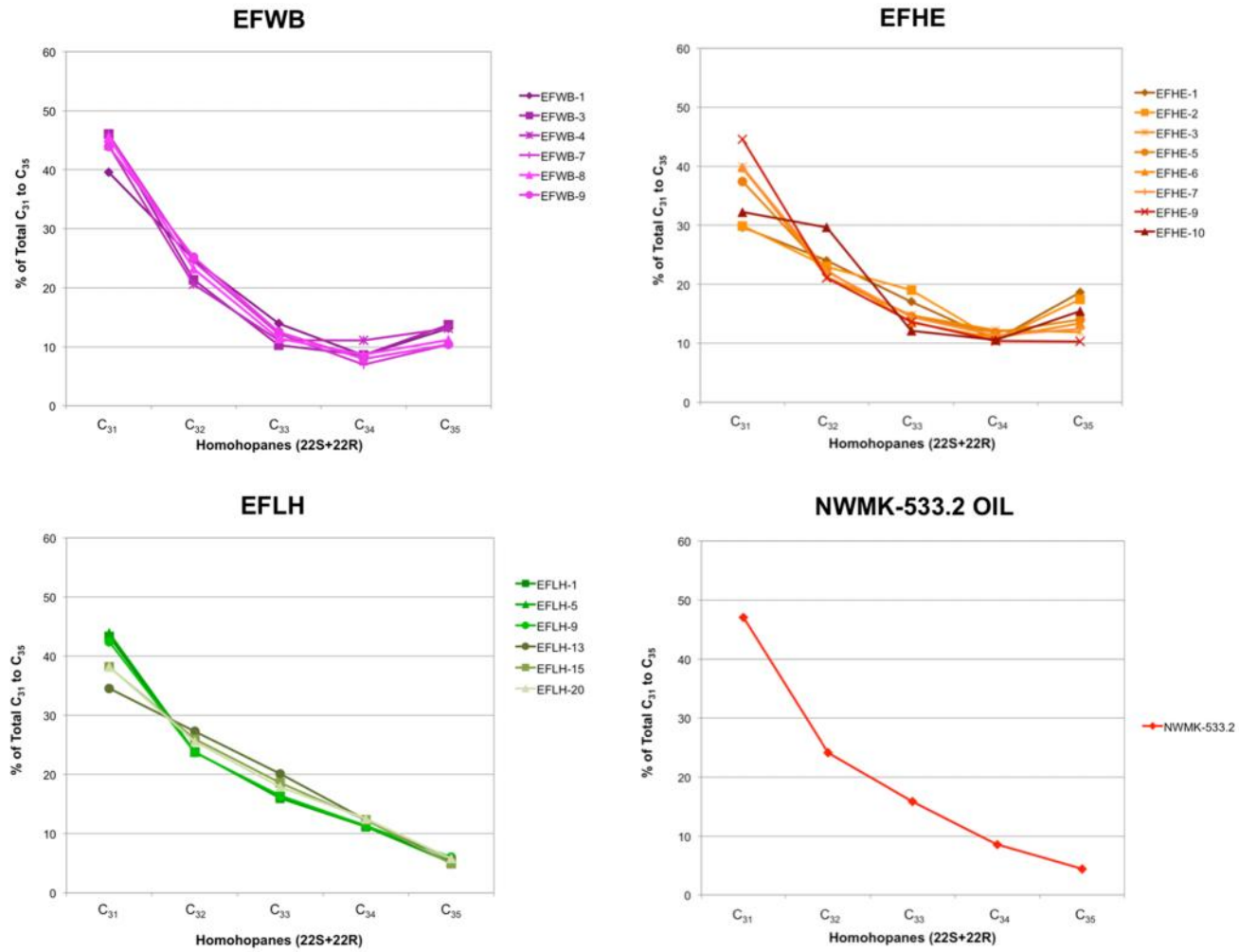


Figure 31. Homohopanes distribution for bitumens and oils from the Eagle Ford Shale (cont.)

2.2.3 Aryl Isoprenoids

Aryl isoprenoids, or trimethylbenzenes, are compounds derived from isorenieratene, a C₄₀ diaromatic carotenoid pigment produced by the green sulfur bacteria *Chlorobiaceae* (Brown and Kenig, 2004; Schwark and Frimmel, 2004). *Chlorobiaceae* performs photosynthesis under light-penetrated, H₂S-saturated, anoxic waters; thus, compounds derived from these bacteria can be specific indicators of photic zone anoxia (PZA; Schwark and Frimmel, 2004).

Analysis of aryl isoprenoids was performed in the B&C fractions of bitumens and oils by SIM/GC-MS using ions m/z 133.1 and 134.1. Results from aryl isoprenoids quantitation are in Appendix N. In all the analyzed samples, only the 1-alkyl-2,3,6-trimethylbenzenes (m/z = 133.1) were identified. The 1-alkyl-3,4,5-trimethylbenzenes (m/z = 134.1) were absent or in very low concentrations hampering detection of these compounds. The identified aryl isoprenoids were a series of C₁₃-C₂₂ homologs, and detection beyond the C₂₆ homolog was problematic (Figure 32). Evaluation of these compounds to understand possible variability of PZA in the Eagle Ford Shale was performed through the calculation of the aryl isoprenoid ratio (AIR; Appendix A). Schwark and Frimmel (2004) introduced this ratio by analyzing the proportions of the short-chain (C₁₃-C₁₇) and intermediate-chain (C₁₈-C₂₂) aryl isoprenoids. High AIR (3.0) are associated with episodic PZA, which leads to alteration of the long- and intermediate-chain aryl isoprenoids. On the contrary, low AIR (0.5) indicates persistent PZA, which contributes to preservation of the long-chain aryl isoprenoids (Schwark and Frimmel, 2004).

Most of the analyzed samples show AIR greater than 1 (Appendix J) suggesting that the Eagle Ford Shale experienced episodic periods of PZA, where shifts of the chemocline occurred periodically. Koopmans et al. (1996) determined that α -isorenieratane and β -carotane are also precursors for the 2,3,6 trimethyl-substituted aryl isoprenoids. When derived from these compounds, the 2,3,6 trimethyl-substituted aryl isoprenoids are depleted in ^{13}C and cannot be used as specific biomarkers for PZA. In the present study, the ^{13}C values for the aryl isoprenoids were not determined. Nevertheless, geochemical logs of AIR for the Eagle Ford Shale samples show a positive correlation with Pr/Ph (Figure 33; Appendix C), and aryl isoprenoids were used to tentatively evaluate variations and degree of PZA. A graph of AIR versus Pr/Ph (Figure 34) helps to better assess redox conditions and PZA during sediments deposition (Schwark and Frimmel, 2004). From this diagram it can be observed that samples from EFLC and EFBC outcrops, and the EFAC core were possibly deposited under anoxic-suboxic conditions and persistent PZA. Samples EFWB and EFHE seem to have experienced episodic PZA. High AIR in samples EFMK, EFFA, and oils could be consequence of their high thermal maturity, which tend to generate short-chain aryl isoprenoids from the long-chain homologs.

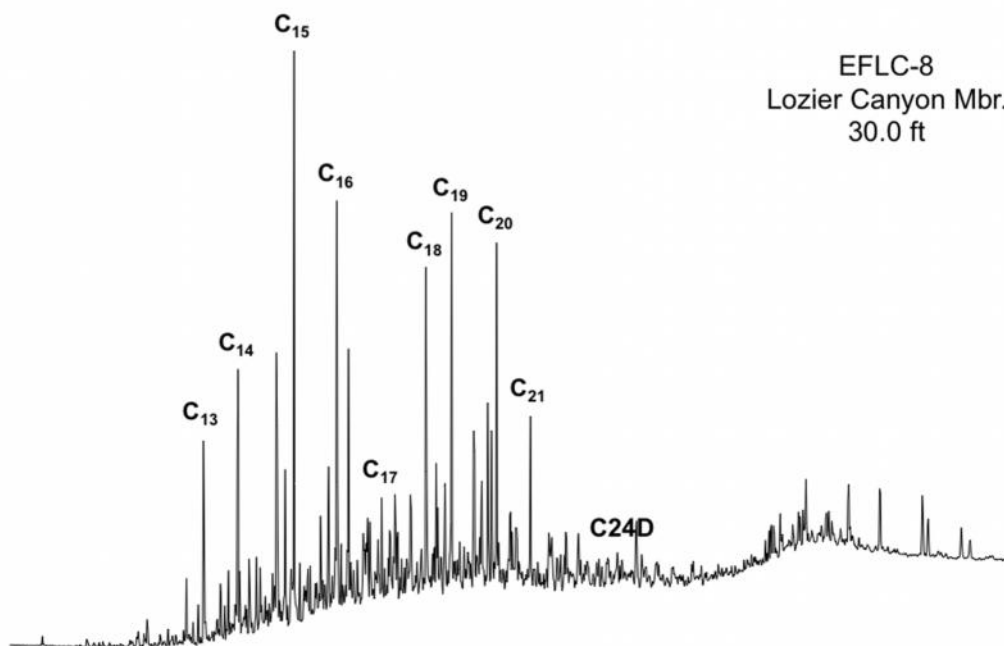
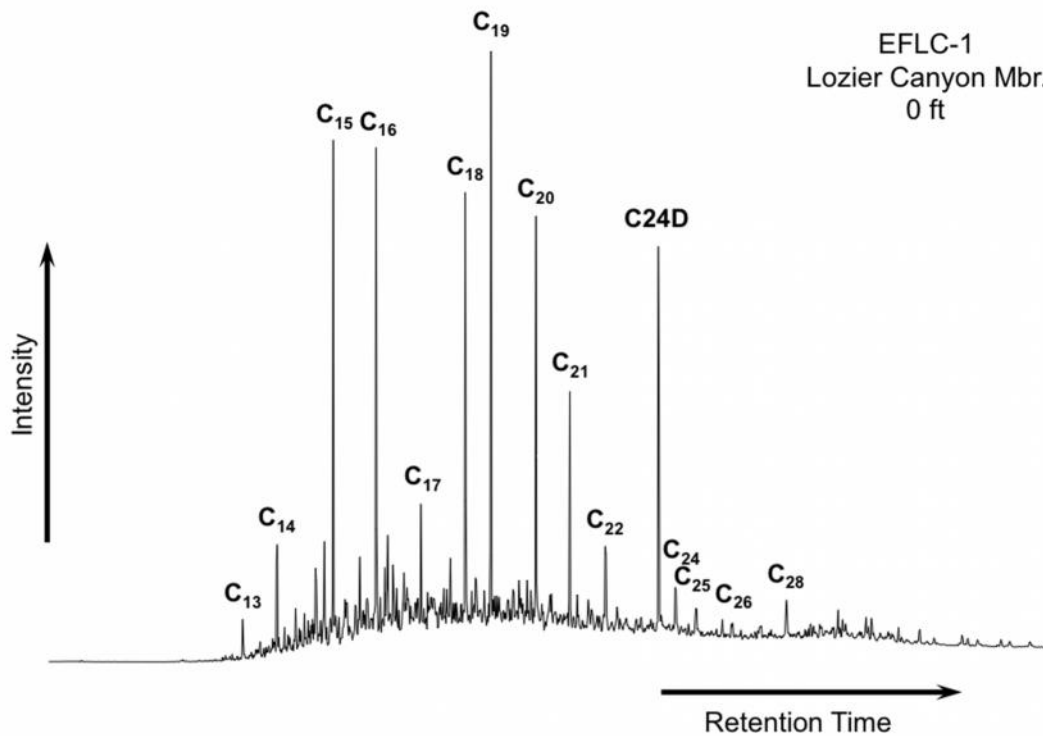


Figure 32. Partial fragmentograms of the m/z 133.1 ion showing the aryl isoprenoids distribution in the B&C fractions of the EFLC samples. Number of carbon atoms for the 2,3,6-trimethyl substituted aryl isoprenoids is indicated above each peak (C_{24D} = internal standard)

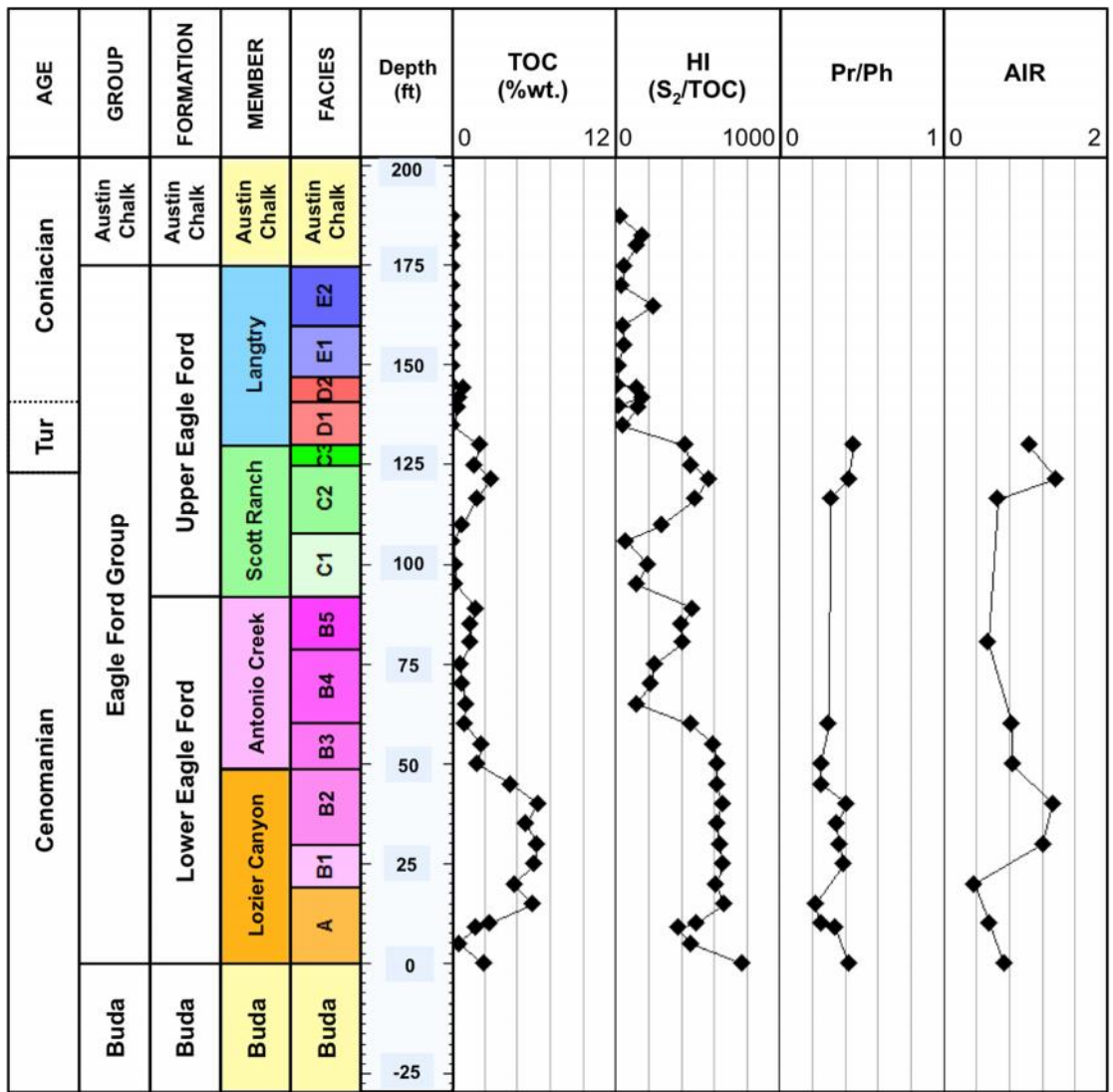


Figure 33. Geochemical logs of biomarker ratios of aryl isoprenoids for Eagle Ford Shale samples at Lozier Canyon. Formulas for calculation of ratios are in Appendix A. Geochemical logs from other locations analyzed are in Appendix C. Numerical values of AIR are in Appendix J

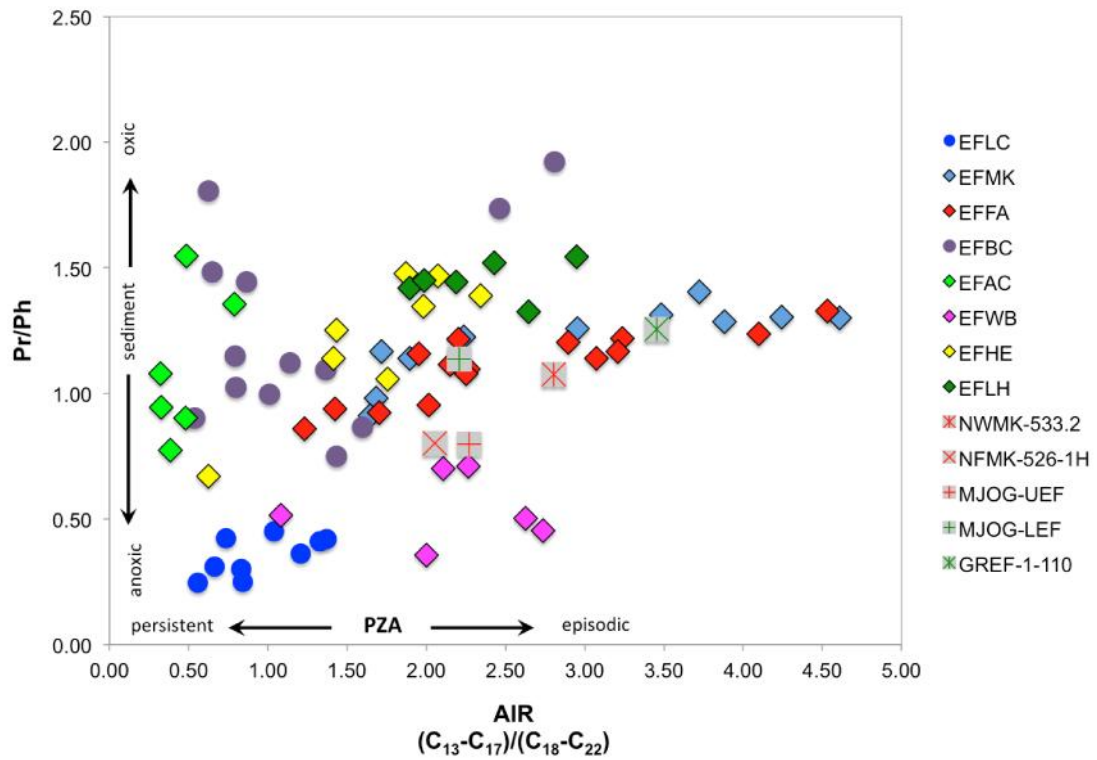


Figure 34. AIR versus Pr/Ph plot for the Eagle Ford Shale samples (Dotted line is used as a guide and does not represent fixed fields on the diagram. Plot template from Schwark and Frimmel, 2004)

2.2.4 Monoaromatic Steroids

Monoaromatic steroids (MAS) are a group of biomarkers derived from sterols containing more than one double bond (Peters et al., 2005). These biomarkers were determined in the B&C fraction of Eagle Ford Shale bitumens and oils through SIM/GC-MS by analysis of the m/z 253.3 ion (Figure 35, Table 8). Compared to the regular steranes ternary plot, the proportions of the C_{27} , C_{28} , and C_{29} MAS plotted in a ternary diagram provide a better characterization of the variations in depositional environments, and are particularly useful for distinguishing marine from non-marine oils (Moldowan et al., 1985). These authors indicated that oils and bitumens derived from marine organic matter contain greater amounts of C_{28} MAS than those of non-marine origin. In addition, Moldowan et al. (1985) observed that some oils derived from marine carbonate source rocks contain higher amounts of C_{29} MAS than oils derived from marine shales. The bitumens and oils from the Eagle Ford Shale have a greater concentration of C_{28} MAS, followed by C_{27} and C_{29} MAS (Appendix O) and their distribution is similar to that of the regular steranes (Figure 36), indicating a marine source for these samples.

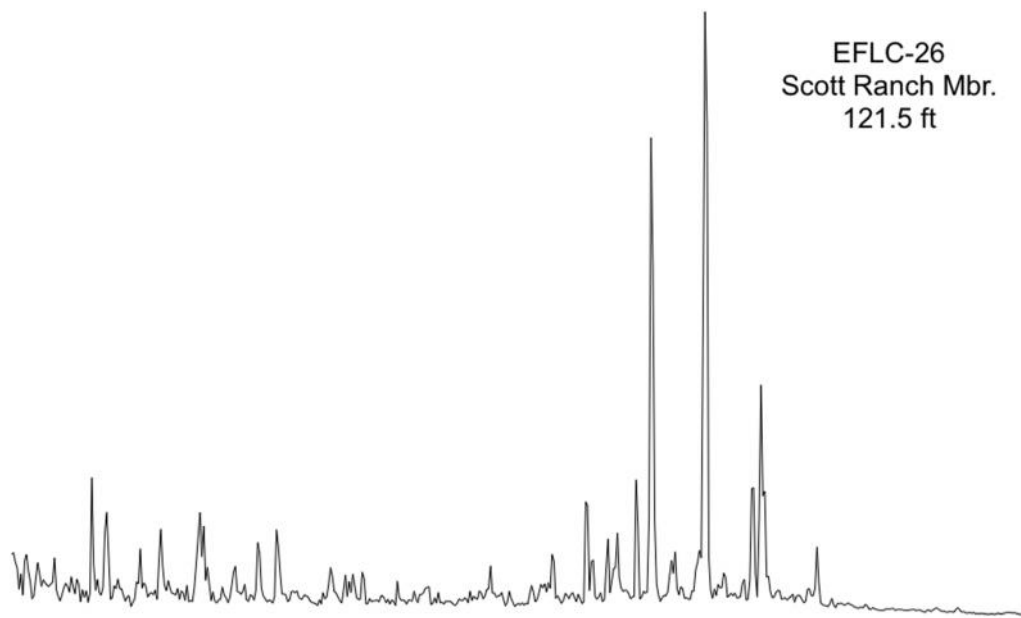
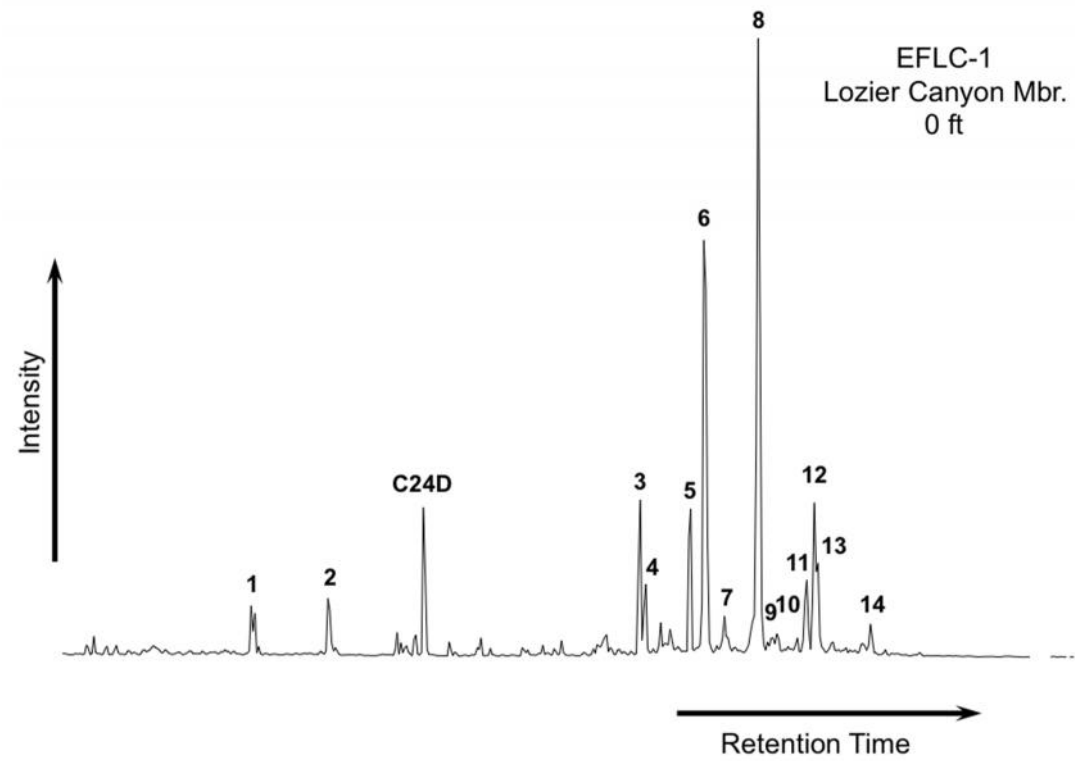


Figure 35. Partial fragmentograms of the m/z 253.3 ion showing distribution of the monoaromatic steroids (MAS) in the B&C fractions of the EFLC samples. Peak identification is presented in Table 8

Table 8. Identification of monoaromatic steroids (MAS) in the partial m/z 253.3 fragmentogram of the B&C fractions

Peak #	Compound
C24D	Deuterated <i>n</i> -tetracosane (ISTD)
1	C ₂₁ Pregnane
2	C ₂₂ 20-Methylpregnane
3	C ₂₇ 5 -Cholestane 20S
4	C ₂₇ Diacholestane 20S
5	C ₂₇ Monoaromatic steroid
6	C ₂₈ 5 -Ergostane 20S + C ₂₈ Diaergostane 20S
7	C ₂₇ 5 -Cholestane 20R
8	C ₂₈ 5 -Ergostane 20S
9	C ₂₈ 5 -Ergostane 20R + C ₂₈ Diaergostane 20R
10	C ₂₉ 5 -Stigmastane 20S + C ₂₉ Diastigmastane 20S
11	C ₂₉ 5 -Stigmastane 20S
12	C ₂₈ 5 -Ergostane 20R
13	C ₂₉ 5 -Stigmastane 20R + C ₂₉ Diastigmastane 20R
14	C ₂₉ 5 -Stigmastane 20R

Peak 5 could be C₂₇ 5 -Cholestane 20R + C₂₇ Diacholestane 20R or C₂₇ 5 -Cholestane 20S

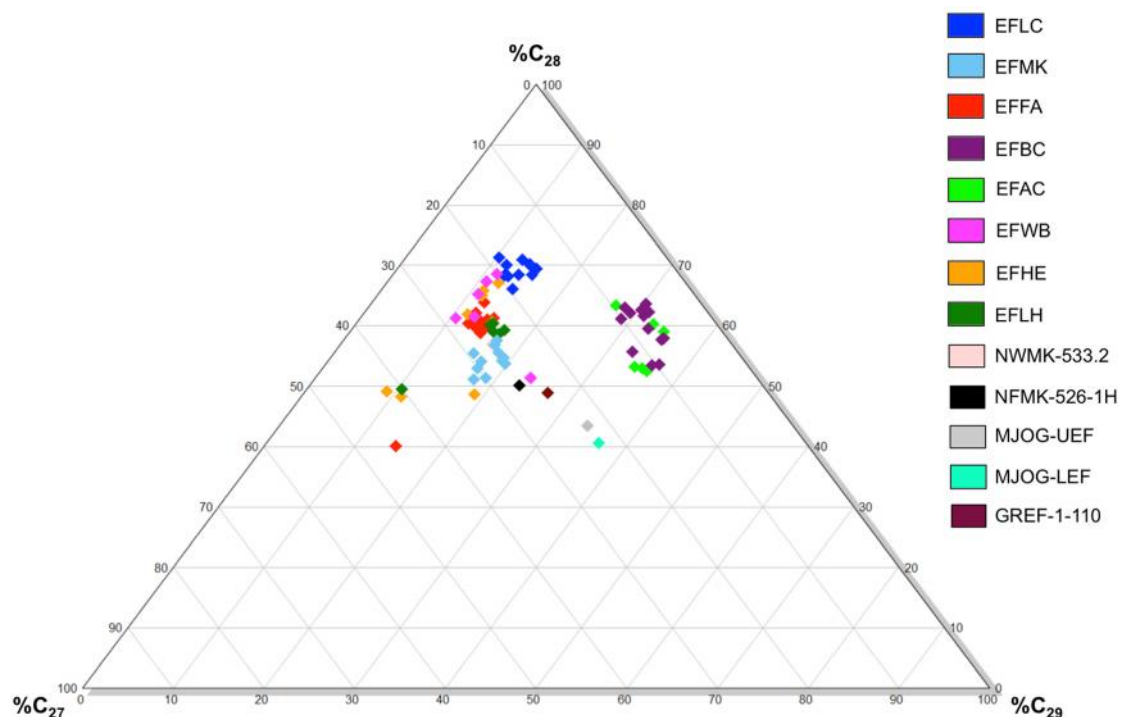


Figure 36. Ternary diagram of C₂₇, C₂₈, and C₂₉ monoaromatic steroids for Eagle Ford Shale bitumens and oils (Plot modified from Moldowan et al., 1985)

2.3 Biomarkers and geochemical parameters for evaluation of thermal maturity

2.3.1 Steranes

Steroid precursors in nature have only an R configuration at C-20, that with increasing maturation are progressively converted to mixture of R (biological) and S (thermal) isomers (Peters et al., 2005). As a result, geochemical ratios of biomarker isomers provide a useful tool for thermal maturity assessment. For example, the C₂₉ 20S/(20S+20R) steranes ratio increases with thermal maturity from 0 to about 0.5, reaching equilibrium at 0.52 to 0.55 (Seifert and Moldowan, 1986; Peters et al., 2005). Organic facies changes, weathering, and biodegradation may affect this maturity

parameter (Peters et al., 2005). The $C_{29} \text{ } \alpha / (\alpha + \beta)$ steranes ratio is not influenced by organic matter source, and its racemic mixture is reached at higher maturity levels compared to the $C_{29} \text{ } 20S/(20S+20R)$ steranes. With increasing thermal maturation isomerization at C-14 and C-17 in the $C_{29} \text{ } 20S$ and $20R$ regular steranes increases the $\alpha / (\alpha + \beta)$ ratio from near-zero values to about 0.7. Equilibrium reaction for this conversion is reached at about 0.67 to 0.71 (Peters et al., 2005). Comparison of the $C_{29} \text{ } \alpha / (\alpha + \beta)$ versus $C_{29} \text{ } 20S/(20S+20R)$ sterane ratios are commonly used to evaluate thermal maturation of oils and bitumens and to perform quality control of these geochemical parameters (Figure 37; Seifert and Moldowan, 1986; Peters et al., 2005). Calculation of these maturity ratios for the Eagle Ford Shale (Appendix A) indicates that all samples are thermally mature, with the exception of samples EFBC and EFAC, which are immature (Table 9; Appendix H).

It is possible that the high $C_{29} \text{ } 20S/(20S+20R)$ sterane ratios for the EFLC samples are attributed to weathering. Clayton and King (1987) found that weathering in shale outcrops produce preferential loss of the $C_{29} \text{ } 20S$ sterane diastereomer. In addition, it has been suggested that this ratio also decreases in high maturity samples (Peters et al., 1990), which could explain why the EFFA samples appear to have low maturity, even though other thermal maturity parameters describe them as high maturity source rocks.

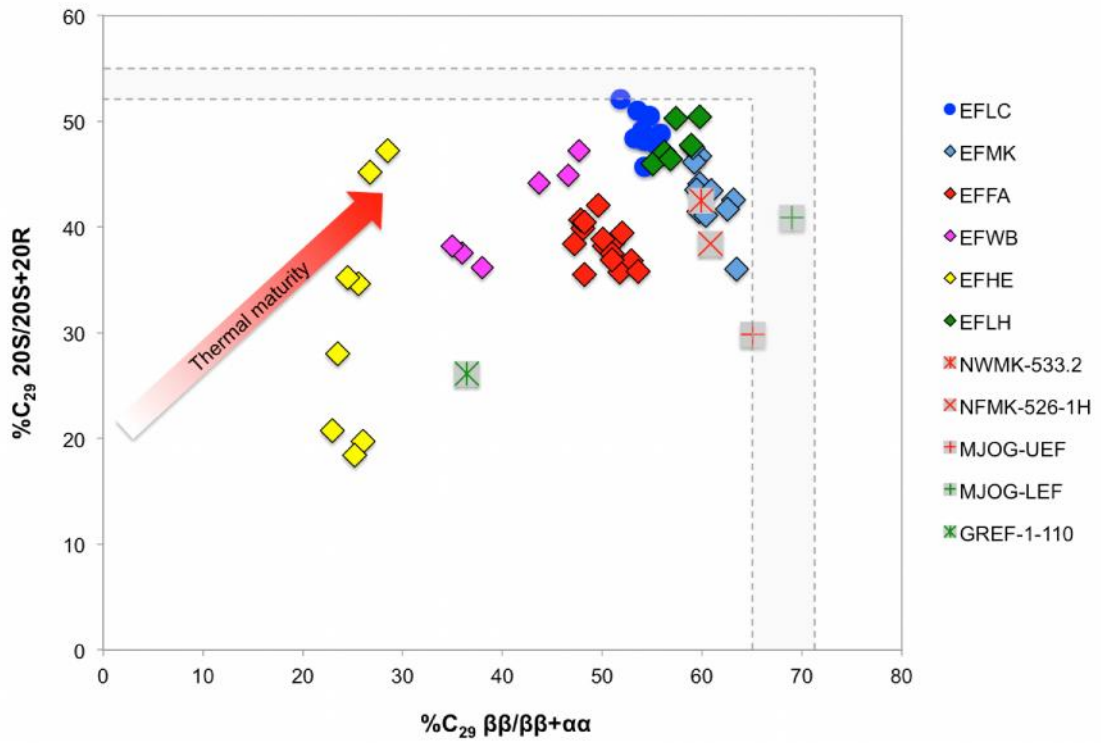


Figure 37. Plot of $C_{29} \beta\beta / (\beta\beta + \alpha\alpha)$ steranes versus $C_{29} \text{ 20S}/(20S+20R)$ steranes showing variations in thermal maturity for the Eagle Ford Shale source rocks and liquids (Gray area represents end points of isomerization reactions. Plot modified from Peters, 1999)

Table 9. Average steranes isomerization ratios for the Eagle Ford Shale bitumens and liquids (ND = not determined; *not averaged)

Sample	# of samples	C ₂₉ 20S/(20S+20R)	C ₂₉ /(+)
EFLC	12	0.49	0.54
EFMK	11	0.43	0.61
EFFA	15	0.38	0.50
EFBC	13	0.01	0.34
EFAC	6	ND	0.26
EFWB	6	0.41	0.41
EFHE	8	0.31	0.25
EFLH	6	0.48	0.57
NWMK-533.2 OIL	*	0.42	0.60
NFMK-526-1H OIL	*	0.38	0.61
MJOG-UEF OIL	*	0.30	0.65
MJOG-LEF OIL	*	0.41	0.69
GREF-1-110 COND.	*	0.26	0.36

2.3.2 Terpanes

2.3.2.1 Hopanes

A maturity parameter based on epimer ratios of 17 -homohopanes is the C₃₁ 22S/(22S+22R) ratio. The biological configuration of the homohopanes is the 22R, and during the early stages of organic matter maturation, isomerization at the C-22 position occurs, gradually converting homohopanes to a mixture of 22R and 22S configurations (Peters and Moldowan, 1991; Peters et al., 2005). The 22S/(22S+22R) ratio increases from 0 to ~0.6 reaching equilibrium at ratios of 0.57 to 0.62 (early oil window) after which no additional thermal maturity information can be obtained (Peters et al., 2005). Samples from the EFBC outcrop and the EFAC core have greater concentrations of the

22R diastereomer and 22S/(22S+22R) ratios of ~0.2 and ~0.3 respectively, pointing towards the low maturity of these samples (Appendices I and M). The bitumens and liquids analyzed from the other outcrops and cores have average 22S/(22S+22R) ratios greater than 0.50, indicating that these samples have reached the main stage of oil generation (Table 10).

Table 10. Average C₃₁ 22S/(22S+22R) hopane ratios for Eagle Ford Shale bitumens and oils (*not averaged)

Sample	# of samples	C ₃₁ 22S/(22S+22R)
EFLC	12	0.61
EFMK	11	0.53
EFFA	15	0.56
EFBC	13	0.23
EFAC	6	0.32
EFWB	6	0.59
EFHE	8	0.58
EFLH	6	0.58
NWMK-533.2 OIL	*	0.50

2.3.2.2 *Moretanes*

Hopanoids in organisms have a 17 (H),21()H-configuration (), which becomes unstable during organic matter burial and easily converts to -moretanes and -hopanes. However, C₂₉ and C₃₀ moretanes are less stable than hopanes and their concentrations decrease relative to hopanes with increasing maturity. As a result, the ratio moretanes/hopanes is used to evaluate thermal maturity for immature to early oil window samples (Peters et al., 2005). This ratio decreases from approximately 0.8 (immature) to <0.15 (minimum 0.05) with increasing maturation (Peters et al., 2005).

High moretanes/hopanes ratios have been associated with hypersaline source rocks (Rullkötter and Marzi, 1988), and in transgressive to highstand systems tracts, with significant terrigenous organic matter input (Isaken and Bohacs, 1995), which suggests that this ratio may also be influenced by depositional environment (Peters et al., 2005). Ratios of C₃₀ moretanes/hopanes calculated for Eagle Ford Shale samples have average values lower than 0.19 (Table 11, Appendix I) indicating that all samples with exception of EFBC and EFAC are thermally mature.

Table 11. Average C₃₀ Moretane/Hopane ratios for Eagle Ford Shale bitumens and oils (*not averaged)

Sample	# of samples	C ₃₀ Moretanes/Hopanes
EFLC	12	0.06
EFMK	11	0.16
EFFA	15	0.19
EFBC	13	1.57
EFAC	6	1.57
EFWB	6	0.13
EFHE	8	0.14
EFLH	6	0.12
NWMK-533.2 OIL	*	0.07

2.3.2.3 *Ts and Tm*

C₂₇ 17 -22,29,30-trisnorhopane (Tm) and C₂₇ 18 -22,29,30-trisnorneohopane (Ts) are terpanes frequently used to assess maturation from immature to postmature stage (Peters et al., 2005). Tm becomes less stable than Ts with increasing thermal maturity; therefore, the Ts/(Ts+Tm) ratio can be used as a maturity indicator,

particularly for samples of similar lithology and organic facies (Moldowan et al., 1994; Peters et al., 2005).

Ts and Tm can be correlated with other parameters to evaluate thermal maturity and depositional environment. For example, the relationship between the Ts/(Ts+Tm) ratio and the diasteranes/(diasteranes+regular) C₂₇ steranes ratio for the Eagle Ford Shale samples reveals three different groups (Figure 38). The first group clusters samples from EFLH, EFMK, EFFA and oils, which are affected by high thermal maturity, indicated by the high Ts/(Ts+Tm) ratios. In the second group, correlation of these ratios gathers samples EFWB, EFHE, and EFLC, suggesting that these bitumens were derived from low maturity, anoxic carbonate source rocks. The third group encompasses four EFLC samples. Ts/(Ts+Tm) ratios for these samples indicate low maturity; however, the diasteranes/(diasteranes+regular) C₂₇ steranes ratios are slightly higher for these samples compared to the second group, suggesting that the rocks that sourced these bitumens are more clay-rich. In fact, three of these samples correspond to the Scott Ranch Member, in which Donovan et al. (2013) identified several bentonite beds. In case of the EFBC samples, the low diasteranes/(diasteranes+regular) C₂₇ steranes ratios could be a result of their low maturity, instead of their lithology. Moreover, in samples from the immature EFAC core, which is closely located to the EFBC outcrop, no C₂₇ diasteranes could be identified.

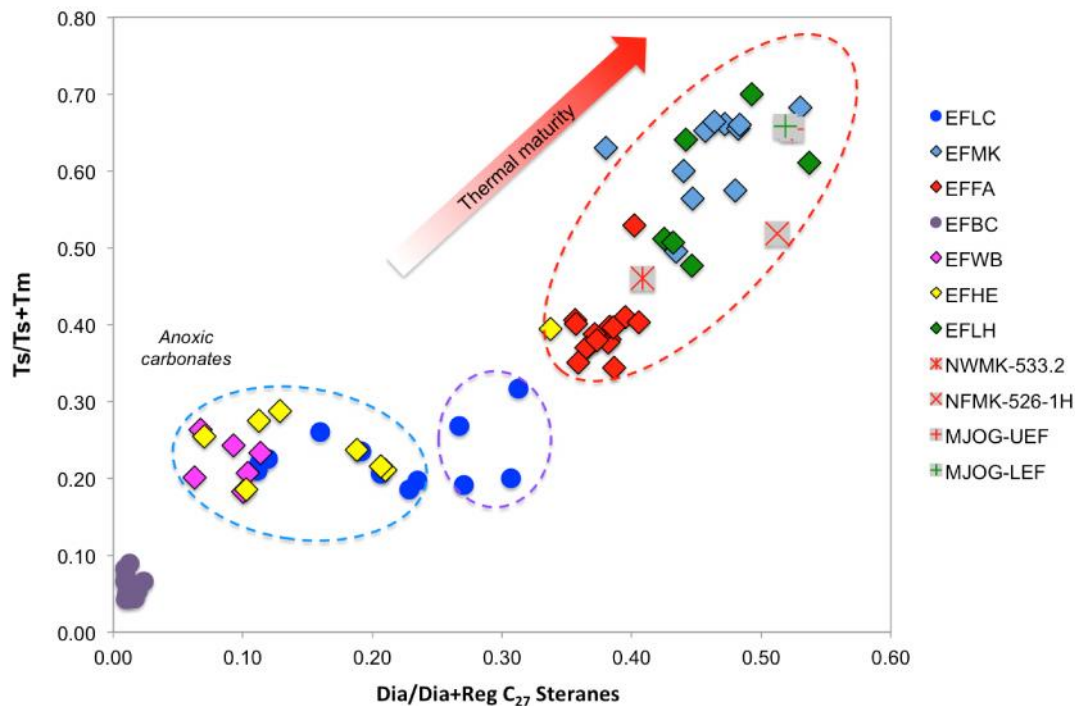


Figure 38. Plot of diasteranes/(diasteranes+regular) C_{27} Steranes versus $Ts/(Ts+Tm)$ ratios shows differences in thermal maturity, source, and redox potential for Eagle Ford Shale samples (Plot modified from Moldowan et al., 1994).

2.3.3 Monoaromatic and Triaromatic Steroids

Monoaromatic and triaromatic steroids were determined in the B&C and aromatic fractions of Eagle Ford source rock extracts and oil samples, respectively. Identification was carried out by analysis of ions m/z 231.3 and 253.3 through SIM/GC-MS. Selected fragmentograms of MAS and TAS are presented in Figures 35 and 39. Peak identifications are in Table 12. Generation of ABC-ring triaromatic steroids (TA) from A-ring monoaromatic (MA) steroids occurs during maturation by aromatization of B and C rings and the loss of both, a methyl group at the A/B junction and the stereocenter at C-5 (Mackenzie, 1984). The ratio $TA_{28}/(MA_{29}+TA_{28})$ increases from 0

to 100% during maturation and provides thermal maturity information from immature to early oil window ($R_o \sim 0.8\%$; Mackenzie, 1984; Peters et al., 2005). The ratios MA(I)/MA(I+II) and TA(I)/TA(I+II) are additional geochemical parameters used to evaluate thermal maturity from immature to late oil generation window ($R_o \sim 1.4\%$; Peters et al., 2005). These ratios increase from 0 to 100% with increase in thermal maturity. This increase may occur due to conversion of long- to short-chain aromatic steroids, preferential thermal degradation of the long- versus short-chain compounds, or both. However, since the TAS are derived from the MAS, the TA(I)/TA(I+II) ratio is more sensitive at high levels of thermal maturity compared to the MAS ratio (Peters et al., 2005).

MAS and TAS ratios (Appendix K) obtained for the Eagle Ford Shale samples indicate that the EFLC, EFBC and EFAC samples are thermally immature as suggested by the other maturity parameters previously described. In addition, these aromatic steroids ratios indicate that samples EFWB and EFHE are marginally mature, while samples EFLH, EFMK, EFFA, and oils are mature, possibly peak oil to late oil window.

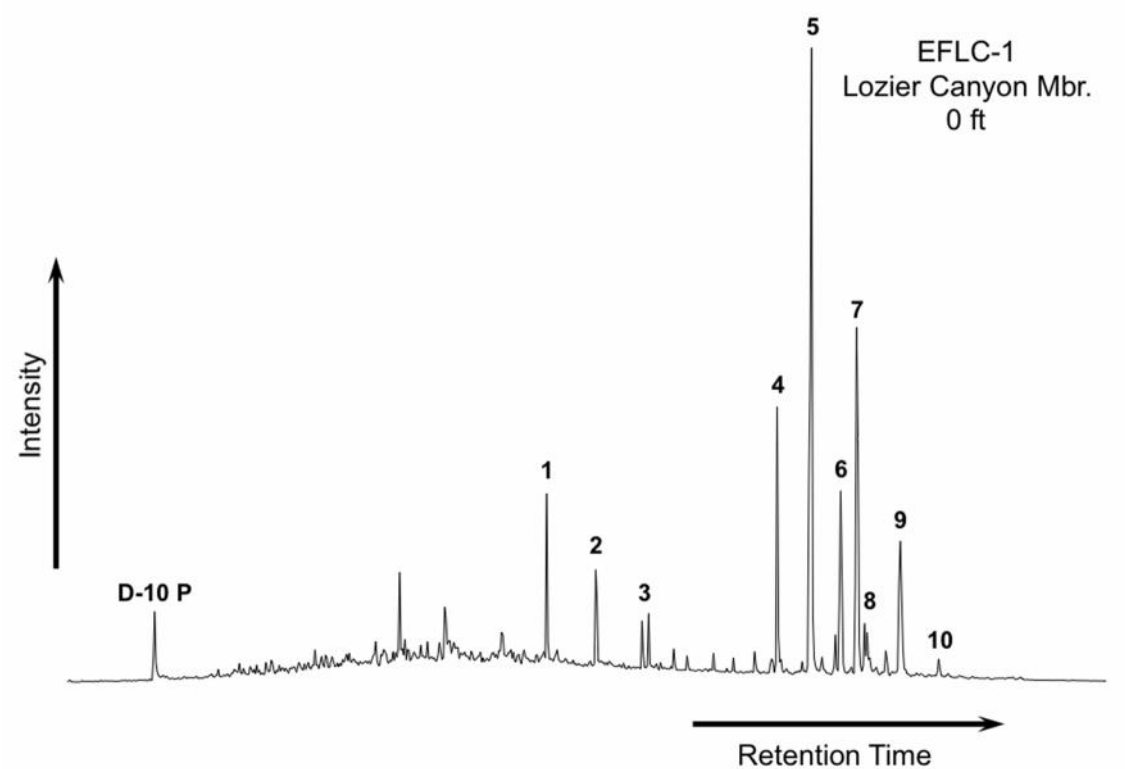


Figure 39. Partial fragmentograms of the m/z 231.3 ion showing distribution of the triaromatic steroids (TAS) in the aromatic fractions of sample EFLC-1. Peak identification is presented in Table 12

Table 12. Identification of triaromatic steroids (TAS) in the partial m/z 231.3 fragmentogram of the aromatic fractions

Peak #	Compound
D-10 P	Deuterated Phenanthrene (ISTD)
1	C ₂₀ Pregnane
2	C ₂₁ 20-Methylpregnane
3	C ₂₂ 20-Ethylpregnanes (a and b are epimeric at C ₂₀)
4	C ₂₆ Cholestane (20S)
5	C ₂₆ Cholestane (20R) + C ₂₇ Ergostane (20S)
6	C ₂₈ Stigmastane (20S) (24-Ethylcholestane 20S)
7	C ₂₇ Ergostane (20R) (24-Methylcholestane 20R)
8	C ₂₉ 24- <i>n</i> -Propylcholestane (20S) (a and b are epimeric at C ₂₀)
9	C ₂₈ Stigmastane (20R)
10	C ₂₉ 24- <i>n</i> -Propylcholestane (20R)

2.3.4 Phenanthrenes

Phenanthrenes, methylphenanthrenes, and dimethylphenanthrenes were identified in the aromatic fraction of bitumens and oils by SIM/GC-MS through the analysis of ions m/z 178.3, 192.3, and 206.3 (Figure 40, Table 13). Results from quantitation of phenanthrenes compounds are in Appendix P. A number of ratios derived from the distribution of these compounds have been used as geochemical maturity parameters (Radke and Welte, 1981; Radke et al., 1984; 1986). In particular, the Methylphenanthrene Indices (MPI-1 and MPI-2) are the most commonly applied. With increasing temperature and burial, an increase of the 2- and 3-methylphenanthrenes relative to the 1- and 9-methylphenanthrenes occurs (Radke et al., 1982). Radke and Welte (1981) recognized a positive correlation between MPI-1 and vitrinite reflectance for samples within the oil window (0.65-1.35%R_o), and a negative correlation for samples of high thermal maturity (1.35-2.00%R_o). From these

observations, the authors derived two equations for R_o calculation, which can be used to estimate thermal maturity for bitumens and oils (Radke and Welte, 1981; Appendix A).

Values of MPI-1 and MPI-2 ratios were obtained for the Eagle Ford Shale samples (Appendix K), and from the MPI-1 parameter, vitrinite reflectance was calculated (Table 14). The equation from Radke and Welte (1981) corresponding to low maturity levels was used for calculating vitrinite reflectance from all samples. Overall, the results are in agreement with the maturity parameters previously described which indicate that most of the Eagle Ford Shale samples analyzed are immature to peak oil window ($R_c \sim 0.58-0.93$).

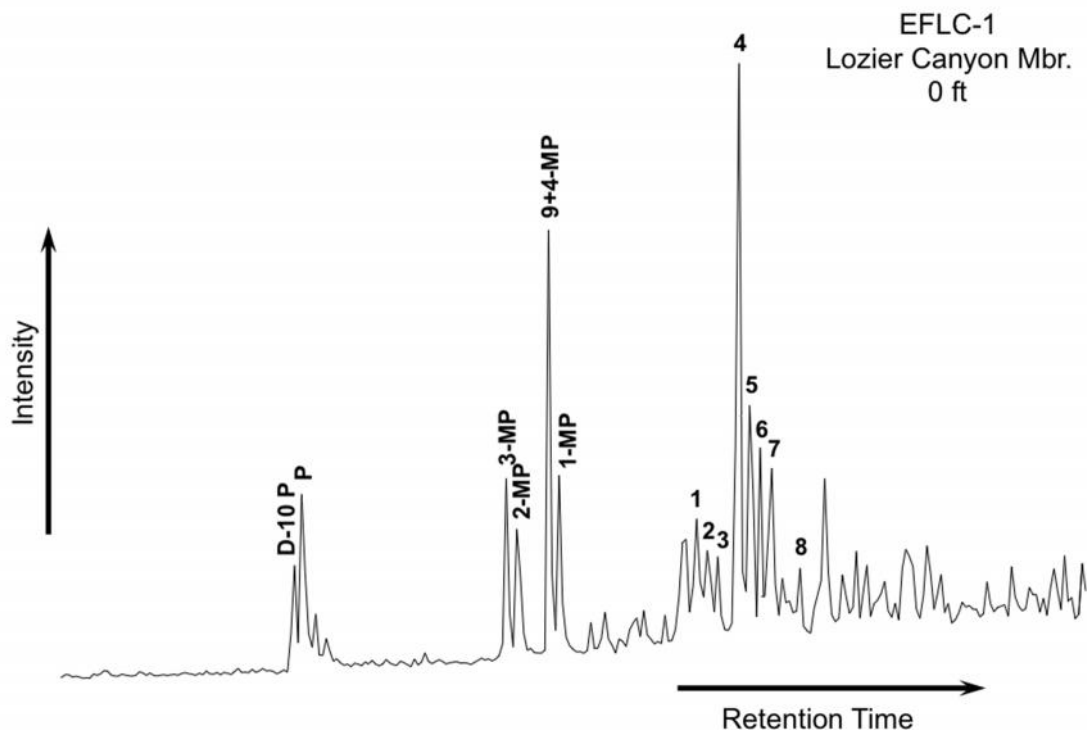


Figure 40. Summed mass chromatograms of m/z 106.3, 192.3, 206.3 ions showing distribution of phenanthrenes compounds in the aromatic fractions of sample EFLC-1. Peak identification is presented in Table 13

Table 13. Identification of phenanthrenes in the partial mass chromatograms of m/z 178.3, 192.3, 206.3 of the aromatic fractions

Peak #	Compound
D-10 P	Deuterated Phenanthrene (ISTD)
P	Phenanthrene
3-MP	3-Methylphenanthrene
2-MP	2-Methylphenanthrene
9+4-MP	9+4-Methylphenanthrene
1-MP	1-Methylphenanthrene
1	Dimethylphenanthrene
2	Dimethylphenanthrene
3	Dimethylphenanthrene
4	Dimethylphenanthrene
5	Dimethylphenanthrene
6	Dimethylphenanthrene
7	Dimethylphenanthrene
8	Dimethylphenanthrene

Table 14. Calculated vitrinite reflectance (R_c) for Eagle Ford Shale bitumens and oils from MPI-1 values

Sample	R_c
EFLC	0.58
EFMK	0.83
EFFA	0.93
EFBC	0.66
EFAC	0.63
EFWB	0.77
EFHE	0.74
EFLH	0.72
NWMK-533.2	0.78
NFMK-526-1H	0.79
MJOG-UEF	0.84
MJOG-LEF	0.89
GREF-1-110	0.92

2.4 Isotope Analysis

Compound specific isotope analysis (CSIA) was performed on the saturate fractions of bitumens and oil samples. At least one sample from each Eagle Ford Shale member per well was selected for CSIA in order to address possible source and maturity variations. Carbon isotope values (^{13}C) for individual *n*-alkanes are reported in per mil (‰) relative to the Vienna Pee Dee Belemnite (VPDB) standard (Appendix Q). Figures 41 to 46 show the *n*-alkane isotope profiles for Eagle Ford Shale samples of different maturity. The carbon isotopic composition of all the samples, including oils and condensates varies from -32.6 to -29.9‰, and in general there is no significant variability on the isotope values for the *n*-alkanes in the range analyzed.

Considering that all the analyzed samples essentially come from the same self-sourced unconventional reservoir, factors such as source rock age, hydrocarbon mixing, and reservoir alteration processes do not have an influence on the isotopic values of the bitumens and liquids analyzed. Therefore, any observed changes in the isotopic composition of the samples would be expected to result from source and thermal maturity variations. Overall, the isotopic profiles for the Eagle Ford Shale samples do not show marked changes in the ^{13}C values. The range of ^{13}C values and the apparently flat and uniform isotopic profiles are in agreement with a marine origin for the samples analyzed (Murray et al., 1994). Even though biomarker data suggest a terrigenous organic matter input for the EFLH well, the *n*-alkanes isotope profiles do not reflect these variations in source. Enrichment in the overall ^{13}C values of *n*-alkanes along with a ^{13}C depletion in the long-chain *n*-alkanes range is commonly observed in samples with terrigenous organic matter contribution (Murray et al., 1994; Dowling et

al., 1995), a characteristic that is not detected in the Eagle Ford Shale samples analyzed from the EFLH core. These observations suggest there are no significant variations in organic facies or depositional environments among the different Eagle Ford Shale members. Likewise, despite the fact that the samples analyzed have different thermal maturity levels, the isotopic data do not reflect variations regarding this effect. This is an interesting observation since usually hydrocarbons become enriched in ^{13}C values with increasing maturity (Clayton, 1991). The similarity in the ^{13}C values between the bitumens and the analyzed liquids indicate that the oils and condensates were generated from the Eagle Ford Shale.

Unfortunately, the lack of published literature regarding CSIA for marine shales using an unconventional reservoir analysis approach prevents making comparisons and correlations of the isotopic data obtained in this research with analog settings.

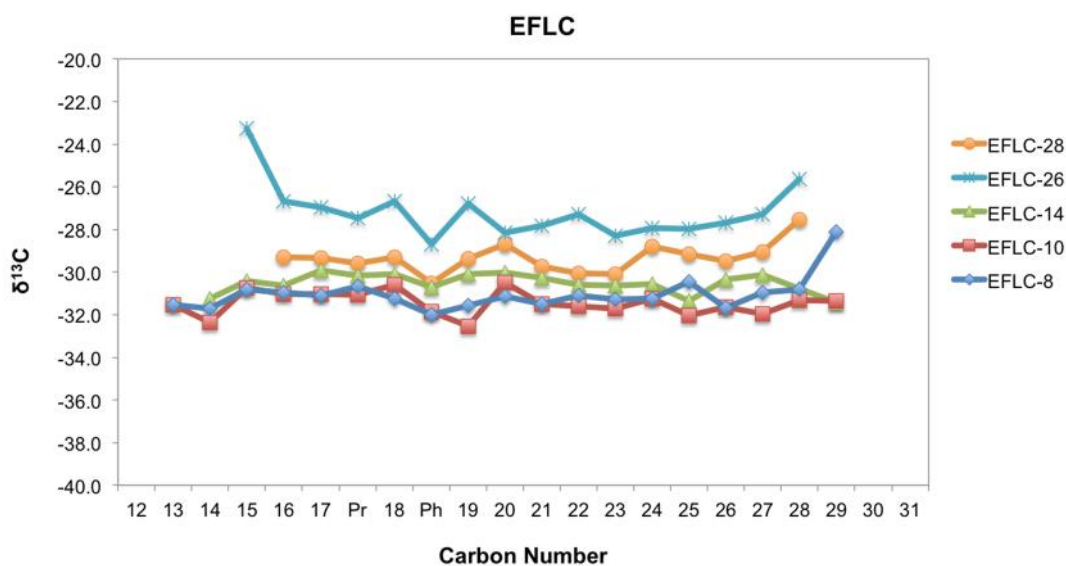


Figure 41. *n*-Alkane isotope profile from the EFLC extracts (immature)

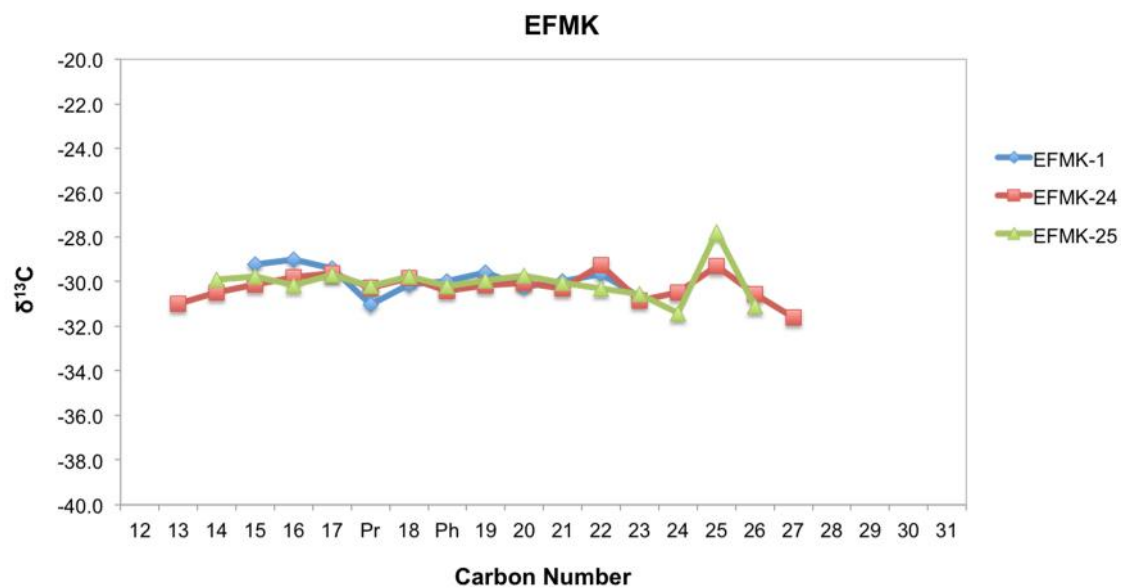


Figure 42. *n*-Alkane isotope profile from the EFMK extracts (oil window)

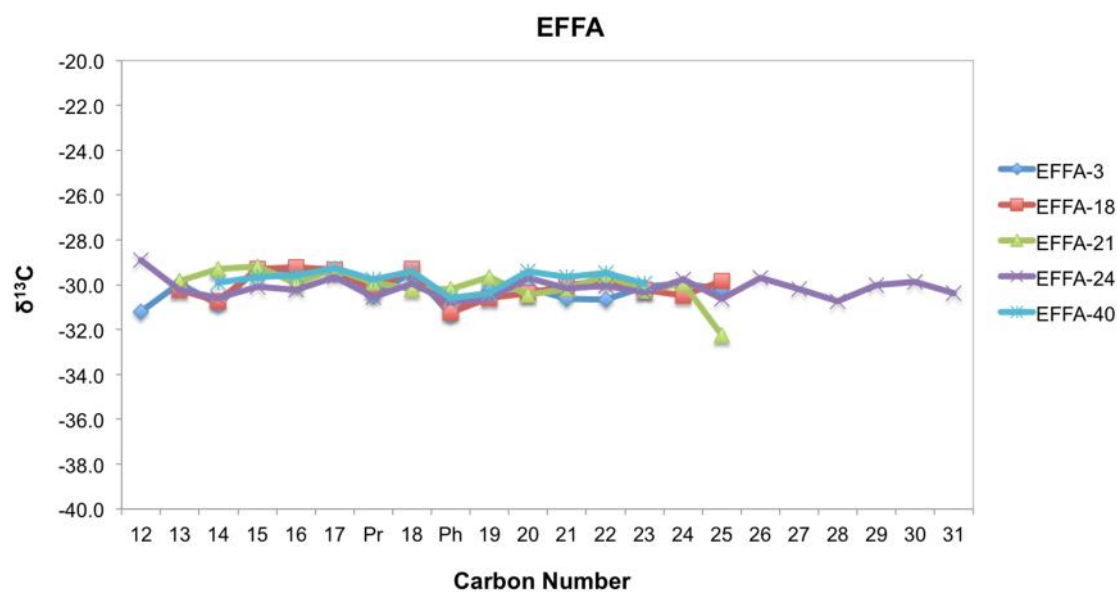


Figure 43. *n*-Alkane isotope profile from the EFFA extracts (peak oil window)

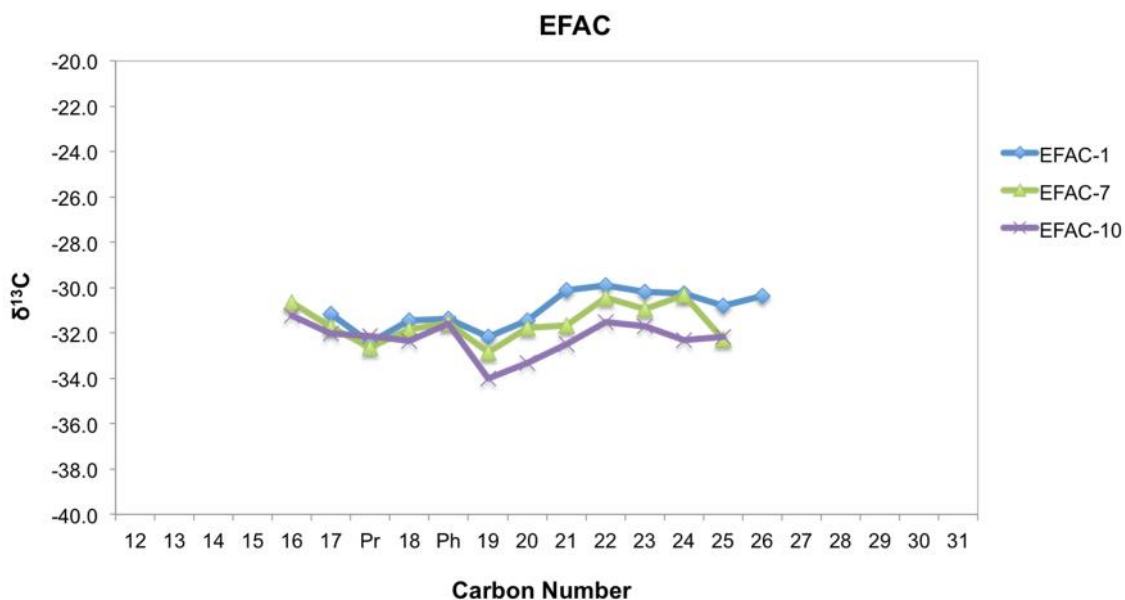


Figure 44. *n*-Alkane isotope profile from the EFAC extracts (immature)

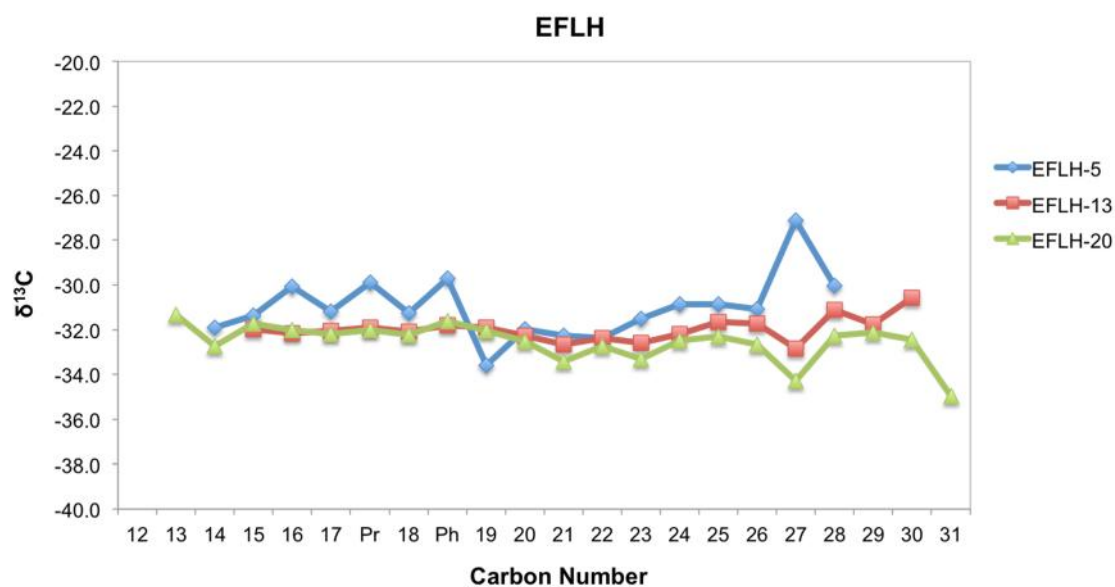


Figure 45. *n*-Alkane isotope profile from the EFLH extracts (oil window)

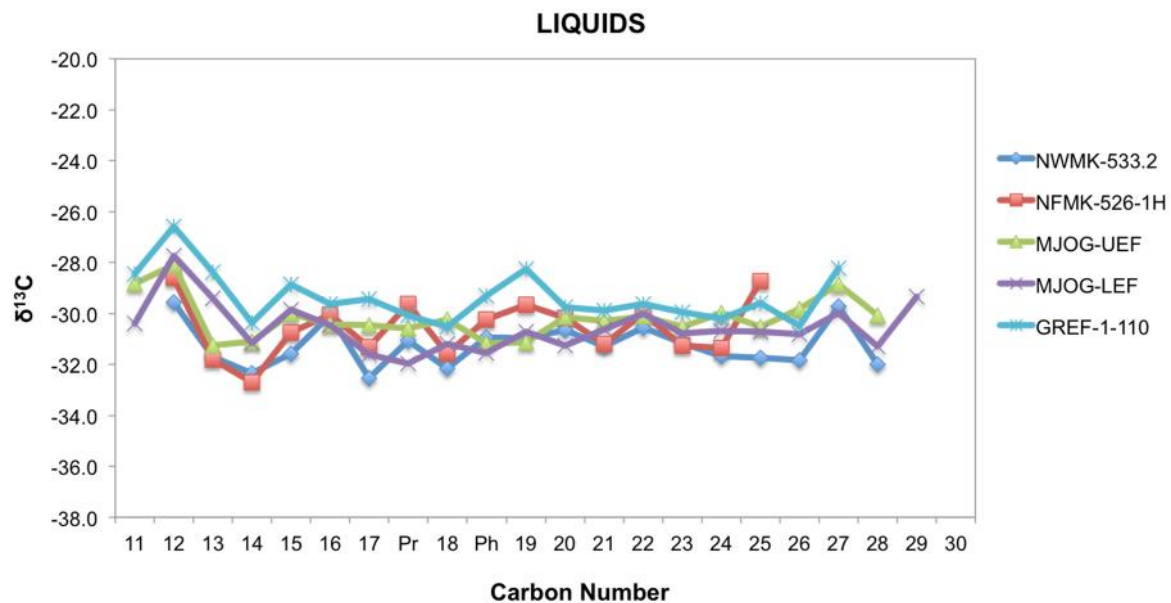


Figure 46. *n*-Alkane isotope profile from oils and condensate samples (condensate/wet gas window)

3. CHAPTER III

3.1 Regional and local organic geochemistry of the Eagle Ford Shale

Geochemical analyses of the Eagle Ford Shale from outcrop and core samples reveal minimal variations in the organic matter type and environmental conditions during deposition of the different Eagle Ford Shale Group members. Detailed geochemical logs of the biomarker ratios calculated for each outcrop, core, and region analyzed in this study are presented in Appendices F through H. Numerical values of these ratios are presented in Appendices I through K. Preliminary analyses of the Eagle Ford Shale samples indicate a similar genetic relationship and generally do not show significant changes in source and depositional settings. Despite the fact that these samples occur in areas separated by important geological structures, they share similar organic geochemical characteristics that point out their common origin. Based on TOC and Rock Eval parameters, the Eagle Ford Shale samples show organic matter quality and good source rock potential. These source rocks are dominated by Type II kerogen and were deposited under marine carbonate anoxic conditions with minimal terrigenous organic matter input, as indicated by low TAR and CPI values. The exception to the latter occurs in the EFLH core (East Texas) where various geochemical parameters, such as high TAR and CPI ratios, Pr/n-C₁₇ and Ph/n-C₁₈ values, low C₃₁R/30H, and low 30-NorH/30H ratios indicate these sediments were possibly influenced by terrigenous material and deposited under suboxic-oxic conditions. This organic matter source variation in East Texas, with respect to the other locations, may be result of the presence of the deltaic depositional system that influenced sedimentation of the Woodbine-Eagle Ford Shale units. Biomarker parameters reflecting the marine organic

matter source for the Eagle Ford Shale sediments include the steroids distributions, and high steranes/17 α -hopanes, C₂₃ tricyclic terpane/C₃₀ hopane, and C₃₁R/30H ratios.

Generally, the Eagle Ford Shale was deposited under anoxic conditions as indicated by their low Pr/Ph values, and the high concentrations of C₃₅ homohopanes relative to their C₃₄ homologs. The presence of aryl isoprenoids suggests the occurrence of PZA conditions during source rock deposition, although AIR values indicate that these conditions were intermittent, pointing towards periodic shifts of the chemocline. The predominance of even numbered *n*-alkanes and steranes, high pregnane/sterane ratios, and tentative identification of gammacerane reveal that the Eagle Ford Shale probably experienced hypersaline conditions and water-column stratification, particularly in West and East Texas. This observation suggests that these regions were possibly more restricted compared to Central Texas.

The Eagle Ford Shale is immature to marginally mature in the EFLC, EFBC, and EFAC wells (West and Central Texas), as determined by T_{max} values, Ts and Tm ratios, and steranes and hopanes isomer ratios. These same thermal maturity parameters indicate that in the EFMK and EFFA wells (Southwest Texas) the Eagle Ford Shale is in the oil window. Based on these maturity parameters and measured R_o, the Eagle Ford Shale is in the main stage of oil generation at the EFLH well (East Texas).

Organic facies variations within the Eagle Ford Shale were determined through correlation of bulk geochemical parameters and biomarker ratios calculated for each member. Tables 15 and 16 show average values of these parameters for each Eagle Ford Shale member for every geographic location considered. Limited data availability for the East Texas EFLH core prevented identification of the different Eagle Ford Shale

members. However, a general characterization of the Lower Eagle Ford Shale based on the same geochemical parameters was carried out on this location. Geochemical data obtained from the EFLC outcrop was the only one considered for the evaluation of vertical variations in West Texas due to the low maturity of these samples compared to the EFMK and EFFA cores.

Across the three regions considered, the Lower Eagle Ford Shale has the highest TOC and HI values. In particular, the Lozier Canyon Member is the most organic-rich (TOC>4.0%; HI>270) followed by the Antonio Creek Member (Table 15). In Central Texas, all the Eagle Ford Shale members show high HI values, suggesting their good source rock potential. Depositional conditions of the different Eagle Ford Shale members seem to have been fairly constant in the West Texas area. Low Pr/Ph ratios suggest the prevalence of anoxic conditions throughout the entire deposition of the Eagle Ford Shale in this area. In addition, variations in Pr/Ph ratios show a positive correlation with AIR. This relationship indicates that variations in PZA occurred during deposition of these units, particularly in the upper interval of the Lozier Canyon Member. Relatively high Pr/Ph ratios and AIR suggest deposition of this member under episodic PZA. This interval contains the greatest amount of organic carbon throughout the analyzed section, indicating the possibility of significant organic matter production during this time. This observation may be supported by a marked increase in phosphorous concentration within the Lozier Canyon Member (at the contact between Donovan et al.'s, 2013 A and B facies; Aris Pramudito, 2014 personal communication), which suggests an increase in nutrient availability that could have promoted production. In Central Texas, Pr/Ph ratios suggest stronger anoxic depositional conditions for the

Lozier Canyon Member compared to the Antonio Creek and Langtry members, whereas deposition of the Lower Eagle Ford Shale in East Texas possibly took place under oxic-suboxic conditions. These observations are supported by the $C_{35}S/C_{34}S$ ratios, which are also consistent with a marine carbonate depositional environment for the Eagle Ford Shale. The presence of pregnanes suggests the establishment of hypersaline conditions during deposition of the Lower Eagle Ford Shale, particularly in the East and West Texas areas, although an enrichment of these biomarkers in the EFMK, EFFA, and EFLH samples might be related to their higher thermal maturity.

Despite the observed variations in depositional conditions, changes in organic matter source were minimal. The C_{23} tricyclic terpane/ C_{30} hopane and C_{24}/C_{23} tricyclic terpane ratios suggest that the Lower Eagle Ford Shale Formation was deposited in a marine carbonate environment, whereas the Upper Eagle Ford Shale Formation might have received some terrigenous input. This observation correlates with the diasteranes/(diasteranes+regular) C_{27} steranes ratio, which indicates a progressive increase in siliciclastic contribution during deposition of the Antonio Creek and the Upper Eagle Ford Shale members. This ratio shows a relationship with the facies description of Donovan et al. (2012, 2013), which refers to significant bentonite beds in the Antonio Creek, Scott Ranch, and Langtry members. These minimal variations in organic matter source are further confirmed by isotopic data. Isotopic profiles are consistent with a marine organic input for the Eagle Ford Shale; however, ^{13}C values do not show significant differences that could be related to organic facies or depositional environment changes. Furthermore, the isotope data analyzed do not reflect the variations in thermal maturity among the different samples.

Donovan et al., (2012, 2013) developed a sequence stratigraphic framework for the Eagle Ford Shale Group at the Lozier Canyon outcrop (EFLC) and the Fasken “A” #1 well (EFFA). The frameworks at these locations were used to determine possible correlations between organic geochemical parameters and the different stratigraphic sequences proposed by these authors. It was observed that low Pr/Ph, AIR, C₂₄ tricyclic terpanes and C₃₁R/30H ratios; and high TOC, HI, and C₂₃ tricyclic terpanes occurred during the highstand systems tract (HST), whereas the contrary was observed at the transgressive systems tract (TST). However, these interpretations need to be further refined by a high-resolution study in order to determine if these variations are consistent within a sequence stratigraphic context for shale gas systems.

Table 15. Average of bulk and *n*-alkanes geochemical parameters for the Eagle Ford Shale members across West, Southwest, Central and East Texas (ND = not determined)

Eagle Ford Formation	WEST TEXAS				CENTRAL TEXAS				EAST TEXAS
	Lower Eagle Ford Shale		Upper Eagle Ford Shale		Lower Eagle Ford Shale		Upper Eagle Ford Shale		Lower Eagle Ford Shale
	Lozier Canyon	Antonio Creek	Scott Ranch	Langtry	Lozier Canyon	Antonio Creek	Scott Ranch	Langtry	
TOC	4.22	1.34	1.25	0.21	5.34	3.17		3.02	4.18
T_{max}	524	434	381	109	421	423		360	443
HI	278	255	106	62	505	412		407	270
OI	19	51	31	166	32	57	NOT PRESENT	102	8
Pr/Ph	0.33	0.28	0.39		0.84	1.19		1.16	1.44
TAR	2.44	2.37	0.59	ND	0.41	ND		ND	0.36
CPI	1.07	0.96	0.74		0.85	0.73		0.74	1.07

Table 16. Average biomarker ratios for the Eagle Ford Shale members across West, Southwest, Central and East Texas (ND = not determined)

Eagle Ford Formation	WEST TEXAS				CENTRAL TEXAS				EAST TEXAS
	Lower Eagle Ford Shale		Upper Eagle Ford Shale		Lower Eagle Ford Shale		Upper Eagle Ford Shale		Lower Eagle Ford Shale
	Lozier Canyon	Antonio Creek	Scott Ranch	Langtry	Lozier Canyon	Antonio Creek	Scott Ranch	Langtry	
Eagle Ford Shale Member									
C₂₇% reg. steranes	39.17	37.59	42.78		39.13	37.56		39.32	44.74
C₂₈% reg. steranes	28.00	30.22	24.21		34.74	37.72		35.27	29.89
C₂₉% reg. steranes	32.83	32.19	33.01		26.13	24.72		25.41	25.37
C₂₉ 20S/(20S + 20R) steranes	0.50	0.49	0.48		0.19	0.18		0.12	0.47
C₂₉ /(+) steranes	0.54	0.55	0.54	ND	0.32	0.35		0.30	0.57
Pregnane/Steranes	0.64	0.73	0.61		0.21	0.26		0.16	1.18
C₂₇ Steranes (Dia/(Dia + Reg))	0.15	0.22	0.30		0.07	0.06		0.06	0.47
Sterane Index	0.05	0.06	0.05		0.05	0.05		0.06	0.07
Steranes/17 -Hopanes	0.49	0.63	0.60		1.02	1.86	NOT	2.05	0.60
Tricyclics/17 -Hopanes	0.05	0.05	0.03		0.04	0.04	PRESENT	0.04	0.09
C₂₄ T/C₂₃ T	0.42	0.33	0.56		0.55	0.46		0.60	0.49
C₂₄ Tetra/C₃₀ Hopane	0.06	0.06	0.04		0.06	0.14		0.05	0.06
C₂₃ T/C₃₀ Hopane	0.54	0.50	0.21		0.20	0.50		0.19	0.26
C₃₁ 22S/(22S+22R)	0.61	0.61	0.62	ND	0.38	0.35		0.43	0.58
Moretane/C₃₀ Hopane	0.07	0.06	0.06		0.97	0.90		0.98	0.12
Ts/Ts+Tm	0.24	0.21	0.24		0.14	0.14		0.14	0.57
C₃₁R/C₃₀ Hopane	0.46	0.54	0.46		0.52	0.81		0.48	0.21
C₃₅S/C₃₄S	1.40	1.45	1.12		0.86	0.92		0.96	0.39
C₂₉/C₃₀ Hopane	0.79	0.75	0.60		1.04	0.89		0.90	0.49

4. CONCLUSIONS

The Eagle Ford Shale Group is a source rock with excellent potential as demonstrated by TOC and Rock Eval parameters. It is dominated by Type II kerogen indicating a marine organic matter input. Biomarker distributions of regular steranes and monoaromatic steroids (MAS), high steranes/17 α -hopanes, C₂₃ tricyclic terpane/C₃₀ hopane, C₂₄/C₂₃ tricyclic terpane, and C₃₁R/C₃₀ hopane ratios are consistent with a marine carbonate depositional environment for the Eagle Ford Shale sediments. However, high terrigenous-to-aquatic (TAR) and carbon preference index (CPI) ratios, low C₃₁R/30H, and low 30-NorH/30H ratios suggest that in East Texas, the Eagle Ford Shale was possibly sourced by terrigenous material, reflecting the influence of the Harris Delta during deposition of these sediments. Low Pr/Ph values, and high C₃₅S/C₃₄S homohopanes suggest that these source rocks were deposited under anoxic conditions. The presence of aryl isoprenoids and the aryl isoprenoids ratio (AIR) suggest the occurrence of intermittent photic zone anoxia (PZA). The predominance of even numbered *n*-alkanes and steranes, high pregnanes/steranes ratios, and the tentative identification of gammacerane suggest the establishment of hypersaline conditions during Eagle Ford Shale deposition particularly in West and East Texas.

Thermal maturity parameters such as T_{max}, Ts and Tm ratios, and steranes and hopanes isomer ratios indicate that in West and Central Texas the Eagle Ford Shale is immature to marginally mature. These parameters show a progressive increase in thermal maturity towards the southeast, from the Lozier Canyon site in Terrell County (early oil window) to the Fasken "A" #1 well in Webb County (dry gas window). Based

on measured R_o values it was determined that in East Texas, the Eagle Ford Shale is in the main stage of oil generation.

Analysis of geochemical logs suggests a narrow range of vertical variation within the Eagle Ford Shale Group. The Lower Eagle Ford Shale Formation has the highest TOC and HI values and in particular, the Lozier Canyon Member is the most organic-rich. Low Pr/Ph ratios suggest the prevalence of stronger anoxic conditions during deposition of the Lower Eagle Ford Shale compared to the Upper Eagle Ford Shale Formation, where the latter may have received an additional siliciclastic and terrigenous organic matter input. Isotope data also points to a marine organic input for the Eagle Ford Shale, but ^{13}C values do not show significant differences that could be related to organic facies, depositional environment, or thermal maturity changes.

Correlation of geochemical data with an already established sequence stratigraphic framework for the Eagle Ford Shale Group in West and Central Texas allowed observation of organic geochemical trends and stratigraphic sequences. Low Pr/Ph, AIR, C_{24} tricyclic terpanes and $\text{C}_{31}\text{R}/30\text{H}$ ratios; and high TOC, HI, and C_{23} tricyclic terpanes intervals were associated with the highstand systems tract (HST), whereas opposite trends were observed within the transgressive systems tract (TST). These observations result in a general interpretation and it is recommended that a high-resolution organic geochemistry study be performed preferably at the same locations. Higher sampling density from fresh outcrops and/or cores may help enhance lower-order variations in biomarker concentrations, geochemical ratios, and isotopic compositions that could improve the use of organic geochemistry in sequence stratigraphic studies.

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6. APPENDIX

A. Abbreviations and formulas used for calculation of geochemical biomarker ratios

n-Alkanes

Carbon Preference Index (CPI)

$$CPI = \frac{1}{2} \left[\frac{C_{25} + C_{27} + C_{29} + C_{31} + C_{33}}{C_{26} + C_{28} + C_{30} + C_{32} + C_{34}} + \frac{C_{25} + C_{27} + C_{29} + C_{31} + C_{33}}{C_{24} + C_{26} + C_{28} + C_{30} + C_{32}} \right]$$

Terrigenous/Aquatic Ratio (TAR):

$$TAR = \frac{C_{27} + C_{29} + C_{31}}{C_{15} + C_{17} + C_{19}}$$

Long- versus Short-chain *n*-alkanes:

$$\text{long-/short-chain } n\text{-alkanes} = \frac{\sum(C_{21} - C_{31})}{\sum(C_{15} - C_{21})}$$

Steranes

C₂₇%, **C₂₈%**, **C₂₉%** = C₂₇, C₂₈, C₂₉ [14 (H),17 (H)- + 14 (H),17 (H)-Cholestane (20S+ 20R)]

C₂₉ 20S/(20S + 20R) = C₂₉ [14 (H),17 (H)-Cholestane (20S)]/[14 (H),17 (H)-Cholestane (20S+20R)]

C₂₉ / (+) = C₂₉ [14 (H),17 (H)-Cholestane (20S+20R)]/[14 (H),17 (H)- + 14 (H),17 (H)-Cholestane (20S+20R)]

Preg/Ster = C₂₁ 14 (H),17 (H)-Pregnane/C₂₇ 14 (H),17 (H)-Cholestane (20R)

C₂₇ Dia/C₂₇ Sterane = [C₂₇ 13 (H),17 (H)-Diacholestane (20S+20R)]/[C₂₇ 14 (H),17 (H)- + 14 (H),17 (H)-Cholestane (20S+20R)]

C₂₇ Dia/(Dia+Reg) = [C₂₇ 13 (H),17 (H)- + 13 (H),17 (H)-Diacholestane (20S+20R)]/[C₂₇ 13 (H),17 (H)- + 13 (H),17 (H)-Diacholestane (20S+20R) + C₂₇ 14 (H),17 (H)- + 14 (H),17 (H)-Cholestane (20S+20R)]

Sterane Index = C₃₀/(C₂₇-C₃₀) = [C₃₀ 14 (H),17 (H)- + 14 (H),17 (H)-Cholestane (20S+20R)]/[C₂₇, C₂₈, C₂₉, C₃₀ [14 (H),17 (H)- + 14 (H),17 (H)-Cholestane (20S+20R)]]

Hopanes

Sterane/17 -Hopanes = [C₂₇, C₂₈, C₂₉, C₃₀ 14 (H),17 (H)- + 14 (H),17 (H)-Cholestane (20S+20R)]/[C₂₉ 17 (H),21 (H)-30-Norhopane + C₃₀ 17 (H),21 (H)-Hopane + C₃₁+C₃₂+C₃₃ 17 (H),21 (H)-Homohopanes (22S+22R)]

TR/17 -H = C₂₈ + C₂₉ Tricyclic terpanes (20S+20R)/17 -Hopanes

C₂₂/C₂₁TR = C₂₂/C₂₁ Tricyclic terpanes

C₂₄/C₂₃TR = C₂₄/C₂₃ Tricyclic terpanes

C₂₆/C₂₅TR = C₂₆/C₂₅ Tricyclic terpanes

C₂₃TR/30H = C₂₃ Tricyclic terpanes/C₃₀ 17 (H),21 (H)-Hopane

C₂₄TT/30H = C₂₄ Tetracyclic terpane/C₃₀ 17 (H),21 (H)-Hopane

30-NorH/30H = C₂₉ 17 (H),21 (H)-30-Norhopane/C₃₀ 17 (H),21 (H)-Hopane

Mor/30H = C₃₀ 17 (H),21 (H)-Moretane/C₃₀ 17 (H),21 (H)-Hopane

31R/30H = C₃₁ 17 (H),21 (H)-Homohopane (22R)/C₃₀ 17 (H),21 (H)-Hopane

35S/34S = C₃₅ 17 (H),21 (H)-Pentakishomohopane (22S)/C₃₄ 17 (H),21 (H)-Tetrakishomohopane (22S)

HH Index = C₃₅/(C₃₁ - C₃₅) = C₃₅ 17 (H),21 (H)-Pentakishomohopane (22S+22R)/[C₃₁+C₃₂+C₃₃+C₃₄+C₃₅ 17 (H),21 (H)-Homohopanes (22S+22R)]

C31 22S/22S+22R = C₃₁ [17 (H),21 (H)-Homohopanes (22S)]/17 (H),21 (H)-Homohopanes (22S+22R)]

Aryl Isoprenoids

Aryl Isoprenoid ratio (AIR):

AIR = (C₁₃-C₁₇)/(C₁₈-C₂₂) 2,3,6-trimethyl substituted aryl isoprenoids

Aromatics

MPI-1:

$$MPI\ 1 = 1.5 \times \frac{[2 - MP + 3 - MP]}{[P + 1 - MP + 9 - MP]}$$

Calculated vitrinite reflectance:

For 0.65 to 1.35%R_o: R_c = 0.60 MPI-1 + 0.40

For 1.35 to 2.00%R_o: R_c = -0.60 MPI-1 + 2.30

MPI-2:

$$MPI\ 2 = 3 \times \frac{[2 - MP]}{P + 1 - MP + 9 - MP}$$

MA(I):

$$MA(I) = \sum C_{27} - C_{29} MAS$$

MA(II):

$$MA(II) = \sum C_{21} - C_{22} MAS$$

TA(I):

$$TA(I) = \sum C_{26} - C_{28} (20S + 20R) TAS$$

TA(II):

$$TA(II) = \sum C_{20} - C_{21} TAS$$

C26TA 20S/20S+20R = C₂₆ Cholestane 20S/20S+20R (Triaromatic steroids)

TA28 = C₂₈ Stigmastane (20S+20R) (Triaromatic steroids)

MA29 = C₂₉ [5 - + 5 -Stigmastane (20S+20R)]

C27R/C28R TAS = C₂₇ Ergostane (20R)/C₂₈ Stigmastane (20R)

C26S/C28S TAS = C₂₆ Cholestane (20S)/C₂₈ Stigmastane (20S)

%C₂₇, %C₂₈, %C₂₉ MAS = C₂₇, C₂₈, C₂₉ MAS

B. Total Organic Carbon (TOC) and Rock Eval (RE) parameters for the Eagle Ford Shale samples

Lozier Canyon outcrop

Sample	Depth (ft)	TOC	S ₁	S ₂	S ₃	T _{max}	HI	OI	S ₂ /S ₃	S ₁ /TOC	PI	S ₁ + S ₂
EFLC-43	187.5	0.04	0.02	0.01	0.05	0	25	125	0	50	0.67	0.03
EFLC-42	182.5	0.05	0.02	0.08	0.11	0	160	220	1	40	0.20	0.10
EFLC-41	180.0	0.04	0.03	0.05	0.07	0	125	175	1	75	0.38	0.08
EFLC-40	175.0	0.07	0.04	0.04	0.37	0	57	529	0	57	0.50	0.08
EFLC-39	170.0	0.08	0.02	0.03	0.06	0	38	75	1	25	0.40	0.05
EFLC-38	165.0	0.06	0.03	0.14	0.06	0	233	100	2	50	0.18	0.17
EFLC-37	160.0	0.11	0.02	0.05	0.09	0	45	82	1	18	0.29	0.07
EFLC-36	155.0	0.06	0.04	0.03	0.25	0	50	417	0	67	0.57	0.07
EFLC-35	150.0	0.06	0.01	0.01	0.21	0	17	350	0	17	0.50	0.02
EFLC-34	145.0	0.07	0.04	0.01	0.23	0	14	329	0	57	0.80	0.05
EFLC-33	144.5	0.89	0.03	1.18	1.41	433	133	158	1	3	0.02	1.21
EFLC-32	142.0	0.57	0.06	0.93	0.92	438	163	161	1	11	0.06	0.99
EFLC-31	140.0	0.05	0.02	0.01	0.09	0	20	180	0	40	0.67	0.03
EFLC-30	139.5	0.48	0.07	0.67	0.27	437	140	56	2	15	0.09	0.74
EFLC-29	135.0	0.07	0.03	0.03	0.11	0	43	157	0	43	0.50	0.06
EFLC-28	130.0	2.07	0.27	8.77	0.64	437	423	31	14	13	0.03	9.04
EFLC-27	125.0	1.68	0.34	7.76	0.60	436	462	36	13	20	0.04	8.10
EFLC-26	121.5	2.92	0.45	16.52	0.91	432	566	31	18	15	0.03	16.97
EFLC-25	116.5	1.84	0.15	8.92	0.65	434	485	35	14	8	0.02	9.07
EFLC-24	110.0	0.78	0.08	2.18	0.40	437	279	51	5	10	0.04	2.26
EFLC-23	106.0	0.08	0.08	0.05	0.17	0	63	213	0	100	0.62	0.13
EFLC-22	100.0	0.29	0.11	0.56	0.24	438	193	83	2	38	0.16	0.67

Lozier Canyon outcrop (cont.)

Sample	Depth (ft)	TOC	S ₁	S ₂	S ₃	T _{max}	HI	OI	S ₂ /S ₃	S ₁ /TOC	PI	S ₁ + S ₂
EFLC-21	95.0	0.30	0.05	0.39	0.28	437	130	93	1	17	0.11	0.44
EFLC-20	89.0	1.80	0.30	8.43	0.55	433	468	31	15	17	0.03	8.73
EFLC-19	85.0	1.40	0.16	5.60	0.43	434	400	31	13	11	0.03	5.76
EFLC-18	80.5	1.40	0.17	5.73	0.50	435	409	36	11	12	0.03	5.90
EFLC-17	75.0	0.63	0.02	1.48	0.29	436	235	46	5	3	0.01	1.50
EFLC-16	70.0	0.77	0.05	1.67	1.31	434	217	170	1	6	0.03	1.72
EFLC-15	65.0	1.03	0.08	1.31	2.34	437	127	227	1	8	0.06	1.39
EFLC-14	60.0	0.92	0.12	4.19	0.51	434	455	55	8	13	0.03	4.31
EFLC-13	55.0	2.19	0.19	13.07	0.92	430	597	42	14	9	0.01	13.26
EFLC-12	50.0	1.93	0.78	11.91	0.76	430	617	39	16	40	0.06	12.69
EFLC-11	45.0	4.32	1.01	26.69	1.22	431	618	28	22	23	0.04	27.70
EFLC-10	40.0	6.32	3.10	41.11	0.85	430	650	13	48	49	0.07	44.21
EFLC-9	35.0	5.47	2.54	33.86	0.98	431	619	18	35	46	0.07	36.40
EFLC-8	30.0	6.29	1.87	39.88	0.92	431	634	15	43	30	0.04	41.75
EFLC-7	25.0	6.05	2.64	39.61	1.13	430	655	19	35	44	0.06	42.25
EFLC-6	20.0	4.58	1.07	28.08	1.12	431	613	24	25	23	0.04	29.15
EFLC-5	15.0	5.91	2.03	39.05	1.95	430	661	33	20	34	0.05	41.08
EFLC-4	10.0	2.79	0.55	13.68	1.78	431	490	64	8	20	0.04	14.23
EFLC-3	9.0	1.77	0.23	6.70	1.44	430	379	81	5	13	0.03	6.93
EFLC-2	5.0	0.54	0.05	2.47	0.32	430	457	59	8	9	0.02	2.52
EFLC-1	0.0	2.36	0.66	18.27	0.52	428	774	22	35	28	0.03	18.93

Ferguson McKnight #526-1H core

Sample	Depth (ft)	TOC	S ₁	S ₂	S ₃	T _{max}	HI	OI	S ₂ /S ₃	S ₁ /TOC	PI	S ₁ + S ₂
EFMK-1	5793.0	1.02	3.01	1.90	0.33	436	186	32	6	295	0.61	4.91
EFMK-2	5801.0	1.23	2.56	2.59	0.49	443	211	40	5	208	0.50	5.15
EFMK-3	5809.0	1.83	3.83	4.08	0.45	440	223	25	9	209	0.48	7.91
EFMK-4	5817.5	2.79	3.28	5.93	0.51	445	213	18	12	118	0.36	9.21
EFMK-5	5825.5	1.32	2.69	3.43	0.40	440	260	30	9	204	0.44	6.12
EFMK-6	5833.0	0.57	1.01	0.76	0.49	443	134	87	2	178	0.57	1.77
EFMK-7	5841.0	0.69	1.54	1.02	0.43	443	147	62	2	222	0.60	2.56
EFMK-8	5849.0	1.70	2.44	3.68	0.60	446	216	35	6	144	0.40	6.12
EFMK-9	5859.0	1.70	4.86	4.06	0.63	442	239	37	6	286	0.54	8.92
EFMK-10	5864.5	2.18	4.37	4.63	0.68	447	212	31	7	200	0.49	9.00
EFMK-11	5873.5	1.76	2.39	3.76	0.61	450	214	35	6	136	0.39	6.15
EFMK-12	5881.0	3.06	3.93	5.88	0.68	453	192	22	9	128	0.40	9.81
EFMK-13	5889.0	2.33	2.91	5.19	0.56	450	223	24	9	125	0.36	8.10
EFMK-14	5897.5	2.44	3.33	5.55	0.45	449	227	18	12	136	0.38	8.88
EFMK-15	5905.5	2.45	3.71	5.66	0.45	448	231	18	13	151	0.40	9.37
EFMK-16	5913.5	1.91	4.33	5.16	0.46	441	270	24	11	227	0.46	9.49
EFMK-17	5921.0	1.14	2.67	2.12	0.53	447	186	46	4	234	0.56	4.79
EFMK-18	5929.0	1.76	3.44	3.47	0.62	447	197	35	6	195	0.50	6.91
EFMK-19	5937.5	1.66	4.01	3.36	0.61	447	202	37	6	242	0.54	7.37
EFMK-20	5945.5	2.24	4.30	4.73	0.52	442	211	23	9	192	0.48	9.03
EFMK-21	5953.0	1.94	4.13	4.42	0.49	443	228	25	9	213	0.48	8.55
EFMK-22	5961.0	2.78	6.12	6.05	0.67	449	218	24	9	220	0.50	12.17
EFMK-23	5969.0	2.28	6.66	5.23	0.59	442	229	26	9	292	0.56	11.89
EFMK-24	5977.5	2.84	7.04	6.40	0.82	447	225	29	8	248	0.52	13.44
EFMK-25	6072.0	3.42	6.76	7.85	1.00	444	230	29	8	198	0.46	14.61
EFMK-26	6174.0	4.99	8.78	11.87	0.73	444	238	15	16	176	0.43	20.65

Fasken "A" #1 core

Sample	Depth (ft)	TOC	S ₁	S ₂	S ₃	T _{max}	HI	OI	S ₂ /S ₃	S ₁ /TOC	PI	S ₁ + S ₂
EFFA-1	9430.5	3.20	0.08	0.28	0.30	553	9	9	1	3	0.22	0.36
EFFA-2	9440.0	1.20	0.09	0.13	0.32	593	11	27	0	7	0.41	0.22
EFFA-3	9450.0	3.72	0.16	0.41	0.31	588	11	8	1	4	0.28	0.57
EFFA-4	9460.5	1.28	0.14	0.17	0.31	587	13	24	1	11	0.45	0.31
EFFA-5	9470.5	2.30	0.11	0.25	0.88	587	11	38	0	5	0.31	0.36
EFFA-6	9479.5	2.18	0.11	0.18	0.24	588	8	11	1	5	0.38	0.29
EFFA-7	9489.0	2.66	0.14	0.29	0.23	599	11	9	1	5	0.33	0.43
EFFA-8	9499.0	1.16	0.18	0.15	0.32	600	13	28	0	16	0.55	0.33
EFFA-9	9509.0	2.14	0.09	0.18	0.27	590	8	13	1	4	0.33	0.27
EFFA-10	9519.5	1.82	0.15	0.20	0.39	599	11	21	1	8	0.43	0.35
EFFA-11	9530.0	1.89	0.11	0.16	0.28	600	8	15	1	6	0.41	0.27
EFFA-12	9540.0	2.07	0.10	0.14	0.35	601	7	17	0	5	0.42	0.24
EFFA-13	9550.0	3.09	0.11	0.25	0.35	593	8	11	1	4	0.31	0.36
EFFA-14	9560.5	2.16	0.10	0.15	0.31	598	7	14	0	5	0.40	0.25
EFFA-15	9570.5	3.32	0.18	0.37	0.31	599	11	9	1	5	0.33	0.55
EFFA-16	9578.5	3.71	0.17	0.37	0.31	596	10	8	1	5	0.31	0.54
EFFA-17	9589.0	2.70	0.15	0.30	0.27	597	11	10	1	6	0.33	0.45
EFFA-18	9599.5	3.70	0.20	0.39	0.30	599	11	8	1	5	0.34	0.59
EFFA-19	9610.0	4.30	0.16	0.35	0.34	598	8	8	1	4	0.31	0.51

Fasken "A" #1 core (cont.)

Sample	Depth (ft)	TOC	S ₁	S ₂	S ₃	T _{max}	HI	OI	S ₂ /S ₃	S ₁ /TOC	PI	S ₁ + S ₂
EFFA-21	9629.5	4.35	0.19	0.38	0.29	602	9	7	1	4	0.33	0.57
EFFA-22	9639.5	4.03	0.21	0.29	0.31	600	7	8	1	5	0.42	0.50
EFFA-23	9650.5	3.91	0.13	0.25	0.29	600	6	7	1	3	0.34	0.38
EFFA-24	9655.5	5.90	0.29	0.57	0.35	604	10	6	2	5	0.34	0.86
EFFA-25	9665.5	5.76	0.30	0.43	0.31	331	7	5	1	5	0.41	0.73
EFFA-26	9670.5	4.03	0.17	0.35	0.29	603	9	7	1	4	0.33	0.52
EFFA-27	9680.5	4.86	0.24	0.48	0.30	604	10	6	2	5	0.33	0.72
EFFA-28	9685.5	5.49	0.23	0.56	0.28	605	10	5	2	4	0.29	0.79
EFFA-29	9695.0	5.42	0.19	0.36	0.28	600	7	5	1	4	0.35	0.55
EFFA-30	9700.5	5.04	0.30	0.54	0.36	604	11	7	2	6	0.36	0.84
EFFA-31	9711.5	5.95	0.23	0.48	0.31	604	8	5	2	4	0.32	0.71
EFFA-32	9717.0	6.20	0.26	0.54	0.41	604	9	7	1	4	0.33	0.80
EFFA-33	9728.5	6.31	0.20	0.51	0.35	603	8	6	1	3	0.28	0.71
EFFA-34	9733.5	5.60	0.29	0.58	0.32	600	10	6	2	5	0.33	0.87
EFFA-35	9742.5	6.42	0.20	0.54	0.30	602	8	5	2	3	0.27	0.74
EFFA-36	9746.5	6.29	0.23	0.60	0.31	602	10	5	2	4	0.28	0.83
EFFA-37	9756.5	5.59	0.19	0.39	0.36	601	7	6	1	3	0.33	0.58
EFFA-38	9761.5	5.39	0.33	0.54	0.35	604	10	6	2	6	0.38	0.87
EFFA-39	9770.5	5.49	0.26	0.67	0.28	604	12	5	2	5	0.28	0.93
EFFA-40	9776.0	5.44	0.20	0.34	0.36	603	6	7	1	4	0.37	0.54

Bouldin Creek outcrop

Sample	Depth (ft)	TOC	S₁	S₂	S₃	T_{max}	HI	OI	S₂/S₃	S₁/TOC	PI	S₁ + S₂
EFBC_Aus2	34.0	0.06	0.01	0.01	0.47	0	17	783	0	17	0.50	0.02
EFBC-Aus	33.0	0.07	0.01	0.02	0.52	0	28	716	0	14	0.33	0.03
EFBC-1	32.0	0.07	0.01	0.01	0.42	0	15	615	0	15	0.50	0.02
EFBC-2	31.0	0.14	0.01	0.02	0.50	0	15	365	0	7	0.33	0.03
EFBC-3	30.0	0.19	0.02	0.07	0.59	0	37	316	0	11	0.22	0.09
EFBC-4	29.0	0.13	0.01	0.09	0.42	0	67	313	0	7	0.10	0.10
EFBC-5	28.0	3.68	0.35	15.99	2.01	423	435	55	8	10	0.02	16.34
EFBC-6	27.0	4.48	0.37	23.18	1.51	414	517	34	15	8	0.02	23.55
EFBC-7	26.0	3.69	0.17	15.92	2.72	422	431	74	6	5	0.01	16.09
EFBC-8	25.0	3.13	0.22	14.08	2.33	425	450	74	6	7	0.02	14.30
EFBC-9	24.0	4.15	0.30	20.70	2.64	420	499	64	8	7	0.01	21.00
EFBC-10	23.0	2.90	0.17	12.89	2.04	422	444	70	6	6	0.01	13.06
EFBC-11	22.0	4.68	0.27	20.67	2.62	422	442	56	8	6	0.01	20.94
EFBC-12	21.0	3.13	0.23	12.37	1.70	425	395	54	7	7	0.02	12.60
EFBC-13	20.0	3.85	0.23	18.58	1.66	417	483	43	11	6	0.01	18.81
EFBC-14	19.0	3.87	0.36	18.37	1.73	418	475	45	11	9	0.02	18.73
EFBC-15	18.0	3.84	0.32	17.29	2.64	420	450	69	7	8	0.02	17.61
EFBC-16	17.0	4.05	0.25	16.05	2.62	422	396	65	6	6	0.02	16.30
EFBC-17	16.0	3.13	0.21	12.41	2.31	424	396	74	5	7	0.02	12.62

Bouldin Creek outcrop (cont.)

Sample	Depth (ft)	TOC	S₁	S₂	S₃	T_{max}	HI	OI	S₂/S₃	S₁/TOC	PI	S₁ + S₂
EFBC-18	15.0	3.45	0.19	15.57	1.64	416	451	48	9	6	0.01	15.76
EFBC-19	14.0	2.14	0.16	7.74	1.07	421	362	50	7	7	0.02	7.90
EFBC-20	13.0	1.12	0.04	2.27	0.58	426	203	52	4	4	0.02	2.31
EFBC-21	12.0	0.17	0.02	0.05	0.60	0	30	355	0	12	0.29	0.07
EFBC-22	11.0	0.39	0.04	0.40	0.74	434	103	191	1	10	0.09	0.44
EFBC-23	10.0	1.35	0.08	3.49	0.87	422	259	64	4	6	0.02	3.57
EFBC-24	9.0	1.02	0.10	2.76	0.67	424	271	66	4	10	0.03	2.86
EFBC-25	8.0	4.83	0.27	20.09	2.18	414	416	45	9	6	0.01	20.36
EFBC-26	7.0	0.95	0.02	2.57	0.70	425	269	73	4	2	0.01	2.59
EFBC-27	6.0	2.02	0.08	6.79	1.06	416	336	52	6	4	0.01	6.87
EFBC-28	5.0	0.59	0.05	1.47	0.54	424	250	92	3	9	0.03	1.52
EFBC-29	4.0	0.91	0.05	2.00	0.86	430	220	95	2	6	0.02	2.05
EFBC-30	3.0	5.95	0.46	35.91	1.56	418	604	26	23	8	0.01	36.37
EFBC-31	2.0	7.22	0.67	48.76	1.66	416	675	23	29	9	0.01	49.43
EFBC-32	1.0	7.13	0.77	48.63	1.51	416	682	21	32	11	0.02	49.40
EFBC-33	0.0	5.59	0.53	37.70	1.42	411	674	25	27	9	0.01	38.23
EFBC-34	-1.0	7.06	1.04	50.59	1.57	412	717	22	32	15	0.02	51.63

ACC#1 core

Sample	Depth (ft)	TOC	S ₁	S ₂	S ₃	T _{max}	HI	OI	S ₂ /S ₃	S ₁ /TOC	PI	S ₁ + S ₂
EFAC-1	80.0	2.12	0.07	6.34	1.18	428	299	56	5	3	0.01	6.41
EFAC-2	85.0	2.80	0.09	10.48	1.39	428	374	50	8	3	0.01	10.57
EFAC-3	90.0	4.04	0.17	17.86	1.71	421	442	42	10	4	0.01	18.03
EFAC-4	95.0	4.55	0.23	20.63	1.94	417	453	43	11	5	0.01	20.86
EFAC-5	100.5	3.88	0.12	14.58	1.76	422	376	45	8	3	0.01	14.70
EFAC-6	105.0	3.89	0.17	15.84	1.93	420	407	50	8	4	0.01	16.01
EFAC-7	110.0	5.12	0.28	26.27	2.00	421	513	39	13	5	0.01	26.55
EFAC-8	114.8	8.07	0.59	45.91	2.77	424	569	34	17	7	0.01	46.50
EFAC-9	120.0	1.20	0.04	0.44	0.71	417	37	59	1	3	0.08	0.48
EFAC-10	124.8	2.22	0.08	6.06	0.79	429	273	36	8	4	0.01	6.14

W. Brechtel #1 core

Sample	Depth (ft)	TOC	S ₁	S ₂	S ₃	T _{max}	HI	OI	S ₂ /S ₃	S ₁ /TOC	PI	S ₁ + S ₂
EFWB-1	3280.1	4.36	0.72	31.11	1.26	423	714	29	25	17	0.02	31.83
EFWB-2	3285.0	3.01	0.34	18.15	1.31	420	603	44	14	11	0.02	18.49
EFWB-3	3289.9	4.11	0.60	25.22	1.37	422	614	33	18	15	0.02	25.82
EFWB-4	3295.0	5.47	0.67	33.31	1.37	422	609	25	24	12	0.02	33.98
EFWB-5	3300.0	4.97	0.69	30.39	1.25	424	611	25	24	14	0.02	31.08
EFWB-6	3304.9	3.83	1.31	24.52	0.97	431	640	25	25	34	0.05	25.83
EFWB-7	3309.8	4.00	0.59	24.61	1.15	421	615	29	21	15	0.02	25.20
EFWB-8	3313.0	5.56	1.24	35.07	0.90	425	631	16	39	22	0.03	36.31
EFWB-9	3315.1	5.95	1.99	40.69	1.17	426	684	20	35	33	0.05	42.68

C.J. Hendershot #1 core

Sample	Depth (ft)	TOC	S₁	S₂	S₃	T_{max}	HI	OI	S₂/S₃	S₁/TOC	PI	S₁ + S₂
EFHE-1	4734.1	0.77	0.07	2.40	0.79	433	313	103	3	9	0.03	2.47
EFHE-2	4736.8	5.10	0.96	36.03	1.45	427	706	28	25	19	0.03	36.99
EFHE-3	4742.1	2.81	0.36	16.19	1.40	428	576	50	12	13	0.02	16.55
EFHE-4	4747.0	3.48	0.40	20.22	1.47	425	581	42	14	11	0.02	20.62
EFHE-5	4751.9	4.39	0.73	26.14	1.35	424	595	31	19	17	0.03	26.87
EFHE-6	4756.9	5.17	0.61	28.93	1.47	422	560	28	20	12	0.02	29.54
EFHE-7	4762.2	7.22	1.40	43.69	1.50	421	605	21	29	19	0.03	45.09
EFHE-8	4768.9	6.86	1.40	41.67	1.43	427	607	21	29	20	0.03	43.07
EFHE-9	4771.0	9.74	2.70	58.56	1.41	425	601	14	42	28	0.04	61.26
EFHE-10	4773.8	1.48	0.20	1.39	0.41	420	94	28	3	14	0.13	1.59

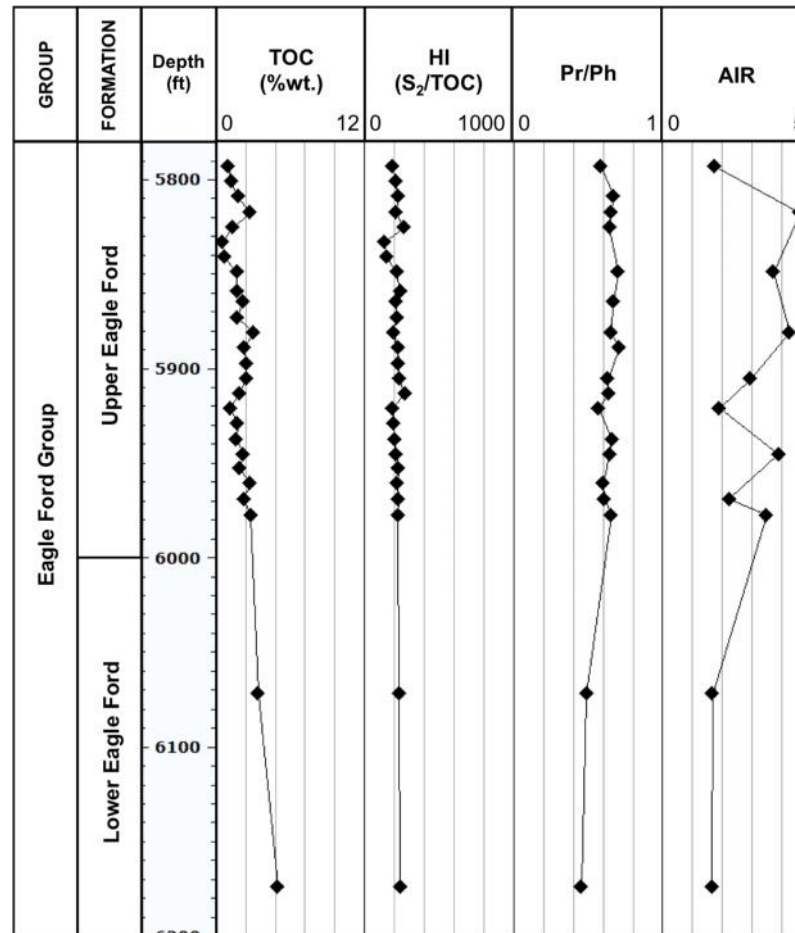
Lily Hoppess #1 core

159

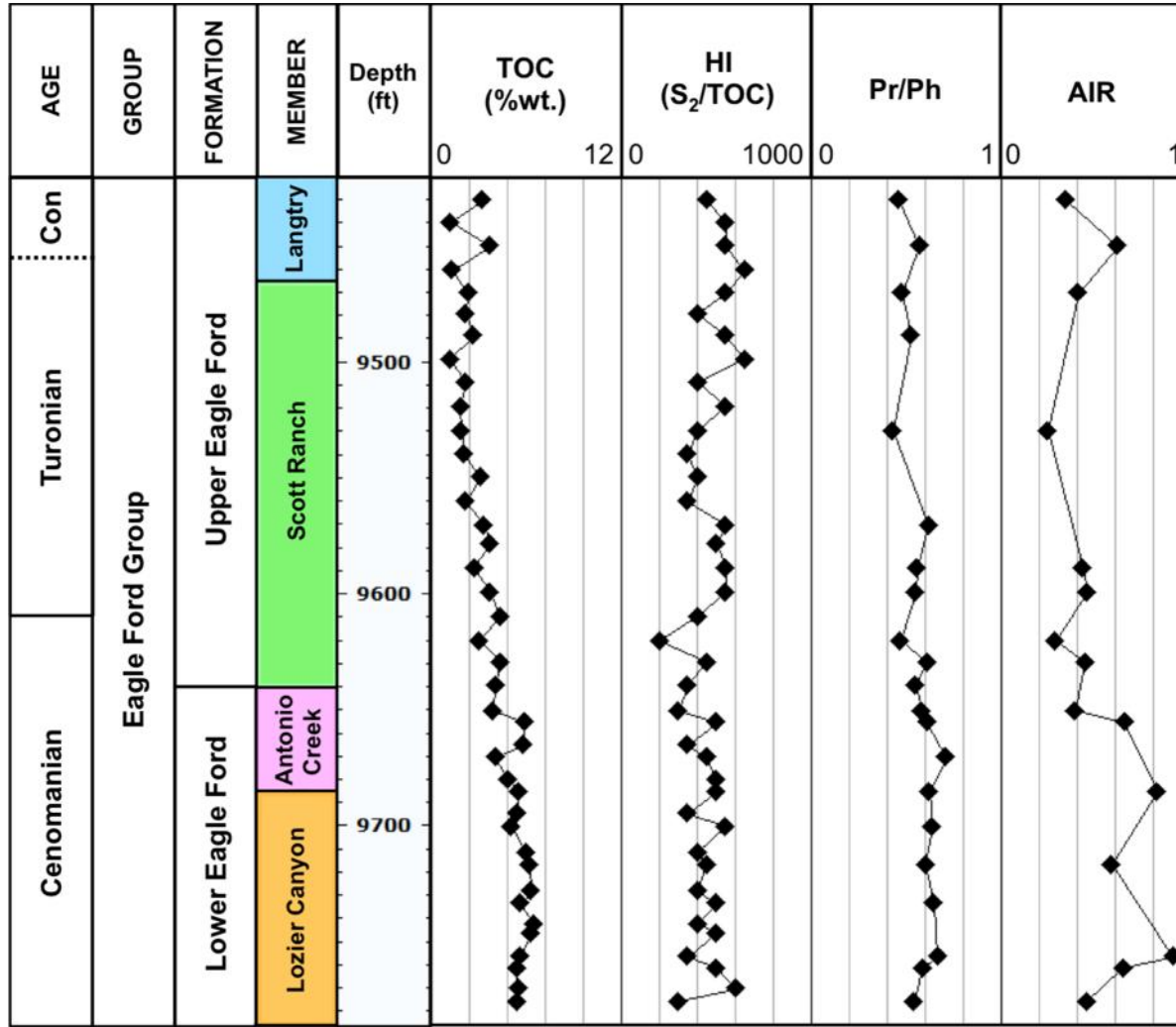
Sample	Depth (ft)	TOC	S ₁	S ₂	S ₃	T _{max}	HI	OI	S ₂ /S ₃	S ₁ /TOC	PI	S ₁ + S ₂
EFLH-1	6640.0	5.74	3.10	19.25	0.31	444	335	5	62	54	0.14	22.35
EFLH-2	6648.0	5.75	2.31	19.08	0.30	444	332	5	64	40	0.11	21.39
EFLH-3	6656.0	4.15	1.99	13.34	0.29	443	322	7	46	48	0.13	15.33
EFLH-4	6664.0	4.25	2.04	13.11	0.44	446	308	10	30	48	0.13	15.15
EFLH-5	6672.0	4.71	2.57	15.95	0.43	442	338	9	37	55	0.14	18.52
EFLH-6	6680.0	3.54	1.09	6.35	0.25	443	180	7	25	31	0.15	7.44
EFLH-7	6688.0	5.87	3.17	24.83	0.33	442	423	6	75	54	0.11	28.00
EFLH-8	6695.0	4.97	2.66	19.26	0.31	442	387	6	62	53	0.12	21.92
EFLH-9	6704.0	4.56	2.27	15.77	0.31	444	346	7	51	50	0.13	18.04
EFLH-10	6711.5	3.67	1.19	8.94	0.27	442	244	7	33	32	0.12	10.13
EFLH-11	6719.0	4.35	1.58	13.60	0.31	443	312	7	44	36	0.10	15.18
EFLH-12	6727.0	3.06	0.57	4.42	0.25	442	145	8	18	19	0.11	4.99
EFLH-13	6735.0	2.28	0.15	0.35	0.17	437	15	7	2	7	0.30	0.50
EFLH-14	6744.0	1.65	0.18	1.43	0.24	441	87	15	6	11	0.11	1.61
EFLH-15	6754.0	1.99	0.37	2.78	0.22	442	140	11	13	19	0.12	3.15
EFLH-16	6764.0	2.49	0.33	1.74	0.17	442	70	7	10	13	0.16	2.07
EFLH-17	6774.0	2.16	0.17	1.31	0.18	444	61	8	7	8	0.11	1.48
EFLH-18	6784.0	2.02	0.13	0.45	0.24	439	22	12	2	6	0.22	0.58
EFLH-19	6794.0	3.56	1.35	10.12	0.37	444	284	10	27	38	0.12	11.47
EFLH-20	6804.0	2.61	0.26	1.97	0.31	445	75	12	6	10	0.12	2.23
EFLH-21	6815.0	0.23	0.04	0.08	0.32	440	35	139	0	17	0.33	0.12
EFLH-22	6820.5	0.34	0.04	0.10	0.40	446	30	119	0	12	0.29	0.14

C. Geochemical logs of TOC, HI, Pr/Ph and AIR for the Eagle Ford Shale bitumens analyzed

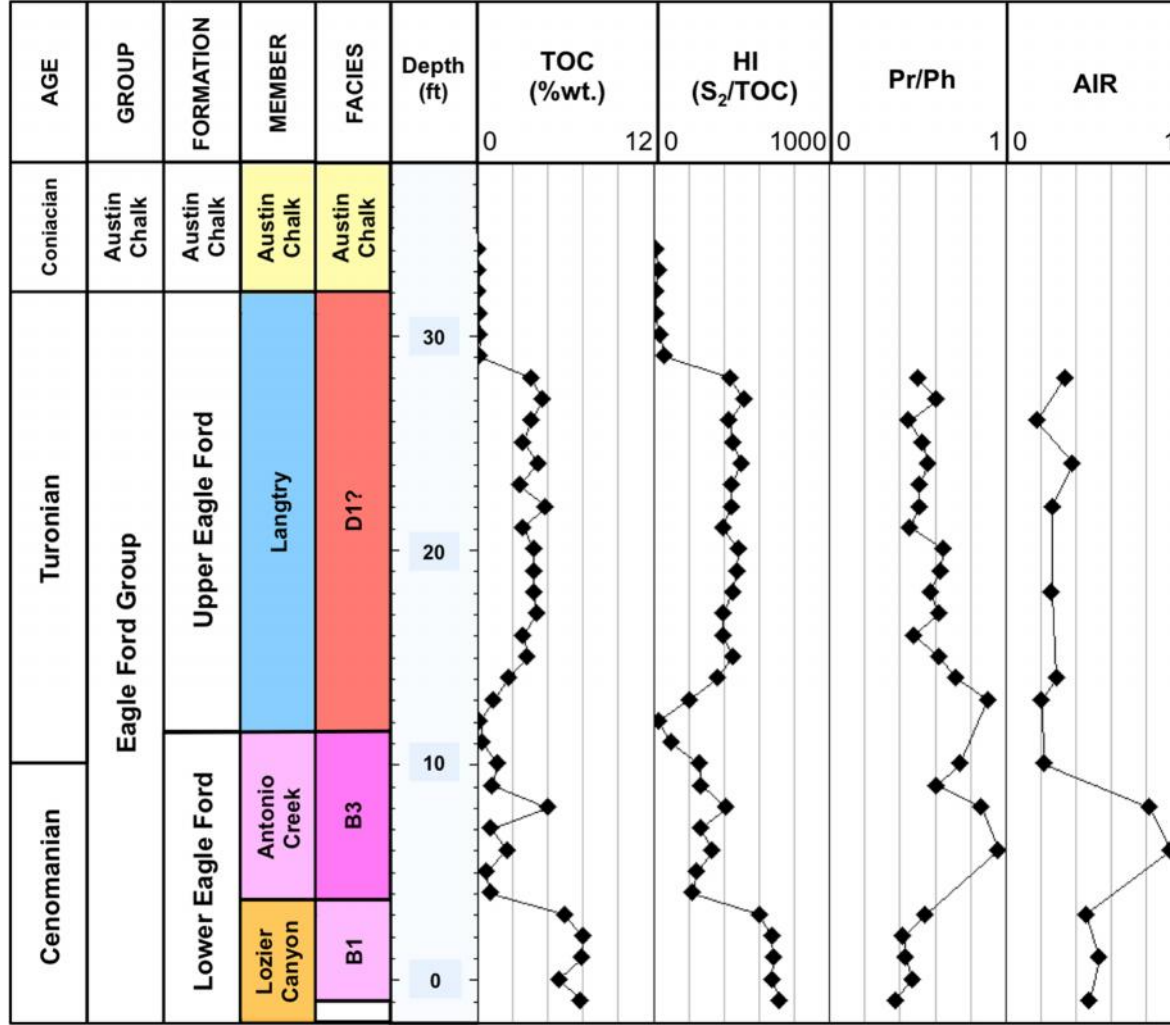
Ferguson McKnight #526-1H core



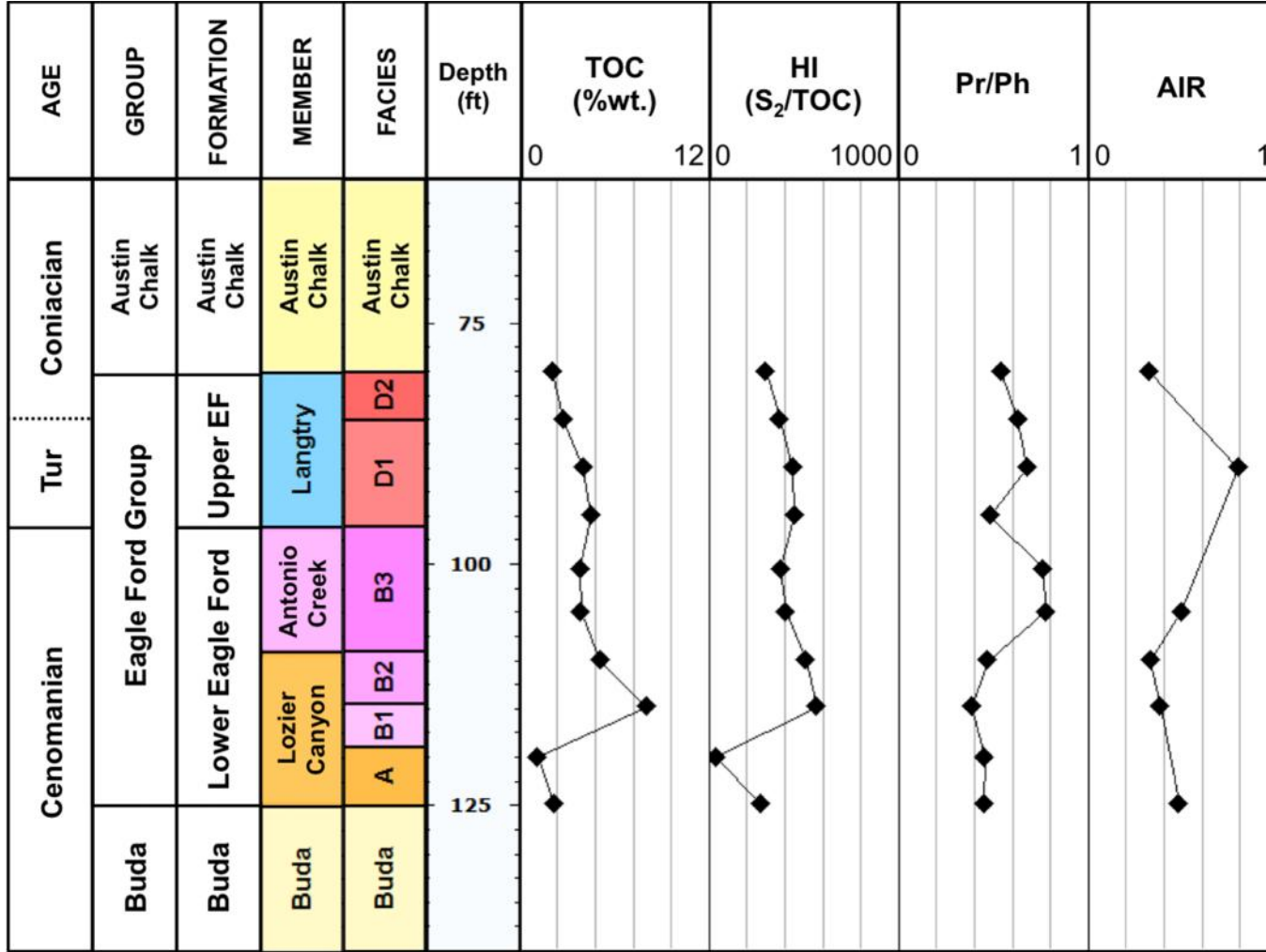
Fasken "A" #1 core



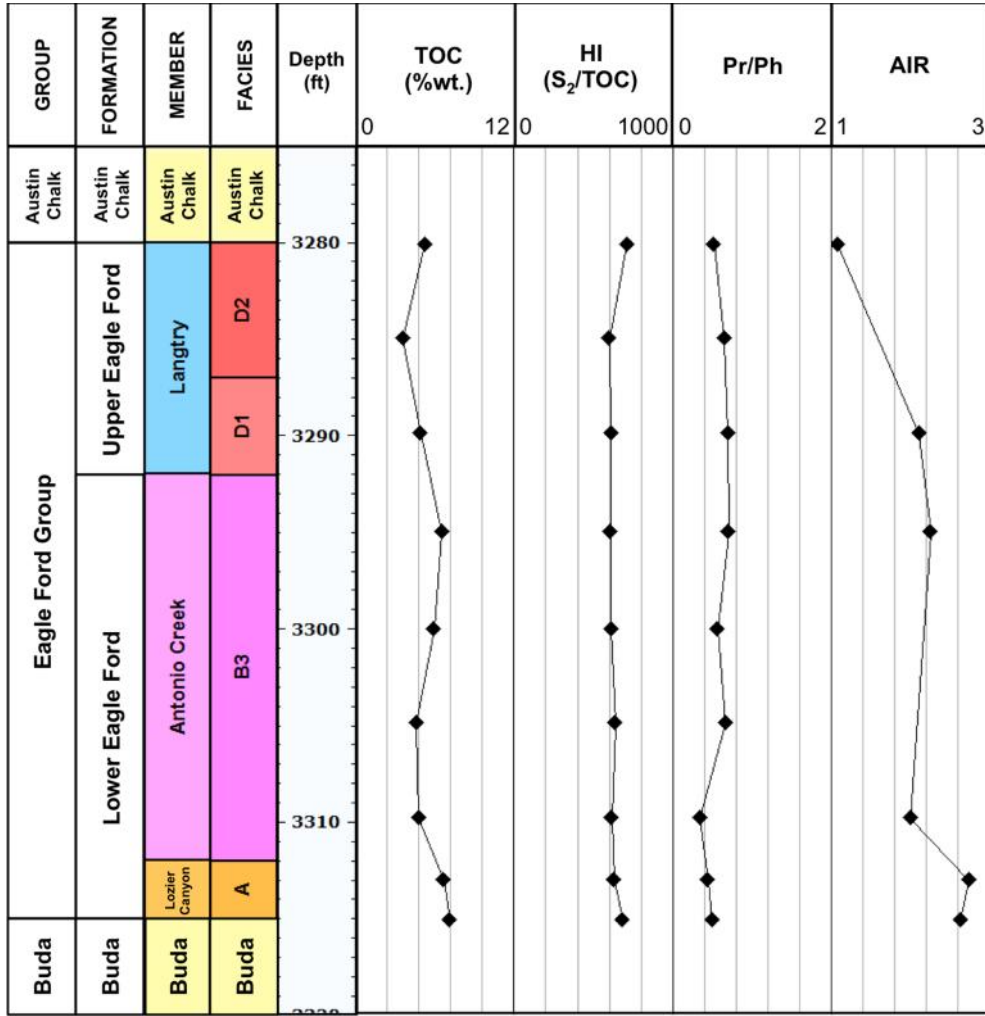
Bouldin Creek outcrop



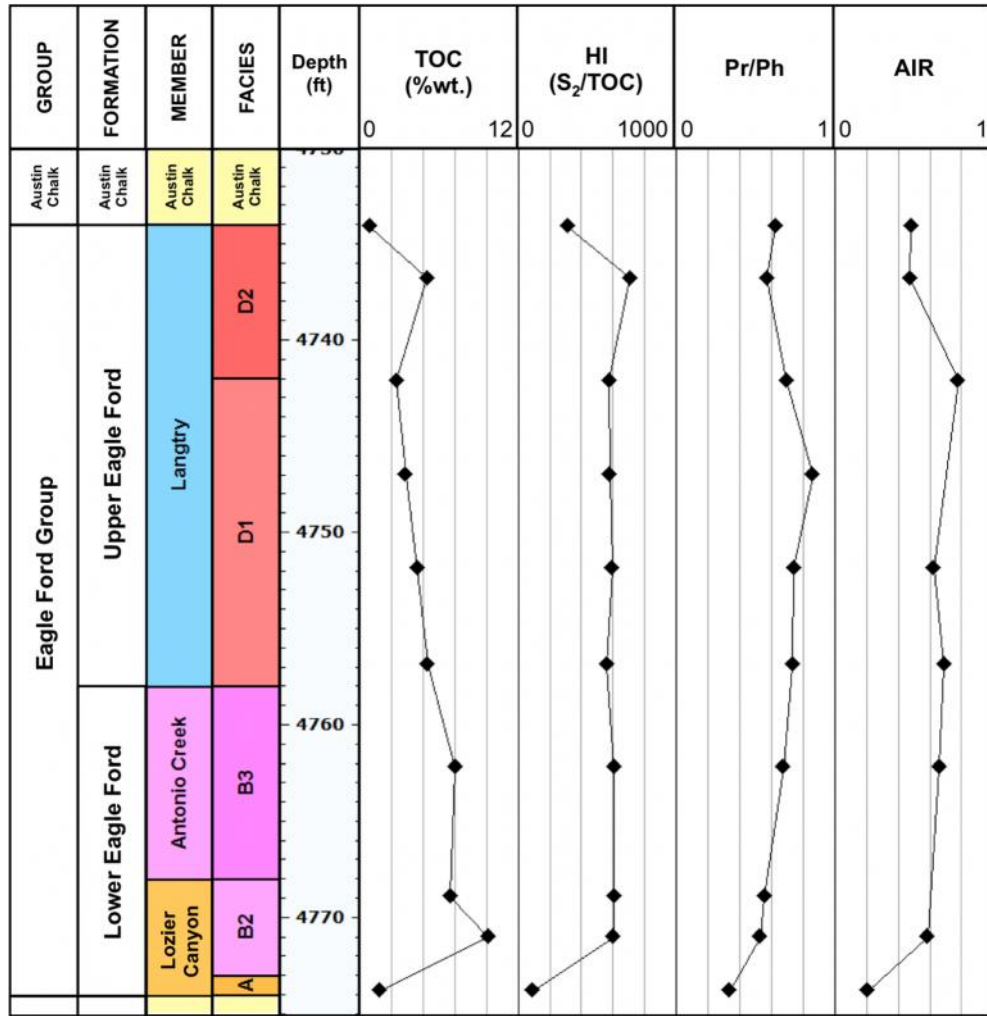
ACC #1 core



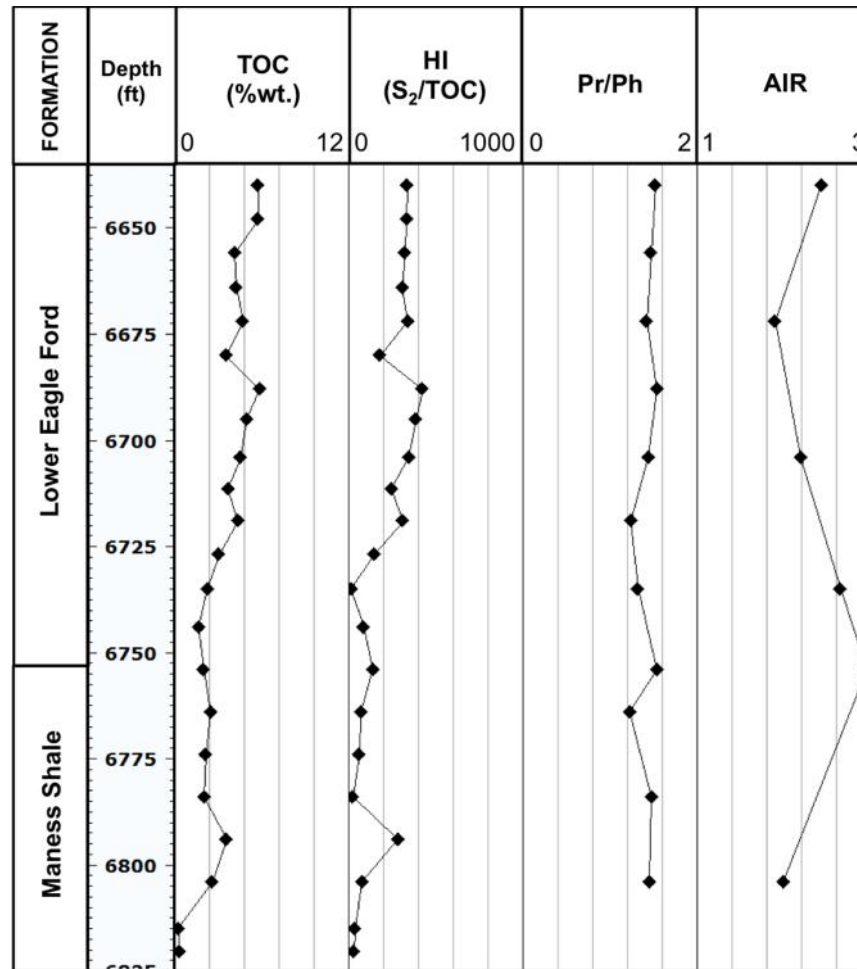
W. Brechtel #1 core



C. J. Hendershot #1 core

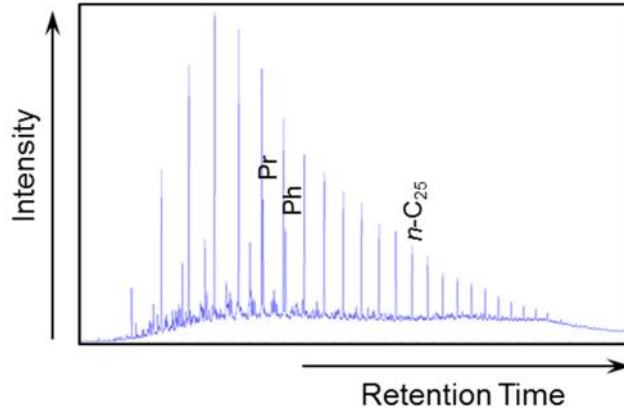


Lily Hoppess #1 core

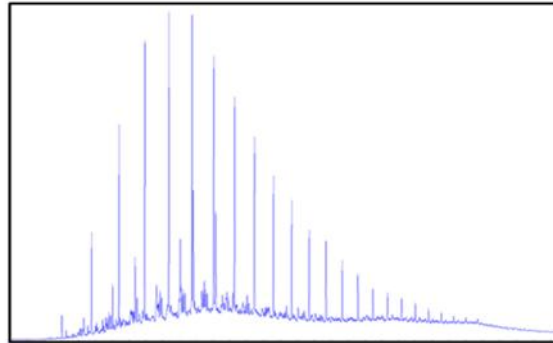


D. Gas chromatograms of the saturate fractions for the Eagle Ford Shale samples analyzed in this study (Pr = pristane; Ph = phytane, n-C₂₅ = C₂₅ normal alkane)

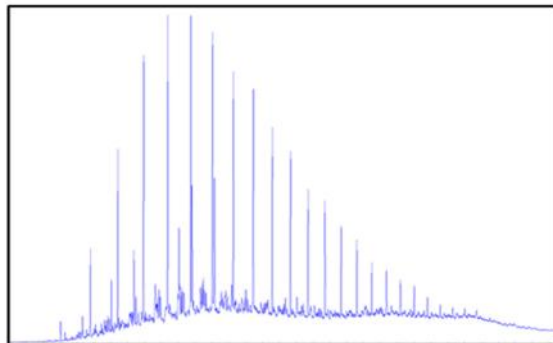
Ferguson McKnight #526-1H core



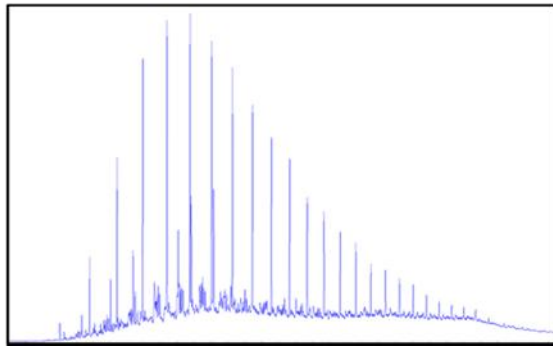
EFMK-4
Upper Eagle Ford Fm.
Depth = 5817.5 ft



EFMK17
Upper Eagle Ford Fm.
Depth = 5921 ft

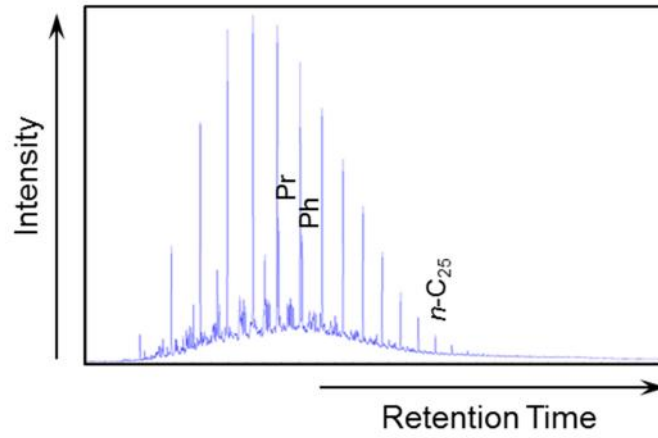


EFMK-25
Lower Eagle Ford Fm.
Depth = 6072 ft

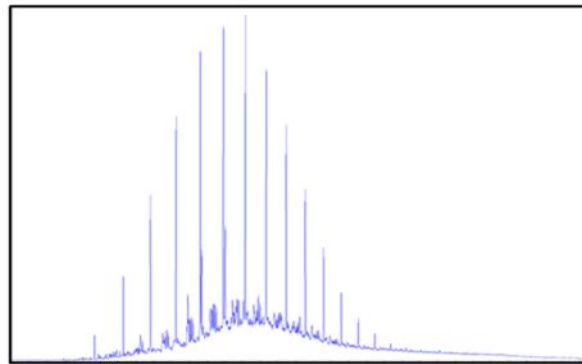


EFMK-26
Lower Eagle Ford Fm.
Depth = 6174 ft

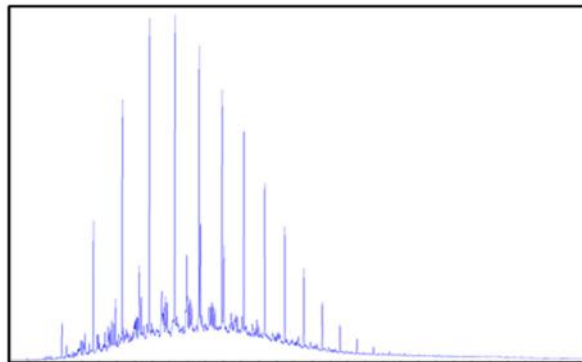
Fasken "A" #1 core



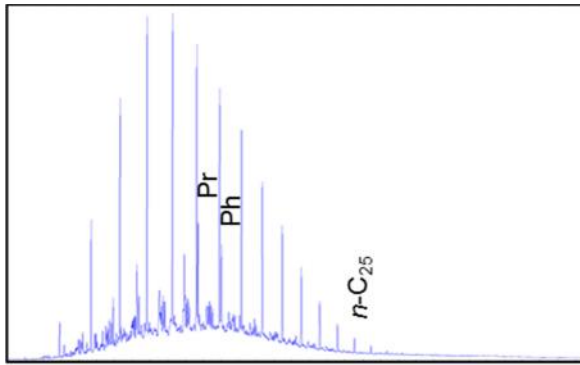
EFA-3
Langtry Mbr.
Depth = 9450.0 ft



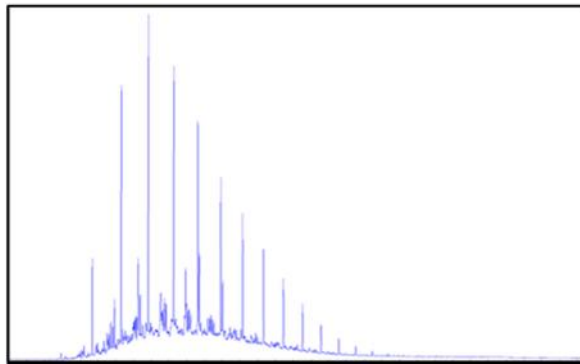
EFA-11
Scott Ranch Mbr.
Depth = 9530.0 ft



EFA-21
Scott Ranch Mbr.
Depth = 9629.5 ft

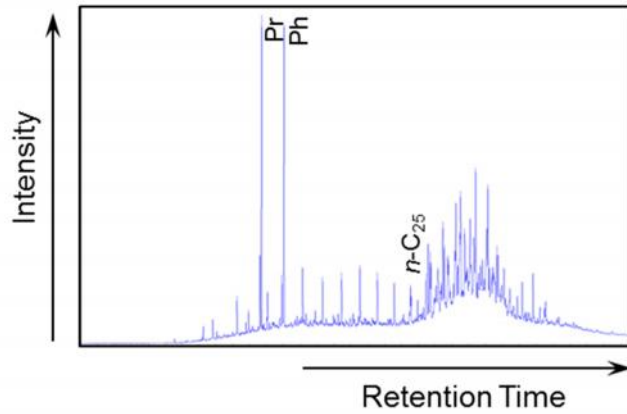


EFFA-24
Antonio Creek Mbr.
Depth = 9655.5 ft

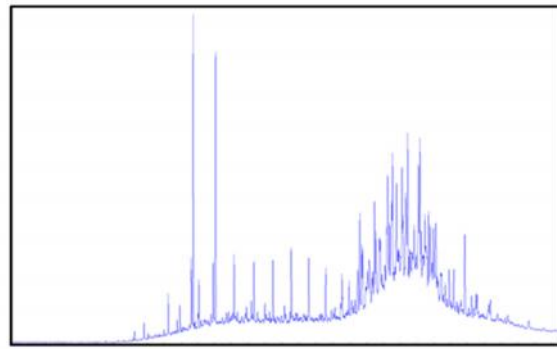


EFFA-37
Lozier Canyon Mbr.
Depth = 9756.5 ft

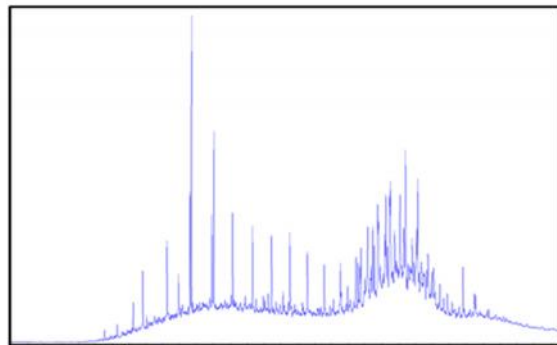
Bouldin Creek outcrop



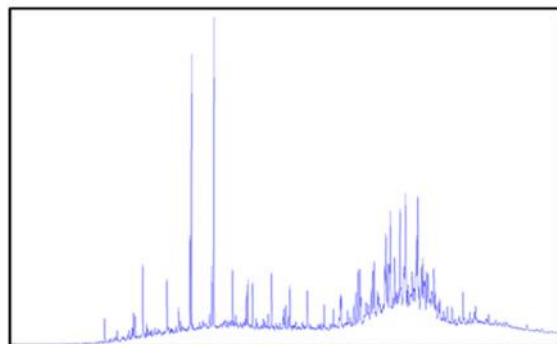
EFBC-5
Langtry Mbr.
Depth = 28.0 ft



EFBC-15
Langtry Mbr.
Depth = 18.0 ft

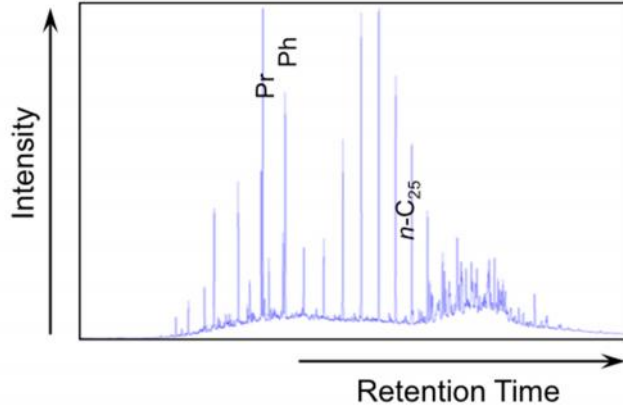


EFBC-25
Antonio Creek Mbr.
Depth = 8.0 ft

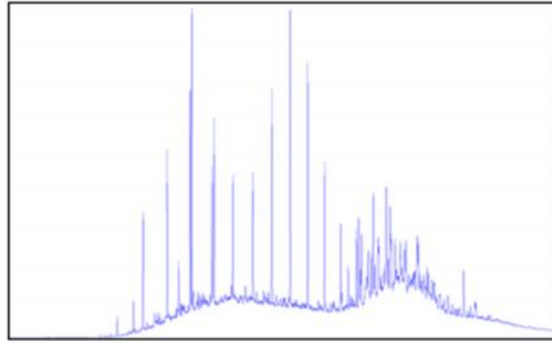


EFBC-32
Lozier Canyon Mbr.
Depth = 1.0 ft

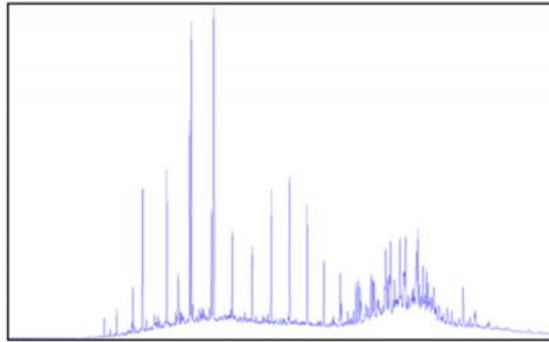
ACC #1 core



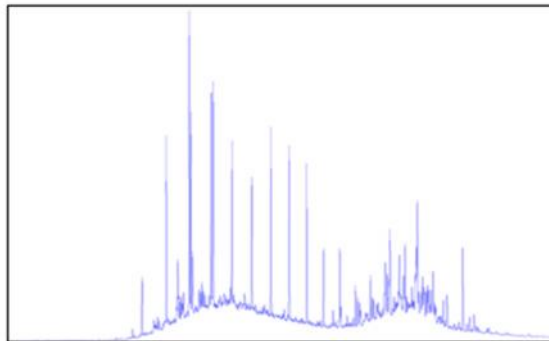
EFAC-3
Langtry Mbr.
Depth = 90.0 ft



EFAC-6
Antonio Creek Mbr.
Depth = 105.0 ft

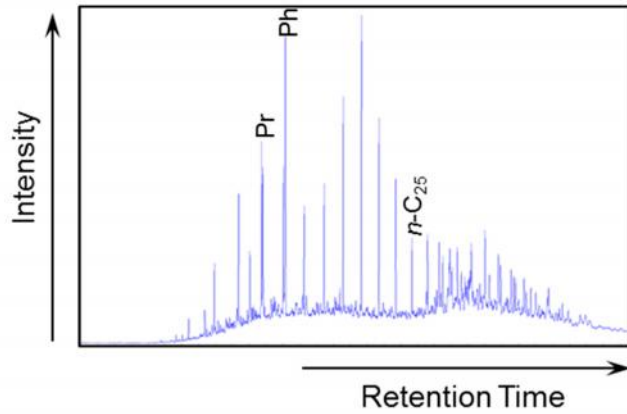


EFAC-7
Lozier Canyon Mbr.
Depth = 110.0 ft

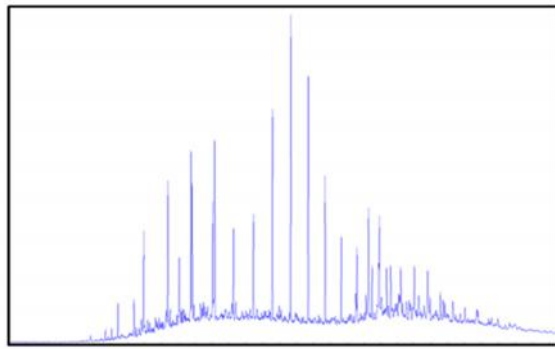


EFAC-10
Lozier Canyon Mbr.
Depth = 124.8 ft

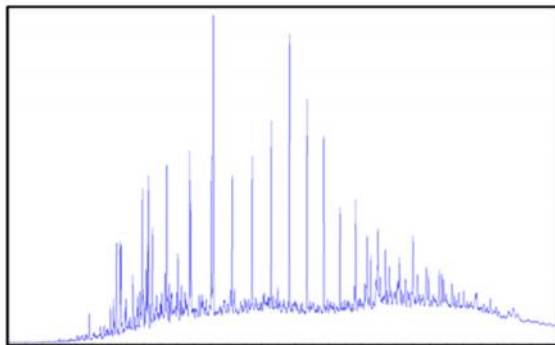
W. Brechtel #1 core



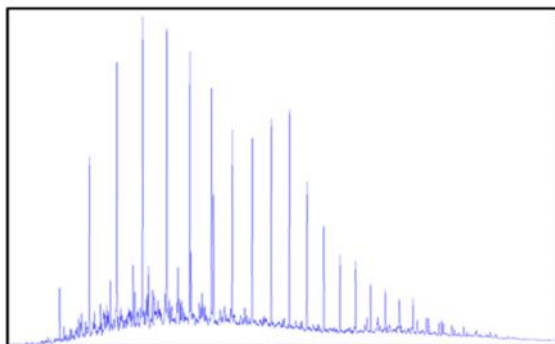
EFWB-1
Langtry Mbr.
Depth = 3280.1 ft



EFWB-3
Langtry Mbr.
Depth = 3289.9 ft

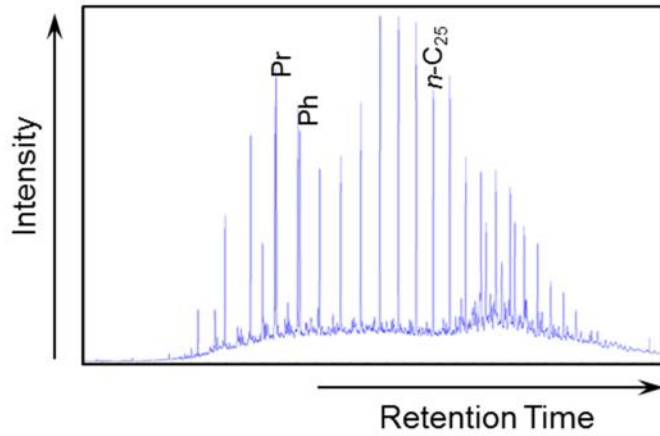


EFWB-7
Antonio Creek Mbr.
Depth = 3309.8 ft

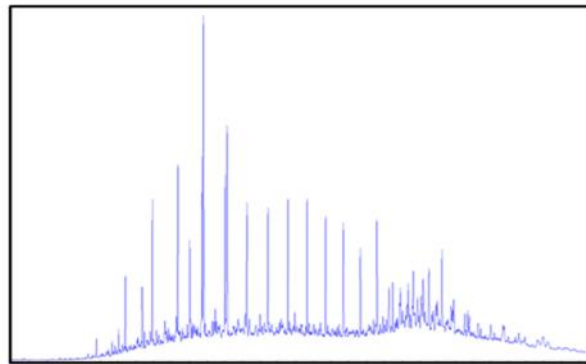


EFWB-9
Lozier Canyon Mbr.
Depth = 3315.1 ft

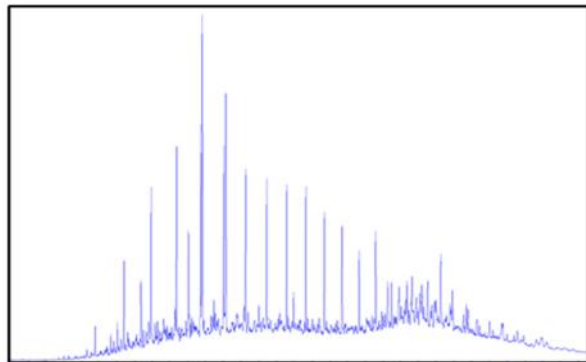
C. J. Hendershot #1 core



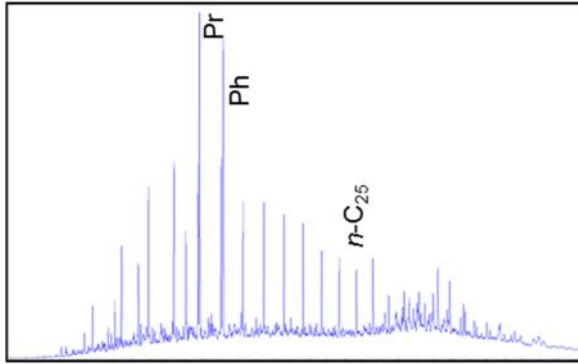
EFHE-1
Langtry Mbr.
Depth = 4734.1 ft



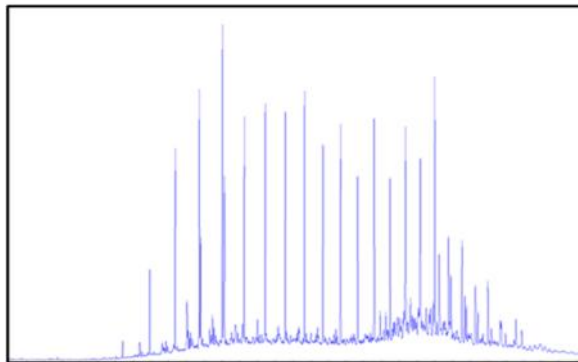
EFHE-5
Scott Ranch Mbr.
Depth = 4751.9 ft



EFHE-7
Antonio Creek Mbr.
Depth = 4762.2 ft

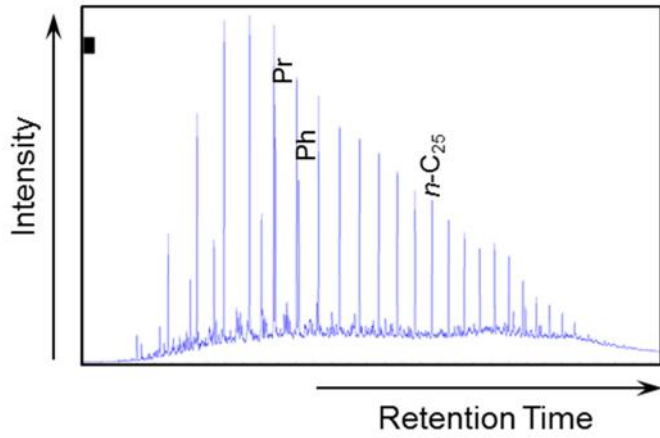


EFHE-9
Lozier Canyon Mbr.
Depth = 4771.0 ft

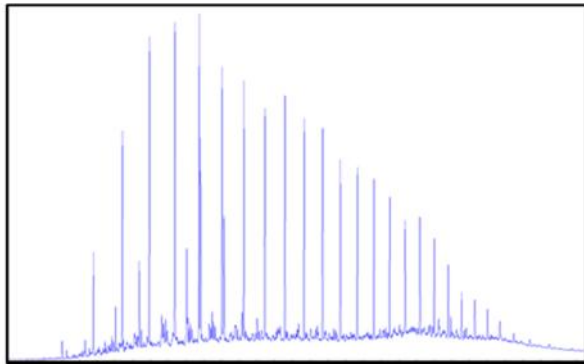


EFHE-10
Lozier Canyon Mbr.
Depth = 4773.8 ft

Lily Hoppess #1 core

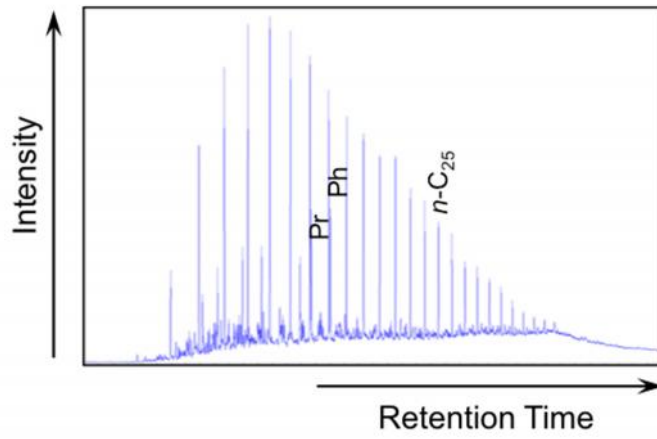


EFLH-9
Lower Eagle Ford Fm.
Depth = 6709 ft



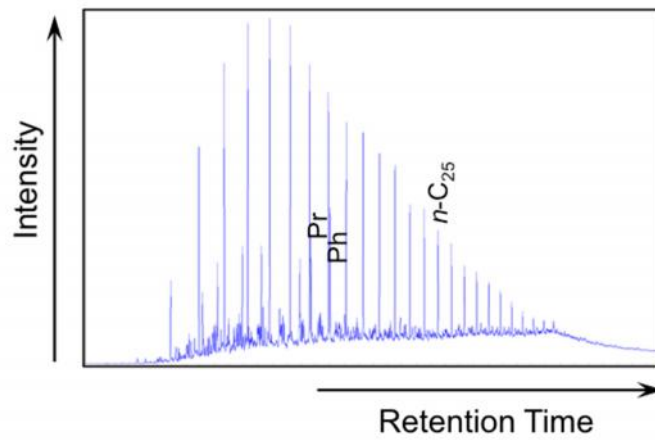
EFLH-15
Maness Shale
Depth = 6754 ft

NWMK-533.2 oil



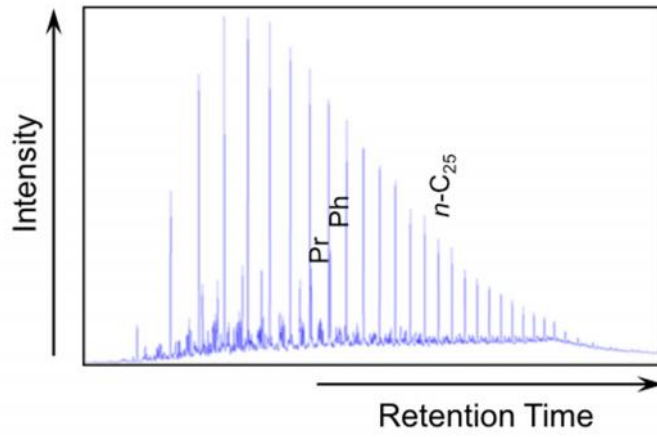
NWMK-533.2
Oil

NFMK 526-1H oil



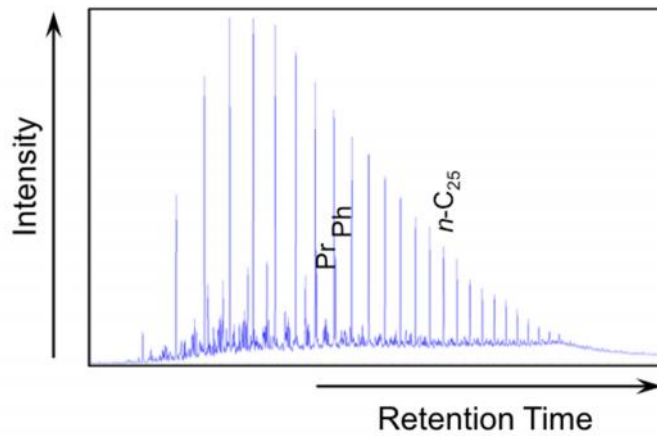
NFMK-526-1H
Oil

MJOG-UEF oil



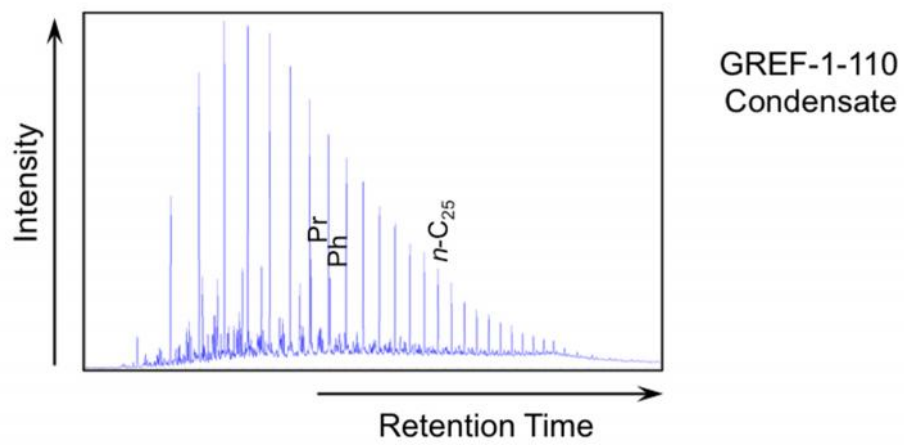
MJOG-UEF
Oil

MJOG-LEF oil



MJOG-LEF
Oil

GRAF-1-110 condensate



E. Geochemical ratios of *n*-alkanes and isoprenoids for the saturate fractions of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined)

Lozier Canyon outcrop

181

Sample	Depth (ft)	Pr/Ph	Pr/ <i>n</i> -C ₁₇	Pr/ <i>n</i> -C ₁₈	Pr/ (Pr+Ph)	TAR	CPI	long/ short chain <i>n</i> -alkanes
EFLC-28	130.0	0.45	1.50	2.52	0.31	1.39	0.84	2.60
EFLC-26	121.5	0.42	0.94	1.78	0.30	0.38	0.82	1.23
EFLC-25	116.5	0.31	1.93	3.68	0.24	0.00	0.56	2.41
EFLC-20	89.0	ND	ND	3.57	ND	1.98	0.41	5.06
EFLC-19	85.0	ND	ND	ND	ND	ND	1.12	13.97
EFLC-18	80.5	ND	ND	ND	ND	ND	1.48	ND
EFLC-16	70.0	ND	ND	ND	ND	27.07	1.20	15.67
EFLC-14	60.0	0.30	0.78	2.18	0.23	1.45	1.06	2.11
EFLC-13	55.0	ND	ND	ND	ND	ND	0.57	33.61
EFLC-12	50.0	0.25	0.88	2.06	0.20	3.70	0.86	4.17
EFLC-11	45.0	0.25	1.36	4.08	0.20	1.86	1.02	2.52
EFLC-10	40.0	0.41	0.45	1.08	0.29	0.20	0.93	0.73
EFLC-9	35.0	0.35	0.60	1.59	0.26	0.51	1.03	1.13
EFLC-8	30.0	0.36	0.40	1.05	0.27	0.22	0.95	0.76
EFLC-7	25.0	0.39	0.46	1.06	0.28	0.23	1.00	0.77
EFLC-6	20.0	ND	ND	5.67	ND	15.41	1.13	9.05
EFLC-5	15.0	0.22	0.75	3.10	0.18	0.99	1.15	2.88
EFLC-4	10.0	0.25	2.15	5.56	0.20	1.04	1.19	2.46
EFLC-3	9.0	0.34	0.73	1.69	0.25	2.42	1.22	2.84
EFLC-1	0.0	0.42	1.13	2.42	0.30	1.48	1.04	2.49

Ferguson McKnight #526-1H core

Sample	Depth (ft)	Pr/Ph	Pr/ <i>n</i> -C ₁₇	Pr/ <i>n</i> -C ₁₈	Pr/ (Pr+Ph)	TAR	CPI	long/ short chain <i>n</i> -alkanes
EFMK-1	5793.0	1.17	0.53	0.51	0.54	0.11	1.01	0.45
EFMK-3	5809.0	1.34	0.51	0.48	0.57	0.11	0.96	0.44
EFMK-4	5817.5	1.30	0.58	0.58	0.57	0.16	0.97	0.54
EFMK-5	5825.5	1.29	0.54	0.52	0.56	0.12	0.98	0.44
EFMK-8	5849.0	1.40	0.55	0.50	0.58	0.15	0.98	0.53
EFMK-10	5864.5	1.34	0.57	0.52	0.57	0.18	0.96	0.60
EFMK-12	5881.0	1.30	0.52	0.51	0.57	0.10	0.96	0.41
EFMK-13	5889.0	1.42	0.58	0.53	0.59	0.16	0.98	0.56
EFMK-15	5905.5	1.26	0.58	0.54	0.56	0.14	0.98	0.51
EFMK-16	5913.5	1.27	0.58	0.55	0.56	0.13	0.97	0.50
EFMK-17	5921.0	1.14	0.49	0.51	0.53	0.10	0.99	0.41
EFMK-19	5937.5	1.32	0.54	0.50	0.57	0.13	0.96	0.52
EFMK-20	5945.5	1.29	0.54	0.53	0.56	0.10	0.97	0.44
EFMK-22	5961.0	1.20	0.54	0.54	0.55	0.12	0.99	0.49
EFMK-23	5969.0	1.22	0.56	0.56	0.55	0.13	0.95	0.51
EFMK-24	5977.5	1.31	0.55	0.53	0.57	0.14	0.99	0.53
EFMK-25	6072.0	0.98	0.54	0.60	0.50	0.15	0.95	0.57
EFMK-26	6174.0	0.91	0.47	0.58	0.48	0.16	0.98	0.54

Fasken "A" #1 core

Sample	Depth (ft)	Pr/Ph	Pr/ <i>n</i> -C ₁₇	Pr/ <i>n</i> -C ₁₈	Pr/ (Pr+Ph)	TAR	CPI	long/ short chain <i>n</i> -alkanes
EFFA-1	9430.5	0.92	0.30	0.34	0.48	0.01	1.47	0.23
EFFA-3	9450.0	1.14	0.42	0.44	0.53	ND	1.44	0.20
EFFA-5	9470.5	0.95	0.36	0.41	0.49	ND	1.55	0.23
EFFA-7	9489.0	1.05	0.38	0.42	0.51	ND	1.50	0.19
EFFA-11	9530.0	0.86	0.36	0.39	0.46	0.01	1.37	0.31
EFFA-15	9570.5	1.23	0.43	0.43	0.55	0.01	1.27	0.20
EFFA-17	9589.0	1.11	0.41	0.43	0.53	ND	1.46	0.22
EFFA-18	9599.5	1.10	0.42	0.44	0.52	ND	1.55	0.20
EFFA-20	9620.5	0.94	0.44	0.48	0.48	0.01	1.28	0.32
EFFA-21	9629.5	1.22	0.43	0.44	0.55	ND	1.69	0.20
EFFA-22	9639.5	1.10	0.39	0.41	0.52	ND	1.52	0.19
EFFA-23	9650.5	1.16	0.38	0.39	0.54	ND	1.50	0.21
EFFA-24	9655.5	1.22	0.43	0.43	0.55	ND	1.33	0.17
EFFA-26	9670.5	1.42	0.45	0.42	0.59	ND	1.29	0.15
EFFA-28	9685.5	1.24	0.42	0.41	0.55	ND	1.63	0.18
EFFA-30	9700.5	1.27	0.43	0.42	0.56	ND	1.53	0.19
EFFA-32	9717.0	1.20	0.40	0.40	0.55	ND	1.44	0.18
EFFA-34	9733.5	1.29	0.44	0.43	0.56	ND	1.21	0.16
EFFA-37	9756.5	1.33	0.45	0.46	0.57	ND	1.28	0.14
EFFA-38	9761.5	1.17	0.43	0.44	0.54	ND	1.52	0.20
EFFA-40	9776.0	1.08	0.42	0.44	0.52	ND	1.49	0.23

Bouldin Creek outcrop

183

Sample	Depth (ft)	Pr/Ph	Pr/ <i>n</i> -C ₁₇	Pr/ <i>n</i> -C ₁₈	Pr/ (Pr+Ph)	TAR	CPI	long/ short chain <i>n</i> -alkanes
EFBC-5	28.0	1.00	4.27	7.12	0.50	ND	0.56	1.01
EFBC-6	27.0	1.22	3.52	5.73	0.55	ND	ND	0.67
EFBC-7	26.0	0.90	4.30	7.49	0.47	ND	ND	0.93
EFBC-8	25.0	1.05	3.36	5.16	0.51	ND	ND	0.74
EFBC-9	24.0	1.12	3.56	5.61	0.53	ND	ND	0.76
EFBC-10	23.0	1.02	4.39	5.60	0.51	ND	ND	0.91
EFBC-11	22.0	1.02	4.52	7.48	0.51	ND	ND	1.00
EFBC-12	21.0	0.92	4.41	6.83	0.48	ND	ND	0.83
EFBC-13	20.0	1.30	4.42	5.76	0.56	ND	ND	0.70
EFBC-14	19.0	1.26	4.21	5.48	0.56	ND	ND	0.76
EFBC-15	18.0	1.15	4.80	5.23	0.53	ND	ND	0.89
EFBC-16	17.0	1.24	4.76	5.47	0.55	ND	ND	0.94
EFBC-17	16.0	0.96	5.62	6.94	0.49	ND	ND	1.01
EFBC-18	15.0	1.24	3.75	5.29	0.55	ND	ND	0.65
EFBC-19	14.0	1.44	4.21	3.55	0.59	ND	ND	0.79
EFBC-20	13.0	1.80	3.22	2.11	0.64	ND	ND	0.82
EFBC-23	10.0	1.48	2.35	1.63	0.60	ND	ND	0.62
EFBC-24	9.0	1.22	4.30	3.72	0.55	ND	0.61	1.09
EFBC-25	8.0	1.73	2.59	2.15	0.63	ND	ND	0.55
EFBC-27	6.0	1.92	1.97	1.47	0.66	ND	ND	0.52
EFBC-30	3.0	1.09	2.23	3.62	0.52	ND	ND	0.47
EFBC-31	2.0	0.84	2.80	6.09	0.46	ND	ND	0.49
EFBC-32	1.0	0.87	3.02	5.73	0.46	ND	ND	0.47
EFBC-33	0.0	0.94	3.51	6.04	0.49	ND	ND	0.43
EFBC-34	-1.0	0.75	3.33	6.30	0.43	ND	ND	0.41

ACC#1 core

Sample	Depth (ft)	Pr/Ph	Pr/ <i>n</i>-C₁₇	Pr/ <i>n</i>-C₁₈	Pr/ (Pr+Ph)	TAR	CPI	long/ short chain <i>n</i>-alkanes
EFAC-1	80.0	1.08	1.30	1.49	0.52	ND	1.07	3.87
EFAC-2	85.0	1.26	1.87	2.20	0.56	ND	0.99	2.33
EFAC-3	90.0	1.36	2.18	3.05	0.58	ND	1.03	1.81
EFAC-4	95.0	0.97	2.52	4.64	0.49	ND	0.98	2.62
EFAC-5	100.5	1.53	1.83	1.91	0.60	ND	0.92	1.80
EFAC-6	105.0	1.55	1.52	1.64	0.61	ND	0.80	1.06
EFAC-7	110.0	0.94	1.57	3.41	0.49	ND	0.90	0.68
EFAC-8	114.8	0.77	2.38	5.96	0.44	ND	0.92	0.68
EFAC-9	120.0	0.91	0.34	0.46	0.48	ND	1.10	0.63
EFAC-10	124.8	0.90	0.78	1.25	0.47	ND	1.29	0.61

184

W. Brechtel #1 core

Sample	Depth (ft)	Pr/Ph	Pr/ <i>n</i>-C₁₇	Pr/ <i>n</i>-C₁₈	Pr/ (Pr+Ph)	TAR	CPI	long/ short chain <i>n</i>-alkanes
EFWB-1	3280.1	0.52	1.06	2.46	0.34	ND	0.67	1.11
EFWB-2	3285.0	0.65	1.29	2.71	0.40	ND	0.83	1.94
EFWB-3	3289.9	0.70	0.99	1.99	0.41	ND	0.78	1.22
EFWB-4	3295.0	0.71	0.90	1.74	0.41	ND	0.77	1.32
EFWB-5	3300.0	0.57	1.02	2.51	0.36	ND	0.76	1.45
EFWB-6	3304.9	0.67	0.90	1.85	0.40	ND	0.76	1.47
EFWB-7	3309.8	0.35	0.80	2.51	0.26	ND	0.64	0.95
EFWB-8	3313.0	0.45	1.10	2.70	0.31	ND	0.61	0.99
EFWB-9	3315.1	0.50	0.30	0.70	0.33	0.10	0.82	0.60

C.J. Hendershot #1 core

Sample	Depth (ft)	Pr/Ph	Pr/ <i>n</i>-C₁₇	Pr/ <i>n</i>-C₁₈	Pr/ (Pr+Ph)	TAR	CPI	long/ short chain <i>n</i>-alkanes
EFHE-1	4734.1	1.25	1.40	1.10	0.35	N/D	0.66	1.31
EFHE-2	4736.8	1.14	1.56	1.20	0.50	N/D	0.49	0.73
EFHE-3	4742.1	1.39	2.01	1.77	0.48	N/D	0.53	0.63
EFHE-4	4747.0	1.72	2.76	1.90	0.42	N/D	0.55	0.89
EFHE-5	4751.9	1.48	2.16	1.75	0.47	N/D	0.56	0.71
EFHE-6	4756.9	1.47	1.86	1.58	0.52	N/D	0.59	0.53
EFHE-7	4762.2	1.35	1.85	1.61	0.49	N/D	0.62	0.61
EFHE-8	4768.9	1.11	1.95	1.89	0.53	N/D	0.67	0.50
EFHE-9	4771.0	1.06	2.22	2.08	0.53	N/D	0.67	0.52
EFHE-10	4773.8	0.67	0.52	0.63	0.48	0.71	0.65	1.51

Lily Hoppess #1 core

Sample	Depth (ft)	Pr/Ph	Pr/ <i>n</i>-C₁₇	Pr/ <i>n</i>-C₁₈	Pr/ (Pr+Ph)	TAR	CPI	long/ short chain <i>n</i>-alkanes
EFLH-1	6640.0	1.52	1.04	0.89	0.60	0.26	1.05	0.72
EFLH-3	6656.0	1.47	1.04	0.88	0.59	0.29	1.10	0.75
EFLH-5	6672.0	1.42	0.95	0.85	0.59	0.31	1.08	0.79
EFLH-7	6688.0	1.54	0.85	0.72	0.61	0.25	1.05	0.68
EFLH-9	6704.0	1.44	0.87	0.73	0.59	0.32	1.06	0.80
EFLH-11	6719.0	1.24	0.70	0.71	0.55	0.42	1.05	0.93
EFLH-13	6735.0	1.32	0.52	0.44	0.57	0.58	1.12	1.12
EFLH-15	6754.0	1.54	0.70	0.55	0.61	0.43	1.06	0.97
EFLH-16	6764.0	1.23	0.50	0.46	0.55	0.56	1.13	1.14
EFLH-18	6784.0	1.48	0.62	0.49	0.60	0.57	1.12	1.16
EFLH-20	6804.0	1.45	0.68	0.58	0.59	0.36	1.08	0.85

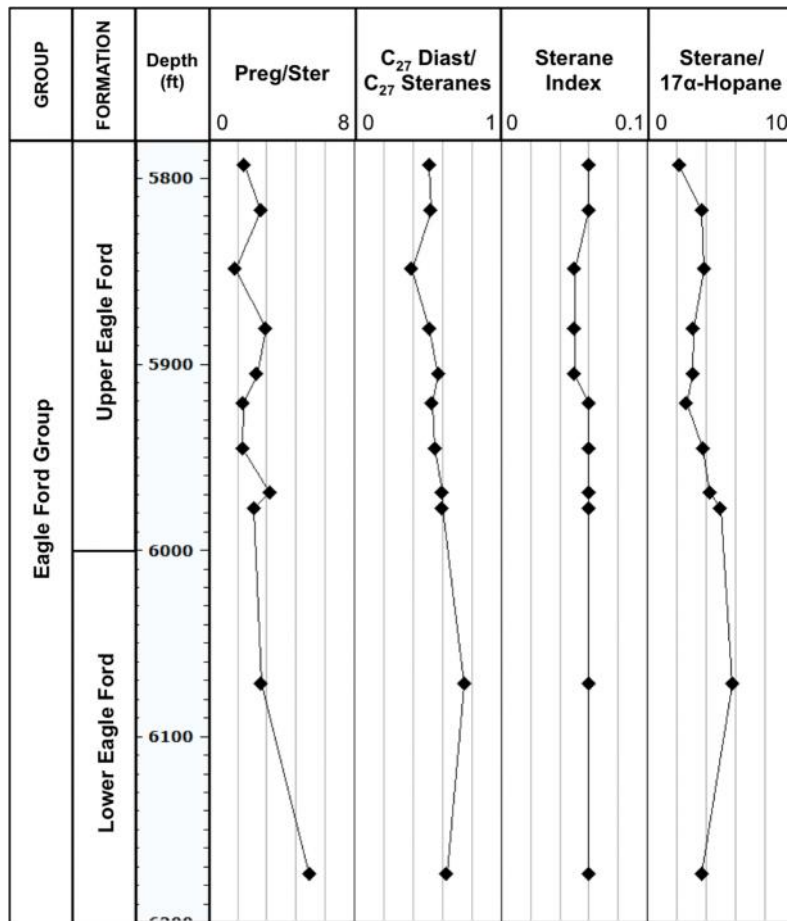
181

Oils and condensates

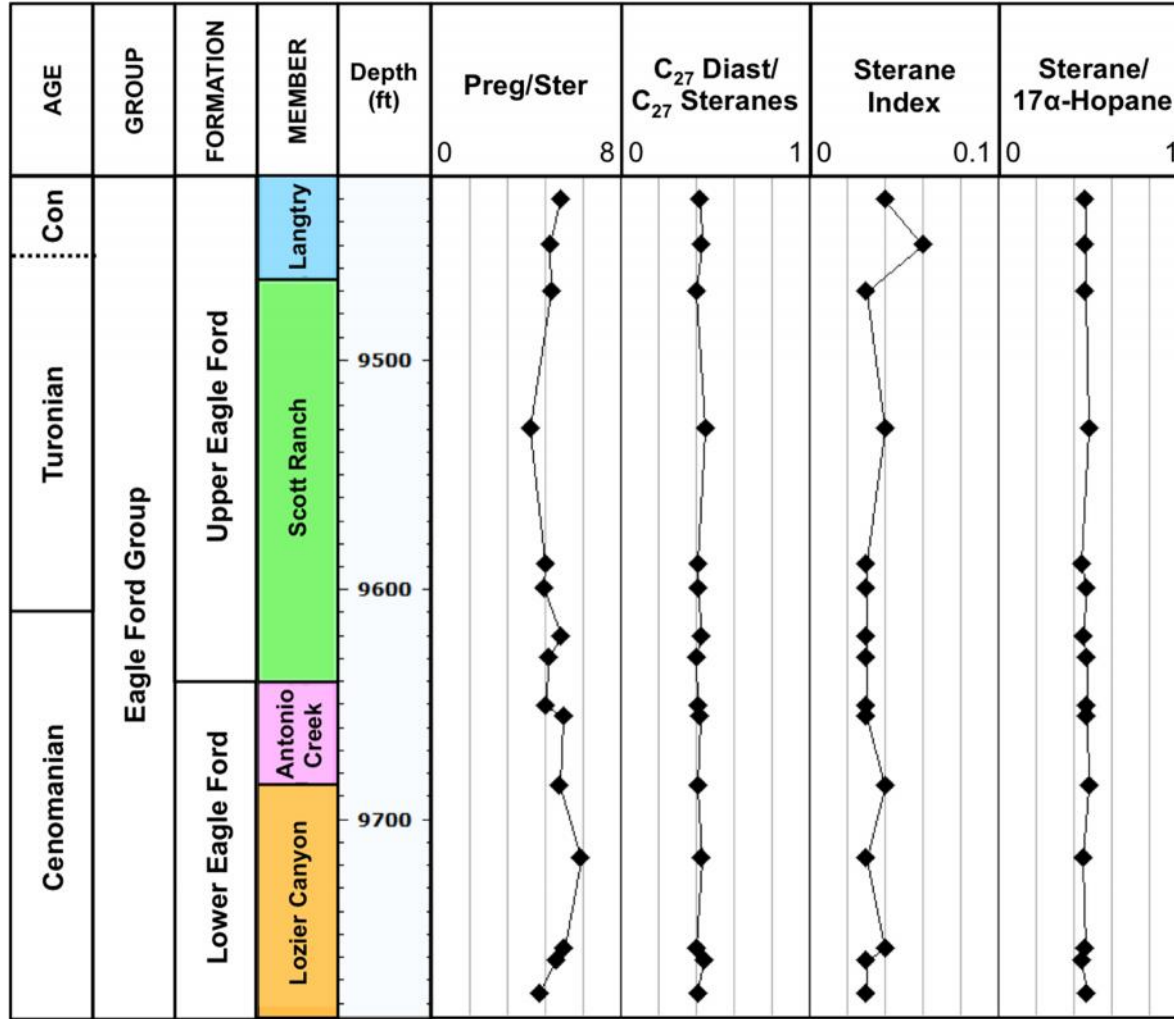
Sample	Pr/ Ph	Pr/ <i>n</i>-C₁₇	Pr/ <i>n</i>-C₁₈	Pr/ (Pr+Ph)	TAR	CPI	long/ short chain <i>n</i>-alkanes
NWMK-533.2	1.07	0.56	0.64	0.74	0.23	0.99	0.71
NFMK-526-1H	0.80	0.46	0.66	0.74	0.21	0.98	0.66
MJOG-UEF	0.80	0.35	0.49	0.78	0.23	0.97	0.68
MJOG-LEF	1.14	0.43	0.46	0.77	0.21	1.00	0.65
GRES-1-110	1.25	0.45	0.43	0.77	0.18	1.04	0.57

F. Geochemical logs of biomarker ratios of steranes for the Eagle Ford Shale bitumens analyzed

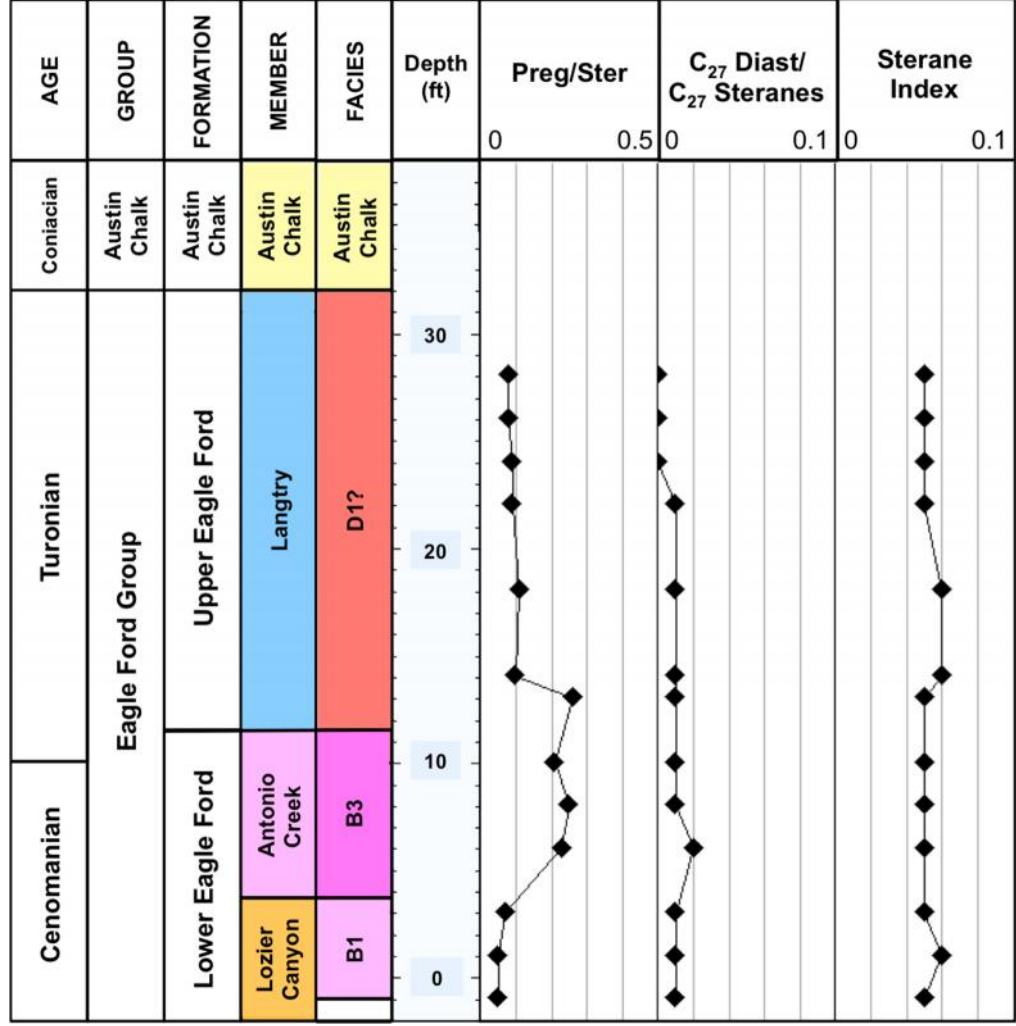
Ferguson McKnight #526-1H core



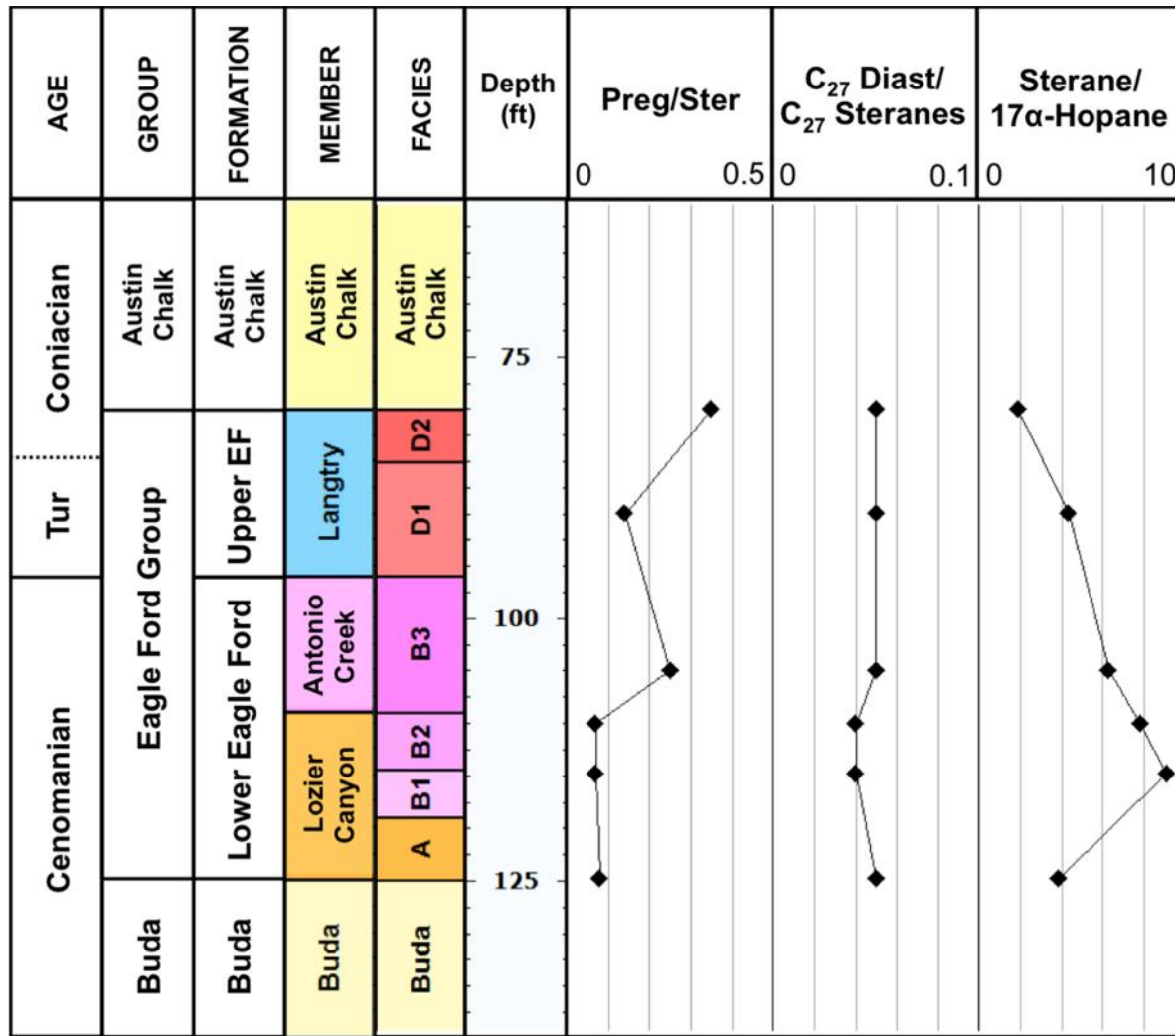
Fasken "A" #1



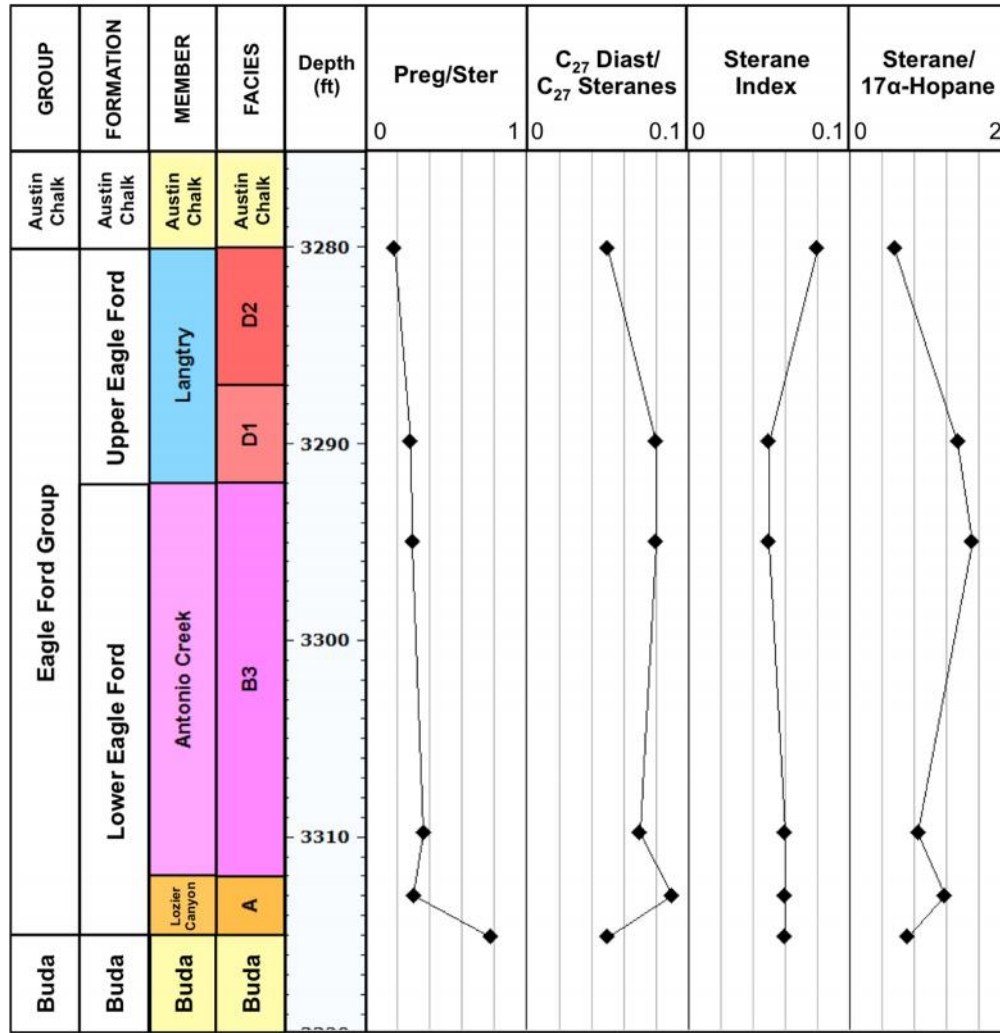
Bouldin Creek outcrop



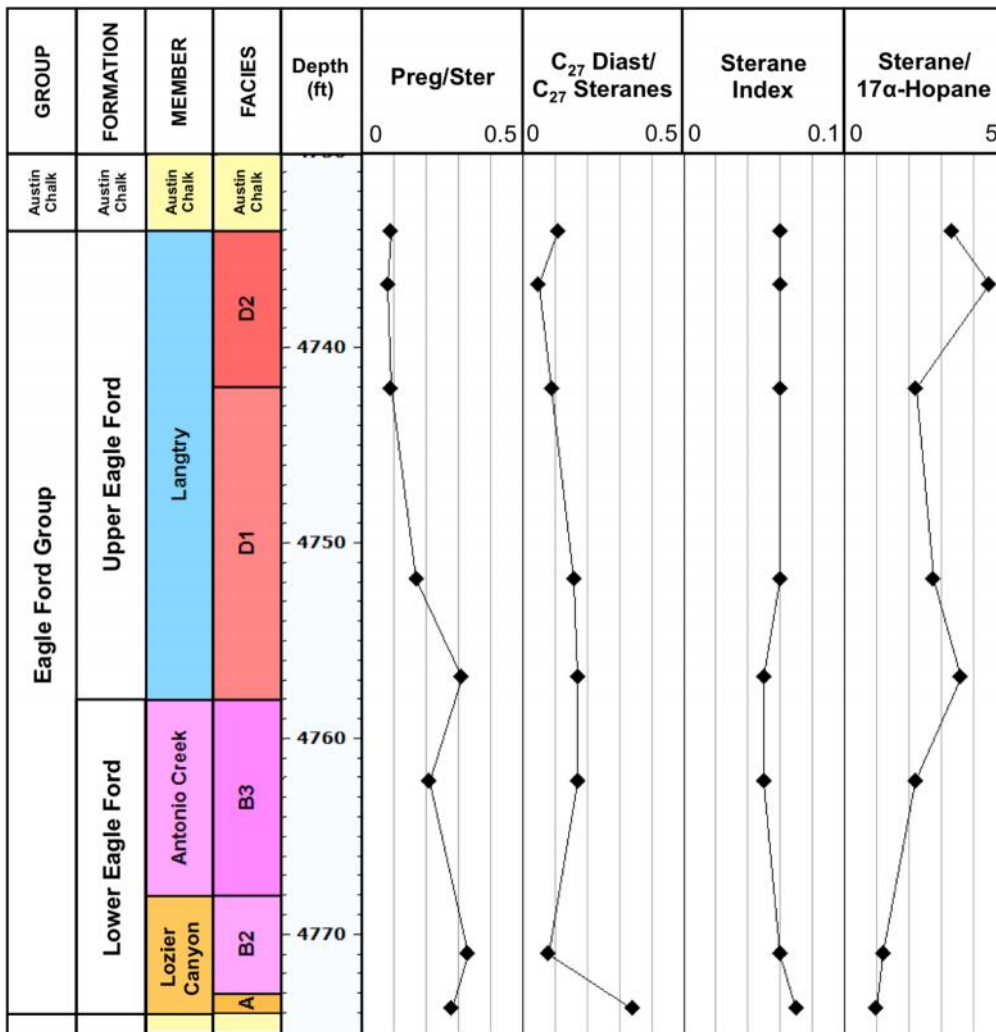
ACC #1 core



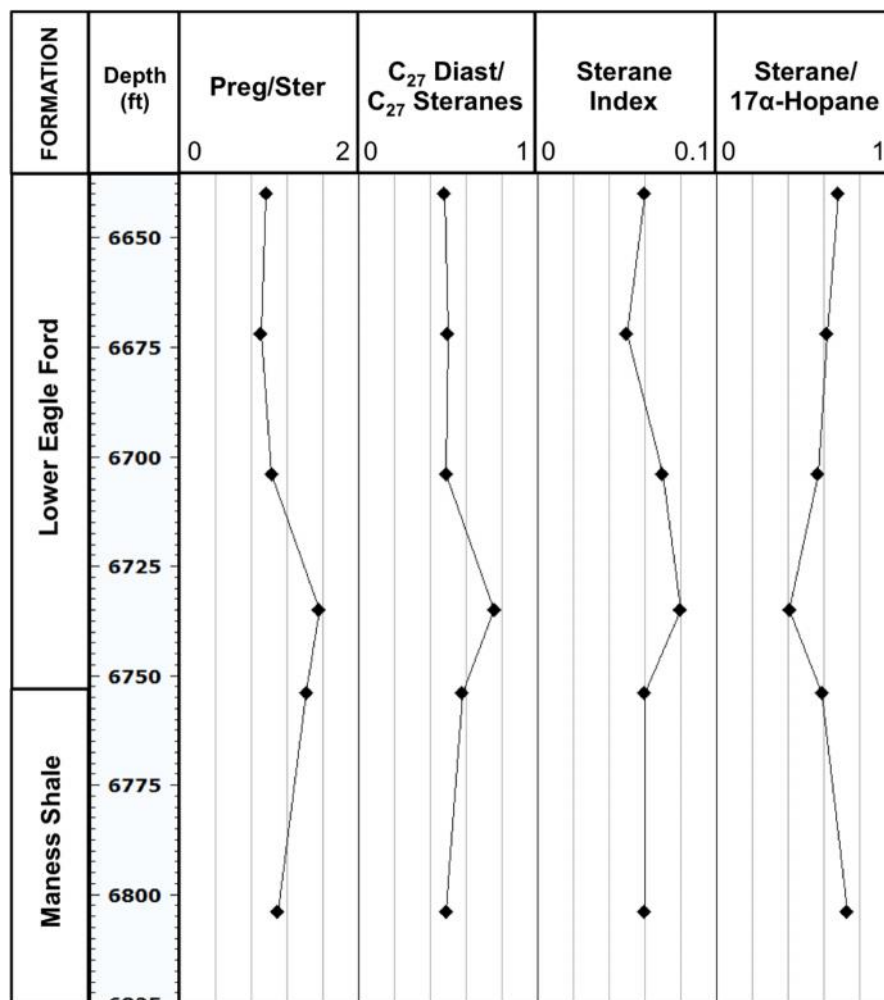
W. Brechtel #1 core



C. J. Hendershot #1 core

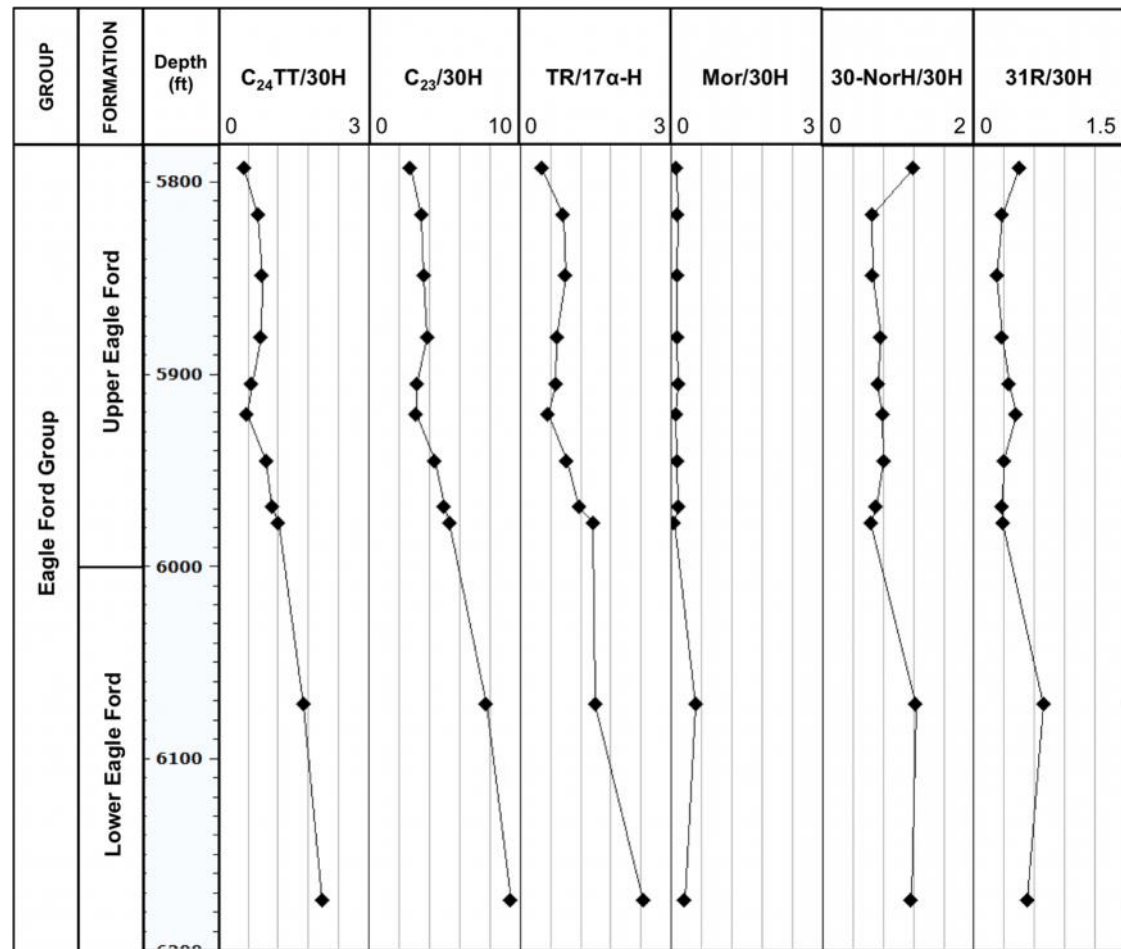


Lily Hoppess #1 core

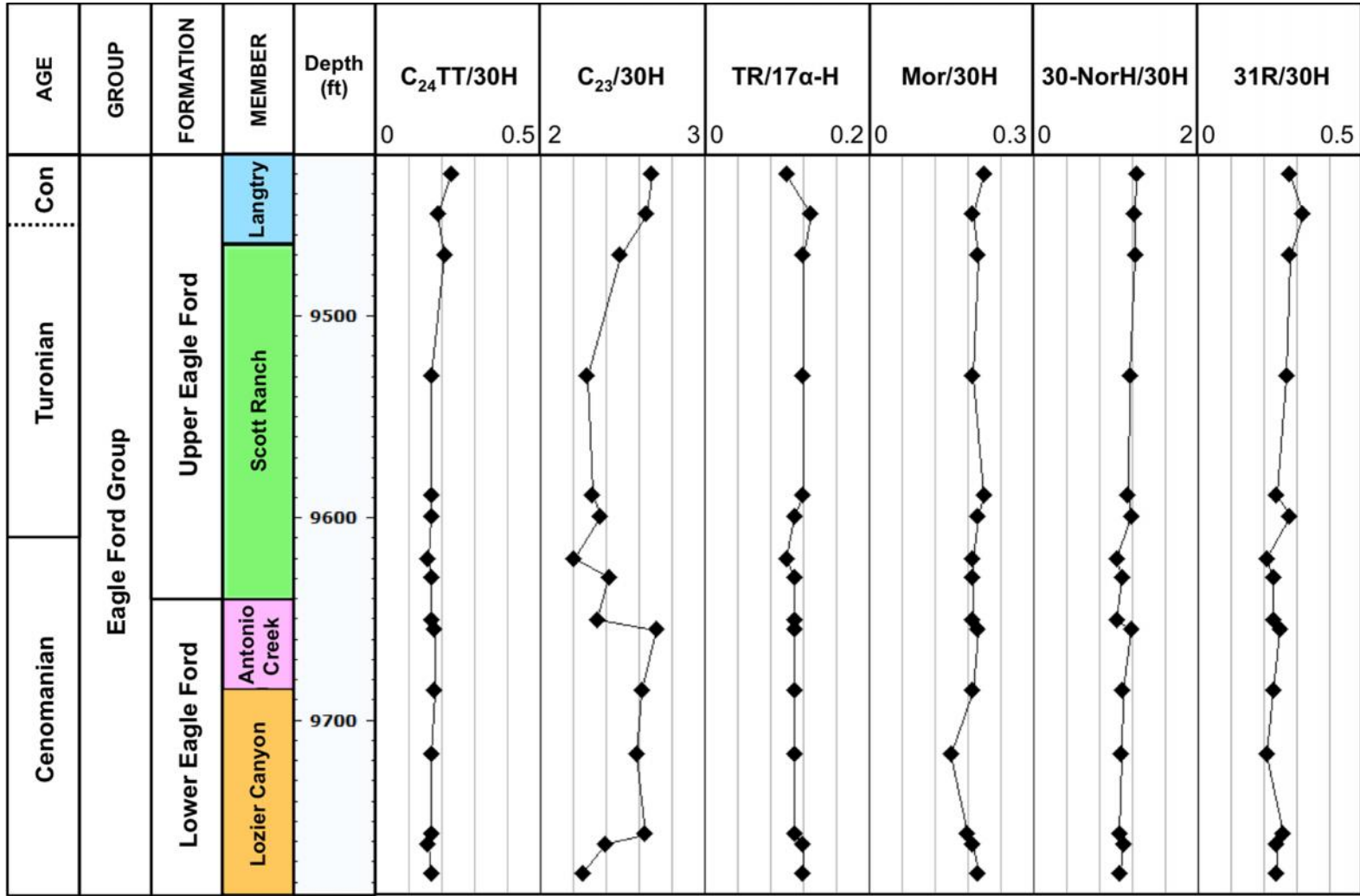


G. Geochemical logs of biomarker ratios of terpanes for the Eagle Ford Shale bitumens analyzed

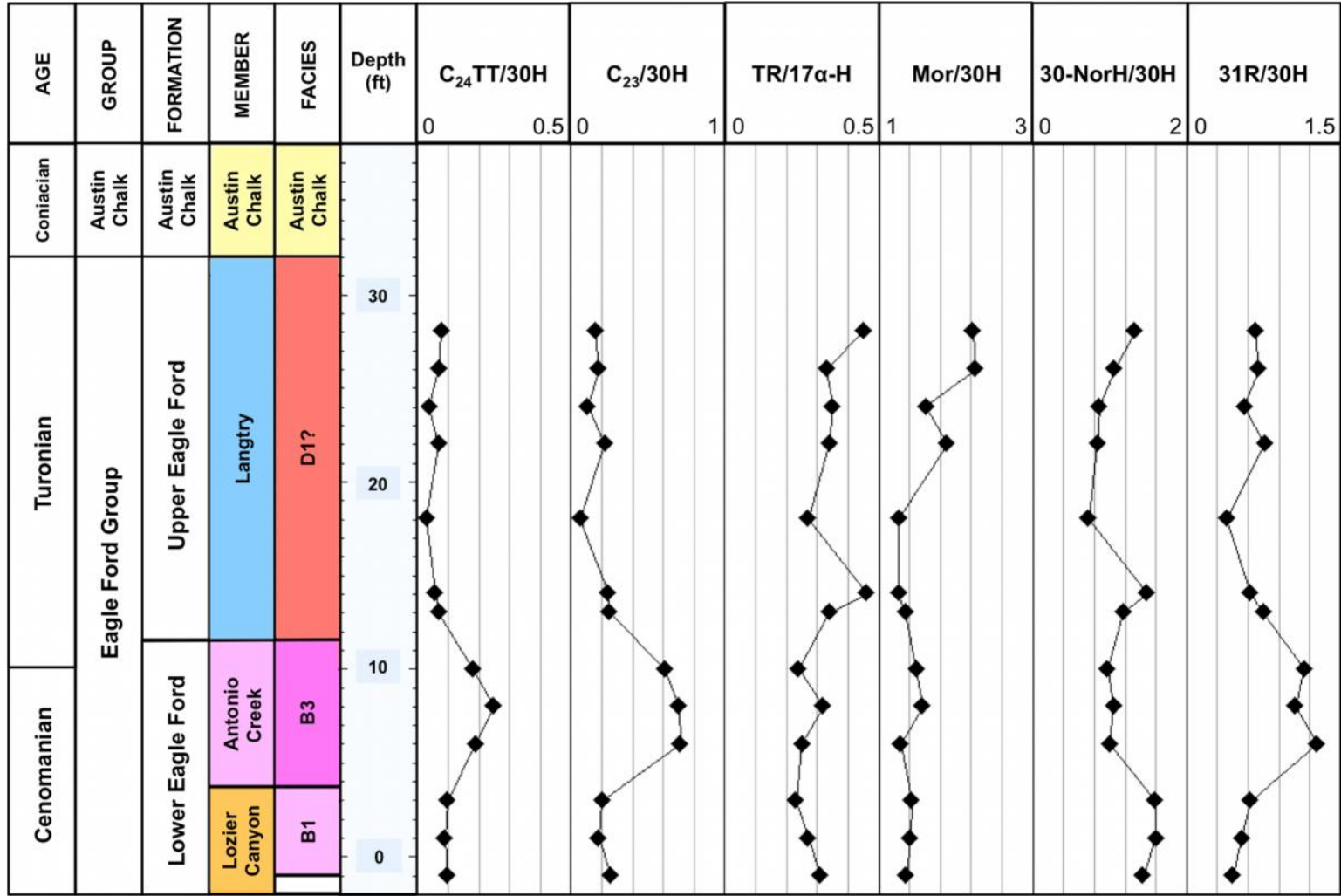
Ferguson McKnight #526-1H core



Fasken "A" #1

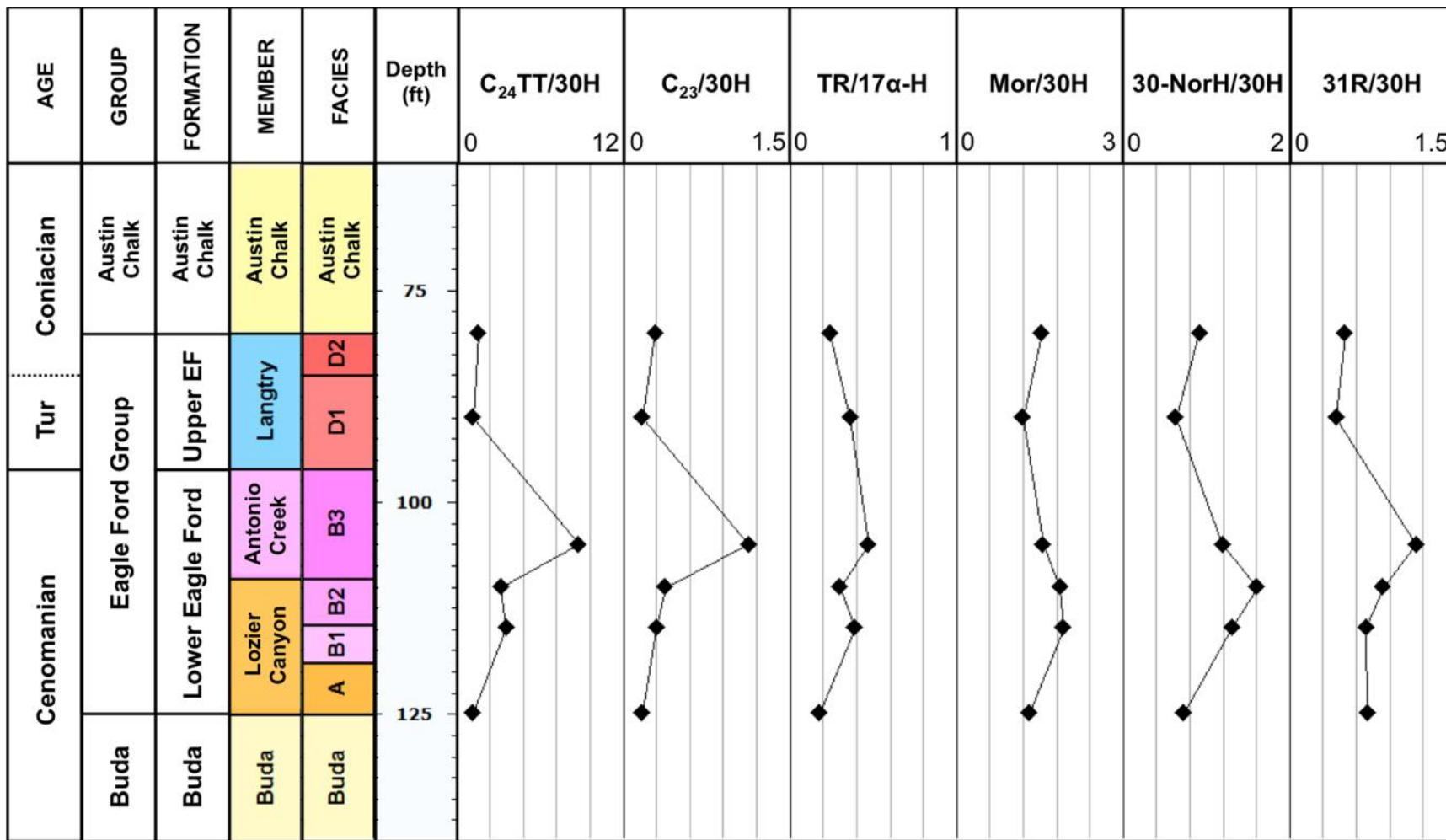


Bouldin Creek outcrop



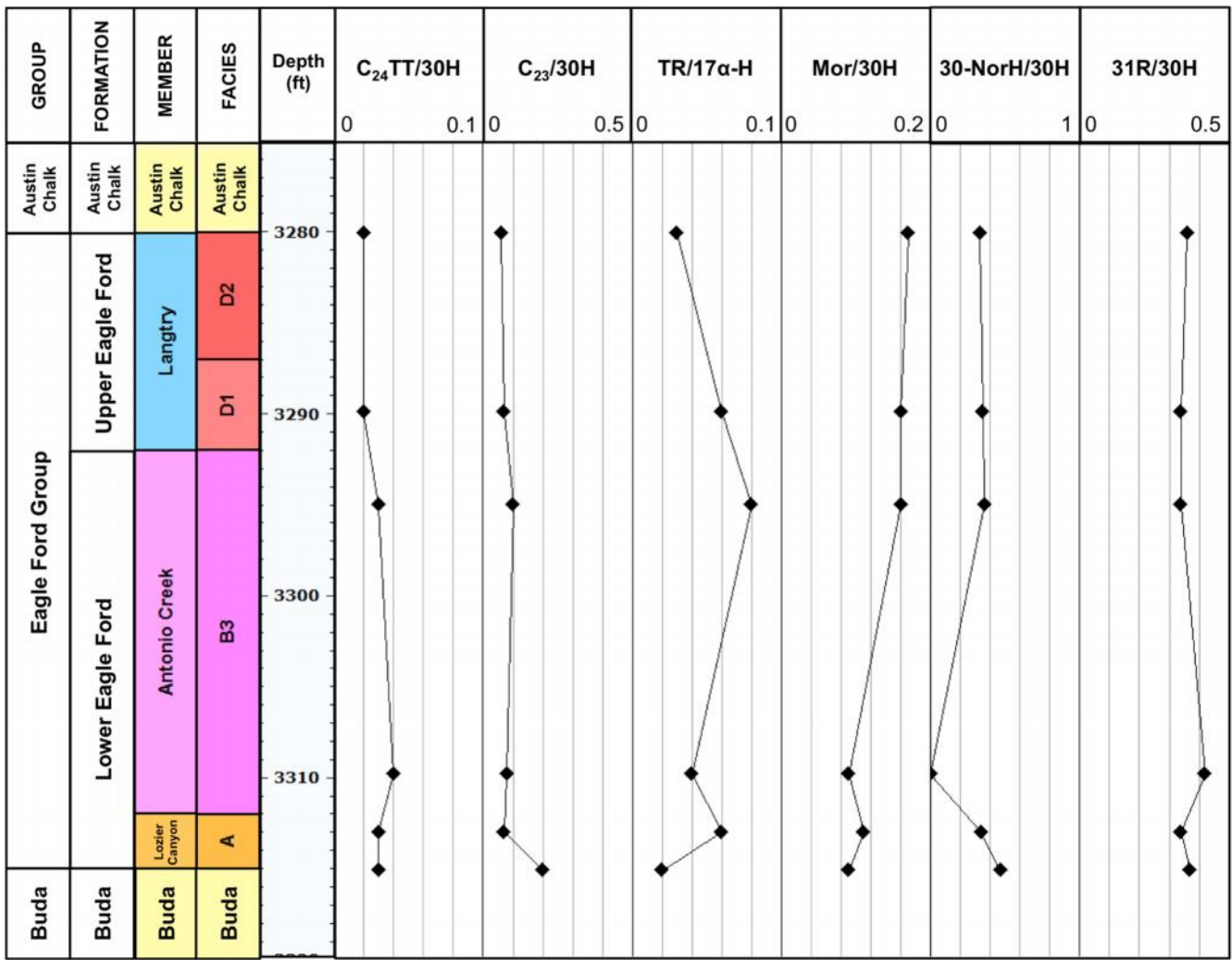
ACC #1 core

191

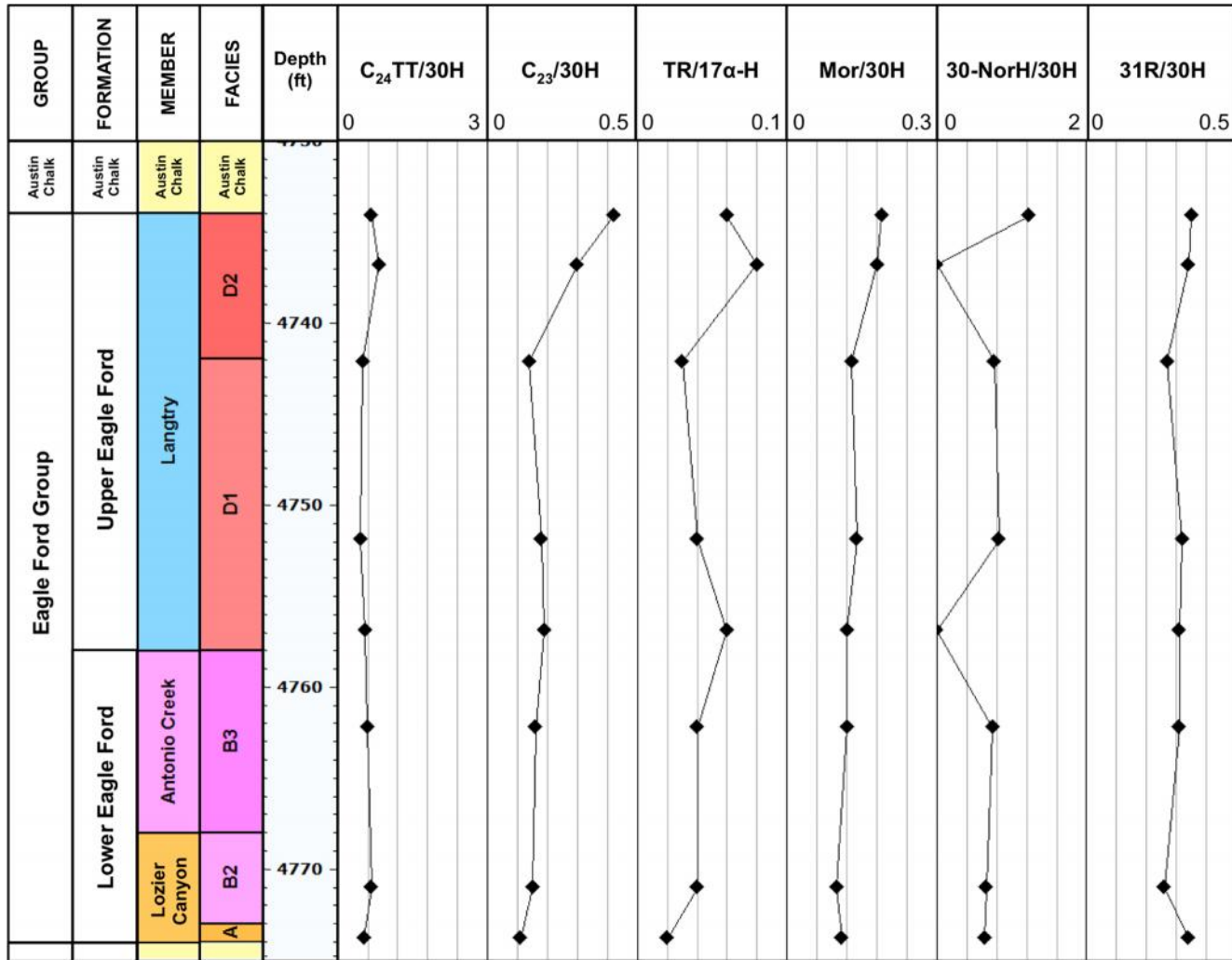


W. Brechtel #1 core

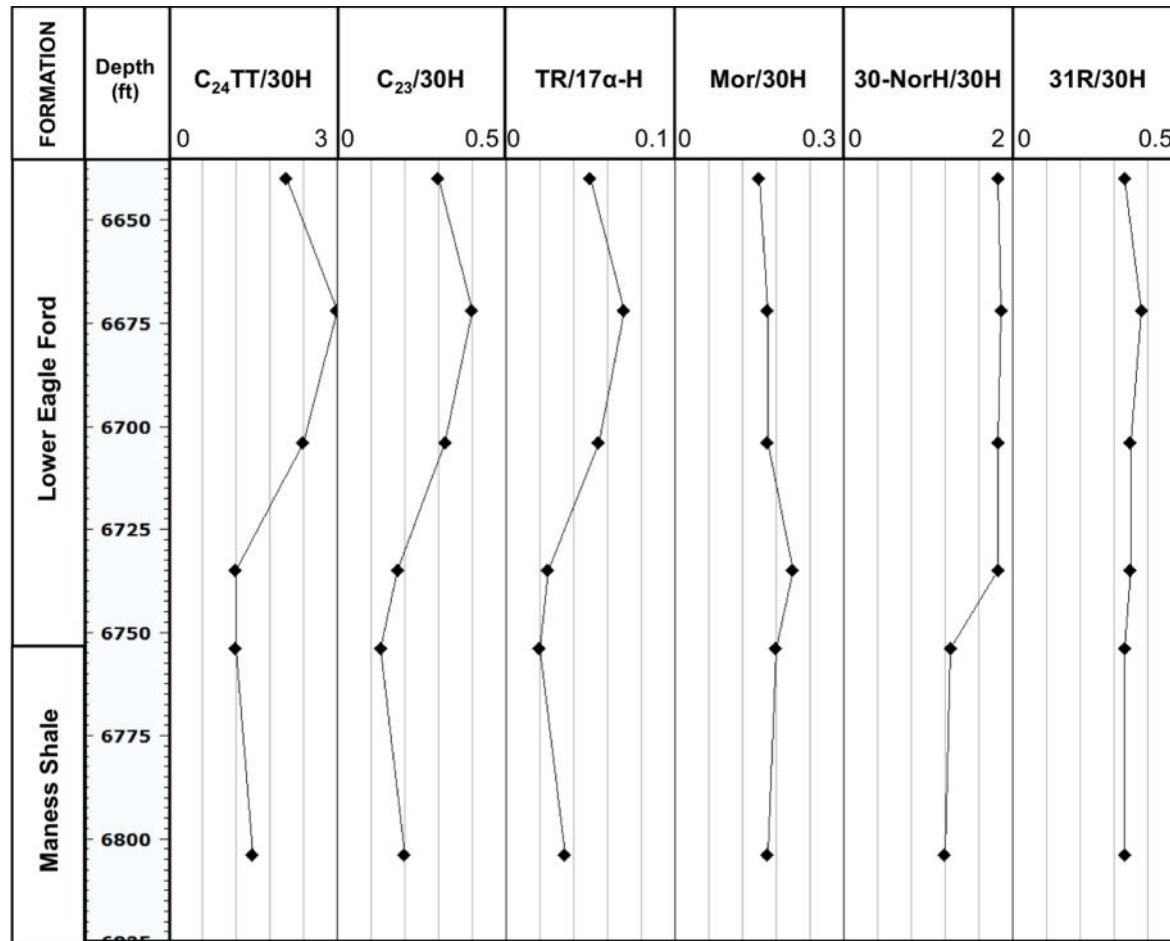
861



C. J. Hendershot #1 core



Lily Hoppess #1 core



H. Geochemical ratios of steranes for the branched and cyclic fractions (B&C) of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined)

Lozier Canyon outcrop

SAMPLE	C₂₇%	C₂₈%	C₂₉%	C₂₉ 20S/ (20S+20R)	C₂₉ / (+)	Preg/ Steranes	C₂₇ Dia/ C₂₇ Ster	C₂₇ Dia/ (Dia+Reg)	Sterane Index
EFLC-28	43.57	24.24	32.19	0.48	0.55	0.67	0.33	0.31	0.05
EFLC-26	42.55	24.17	33.29	0.48	0.54	0.70	0.32	0.31	0.05
EFLC-25	42.23	24.22	33.55	0.48	0.53	0.46	0.27	0.27	0.05
EFLC-20	37.68	30.06	32.26	0.49	0.54	0.61	0.20	0.23	0.05
EFLC-18	36.73	30.98	32.29	0.49	0.54	0.66	0.20	0.23	0.06
EFLC-14	36.87	31.62	31.51	0.48	0.54	0.67	0.17	0.21	0.06
EFLC-12	39.10	28.22	32.68	0.49	0.56	0.98	0.16	0.19	0.05
EFLC-10	39.27	27.22	33.51	0.51	0.55	1.03	0.09	0.11	0.05
EFLC-8	38.84	27.79	33.37	0.50	0.55	1.14	0.09	0.11	0.05
EFLC-6	38.59	28.35	33.05	0.51	0.54	0.95	0.10	0.12	0.06
EFLC-4	40.67	28.52	30.81	0.46	0.54	0.00	0.13	0.16	0.05
EFLC-1	38.47	28.11	33.42	0.52	0.52	0.07	0.27	0.27	0.06

Ferguson McKnight #526-1H core

SAMPLE	C₂₇%	C₂₈%	C₂₉%	C₂₉ 20S/ (20S+20R)	C₂₉ / (+)	Preg/ Steranes	C₂₇ Dia/ C₂₇ Ster	C₂₇ Dia/ (Dia+Reg)	Sterane Index
EFMK-1	44.91	28.47	26.62	0.44	0.60	1.99	0.51	0.43	0.06
EFMK-4	43.64	28.67	27.69	0.44	0.59	2.88	0.52	0.45	0.06
EFMK-8	48.09	27.22	24.68	0.47	0.60	1.49	0.39	0.38	0.05
EFMK-12	44.02	28.94	27.04	0.46	0.59	3.16	0.51	0.44	0.05
EFMK-15	45.60	27.88	26.52	0.43	0.61	2.67	0.57	0.47	0.05
EFMK-17	48.55	25.85	25.60	0.41	0.60	1.93	0.53	0.46	0.06
EFMK-20	48.24	25.72	26.05	0.41	0.60	1.88	0.55	0.46	0.06
EFMK-23	47.25	25.63	27.12	0.43	0.63	3.37	0.60	0.48	0.06
EFMK-24	47.46	25.95	26.59	0.42	0.63	2.50	0.60	0.48	0.06
EFMK-25	49.86	27.15	23.00	0.36	0.63	2.92	0.75	0.53	0.06
EFMK-26	44.86	29.52	25.62	0.48	0.59	5.52	0.63	0.48	0.06

Fasken "A" #1 core

SAMPLE	C ₂₇ %	C ₂₈ %	C ₂₉ %	C ₂₉ 20S/ (20S+20R)	C ₂₉ / (+)	Preg/ Steranes	C ₂₇ Dia/ C ₂₇ Ster	C ₂₇ Dia/ (Dia+Reg)	Sterane Index
EFFA-1	44.65	27.48	27.87	0.40	0.48	5.46	0.42	0.40	0.04
EFFA-3	44.88	26.65	28.47	0.37	0.53	5.00	0.43	0.39	0.06
EFFA-5	45.51	25.89	28.59	0.39	0.51	5.08	0.40	0.37	0.03
EFFA-11	44.61	27.43	27.95	0.36	0.52	4.18	0.45	0.41	0.04
EFFA-17	43.94	26.29	29.77	0.38	0.47	4.82	0.41	0.38	0.03
EFFA-18	42.11	29.24	28.65	0.41	0.48	4.73	0.41	0.38	0.03
EFFA-20	42.78	27.62	29.60	0.38	0.50	5.49	0.43	0.38	0.03
EFFA-21	43.71	28.29	28.00	0.39	0.50	4.96	0.40	0.36	0.03
EFFA-23	43.79	28.93	27.28	0.39	0.52	4.82	0.41	0.37	0.03
EFFA-24	42.82	27.74	29.44	0.40	0.48	5.59	0.42	0.36	0.03
EFFA-28	44.33	27.18	28.49	0.42	0.50	5.42	0.41	0.36	0.04
EFFA-32	42.89	28.89	28.22	0.36	0.48	6.32	0.43	0.39	0.03
EFFA-37	43.96	29.96	26.08	0.36	0.54	5.62	0.40	0.36	0.04
EFFA-38	45.53	26.16	28.31	0.37	0.51	5.27	0.44	0.39	0.03
EFFA-40	45.71	25.90	28.39	0.37	0.51	4.56	0.41	0.37	0.03

Bouldin Creek outcrop

SAMPLE	C₂₇%	C₂₈%	C₂₉%	C₂₉ 20S/ (20S+20R)	C₂₉ / (+)	Preg/ Steranes	C₂₇ Dia/ C₂₇ Ster	C₂₇ Dia/ (Dia+Reg)	Sterane Index
EFBC-5	36.50	40.67	22.83	ND	0.31	0.08	ND	0.01	0.05
EFBC-7	37.59	42.03	20.38	ND	0.30	0.08	ND	0.01	0.05
EFBC-9	34.98	43.15	21.87	ND	0.31	0.09	ND	0.01	0.05
EFBC-11	36.11	43.30	20.59	ND	0.34	0.09	0.01	0.01	0.05
EFBC-15	36.58	38.78	24.64	ND	0.33	0.11	0.01	0.01	0.06
EFBC-19	36.15	33.58	30.28	0.04	0.34	0.10	0.01	0.01	0.06
EFBC-20	38.47	34.73	26.80	ND	0.35	0.26	0.01	0.02	0.05
EFBC-23	34.05	42.42	23.54	0.04	0.36	0.21	0.01	0.02	0.05
EFBC-25	32.88	44.76	22.36	ND	0.38	0.25	0.01	0.02	0.05
EFBC-27	36.35	39.13	24.52	ND	0.37	0.23	0.02	0.02	0.05
EFBC-30	38.99	37.58	23.43	ND	0.34	0.07	0.01	0.01	0.05
EFBC-32	42.18	31.28	26.54	0.03	0.32	0.05	0.01	0.01	0.06
EFBC-34	33.10	43.50	23.40	0.03	0.32	0.05	0.01	0.01	0.05

ACC#1 core

SAMPLE	C₂₇%	C₂₈%	C₂₉%	C₂₉ 20S/ (20S+20R)	C₂₉ / (+)	Preg/ Steranes	C₂₇ Dia/ C₂₇ Ster	C₂₇ Dia/ (Dia+Reg)	Sterane Index
EFAC-1	44.88	28.34	26.78	ND	0.28	0.35	ND	ND	0.05
EFAC-3	38.73	36.51	24.76	ND	0.27	0.14	ND	ND	0.05
EFAC-6	42.59	34.92	22.49	ND	0.26	0.25	ND	ND	0.05
EFAC-7	38.98	38.56	22.46	ND	0.26	0.07	ND	ND	0.04
EFAC-8	39.98	39.03	20.99	ND	0.26	0.07	ND	ND	0.04
EFAC-10	36.30	31.90	31.80	ND	0.22	0.08	ND	ND	0.05

W. Brechtel #1 core

SAMPLE	C₂₇%	C₂₈%	C₂₉%	C₂₉ 20S/ (20S+20R)	C₂₉ / (+)	Preg/ Steranes	C₂₇ Dia/ C₂₇ Ster	C₂₇ Dia/ (Dia+Reg)	Sterane Index
EFWB-1	44.00	29.54	26.46	0.36	0.38	0.18	0.05	0.06	0.08
EFWB-3	37.05	38.00	24.95	0.38	0.36	0.28	0.08	0.10	0.05
EFWB-4	36.55	37.41	26.04	0.38	0.35	0.29	0.08	0.10	0.05
EFWB-7	40.68	33.62	25.70	0.45	0.47	0.36	0.07	0.09	0.06
EFWB-8	40.96	34.76	24.28	0.44	0.44	0.30	0.09	0.11	0.06
EFWB-9	38.92	32.76	28.31	0.47	0.48	0.78	0.05	0.07	0.06

C.J. Hendershot #1 core

SAMPLE	C₂₇%	C₂₈%	C₂₉%	C₂₉ 20S/ (20S+20R)	C₂₉ / (+)	Preg/ Steranes	C₂₇ Dia/ C₂₇ Ster	C₂₇ Dia/ (Dia+Reg)	Sterane Index
EFHE-1	46.16	27.62	26.22	0.20	0.26	0.09	0.11	0.13	0.06
EFHE-2	42.57	29.58	27.85	0.18	0.25	0.08	0.05	0.07	0.06
EFHE-3	40.76	31.23	28.01	0.21	0.23	0.09	0.09	0.11	0.06
EFHE-5	37.72	35.91	26.37	0.28	0.23	0.17	0.16	0.21	0.06
EFHE-6	40.85	31.32	27.83	0.35	0.26	0.31	0.17	0.21	0.05
EFHE-7	39.82	31.77	28.41	0.35	0.24	0.21	0.17	0.19	0.05
EFHE-9	39.53	31.93	28.55	0.47	0.29	0.33	0.08	0.10	0.06
EFHE-10	42.37	26.13	31.50	0.45	0.27	0.28	0.34	0.34	0.07

Oils and condensates

SAMPLE	C₂₇%	C₂₈%	C₂₉%	C₂₉ 20S/ (20S+20R)	C₂₉ / (+)	Preg/ Steranes	C₂₇ Dia/ C₂₇ Ster	C₂₇ Dia/ (Dia+Reg)	Sterane Index
NWMK-533.2	37.52	39.45	23.03	0.42	0.60	2.51	0.47	0.41	0.06
NFMK-526-1H	42.73	32.12	25.15	0.38	0.61	4.56	0.70	0.51	0.06
MJOG-UEF	59.99	19.28	20.74	0.30	0.65	2.40	0.76	0.52	0.01
MJOG-LEF	62.16	20.28	17.56	0.41	0.69	2.06	0.72	0.52	0.04
GRAF-1-110	54.40	22.71	22.89	0.26	0.36	1.36	0.96	0.58	0.09

I. Geochemical ratios of terpanes for the branched and cyclic fractions (B&C) of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined)

Lozier Canyon outcrop

Sample	Steranes/ 17 -H	TR/ 17 -H	C ₂₂ /C ₂₁ TR	C ₂₄ /C ₂₃ TR	C ₂₆ /C ₂₅ TR	C ₂₃ / 30H	C ₂₄ TT/ 30H	30-NorH/ 30H	Mor/ 30H	31R/ 30H	35S/ 34S	HH Index	Ts/ Ts+Tm	C ₃₁ 22S/ (22S+22R)
EFLC-28	0.66	0.03	0.24	0.57	0.96	0.23	0.05	0.6	0.06	0.46	1.20	0.11	0.32	0.62
EFLC-26	0.63	0.03	0.32	0.53	0.76	0.27	0.04	0.62	0.06	0.47	1.01	0.09	0.20	0.61
EFLC-25	0.51	0.02	0.36	0.59	0.89	0.14	0.03	0.58	0.06	0.46	1.16	0.11	0.19	0.61
EFLC-20	0.7	0.04	0.54	0.43	0.71	0.28	0.04	0.65	0.06	0.53	1.46	0.13	0.19	0.61
EFLC-18	0.68	0.04	0.63	0.33	0.64	0.39	0.05	0.69	0.06	0.57	1.42	0.14	0.20	0.62
EFLC-14	0.56	0.05	0.84	0.29	0.52	0.55	0.06	0.84	0.06	0.56	1.44	0.14	0.21	0.62
EFLC-12	0.58	0.07	0.88	0.29	0.52	0.77	0.08	0.83	0.06	0.52	1.46	0.14	0.23	0.61
EFLC-10	0.42	0.06	0.92	0.29	0.59	0.59	0.06	0.82	0.06	0.46	1.39	0.14	0.21	0.6
EFLC-8	0.42	0.06	0.88	0.29	0.61	0.65	0.06	0.84	0.06	0.46	1.36	0.13	0.22	0.61
EFLC-6	0.42	0.06	0.9	0.32	0.62	0.56	0.07	0.84	0.06	0.47	1.42	0.14	0.22	0.61
EFLC-4	0.35	0.05	0.81	0.26	0.63	0.75	0.07	0.88	0.06	0.49	1.38	0.15	0.26	0.61
EFLC-1	0.82	0.03	0.35	0.95	1.11	0.14	0.04	0.56	0.08	0.43	1.44	0.21	0.27	0.6

Ferguson McKnight #526-1H core

Sample	Steranes/ 17 -H	TR/ 17 -H	C₂₂/C₂₁ TR	C₂₄/C₂₃ TR	C₂₆/C₂₅ TR	C₂₃/ 30H	C₂₄TT/ 30H	30-NorH/ 30H	Mor/ 30H	31R/ 30H	35S/ 34S	HH Index	Ts/ Ts+Tm	C₃₁ 22S/ (22S+22R)
EFMK-1	2.15	0.43	0.30	0.62	0.48	2.74	0.53	1.20	0.10	0.45	ND	ND	0.49	0.46
EFMK-4	3.68	0.86	0.22	0.65	0.55	3.49	0.80	0.65	0.13	0.28	ND	ND	0.56	0.57
EFMK-8	3.84	0.90	0.24	0.62	0.64	3.62	0.88	0.66	0.11	0.23	ND	ND	0.63	0.59
EFMK-12	3.12	0.73	0.27	0.60	0.67	3.85	0.86	0.77	0.11	0.28	ND	ND	0.60	0.60
EFMK-15	3.10	0.71	0.29	0.63	0.70	3.19	0.68	0.73	0.14	0.35	ND	ND	0.66	0.54
EFMK-17	2.67	0.56	0.26	0.57	0.79	3.13	0.58	0.79	0.09	0.42	ND	ND	0.65	0.49
EFMK-20	3.79	0.93	0.25	0.64	0.65	4.35	0.96	0.81	0.12	0.30	ND	ND	0.66	0.60
EFMK-23	4.25	1.17	0.25	0.64	0.66	4.98	1.08	0.71	0.14	0.28	ND	ND	0.66	0.56
EFMK-24	4.98	1.45	0.25	0.67	0.64	5.35	1.21	0.64	0.06	0.29	1.11	0.12	0.66	0.53
EFMK-25	5.80	1.49	0.37	0.72	0.68	7.76	1.71	1.23	0.48	0.69	ND	ND	0.68	0.52
EFMK-26	3.68	2.44	0.41	0.74	0.65	9.38	2.07	1.17	0.26	0.53	ND	ND	0.57	0.37

Fasken "A" #1 core

Sample	Steranes/ 17 -H	TR/ 17 -H	C ₂₂ /C ₂₁ TR	C ₂₄ /C ₂₃ TR	C ₂₆ /C ₂₅ TR	C ₂₃ / 30H	C ₂₄ TT/ 30H	30-NorH/ 30H	Mor/ 30H	31R/ 30H	35S/ 34S	HH Index	Ts/ Ts+Tm	C ₃₁ 22S/ (22S+22R)
EFFA-1	0.46	0.10	0.36	0.45	0.94	2.68	0.23	1.26	0.21	0.28	ND	ND	0.53	0.57
EFFA-3	0.46	0.13	0.35	0.44	0.91	2.65	0.19	1.23	0.19	0.32	ND	ND	0.41	0.53
EFFA-5	0.46	0.12	0.36	0.45	0.92	2.49	0.21	1.24	0.20	0.28	ND	ND	0.37	0.57
EFFA-11	0.48	0.12	0.38	0.45	0.92	2.29	0.17	1.18	0.19	0.27	ND	ND	0.40	0.54
EFFA-17	0.44	0.12	0.35	0.47	0.92	2.32	0.17	1.15	0.21	0.24	ND	ND	0.38	0.57
EFFA-18	0.47	0.11	0.36	0.43	0.96	2.37	0.17	1.19	0.20	0.28	ND	ND	0.38	0.55
EFFA-20	0.45	0.10	0.36	0.43	0.87	2.21	0.16	1.02	0.19	0.21	ND	ND	0.40	0.58
EFFA-21	0.47	0.11	0.31	0.44	1.05	2.42	0.17	1.08	0.19	0.23	ND	ND	0.41	0.56
EFFA-23	0.47	0.11	0.34	0.43	0.93	2.35	0.17	1.02	0.19	0.23	ND	ND	0.39	0.57
EFFA-24	0.47	0.11	0.33	0.42	0.95	2.71	0.18	1.19	0.20	0.25	ND	ND	0.37	0.58
EFFA-28	0.48	0.11	0.32	0.43	0.92	2.62	0.18	1.09	0.19	0.23	ND	ND	0.35	0.54
EFFA-32	0.45	0.11	0.32	0.44	0.96	2.59	0.17	1.07	0.15	0.21	ND	ND	0.40	0.57
EFFA-37	0.46	0.11	0.30	0.43	0.95	2.64	0.17	1.05	0.18	0.26	ND	ND	0.40	0.51
EFFA-38	0.44	0.12	0.31	0.45	0.96	2.40	0.16	1.10	0.19	0.24	ND	ND	0.34	0.57
EFFA-40	0.47	0.12	0.34	0.45	0.87	2.26	0.17	1.06	0.20	0.24	ND	ND	0.38	0.56

Bouldin Creek outcrop

Sample	Steranes/ 17 -H	TR/ 17 -H	C₂₂/C₂₁ TR	C₂₄/C₂₃ TR	C₂₆/C₂₅ TR	C₂₃/ 30H	C₂₄TT/ 30H	30-NorH/ 30H	Mor/ 30H	31R/ 30H	35S/ 34S	HH Index	Ts/ Ts+Tm	C₃₁ 22S/ (22S+22R)
EFBC-5	13.23	0.45	ND	0.47	0.82	0.16	0.08	1.33	2.22	0.67	ND	ND	0.08	0.42
EFBC-7	13.69	0.33	ND	0.50	0.98	0.18	0.07	1.06	2.26	0.70	ND	ND	0.07	0.31
EFBC-9	10.01	0.35	ND	0.52	0.86	0.11	0.04	0.87	1.61	0.56	ND	ND	0.07	0.31
EFBC-11	13.90	0.34	ND	0.40	0.70	0.22	0.07	0.84	1.89	0.76	ND	ND	0.06	0.34
EFBC-15	7.53	0.27	ND	0.71	1.06	0.06	0.03	0.73	1.26	0.39	ND	ND	0.06	0.30
EFBC-19	11.00	0.46	ND	0.37	1.16	0.24	0.06	1.49	1.26	0.62	ND	ND	0.09	0.35
EFBC-20	8.03	0.34	ND	0.25	1.42	0.25	0.07	1.18	1.35	0.75	0.06	0.11	0.04	0.23
EFBC-23	17.80	0.24	ND	0.21	1.10	0.61	0.18	0.97	1.49	1.15	ND	ND	0.06	0.15
EFBC-25	20.06	0.32	ND	0.17	0.90	0.70	0.25	1.05	1.57	1.06	ND	ND	0.05	0.15
EFBC-27	13.95	0.25	ND	0.15	0.89	0.71	0.19	1.00	1.29	1.27	ND	ND	0.07	0.11
EFBC-30	10.83	0.23	ND	0.45	1.21	0.20	0.10	1.59	1.43	0.62	ND	ND	0.05	0.12
EFBC-32	10.61	0.27	ND	0.33	0.87	0.18	0.09	1.60	1.41	0.54	ND	ND	0.05	0.10
EFBC-34	16.68	0.31	ND	0.27	0.60	0.26	0.10	1.43	1.36	0.45	0.06	0.13	0.04	0.13

ACC#1 core

Sample	Steranes/ 17 -H	TR/ 17 -H	C₂₂/C₂₁ TR	C₂₄/C₂₃ TR	C₂₆/C₂₅ TR	C₂₃/ 30H	C₂₄TT/ 30H	30-NorH/ 30H	Mor/ 30H	31R/ 30H	35S/ 34S	HH Index	Ts/ Ts+Tm	C₃₁ 22S/ (22S+22R)
EFAC-1	1.89	0.24	0.19	0.50	1.13	0.29	0.04	0.92	1.54	0.50	ND	ND	0.05	0.30
EFAC-3	4.35	0.36	0.17	0.52	0.69	0.17	0.03	0.63	1.19	0.42	ND	ND	0.07	0.31
EFAC-6	6.31	0.47	0.16	0.34	0.69	1.13	0.22	1.19	1.56	1.14	ND	ND	0.09	0.26
EFAC-7	7.82	0.30	0.17	0.38	0.75	0.38	0.08	1.61	1.86	0.84	ND	ND	0.04	0.33
EFAC-8	9.11	0.39	0.22	0.59	0.53	0.30	0.09	1.31	1.92	0.69	ND	ND	0.03	0.40
EFAC-10	3.86	0.18	0.18	0.36	1.68	0.17	0.03	0.73	1.32	0.70	ND	ND	0.08	0.31

W. Brechtel #1 core

Sample	Steranes/ 17 -H	TR/ 17 -H	C₂₂/C₂₁ TR	C₂₄/C₂₃ TR	C₂₆/C₂₅ TR	C₂₃/ 30H	C₂₄TT/ 30H	30-NorH/ 30H	Mor/ 30H	31R/ 30H	35S/ 34S	HH Index	Ts/ Ts+Tm	C₃₁ 22S/ (22S+22R)
EFWB-1	0.56	0.03	1.57	1.50	1.16	0.06	0.02	0.33	0.17	0.36	1.19	0.13	0.20	0.56
EFWB-3	1.34	0.06	0.39	0.89	0.84	0.07	0.02	0.35	0.16	0.34	0.87	0.14	0.18	0.59
EFWB-4	1.52	0.08	0.38	0.79	0.91	0.10	0.03	0.36	0.16	0.34	0.60	0.13	0.21	0.59
EFWB-7	0.85	0.04	2.77	0.96	1.13	0.08	0.04	0.00	0.09	0.42	1.23	0.10	0.24	0.61
EFWB-8	1.18	0.06	0.26	1.32	0.64	0.07	0.03	0.34	0.11	0.34	0.91	0.11	0.23	0.61
EFWB-9	0.72	0.02	0.18	0.59	1.15	0.20	0.03	0.47	0.09	0.37	1.11	0.10	0.26	0.61

C.J. Hendershot #1 core

Sample	Steranes/ 17 -H	TR/ 17 -H	C ₂₂ /C ₂₁ TR	C ₂₄ /C ₂₃ TR	C ₂₆ /C ₂₅ TR	C ₂₃ / 30H	C ₂₄ TT/ 30H	30-NorH/ 30H	Mor/ 30H	31R/ 30H	35S/ 34S	HH Index	Ts/ Ts+Tm	C ₃₁ 22S/ (22S+22R)
EFHE-1	3.34	0.06	0.25	0.68	0.72	0.42	0.10	1.23	0.19	0.35	1.56	0.19	0.29	0.53
EFHE-2	4.51	0.08	0.15	0.82	0.69	0.30	0.10	0.00	0.18	0.34	1.40	0.17	0.25	0.53
EFHE-3	2.23	0.03	0.14	0.50	0.97	0.14	0.04	0.77	0.13	0.27	0.76	0.12	0.28	0.56
EFHE-5	2.77	0.04	0.18	0.46	0.68	0.18	0.04	0.83	0.14	0.32	0.94	0.14	0.21	0.59
EFHE-6	3.61	0.06	0.15	0.55	0.71	0.19	0.05	0.00	0.12	0.31	0.92	0.13	0.22	0.61
EFHE-7	2.20	0.04	0.16	0.59	0.38	0.16	0.04	0.75	0.12	0.31	0.94	0.12	0.24	0.60
EFHE-9	1.21	0.04	0.13	0.68	0.61	0.15	0.05	0.66	0.10	0.26	0.80	0.10	0.19	0.61
EFHE-10	0.99	0.02	0.27	0.53	1.34	0.11	0.03	0.64	0.11	0.34	1.41	0.15	0.39	0.61

212

Lily Hoppess #1 core

Sample	Steranes/ 17 -H	TR/ 17 -H	C ₂₂ /C ₂₁ TR	C ₂₄ /C ₂₃ TR	C ₂₆ /C ₂₅ TR	C ₂₃ / 30H	C ₂₄ TT/ 30H	30-NorH/ 30H	Mor/ 30H	31R/ 30H	35S/ 34S	HH Index	Ts/ Ts+Tm	C ₃₁ 22S/ (22S+22R)
EFLH-1	0.68	0.10	0.20	0.46	0.86	0.30	0.07	0.55	0.10	0.20	0.34	0.05	0.51	0.57
EFLH-5	0.62	0.14	0.23	0.49	0.72	0.40	0.10	0.56	0.11	0.23	0.41	0.05	0.48	0.55
EFLH-9	0.57	0.11	0.21	0.48	0.94	0.32	0.08	0.55	0.11	0.21	0.42	0.06	0.51	0.58
EFLH-13	0.41	0.05	0.29	0.47	1.40	0.18	0.04	0.55	0.14	0.21	0.40	0.06	0.61	0.61
EFLH-15	0.59	0.04	0.30	0.54	1.44	0.13	0.04	0.38	0.12	0.20	0.34	0.05	0.70	0.60
EFLH-20	0.73	0.07	0.22	0.50	1.02	0.20	0.05	0.36	0.11	0.20	0.40	0.06	0.64	0.59

Oils and condensates

Sample	Steranes/ 17 -H	TR/ 17 -H	C₂₂/C₂₁ TR	C₂₄/C₂₃ TR	C₂₆/C₂₅ TR	C₂₃/ 30H	C₂₄TT/ 30H	30-NorH/ 30H	Mor/ 30H	31R/ 30H	35S/ 34S	HH Index	Ts/ Ts+Tm	C₃₁ 22S/ (22S+22R)
NWMK-533.2	3.19	0.61	0.22	0.76	0.63	1.98	0.42	0.30	0.07	0.37	0.58	0.04	0.46	0.50
NFMK-526-1H	7.87	3.71	0.28	0.86	0.58	7.75	1.99	0.88	ND	0.31	ND	ND	0.52	ND
MJOG-UEF	13.20	8.13	0.70	0.66	0.63	ND	ND	ND	ND	ND	ND	ND	0.65	ND
MJOG-LEF	10.57	3.53	0.24	0.72	0.64	12.87	3.27	2.29	ND	ND	ND	ND	0.66	ND
GRES-1-110	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND

J. Aryl isoprenoids ratio (AIR) for the branched and cyclic fractions (B&C) of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined)

Lozier Canyon outcrop

Sample	AIR
EFLC-28	1.04
EFLC-26	1.37
EFLC-25	0.66
EFLC-18	0.54
EFLC-14	0.83
EFLC-12	0.84
EFLC-10	1.33
EFLC-8	1.21
EFLC-6	0.37
EFLC-4	0.56
EFLC-1	0.74

Ferguson McKnight #526-1H core

Sample	AIR
EFMK-1	1.72
EFMK-4	4.61
EFMK-8	3.72
EFMK-12	4.25
EFMK-15	2.95
EFMK-17	1.90
EFMK-20	3.88
EFMK-23	2.24
EFMK-24	3.48
EFMK-25	1.68
EFMK-26	1.65

Fasken "A" #1 core

Sample	AIR
EFFA-1	1.70
EFFA-3	3.07
EFFA-5	2.02
EFFA-11	1.23
EFFA-17	2.15
EFFA-18	2.26
EFFA-20	1.42
EFFA-21	2.20
EFFA-23	1.95
EFFA-24	3.24
EFFA-28	4.10
EFFA-32	2.90
EFFA-37	4.53
EFFA-38	3.21
EFFA-40	2.25

Bouldin Creek outcrop

Sample	AIR
EFBC-5	1.01
EFBC-7	0.54
EFBC-9	1.14
EFBC-11	0.80
EFBC-15	0.79
EFBC-19	0.87
EFBC-20	0.62
EFBC-23	0.65
EFBC-25	2.46
EFBC-27	2.81
EFBC-30	1.37
EFBC-32	1.60
EFBC-34	1.43

ACC #1 core

Sample	AIR
EFAC-1	0.32
EFAC-3	0.79
EFAC-6	0.49
EFAC-7	0.33
EFAC-8	0.38
EFAC-10	0.48

W. Brechtel #1 core

Sample	AIR
EFWB-1	1.08
EFWB-3	2.11
EFWB-4	2.26
EFWB-7	2.00
EFWB-8	2.74
EFWB-9	2.63

C.J. Hendershot #1 core

Sample	AIR
EFHE-1	1.44
EFHE-2	1.42
EFHE-3	2.34
EFHE-5	1.87
EFHE-6	2.07
EFHE-7	1.98
EFHE-9	1.76
EFHE-10	0.62

Lily Hoppess #1 core

Sample	AIR
EFLH-1	2.43
EFLH-5	1.90
EFLH-9	2.19
EFLH-13	2.65
EFLH-15	2.95
EFLH-20	1.99

Oils and condensates

Sample	AIR
NWMK-533.2	2.80
NFMK-526-1H	2.05
MJOG-UEF	2.27
MJOG-LEF	2.21
GRES-1-110	3.45

K. Geochemical ratios for aromatic biomarkers of the Eagle Ford Shale bitumen, oil, and condensate samples (ND = not determined)

Lozier Canyon outcrop

Sample	MPI 1	MPI 2	MA(I)/ MA(I+II)	TA(I)/ TA(I+II)	C₂₆ TA (20S/20S+20R)	TA28/ (MA29+TA28)	%C₂₇ MAS	%C₂₈ MAS	%C₂₉ MAS
EFLC-28	0.22	0.28	0.06	0.10	0.17	0.79	15.58	70.17	14.25
EFLC-26	0.13	0.18	0.10	0.10	0.17	0.78	15.68	69.81	14.51
EFLC-25	0.18	0.25	0.07	0.08	0.17	0.74	15.28	69.41	15.31
EFLC-20	0.18	0.24	0.04	0.06	0.16	0.78	16.04	70.91	13.05
EFLC-18	0.10	0.15	0.10	0.16	0.15	0.55	16.26	68.33	15.41
EFLC-14	0.16	0.22	0.18	0.29	0.15	0.33	17.72	68.37	13.91
EFLC-12	0.52	0.56	0.13	0.14	0.16	0.71	18.27	70.02	11.71
EFLC-10	0.56	0.63	0.18	0.14	0.17	0.76	19.03	68.22	12.75
EFLC-8	0.39	0.48	0.21	0.29	0.18	0.71	19.05	68.11	12.83
EFLC-6	0.43	0.52	0.14	0.17	0.19	0.63	19.59	66.03	14.38
EFLC-4	0.12	0.17	0.16	0.35	0.17	0.47	19.39	68.21	12.40
EFLC-1	0.59	0.57	0.05	0.11	0.21	0.30	18.49	71.25	10.26

Ferguson McKnight #526-1H core

Sample	MPI 1	MPI 2	MA(I)/ MA(I+II)	TA(I)/ TA(I+II)	C₂₆ TA (20S/20S+20R)	TA28/ (MA29+TA28)	%C₂₇ MAS	%C₂₈ MAS	%C₂₉ MAS
EFMK-1	0.72	0.78	0.46	0.53	0.21	0.96	30.03	52.86	17.11
EFMK-4	0.68	0.77	0.53	0.64	0.20	0.96	31.38	51.03	17.59
EFMK-8	0.73	0.80	0.57	0.69	0.20	0.96	29.90	51.33	18.77
EFMK-12	0.68	0.78	0.55	0.68	0.20	0.94	29.19	55.34	15.47
EFMK-15	0.72	0.81	0.43	0.62	0.17	0.96	26.19	56.77	17.04
EFMK-17	0.72	0.80	0.37	0.61	0.19	0.95	26.21	54.68	19.11
EFMK-20	0.72	0.82	0.42	0.63	0.17	0.96	25.54	57.54	16.92
EFMK-23	0.74	0.84	0.42	0.61	0.19	0.97	26.72	54.10	19.18
EFMK-24	0.71	0.81	0.42	0.63	0.18	0.96	26.45	55.46	18.09
EFMK-25	0.72	0.84	0.44	0.66	0.19	0.97	26.57	53.63	19.80
EFMK-26	0.73	0.84	0.49	0.69	0.19	0.98	29.02	54.04	16.94

Fasken "A" #1 core

Sample	MPI 1	MPI 2	MA(I)/ MA(I+II)	TA(I)/ TA(I+II)	C ₂₆ TA (20S/20S+20R)	TA28/ (MA29+TA28)	%C ₂₇ MAS	%C ₂₈ MAS	%C ₂₉ MAS
EFFA-1	0.93	1.03	0.59	0.74	0.21	0.50	45.42	40.05	14.53
EFFA-3	0.89	0.93	0.55	0.70	0.22	0.63	25.40	60.54	14.06
EFFA-5	0.91	0.96	0.50	0.72	0.21	0.62	26.16	60.05	13.79
EFFA-11	0.96	1.04	0.46	0.71	0.22	0.65	24.93	60.99	14.09
EFFA-17	0.89	0.95	0.50	0.71	0.21	0.60	25.69	59.85	14.47
EFFA-18	0.91	0.97	0.49	0.76	0.23	0.58	26.78	58.71	14.51
EFFA-20	0.90	0.97	0.49	0.75	0.25	0.41	23.84	63.85	12.31
EFFA-21	0.82	0.86	0.62	0.66	0.22	0.58	26.90	59.66	13.44
EFFA-23	0.89	0.95	0.59	0.72	0.21	0.59	27.30	60.39	12.31
EFFA-24	0.87	0.88	0.58	0.67	0.19	0.62	25.87	61.83	12.31
EFFA-28	0.87	0.89	0.58	0.70	0.21	0.61	24.05	61.25	14.70
EFFA-32	0.91	0.94	0.53	0.69	0.21	0.64	26.69	60.82	12.49
EFFA-37	0.77	0.82	0.64	0.73	0.21	0.59	25.61	62.11	12.29
EFFA-38	0.93	0.95	0.52	0.66	0.21	0.62	25.86	60.97	13.17
EFFA-40	0.86	0.90	0.50	0.71	0.23	0.59	26.49	60.92	12.58

Bouldin Creek outcrop

Sample	MPI 1	MPI 2	MA(I)/ MA(I+II)	TA(I)/ TA(I+II)	C₂₆ TA (20S/20S+20R)	TA28/ (MA29+TA28)	%C₂₇ MAS	%C₂₈ MAS	%C₂₉ MAS
EFBC-5	0.44	0.36	0.03	0.13	0.11	ND	7.31	61.62	31.07
EFBC-7	0.47	0.46	0.03	0.14	0.11	ND	7.14	62.56	30.31
EFBC-9	0.47	0.43	0.03	0.12	0.10	ND	6.51	62.17	31.32
EFBC-11	0.49	0.44	0.02	0.13	0.09	ND	6.15	63.50	30.35
EFBC-15	0.41	0.44	0.03	0.12	0.11	ND	7.36	57.59	35.05
EFBC-19	0.22	0.23	0.03	0.11	0.10	ND	7.01	57.90	35.09
EFBC-20	0.31	0.31	0.04	0.10	0.12	0.01	7.96	59.44	32.60
EFBC-23	0.30	0.31	0.04	0.11	0.12	ND	8.57	62.08	29.35
EFBC-25	0.36	0.37	0.04	0.11	0.12	0.01	8.77	62.88	28.35
EFBC-27	0.45	0.47	0.05	0.13	0.14	0.01	10.04	61.12	28.84
EFBC-30	0.49	0.50	0.07	0.23	0.15	ND	11.62	55.58	32.80
EFBC-32	0.55	0.60	0.05	0.15	0.17	ND	10.56	53.41	36.03
EFBC-34	0.67	0.74	0.05	0.15	0.15	ND	9.67	53.50	36.83

ACC#1 core

Sample	MPI 1	MPI 2	MA(I)/ MA(I+II)	TA(I)/ TA(I+II)	C₂₆ TA (20S/20S+20R)	TA28/ (MA29+TA28)	%C₂₇ MAS	%C₂₈ MAS	%C₂₉ MAS
EFAC-1	0.32	0.32	0.03	0.13	0.12	0.01	7.00	60.22	32.78
EFAC-3	0.33	0.34	0.03	0.15	0.10	ND	6.46	58.97	34.57
EFAC-6	0.39	0.40	0.05	0.19	0.13	0.01	9.57	63.28	27.15
EFAC-7	0.37	0.39	0.06	0.23	0.13	0.01	12.60	53.11	34.29
EFAC-8	0.42	0.44	0.05	0.21	0.13	ND	11.64	52.45	35.91
EFAC-10	0.45	0.45	0.04	0.25	0.13	0.01	11.89	52.94	35.17

W. Brechtel #1 core

Sample	MPI 1	MPI 2	MA(I)/ MA(I+II)	TA(I)/ TA(I+II)	C₂₆ TA (20S/20S+20R)	TA28/ (MA29+TA28)	%C₂₇ MAS	%C₂₈ MAS	%C₂₉ MAS
EFWB-1	0.68	0.79	0.08	0.09	0.22	0.17	26.07	61.47	12.46
EFWB-3	0.66	0.77	0.07	0.07	0.16	0.13	24.99	51.28	23.72
EFWB-4	0.65	0.74	0.05	0.07	0.17	0.22	20.09	68.54	11.38
EFWB-7	0.58	0.67	0.06	0.09	0.20	0.26	23.80	65.21	10.99
EFWB-8	0.48	0.56	0.05	0.08	0.19	0.19	21.91	67.22	10.87
EFWB-9	0.63	0.74	0.10	0.34	0.20	0.35	28.27	61.19	10.54

C.J. Hendershot #1 core

Sample	MPI 1	MPI 2	MA(I)/ MA(I+II)	TA(I)/ TA(I+II)	C₂₆ TA (20S/20S+20R)	TA28/ (MA29+TA28)	%C₂₇ MAS	%C₂₈ MAS	%C₂₉ MAS
EFHE-1	0.59	0.56	0.11	0.12	0.22	0.26	32.52	48.57	18.91
EFHE-2	0.58	0.52	0.09	0.12	0.21	0.26	24.55	60.57	14.88
EFHE-3	0.59	0.55	0.06	0.12	0.19	0.31	40.79	48.21	10.99
EFHE-5	0.55	0.49	0.05	0.08	0.17	0.31	20.68	67.03	12.28
EFHE-6	0.55	0.47	0.07	0.09	0.19	0.44	23.55	64.91	11.54
EFHE-7	0.55	0.48	0.06	0.09	0.19	0.41	22.92	65.82	11.26
EFHE-9	0.54	0.46	0.05	0.05	0.21	0.46	41.96	49.02	9.02
EFHE-10	0.53	0.53	0.08	0.13	0.25	0.47	26.60	61.83	11.57

Lily Hoppess #1 core

Sample	MPI 1	MPI 2	MA(I)/ MA(I+II)	TA(I)/ TA(I+II)	C ₂₆ TA (20S/20S+20R)	TA28/ (MA29+TA28)	%C ₂₇ MAS	%C ₂₈ MAS	%C ₂₉ MAS
EFLH-1	0.55	0.52	0.19	0.20	0.21	0.94	25.06	60.04	14.90
EFLH-5	0.55	0.52	0.18	0.20	0.19	0.93	24.63	60.29	15.08
EFLH-9	0.52	0.50	0.20	0.15	0.20	0.94	25.24	59.01	15.75
EFLH-13	0.53	0.51	0.15	0.35	0.22	0.92	40.12	49.39	10.49
EFLH-15	0.54	0.53	0.14	0.25	0.19	0.94	23.92	59.25	16.83
EFLH-20	0.53	0.51	0.18	0.27	0.19	0.93	24.80	58.55	16.65

Oils and condensates

Sample	MPI 1	MPI 2	MA(I)/ MA(I+II)	TA(I)/ TA(I+II)	C ₂₆ TA (20S/ 20S+20R)	TA28/ (MA29+TA28)	%C ₂₇ MAS	%C ₂₈ MAS	%C ₂₉ MAS
NWMK-533.2	0.63	0.66	0.44	0.70	0.16	0.78	26.46	56.81	16.73
NFMK-526-1H	0.65	0.73	0.54	0.79	0.17	0.76	26.82	50.03	23.16
MJOG-UEF	0.74	0.81	0.56	0.91	0.40	0.52	22.63	43.42	33.95
MJOG-LEF	0.81	0.87	0.64	0.94	0.50	0.43	22.83	40.49	36.68
GRES-1-110	0.86	0.89	0.53	ND	ND	ND	24.29	48.78	26.93

L. Quantitative biomarker analysis results for steranes (Concentrations are expressed as µg biomarkers/g TOC; ND = not determined)

Lozier Canyon outcrop

Sample Name	Diapreg	Preg	Diahomopreg	Homopreg	C ₂₇ S -Dia (20S)	C ₂₇ S -Dia (20R)	C ₂₇ S-Dia (20S)	C ₂₇ S-Dia (20R)	C ₂₈ S -Dia (20S)	C ₂₈ S -Dia (20R)	C ₂₈ S-Dia (20S) + C ₂₇ (20S)	C ₂₈ S-Dia (20S)	C ₂₇ SS (20R) + C ₂₉ S -Dia (20S)	C ₂₇ SS (20R) + C ₂₈ S-Dia (20R)	C ₂₇ (20R)
EFLC-28	52	78	2	31	97	53	19	36	17	41	88	144	103	117	
EFLC-26	91	157	6	71	176	95	28	75	30	61	167	267	188	223	
EFLC-25	21	38	1	18	57	32	9	26	11	25	68	102	81	83	
EFLC-20	45	114	3	57	100	47	15	58	22	46	155	217	185	186	
EFLC-18	12	29	1	15	23	11	4	14	6	11	36	47	41	43	
EFLC-14	9	26	1	13	16	7	2	11	3	8	31	37	34	38	
EFLC-12	23	77	4	38	32	17	5	17	6	17	62	84	75	79	
EFLC-10	20	103	4	51	23	10	4	11	8	6	78	103	101	99	
EFLC-8	33	157	7	79	34	14	5	11	11	10	107	140	124	138	
EFLC-6	9	40	2	20	11	5	2	4	4	3	39	39	39	42	
EFLC-4	3	0	1	6	4	2	1	2	1	2	10	9	9	11	
EFLC-1	122	27	15	113	263	148	52	97	58	141	287	466	386	399	

Lozier Canyon outcrop (cont.)

Sample Name	C ₂₉ S -Dia (20R)	C ₂₉ S-Dia (20S)	C ₂₈ (20S)	C ₂₈ SS (20R) +C ₂₉ S-Dia (20R)	C ₂₈ SS (20S)	C ₂₈ (20R)	C ₂₉ (20S)	C ₂₉ SS (20R)	C ₂₉ SS (20S)	C ₂₉ (20R)	C ₃₀ (20S)	C ₃₀ SS (20R)	C ₃₀ SS (20S)	C ₃₀ (20R)
EFLC-28	44	70	74	69	47	61	71	102	82	78	9	13	20	7
EFLC-26	83	127	140	127	94	119	145	200	158	157	19	27	37	15
EFLC-25	34	49	58	53	34	48	60	79	62	64	8	12	17	6
EFLC-20	93	148	186	168	94	146	143	194	151	150	19	29	37	16
EFLC-18	26	36	44	40	21	36	33	44	35	34	5	8	9	5
EFLC-14	22	31	38	35	16	31	26	37	29	28	5	7	10	5
EFLC-12	36	50	66	61	37	53	54	78	62	57	8	12	15	7
EFLC-10	45	57	83	77	38	67	74	100	78	73	10	15	19	10
EFLC-8	61	78	114	100	56	94	100	132	106	99	14	17	21	13
EFLC-6	19	25	35	32	20	30	32	40	32	31	5	7	9	4
EFLC-4	5	7	8	7	4	7	6	9	7	7	1	2	2	1
EFLC-1	223	261	347	300	218	259	335	405	287	308	35	71	100	32

Ferguson McKnight #526-1H core

Sample Name	Diapreg	Preg	Diahomopreg	Homopreg	C ₂₇ S -Dia (20S)	C ₂₇ S -Dia (20R)	C ₂₇ S-Dia (20S)	C ₂₇ S-Dia (20R)	C ₂₈ S -Dia (20S)	C ₂₈ S -Dia (20R)	C ₂₈ S-Dia (20S) + C ₂₇ (20S)	C ₂₇ SS (20R) + C ₂₉ S -Dia (20S)	C ₂₇ SS (20S) + C ₂₈ S-Dia (20R)	C ₂₇ (20R)
EFMK-1	127	199	113	109	137	83	32	79	79	56	81	140	110	100
EFMK-4	76	145	73	73	79	47	18	50	45	47	47	80	62	50
EFMK-8	166	308	150	150	143	83	33	94	88	106	88	151	129	206
EFMK-12	88	159	76	75	70	43	16	46	45	53	43	72	57	50
EFMK-15	82	139	70	60	78	47	17	54	50	63	40	72	55	52
EFMK-17	269	441	237	194	242	147	56	167	147	190	127	211	160	229
EFMK-20	189	311	155	130	186	112	42	130	111	135	91	164	122	166
EFMK-23	196	435	35	131	202	122	42	135	113	149	68	214	127	129
EFMK-24	168	294	144	114	181	104	40	117	103	128	60	184	110	117
EFMK-25	103	178	83	65	112	67	23	68	66	84	29	91	59	61
EFMK-26	82	180	77	57	53	33	11	28	23	36	30	44	29	33

Ferguson McKnight #526-1H core (cont.)

Sample Name	C ₂₉ S -Dia (20R)	C ₂₉ S-Dia (20S)	C ₂₈ (20S)	C ₂₈ SS (20R) +C ₂₉ S-Dia (20R)	C ₂₈ SS (20S)	C ₂₈ (20R)	C ₂₉ (20S)	C ₂₉ SS (20R)	C ₂₉ SS (20S)	C ₂₉ (20R)	C ₃₀ (20S)	C ₃₀ SS (20R)	C ₃₀ SS (20S)	C ₃₀ (20R)
EFMK-1	98	73	84	67	53	69	45	84	68	58	13	14	19	10
EFMK-4	52	46	52	43	21	41	27	50	41	35	8	9	11	5
EFMK-8	105	87	96	93	65	71	56	96	81	63	15	16	21	10
EFMK-12	49	39	44	42	28	33	26	44	37	30	7	8	10	4
EFMK-15	49	40	43	34	25	32	22	42	35	28	6	8	8	5
EFMK-17	99	121	128	96	60	102	64	124	105	90	23	26	29	13
EFMK-20	69	88	90	70	61	69	48	97	80	68	16	18	22	10
EFMK-23	129	92	90	74	59	69	48	107	88	65	18	19	22	11
EFMK-24	60	81	78	64	58	59	41	92	74	58	14	18	18	9
EFMK-25	36	43	44	41	12	33	15	40	30	26	7	8	10	5
EFMK-26	35	25	28	23	16	23	15	27	19	17	3	7	6	3

Fasken "A" #1 core

Sample Name	Diapreg	Preg	Diahomopreg	Homopreg	C ₂₇ S -Dia (20S)	C ₂₇ S -Dia (20R)	C ₂₇ S-Dia (20S)	C ₂₇ S-Dia (20R)	C ₂₈ S -Dia (20S)	C ₂₈ S -Dia (20R)	C ₂₈ S-Dia (20S) + C ₂₇ (20S)	C ₂₇ SS (20R) + C ₂₉ S -Dia (20S)	C ₂₇ SS (20S) + C ₂₈ S-Dia (20R)	C ₂₇ (20R)
EFFA-1	2	4	0	1	1	0	0	0	0	0	0	1	0	1
EFFA-3	0	0	0	0	0	0	0	0	0	0	0	0	0	0
EFFA-5	9	25	2	6	3	2	1	2	1	2	2	4	2	5
EFFA-11	14	30	2	8	5	3	1	3	1	3	4	5	3	7
EFFA-17	11	23	1	6	3	2	1	2	1	2	2	3	2	5
EFFA-18	11	28	2	7	4	2	1	2	1	2	3	4	2	6
EFFA-20	1	3	0	1	0	0	0	0	0	0	0	0	0	1
EFFA-21	3	7	0	2	1	1	0	0	0	0	1	1	0	1
EFFA-23	6	12	1	3	2	1	0	1	1	1	1	2	1	2
EFFA-24	34	68	4	17	8	5	1	3	3	4	6	8	5	12
EFFA-28	18	44	2	12	5	3	1	2	2	2	4	6	4	8
EFFA-32	38	95	6	23	12	7	3	6	4	6	9	12	7	15
EFFA-37	4	9	0	2	1	1	0	0	0	1	1	1	1	2
EFFA-38	21	43	3	12	6	3	1	3	2	2	4	6	3	8
EFFA-40	5	12	1	3	2	1	0	1	1	1	1	2	1	3

Fasken "A" #1 core (cont.)

Sample Name	C ₂₉ S -Dia (20R)	C ₂₉ S-Dia (20S)	C ₂₈ (20S)	C ₂₈ SS (20R) +C ₂₉ S-Dia (20R)	C ₂₈ SS (20S)	C ₂₈ (20R)	C ₂₉ (20S)	C ₂₉ SS (20R)	C ₂₉ SS (20S)	C ₂₉ (20R)	C ₃₀ (20S)	C ₃₀ SS (20R)	C ₃₀ SS (20S)	C ₃₀ (20R)
EFFA-1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
EFFA-3	0	0	0	0	0	0	0	0	0	0	0	0	0	0
EFFA-5	1	2	2	1	1	3	2	3	2	2	0	0	0	0
EFFA-11	2	3	3	2	2	4	2	4	2	4	0	0	1	0
EFFA-17	1	2	2	1	1	3	2	2	1	3	0	0	0	0
EFFA-18	2	2	2	2	3	4	2	3	2	3	0	0	0	0
EFFA-20	0	0	0	0	0	0	0	0	0	0	0	0	0	0
EFFA-21	0	0	1	0	0	1	0	1	0	1	0	0	0	0
EFFA-23	1	1	1	1	1	1	1	1	1	1	0	0	0	0
EFFA-24	4	5	5	4	3	8	4	7	4	6	0	0	1	0
EFFA-28	2	3	3	2	3	5	3	5	2	4	0	0	1	0
EFFA-32	8	7	6	5	6	12	5	9	5	9	1	1	1	1
EFFA-37	0	1	1	1	1	1	0	1	0	1	0	0	0	0
EFFA-38	3	3	3	2	2	4	2	4	2	4	0	0	1	0
EFFA-40	1	1	1	1	1	1	1	1	1	1	0	0	0	0

Bouldin Creek outcrop

Sample Name	Diapreg	Preg	Diahomopreg	Homopreg	C ₂₇ S -Dia (20S)	C ₂₇ S -Dia (20R)	C ₂₇ S-Dia (20S)	C ₂₇ S-Dia (20R)	C ₂₈ S -Dia (20S)	C ₂₈ S -Dia (20R)	C ₂₈ S-Dia (20S) + C ₂₇ (20S)	C ₂₇ SS (20R) + C ₂₉ S -Dia (20S)	C ₂₇ SS (20S) + C ₂₈ S-Dia (20R)	C ₂₇ (20R)
EFBC-5	0	1	2	0	ND	0	0	ND	ND	0	6	0	0	14
EFBC-7	0	1	2	0	ND	0	0	ND	ND	0	7	1	0	16
EFBC-9	0	1	3	1	ND	0	0	ND	ND	1	7	1	1	16
EFBC-11	0	1	2	0	ND	0	0	ND	ND	1	7	1	1	14
EFBC-15	0	1	2	0	ND	0	0	ND	ND	0	3	0	0	6
EFBC-19	1	3	4	1	ND	0	0	ND	ND	1	16	2	0	33
EFBC-20	1	11	2	3	ND	1	1	ND	ND	2	22	3	2	43
EFBC-23	1	7	1	2	ND	0	0	ND	ND	2	18	3	2	34
EFBC-25	1	6	1	2	ND	0	0	ND	ND	1	12	2	2	24
EFBC-27	1	6	1	2	0	0	0	ND	ND	2	13	2	2	24
EFBC-30	1	3	5	2	0	0	0	ND	ND	2	22	3	2	42
EFBC-32	1	5	10	2	0	1	0	ND	ND	4	47	5	4	89
EFBC-34	1	3	6	1	0	0	0	ND	ND	3	31	3	3	58

Bouldin Creek outcrop (cont.)

Sample Name	C ₂₉ S -Dia (20R)	C ₂₉ S-Dia (20S)	C ₂₈ (20S)	C ₂₈ SS (20R) +C ₂₉ S-Dia (20R)	C ₂₈ SS (20S)	C ₂₈ (20R)	C ₂₉ (20S)	C ₂₉ SS (20R)	C ₂₉ SS (20S)	C ₂₉ (20R)	C ₃₀ (20S)	C ₃₀ SS (20R)	C ₃₀ SS (20S)	C ₃₀ (20R)
EFBC-5	1	ND	8	1	0	15	ND	4	ND	9	0	ND	1	2
EFBC-7	1	ND	9	1	0	17	ND	4	ND	9	0	ND	1	2
EFBC-9	1	ND	10	1	0	19	ND	5	ND	11	0	ND	1	2
EFBC-11	1	ND	10	1	0	16	ND	4	ND	8	0	ND	1	2
EFBC-15	0	ND	4	0	0	6	ND	2	ND	4	0	ND	1	1
EFBC-19	3	ND	17	2	1	28	1	15	ND	27	1	ND	4	5
EFBC-20	2	ND	24	3	2	35	ND	17	ND	32	ND	ND	4	5
EFBC-23	4	ND	28	3	0	39	1	14	ND	24	1	ND	3	4
EFBC-25	3	ND	22	2	1	29	ND	10	ND	17	1	ND	3	3
EFBC-27	3	ND	18	2	1	24	ND	10	ND	17	1	ND	3	3
EFBC-30	3	ND	24	2	1	39	ND	14	ND	27	1	ND	5	4
EFBC-32	8	4	ND	5	2	101	2	30	ND	60	2	ND	10	10
EFBC-34	5	ND	43	4	2	75	2	21	ND	43	2	ND	7	7

ACC #1 core

Sample Name	Diapreg	Preg	Diahomopreg	Homopreg	C ₂₇ S -Dia (20S)	C ₂₇ S -Dia (20R)	C ₂₇ S-Dia (20S)	C ₂₇ S-Dia (20R)	C ₂₈ S -Dia (20S)	C ₂₈ S -Dia (20R)	C ₂₈ S-Dia (20S) + C ₂₇ (20S)	C ₂₇ SS (20R) + C ₂₉ S -Dia (20S)	C ₂₇ SS (20S) + C ₂₈ S-Dia (20R)	C ₂₇ (20R)
EFAC-1	10	56	56	12	ND	ND	ND	ND	ND	ND	61	4	ND	163
EFAC-3	10	37	69	11	ND	ND	ND	ND	ND	ND	108	8	8	273
EFAC-6	9	32	22	10	ND	ND	ND	ND	ND	ND	46	6	ND	128
EFAC-7	3	13	17	4	1	ND	ND	ND	ND	ND	72	4	ND	181
EFAC-8	2	14	21	4	ND	ND	ND	ND	ND	ND	73	4	ND	196
EFAC-10	24	64	50	20	ND	ND	ND	ND	ND	ND	235	22	ND	790

ACC #1 core (cont.)

Sample Name	C ₂₉ S -Dia (20R)	C ₂₉ S-Dia (20S)	C ₂₈ (20S)	C ₂₈ SS (20R) +C ₂₉ S-Dia (20R)	C ₂₈ SS (20S)	C ₂₈ (20R)	C ₂₉ (20S)	C ₂₉ SS (20R)	C ₂₉ SS (20S)	C ₂₉ (20R)	C ₃₀ (20S)	C ₃₀ SS (20R)	C ₃₀ SS (20S)	C ₃₀ (20R)
EFAC-1	6	ND	44	ND	ND	100	ND	38	ND	98	ND	ND	10	17
EFAC-3	9	ND	112	ND	ND	263	ND	70	ND	184	ND	ND	19	36
EFAC-6	4	ND	44	ND	ND	103	ND	25	ND	70	ND	ND	8	12
EFAC-7	6	ND	71	ND	ND	184	ND	38	ND	110	ND	ND	11	14
EFAC-8	6	ND	77	ND	ND	189	ND	37	ND	106	ND	ND	12	15
EFAC-10	24	ND	236	ND	ND	683	ND	205	ND	711	ND	ND	60	86

W. Brechtel #1 core

Sample Name	Diapreg	Preg	Diahomopreg	Homopreg	C ₂₇ S -Dia (20S)	C ₂₇ S -Dia (20R)	C ₂₇ S-Dia (20S)	C ₂₇ S-Dia (20R)	C ₂₈ S -Dia (20S)	C ₂₈ S -Dia (20R)	C ₂₈ S-Dia (20S) + C ₂₇ (20S)	C ₂₇ SS (20R) + C ₂₉ S -Dia (20S)	C ₂₇ SS (20S) + C ₂₈ S-Dia (20R)	C ₂₇ (20R)
EFWB-1	35	43	23	75	16	6	ND	10	12	5	144	57	51	233
EFWB-3	38	67	30	93	27	12	6	12	14	15	149	58	60	239
EFWB-4	29	44	14	70	19	8	4	9	10	11	103	42	50	154
EFWB-7	57	71	19	137	29	10	6	10	13	11	147	96	99	200
EFWB-8	27	34	9	65	18	8	3	7	7	9	83	46	48	111
EFWB-9	38	93	27	91	15	6	2	6	7	ND	107	78	77	119

W. Brechtel #1 core (cont.)

Sample Name	C ₂₉ S -Dia (20R)	C ₂₉ S-Dia (20S)	C ₂₈ (20S)	C ₂₈ SS (20R) +C ₂₉ S-Dia (20R)	C ₂₈ SS (20S)	C ₂₈ (20R)	C ₂₉ (20S)	C ₂₉ SS (20R)	C ₂₉ SS (20S)	C ₂₉ (20R)	C ₃₀ (20S)	C ₃₀ SS (20R)	C ₃₀ SS (20S)	C ₃₀ (20R)
EFWB-1	24	48	101	63	21	140	65	66	45	115	25	12	30	24
EFWB-3	57	93	151	105	28	235	82	75	47	136	21	7	23	22
EFWB-4	39	61	100	73	25	159	62	55	31	100	15	5	17	16
EFWB-7	21	69	142	124	24	158	82	89	70	101	20	11	35	18
EFWB-8	26	41	74	66	14	91	42	42	32	54	11	6	18	10
EFWB-9	26	48	104	96	18	104	69	73	60	77	14	13	24	11

C.J. Hendershot #1 core

Sample Name	Diapreg	Preg	Diahomopreg	Homopreg	C ₂₇ S -Dia (20S)	C ₂₇ S -Dia (20R)	C ₂₇ S-Dia (20S)	C ₂₇ S-Dia (20R)	C ₂₈ S -Dia (20S)	C ₂₈ S -Dia (20R)	C ₂₈ S-Dia (20S) + C ₂₇ (20S)	C ₂₇ SS (20R) + C ₂₉ S -Dia (20S)	C ₂₇ SS (20S) + C ₂₈ S-Dia (20R)	C ₂₇ (20R)
EFHE-1	184	236	171	145	261	191	67	90	116	132	1142	276	224	2487
EFHE-2	151	206	151	133	117	95	35	62	58	82	1196	169	187	2528
EFHE-3	196	244	148	137	214	165	59	124	89	188	1272	231	270	2657
EFHE-5	192	271	147	173	285	194	65	252	168	281	916	273	218	1598
EFHE-6	167	334	334	151	229	143	48	148	93	153	717	207	162	1090
EFHE-7	186	253	128	160	248	160	55	95	107	194	789	232	188	1198
EFHE-9	124	298	29	175	116	66	26	45	39	52	835	241	238	903
EFHE-10	164	187	15	47	364	203	77	193	120	203	538	230	198	677

C.J. Hendershot #1 core (cont.)

Sample Name	C ₂₉ S -Dia (20R)	C ₂₉ S-Dia (20S)	C ₂₈ (20S)	C ₂₈ SS (20R) +C ₂₉ S-Dia (20R)	C ₂₈ SS (20S)	C ₂₈ (20R)	C ₂₉ (20S)	C ₂₉ SS (20R)	C ₂₉ SS (20S)	C ₂₉ (20R)	C ₃₀ (20S)	C ₃₀ SS (20R)	C ₃₀ SS (20S)	C ₃₀ (20R)
EFHE-1	253	289	468	75	181	1747	343	531	79	1392	143	26	98	266
EFHE-2	269	349	520	80	177	2058	368	594	77	1629	160	48	105	305
EFHE-3	298	492	606	138	221	2430	486	629	68	1860	190	39	124	313
EFHE-5	249	557	565	213	175	1907	451	385	108	1156	128	43	101	216
EFHE-6	167	379	345	147	134	1043	382	282	98	720	85	29	62	119
EFHE-7	198	424	407	180	156	1177	457	315	106	840	100	35	79	138
EFHE-9	28	454	421	255	149	966	541	294	163	604	104	46	92	102
EFHE-10	117	283	230	109	147	527	404	226	101	491	74	32	77	89

Lily Hoppess #1 core

Sample Name	Diapreg	Preg	Diahomopreg	Homopreg	C ₂₇ S -Dia (20S)	C ₂₇ S -Dia (20R)	C ₂₇ S-Dia (20S)	C ₂₇ S-Dia (20R)	C ₂₈ S -Dia (20S)	C ₂₈ S -Dia (20R)	C ₂₈ S-Dia (20S) + C ₂₇ (20S)	C ₂₇ SS (20R) + C ₂₉ S -Dia (20S)	C ₂₇ SS (20S) + C ₂₈ S-Dia (20R)	C ₂₇ (20R)
EFLH-1	248	350	205	217	421	256	107	258	229	326	294	413	340	362
EFLH-5	337	473	288	277	606	377	161	456	406	566	407	559	498	522
EFLH-9	281	422	226	248	482	291	126	310	263	385	323	460	400	408
EFLH-13	48	47	31	23	102	64	26	61	45	71	31	99	59	30
EFLH-15	177	232	138	125	358	220	94	289	220	330	141	414	272	163
EFLH-20	149	198	107	99	237	143	64	174	131	195	97	279	224	180

Lily Hoppess #1 core (cont.)

Sample Name	C ₂₉ S -Dia (20R)	C ₂₉ S-Dia (20S)	C ₂₈ (20S)	C ₂₈ SS (20R) +C ₂₉ S-Dia (20R)	C ₂₈ SS (20S)	C ₂₈ (20R)	C ₂₉ (20S)	C ₂₉ SS (20R)	C ₂₉ SS (20S)	C ₂₉ (20R)	C ₃₀ (20S)	C ₃₀ SS (20R)	C ₃₀ SS (20S)	C ₃₀ (20R)
EFLH-1	275	256	245	258	116	227	154	222	188	181	48	57	54	38
EFLH-5	498	409	426	510	270	367	223	336	274	251	62	79	74	52
EFLH-9	365	304	350	296	179	282	184	278	244	213	56	80	76	46
EFLH-13	92	40	34	27	39	24	27	39	32	26	6	14	14	5
EFLH-15	454	199	203	161	178	154	121	190	172	132	33	52	46	27
EFLH-20	183	126	146	116	120	96	88	137	123	87	21	33	34	17

Oils and condensates (Concentrations are expressed as µg biomarkers/g whole oil; ND = not determined)

Sample Name	Diapreg	Preg	Diahomopreg	Homopreg	C ₂₇ S -Dia (20S)	C ₂₇ S -Dia (20R)	C ₂₇ S-Dia (20S)	C ₂₇ S-Dia (20R)	C ₂₈ S -Dia (20S)	C ₂₈ S -Dia (20R)	C ₂₈ S-Dia (20S) + C ₂₇ (20S)	C ₂₇ SS (20R) + C ₂₉ S -Dia (20S)	C ₂₇ SS (20S) + C ₂₈ S-Dia (20R)	C ₂₇ (20R)
NWMK-533.2	343	1133	35	415	435	348	100	273	271	154	262	555	405	452
NFMK-526-1H	286	882	34	214	289	216	70	178	149	88	115	243	166	194
MJOG-UEF	63	85	4	17	32	29	9	19	13	9	12	22	11	35
MJOG-LEF	115	175	8	14	95	57	26	50	41	30	26	67	34	85
REF-1-110	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND

Oils and condensates (Concentrations are expressed as µg biomarkers/g whole oil; ND = not determined; cont.)

Sample Name	C ₂₉ S -Dia (20R)	C ₂₉ S-Dia (20S)	C ₂₈ (20S)	C ₂₈ SS (20R) +C ₂₉ S-Dia (20R)	C ₂₈ SS (20S)	C ₂₈ (20R)	C ₂₉ (20S)	C ₂₉ SS (20R)	C ₂₉ SS (20S)	C ₂₉ (20R)	C ₃₀ (20S)	C ₃₀ SS (20R)	C ₃₀ SS (20S)	C ₃₀ (20R)
NWMK-533.2	290	297	425	692	465	178	175	348	268	237	56	67	121	43
NFMK-526-1H	105	133	162	212	50	116	64	151	106	102	21	28	31	20
MJOG-UEF	6	8	8	7	6	6	3	8	10	7	1	ND	ND	ND
MJOG-LEF	11	22	17	14	23	15	8	25	16	11	3	5	3	4
GRAF-1-110	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND

M. Quantitative biomarker analysis results for terpanes (Concentrations are expressed as μg biomarkers/g TOC; ND = not determined)

Lozier Canyon outcrop

Sample Name	C ₂₀ TR	C ₂₁ TR	C ₂₂ TR	C ₂₃ TR	C ₂₄ TR	C ₂₅ TR	C ₂₄ TT	C ₂₆ TR	C ₂₆ TR	C ₂₈ TR	C ₂₈ TR	C ₂₉ TR	C ₂₉ TR	T _s	T _m
EFLC-28	28	61	15	92	52	44	18	17	25	11	13	8	11	51	111
EFLC-26	52	108	35	216	115	108	35	30	52	26	27	24	28	63	253
EFLC-25	11	25	9	55	32	31	11	10	18	7	10	9	10	25	104
EFLC-20	20	55	30	181	77	87	27	23	39	23	24	25	28	53	232
EFLC-18	6	14	9	58	19	24	7	6	10	6	7	7	8	13	52
EFLC-14	6	15	12	79	23	31	8	7	9	8	9	10	10	15	58
EFLC-12	19	48	42	236	68	83	23	20	23	19	20	26	27	44	143
EFLC-10	30	72	66	336	98	122	32	28	44	27	30	38	37	59	221
EFLC-8	49	112	98	495	146	177	48	40	68	38	42	51	49	88	315
EFLC-6	12	29	26	132	42	52	16	13	19	13	14	16	16	28	97
EFLC-4	5	11	9	47	12	15	4	3	6	3	3	4	4	10	27
EFLC-1	76	113	40	169	161	133	49	87	60	25	25	33	40	169	463

Lozier Canyon outcrop (cont.)

Sample Name	C ₂₉ -30- NorH	C ₂₉ Ts	C ₃₀ D	C ₂₉ -30- NorM	C ₃₀ H	C ₃₀ Mor	C ₃₁ HH (22S)	C ₃₁ HH (22R)	C ₃₂ HH (22S)	C ₃₂ HH (22R)	C ₃₃ HH (22S)	C ₃₃ HH (22R)	C ₃₄ HH (22S)	C ₃₄ HH (22R)	C ₃₅ HH (22S)	C ₃₅ HH (22R)
EFLC-28	236	44	8	18	392	25	300	182	183	121	97	60	72	40	86	48
EFLC-26	504	65	13	34	808	52	606	383	344	227	185	114	123	70	125	71
EFLC-25	231	28	6	15	401	24	286	183	172	116	97	60	68	39	79	45
EFLC-20	425	47	7	30	652	41	548	345	313	205	197	121	126	74	184	104
EFLC-18	101	10	2	7	147	9	133	83	75	49	50	31	33	19	48	27
EFLC-14	121	ND	1	7	144	9	132	81	74	48	50	30	33	18	48	26
EFLC-12	253	ND	3	16	305	18	244	158	132	90	88	56	61	35	89	51
EFLC-10	464	ND	6	29	567	34	389	260	214	147	149	96	102	60	142	82
EFLC-8	646	ND	9	39	766	49	540	351	287	197	204	130	140	82	190	108
EFLC-6	198	ND	3	12	234	15	168	109	91	64	65	41	46	27	65	38
EFLC-4	56	ND	1	3	63	4	49	31	27	18	19	12	14	8	20	11
EFLC-1	696	171	47	68	1233	94	801	535	580	429	375	244	393	228	567	376

Ferguson McKnight #526-1H core

Sample Name	C ₂₀ TR	C ₂₁ TR	C ₂₂ TR	C ₂₃ TR	C ₂₄ TR	C ₂₅ TR	C ₂₄ TT	C ₂₆ TR	C ₂₆ TR	C ₂₈ TR	C ₂₈ TR	C ₂₉ TR	C ₂₉ TR	T _s	T _m
EFMK-1	209	216	65	331	205	184	64	43	46	44	45	52	50	79	81
EFMK-4	92	120	26	178	115	116	41	36	28	30	31	37	30	56	44
EFMK-8	244	272	65	393	243	237	96	83	68	73	68	77	61	160	94
EFMK-12	137	135	37	195	118	110	44	38	36	29	30	33	27	65	43
EFMK-15	111	111	32	158	99	93	34	36	29	28	26	31	25	82	42
EFMK-17	392	381	100	529	299	264	98	115	93	80	89	80	68	261	139
EFMK-20	254	280	69	402	256	241	88	92	64	66	65	81	65	188	95
EFMK-23	255	292	73	432	279	257	94	107	62	77	77	87	72	203	107
EFMK-24	223	243	60	365	243	234	82	95	55	73	69	78	68	183	94
EFMK-25	104	93	34	144	104	91	32	38	24	29	32	34	28	77	36
EFMK-26	145	153	63	257	190	150	57	64	34	53	46	54	47	99	73

Ferguson McKnight #526-1H core (cont.)

Sample Name	C ₂₉ -30-NorH	C ₂₉ Ts	C ₃₀ D	C ₂₉ -30-NorM	C ₃₀ H	C ₃₀ Mor	C ₃₁ HH (22S)	C ₃₁ HH (22R)	C ₃₂ HH (22S)	C ₃₂ HH (22R)	C ₃₃ HH (22S)	C ₃₃ HH (22R)	C ₃₄ HH (22S)	C ₃₄ HH (22R)	C ₃₅ HH (22S)	C ₃₅ HH (22R)
EFMK-1	144	29	12	12	121	12	45	54	29	17	17	18	ND	ND	ND	ND
EFMK-4	33	33	4	6	51	7	19	14	12	8	8	5	ND	ND	ND	ND
EFMK-8	72	62	11	ND	109	12	36	25	21	18	17	13	ND	ND	ND	ND
EFMK-12	39	38	7	8	51	6	22	14	13	8	8	6	ND	ND	ND	ND
EFMK-15	36	37	5	7	49	7	20	17	9	6	10	7	ND	ND	ND	ND
EFMK-17	134	112	18	26	169	16	68	72	41	29	30	18	ND	ND	ND	ND
EFMK-20	74	83	13	17	92	11	40	27	21	13	13	17	ND	ND	ND	ND
EFMK-23	62	90	13	6	87	12	30	24	17	14	21	14	ND	ND	ND	ND
EFMK-24	44	75	8	11	68	4	23	20	11	10	16	6	7	3	8	6
EFMK-25	23	24	7	8	19	9	14	13	9	6	ND	ND	ND	ND	ND	ND
EFMK-26	32	27	ND	ND	27	7	8	14	ND	ND	ND	ND	ND	ND	ND	ND

Fasken "A" #1 core

Sample Name	C ₂₀ TR	C ₂₁ TR	C ₂₂ TR	C ₂₃ TR	C ₂₄ TR	C ₂₅ TR	C ₂₄ TT	C ₂₆ TR	C ₂₆ TR	C ₂₈ TR	C ₂₈ TR	C ₂₉ TR	C ₂₉ TR	T _s	T _m
EFFA-1	7	6	2	7	3	2	1	0	1	0	0	0	0	1	1
EFFA-3	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
EFFA-5	39	36	13	46	21	10	4	3	6	2	2	2	1	5	9
EFFA-11	42	42	16	60	27	14	4	4	9	2	3	2	2	7	10
EFFA-17	39	35	12	44	20	9	3	3	6	2	2	2	1	5	8
EFFA-18	48	42	15	54	23	11	4	3	8	2	2	2	2	6	9
EFFA-20	5	5	2	6	3	1	0	0	1	0	0	0	0	1	1
EFFA-21	12	11	3	13	6	3	1	1	2	1	0	1	0	1	2
EFFA-23	20	18	6	23	10	5	2	1	3	1	1	1	1	2	4
EFFA-24	129	110	36	128	54	27	9	8	18	4	5	3	4	12	21
EFFA-28	81	72	23	86	37	17	6	4	12	3	2	3	2	8	15
EFFA-32	187	159	51	189	83	38	13	11	26	7	6	7	6	19	29
EFFA-37	20	16	5	19	8	4	1	1	2	1	1	1	0	2	3
EFFA-38	82	71	22	82	37	16	6	5	11	4	3	3	2	9	16
EFFA-40	18	17	6	22	10	5	2	1	3	1	1	1	1	2	4

Fasken "A" #1 core (cont.)

Sample Name	C ₂₉ -30-NorH	C ₂₉ Ts	C ₃₀ D	C ₂₉ -30-NorM	C ₃₀ H	C ₃₀ Mor	C ₃₁ HH (22S)	C ₃₁ HH (22R)	C ₃₂ HH (22S)	C ₃₂ HH (22R)	C ₃₃ HH (22S)	C ₃₃ HH (22R)	C ₃₄ HH (22S)	C ₃₄ HH (22R)	C ₃₅ HH (22S)	C ₃₅ HH (22R)
EFFA-1	3	0	0	0	3	1	1	1	1	1	1	0	ND	ND	ND	ND
EFFA-3	0	0	0	0	0	0	0	0	0	0	0	0	ND	ND	ND	ND
EFFA-5	23	1	1	3	18	4	7	5	4	3	2	2	ND	ND	ND	ND
EFFA-11	31	1	1	3	26	5	8	7	4	4	3	3	ND	ND	ND	ND
EFFA-17	22	1	1	2	19	4	6	5	4	3	2	1	ND	ND	ND	ND
EFFA-18	27	1	1	3	23	5	8	6	4	4	2	2	ND	ND	ND	ND
EFFA-20	3	0	0	0	3	1	1	1	0	0	0	0	0	0	ND	ND
EFFA-21	6	0	0	1	5	1	2	1	1	1	1	0	ND	ND	ND	ND
EFFA-23	10	0	0	1	10	2	3	2	1	1	1	1	ND	ND	ND	ND
EFFA-24	56	1	1	5	47	10	16	12	7	6	6	3	5	2	ND	ND
EFFA-28	36	1	1	3	33	6	9	7	5	4	3	2	ND	ND	ND	ND
EFFA-32	78	1	2	8	73	11	20	16	12	10	8	5	ND	ND	ND	ND
EFFA-37	7	0	0	1	7	1	2	2	1	1	1	0	ND	ND	ND	ND
EFFA-38	37	1	1	5	34	7	11	8	5	4	3	2	ND	ND	ND	ND
EFFA-40	10	0	0	1	10	2	3	2	2	1	1	1	ND	ND	ND	ND

Bouldin Creek outcrop

Sample Name	C ₂₀ TR	C ₂₁ TR	C ₂₂ TR	C ₂₃ TR	C ₂₄ TR	C ₂₅ TR	C ₂₄ TT	C ₂₆ TR	C ₂₆ TR	C ₂₈ TR	C ₂₈ TR	C ₂₉ TR	C ₂₉ TR	T _s	T _m
EFBC-5	0	0	ND	0	0	0	0	0	0	ND	ND	2	0	0	2
EFBC-7	0	0	ND	0	0	0	0	0	0	ND	ND	1	0	0	3
EFBC-9	0	0	ND	0	0	0	0	0	0	ND	ND	2	0	0	3
EFBC-11	0	0	ND	0	0	0	0	0	0	ND	ND	1	0	0	2
EFBC-15	0	0	ND	0	0	0	0	0	0	ND	ND	1	0	0	1
EFBC-19	1	1	ND	1	0	1	0	0	1	ND	ND	6	0	0	4
EFBC-20	1	2	ND	2	0	1	0	1	1	ND	ND	7	0	0	11
EFBC-23	1	1	ND	1	0	1	0	0	1	ND	ND	2	0	0	5
EFBC-25	1	1	ND	1	0	1	0	0	0	ND	ND	2	0	0	4
EFBC-27	1	1	ND	1	0	1	0	0	0	ND	ND	2	0	0	4
EFBC-30	1	1	ND	1	0	1	0	0	1	ND	ND	4	0	0	9
EFBC-32	2	2	ND	2	1	3	1	1	2	ND	ND	9	0	1	16
EFBC-34	1	1	ND	1	0	3	0	1	1	ND	ND	5	0	0	9

Bouldin Creek outcrop (cont.)

Sample Name	C ₂₉ -30-NorH	C ₂₉ Ts	C ₃₀ D	C ₂₉ -30-Norm	C ₃₀ H	C ₃₀ Mor	C ₃₁ HH (22S)	C ₃₁ HH (22R)	C ₃₂ HH (22S)	C ₃₂ HH (22R)	C ₃₃ HH (22S)	C ₃₃ HH (22R)	C ₃₄ HH (22S)	C ₃₄ HH (22R)	C ₃₅ HH (22S)	C ₃₅ HH (22R)
EFBC-5	2	ND	0	2	1	3	1	1	ND	0	ND	0	ND	ND	ND	ND
EFBC-7	1	ND	ND	2	1	3	0	1	0	0	ND	ND	ND	ND	ND	ND
EFBC-9	2	ND	ND	3	2	4	1	1	0	0	ND	0	ND	ND	ND	ND
EFBC-11	1	ND	ND	2	1	3	1	1	0	0	ND	0	ND	ND	ND	ND
EFBC-15	1	ND	ND	1	1	2	0	1	0	0	ND	0	ND	ND	ND	ND
EFBC-19	5	ND	ND	4	3	4	1	2	0	1	ND	0	ND	ND	ND	ND
EFBC-20	8	ND	ND	8	6	9	1	5	0	2	ND	0	1	0	0	1
EFBC-23	2	ND	ND	4	2	3	0	3	0	1	ND	0	ND	ND	ND	ND
EFBC-25	2	ND	ND	2	2	2	0	2	0	1	ND	0	ND	ND	ND	ND
EFBC-27	2	ND	ND	3	2	3	0	3	0	1	ND	0	ND	ND	ND	ND
EFBC-30	7	ND	ND	10	4	6	0	3	0	1	ND	0	ND	ND	ND	ND
EFBC-32	14	ND	ND	17	9	13	1	5	0	2	ND	1	ND	ND	ND	ND
EFBC-34	7	ND	ND	8	5	7	0	2	0	1	ND	1	1	0	0	1

ACC #1 core

Sample Name	C ₂₀ TR	C ₂₁ TR	C ₂₂ TR	C ₂₃ TR	C ₂₄ TR	C ₂₅ TR	C ₂₄ TT	C ₂₆ TR	C ₂₆ TR	C ₂₈ TR	C ₂₈ TR	C ₂₉ TR	C ₂₉ TR	T _s	T _m	
EFAC-1	30	22	4	28	14	13	4	6	9	8	4	46	6	4	77	
EFAC-3	28	18	3	17	9	13	3	4	5	8	2	70	6	4	60	
EFAC-6	18	15	2	18	6	9	3	3	3	4	2	22	3	2	22	
EFAC-7	14	9	1	8	3	7	2	2	3	2	2	19	1	1	36	
EFAC-8	7	7	2	6	4	9	2	2	2	4	2	22	2	1	36	
EFAC-10	66	49	9	43	16	19	7	9	22	9	10	106	8	20	214	
Sample Name	C ₂₉ -30-NorH	C ₂₉ Ts	C ₃₀ D	C ₂₉ -30-NorM	C ₃₀ H	C ₃₀ Mor	C ₃₁ HH (22S)	C ₃₁ HH (22R)	C ₃₂ HH (22S)	C ₃₂ HH (22R)	C ₃₃ HH (22S)	C ₃₃ HH (22R)	C ₃₄ HH (22S)	C ₃₄ HH (22R)	C ₃₅ HH (22S)	C ₃₅ HH (22R)
EFAC-1	89	ND	4	83	97	149	21	49	2	12	ND	ND	ND	ND	ND	ND
EFAC-3	63	ND	ND	49	100	119	18	42	2	11	ND	ND	ND	ND	ND	ND
EFAC-6	18	0	ND	12	15	24	6	18	2	7	ND	ND	ND	ND	ND	ND
EFAC-7	32	ND	2	20	20	37	8	17	1	7	ND	ND	ND	ND	ND	ND
EFAC-8	26	ND	2	14	20	38	9	14	1	6	ND	ND	ND	ND	ND	ND
EFAC-10	186	ND	7	230	254	335	80	179	5	41	ND	ND	ND	ND	ND	ND

W. Brechtel #1 core

Sample Name	C ₂₀ TR	C ₂₁ TR	C ₂₂ TR	C ₂₃ TR	C ₂₄ TR	C ₂₅ TR	C ₂₄ TT	C ₂₆ TR	C ₂₆ TR	C ₂₈ TR	C ₂₈ TR	C ₂₉ TR	C ₂₉ TR	T _s	T _m	
EFWB-1	78	41	64	37	56	29	16	14	19	34	ND	5	14	37	149	
EFWB-3	50	35	14	25	22	30	9	9	16	43	ND	5	16	24	106	
EFWB-4	40	26	10	22	17	18	6	7	10	35	ND	3	11	18	70	
EFWB-7	56	28	77	44	42	32	21	16	20	34	ND	10	18	43	134	
EFWB-8	25	20	5	14	19	18	6	6	6	19	1	6	10	18	60	
EFWB-9	70	61	11	84	50	42	13	20	28	7	7	8	11	43	120	
Sample Name	C ₂₉ -30-NorH	C ₂₉ Ts	C ₃₀ D	C ₂₉ -30-NorM	C ₃₀ H	C ₃₀ Mor	C ₃₁ HH (22S)	C ₃₁ HH (22R)	C ₃₂ HH (22S)	C ₃₂ HH (22R)	C ₃₃ HH (22S)	C ₃₃ HH (22R)	C ₃₄ HH (22S)	C ₃₄ HH (22R)	C ₃₅ HH (22S)	C ₃₅ HH (22R)
EFWB-1	223	ND	ND	28	673	112	306	240	189	153	104	88	70	48	83	98
EFWB-3	129	127	ND	ND	370	58	180	127	80	63	41	28	36	22	31	61
EFWB-4	81	ND	ND	7	223	36	112	76	49	39	26	21	33	14	20	36
EFWB-7	ND	ND	157	18	536	47	346	225	178	126	92	60	57	30	70	59
EFWB-8	71	ND	2	5	209	23	110	71	56	36	26	19	21	14	19	25
EFWB-9	197	ND	ND	17	418	36	247	156	138	93	70	44	48	25	53	42

C.J. Hendershot #1 core

Sample Name	C ₂₀ TR	C ₂₁ TR	C ₂₂ TR	C ₂₃ TR	C ₂₄ TR	C ₂₅ TR	C ₂₄ TT	C ₂₆ TR	C ₂₆ TR	C ₂₈ TR	C ₂₈ TR	C ₂₉ TR	C ₂₉ TR	T _s	T _m
EFHE-1	200	167	43	278	190	182	69	79	52	47	40	40	35	121	301
EFHE-2	170	144	21	236	194	183	74	80	45	50	41	41	36	107	314
EFHE-3	126	133	19	224	113	126	60	67	54	28	31	43	45	164	433
EFHE-5	100	77	14	157	72	92	38	38	24	24	24	30	32	66	246
EFHE-6	82	64	10	112	62	73	30	32	19	18	18	29	22	51	187
EFHE-7	93	81	13	137	80	114	38	42	2	24	24	27	29	72	233
EFHE-9	177	137	18	235	159	183	75	77	35	44	48	49	47	91	402
EFHE-10	54	82	22	117	62	67	29	22	67	13	23	19	21	200	307

C.J. Hendershot #1 core (cont.)

Sample Name	C ₂₉ -30- NorH	C ₂₉ Ts	C ₃₀ D	C ₂₉ -30- Norm	C ₃₀ H	C ₃₀ Mor	C ₃₁ HH (22S)	C ₃₁ HH (22R)	C ₃₂ HH (22S)	C ₃₂ HH (22R)	C ₃₃ HH (22S)	C ₃₃ HH (22R)	C ₃₄ HH (22S)	C ₃₄ HH (22R)	C ₃₅ HH (22S)	C ₃₅ HH (22R)
EFHE-1	820	ND	24	143	667	125	263	236	197	207	146	140	96	81	150	164
EFHE-2	ND	25	736	151	776	143	300	261	225	207	184	174	111	88	154	174
EFHE-3	1272	ND	37	184	1655	220	578	445	289	257	198	180	192	122	145	160
EFHE-5	701	ND	25	85	846	116	397	273	224	173	146	114	121	91	113	137
EFHE-6	ND	480	9	57	591	73	282	183	149	113	92	65	77	50	71	85
EFHE-7	640	ND	15	73	851	104	393	260	212	153	138	100	111	76	104	102
EFHE-9	1071	ND	16	98	1620	159	660	427	302	212	199	134	160	93	129	122
EFHE-10	686	188	42	89	1077	119	573	362	508	352	210	142	200	107	281	167

Lily Hoppess #1 core

Sample Name	C ₂₀ TR	C ₂₁ TR	C ₂₂ TR	C ₂₃ TR	C ₂₄ TR	C ₂₅ TR	C ₂₄ TT	C ₂₆ TR	C ₂₆ TR	C ₂₈ TR	C ₂₈ TR	C ₂₉ TR	C ₂₉ TR	T _s	T _m
EFLH-1	149	227	46	541	250	324	130	200	79	119	130	101	111	335	320
EFLH-5	281	475	109	1170	571	740	305	396	134	272	305	231	254	541	595
EFLH-9	218	352	75	795	381	477	194	310	136	179	183	151	166	491	477
EFLH-13	33	46	13	74	35	44	16	23	39	14	15	13	13	107	68
EFLH-15	101	156	47	210	113	116	55	75	92	43	48	38	43	444	190
EFLH-20	66	121	26	191	95	119	47	63	58	40	45	37	39	247	139

Lily Hoppess #1 core (cont.)

Sample Name	C ₂₉ -30- NorH	C ₂₉ Ts	C ₃₀ D	C ₂₉ -30- Norm	C ₃₀ H	C ₃₀ Mor	C ₃₁ HH (22S)	C ₃₁ HH (22R)	C ₃₂ HH (22S)	C ₃₂ HH (22R)	C ₃₃ HH (22S)	C ₃₃ HH (22R)	C ₃₄ HH (22S)	C ₃₄ HH (22R)	C ₃₅ HH (22S)	C ₃₅ HH (22R)
EFLH-1	980	161	71	115	1784	184	483	366	275	190	185	135	148	75	51	52
EFLH-5	1626	220	85	167	2917	315	847	681	493	337	331	225	257	132	105	72
EFLH-9	1388	230	88	167	2510	276	727	537	414	293	290	200	222	117	94	86
EFLH-13	223	51	45	32	403	57	132	86	101	71	75	51	51	27	20	16
EFLH-15	601	420	138	97	1572	193	470	314	313	222	226	156	162	91	55	46
EFLH-20	343	237	89	54	963	108	285	196	189	133	132	93	104	53	41	32

Oils and condensates (Concentrations are expressed as µg biomarkers/g whole oil; ND = not determined)

Sample Name	C ₂₀ TR	C ₂₁ TR	C ₂₂ TR	C ₂₃ TR	C ₂₄ TR	C ₂₅ TR	C ₂₄ TT	C ₂₆ TR	C ₂₆ TR	C ₂₈ TR	C ₂₈ TR	C ₂₉ TR	C ₂₉ TR	T _s	T _m
NWMK-533.2	426	631	139	1028	782	630	219	197	198	189	187	266	216	265	310
NFMK-526-1H	354	446	126	756	651	484	194	158	121	180	187	220	204	199	185
MJOG-UEF	37	21	15	74	49	38	22	18	6	18	23	22	21	43	23
MJOG-LEF	60	79	19	126	91	67	32	29	14	27	27	34	26	63	33
GRES-1-110	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND

Oils and condensates (Concentrations are expressed as µg biomarkers/g whole oil; ND = not determined; cont.)

Sample Name	C ₂₉ -30-NorH	C ₂₉ Ts	C ₃₀ D	C ₂₉ -30-NorM	C ₃₀ H	C ₃₀ Mor	C ₃₁ HH (22S)	C ₃₁ HH (22R)	C ₃₂ HH (22S)	C ₃₂ HH (22R)	C ₃₃ HH (22S)	C ₃₃ HH (22R)	C ₃₄ HH (22S)	C ₃₄ HH (22R)	C ₃₅ HH (22S)	C ₃₅ HH (22R)
NWMK-533.2	156	1	26	38	520	38	197	194	118	83	78	53	44	27	26	11
NFMK-526-1H	85	9	17	15	98	ND	ND	30	ND	ND	ND	ND	ND	ND	ND	ND
MJOG-UEF	10	1	4	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
MJOG-LEF	22	2	18	7	10	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
GRES-1-110	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND

N. Quantitative biomarker analysis results for aryl isoprenoids (Concentrations are expressed as µg biomarkers/g TOC; ND = not determined)

Lozier Canyon outcrop

258

Sample Name	C13 AI	C14 AI	C15 AI	C16 AI	C17 AI	C18 AI	C19 AI	C20 AI	C21 AI	C22 AI	C23 AI	C24 AI	C25 AI	C26 AI	C27 AI	C28 AI	C30 AI	C31 AI
EFLC-28	1	1	1	2	1	2	1	1	0	2	ND	2	ND	ND	ND	ND	ND	ND
EFLC-26	3	2	3	6	2	3	2	2	1	3	ND	4	ND	ND	ND	ND	ND	ND
EFLC-25	0	0	0	1	0	1	0	0	0	1	ND	1	0	0	ND	ND	ND	ND
EFLC-20	ND	ND	ND	ND	ND	3	1	2	2	ND	ND	ND	ND	ND	ND	ND	ND	ND
EFLC-18	0	0	0	0	0	0	0	0	0	0	ND	1	0	0	ND	ND	ND	ND
EFLC-14	0	0	1	1	0	1	1	1	1	0	ND	1	ND	ND	ND	ND	ND	ND
EFLC-12	13	22	44	45	12	48	40	39	19	15	4	18	6	4	4	9	4	5
EFLC-10	81	98	142	115	28	103	94	87	41	26	8	31	16	7	ND	11	ND	ND
EFLC-8	41	52	72	68	15	60	54	50	26	15	6	21	4	4	ND	ND	ND	ND
EFLC-6	1	1	4	6	1	9	9	9	6	3	1	6	2	2	ND	ND	ND	ND
EFLC-4	0	0	0	0	0	1	1	1	0	0	ND	0	0	0	0	ND	ND	ND
EFLC-1	84	242	636	750	164	691	763	549	335	213	ND	156	94	20	ND	104	ND	53

Ferguson McKnight #526-1H core

Sample Name	C₁₃ AI	C₁₄ AI	C₁₅ AI	C₁₆ AI	C₁₇ AI	C₁₈ AI	C₁₉ AI	C₂₀ AI	C₂₁ AI	C₂₂ AI	C₂₃ AI	C₂₄ AI	C₂₅ AI	C₂₆ AI	C₂₇ AI	C₂₈ AI	C₃₀ AI	C₃₁ AI
EFMK-1	290	260	401	519	277	351	279	179	106	100	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-4	330	200	193	263	104	53	67	51	34	33	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-8	654	361	391	612	251	142	171	126	79	92	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-12	441	273	264	362	139	85	104	69	42	49	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-15	251	175	233	386	152	96	116	85	53	55	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-17	1029	664	819	1372	566	996	467	531	178	173	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-20	1043	651	642	854	349	224	273	197	98	120	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-23	788	448	503	862	335	560	296	208	113	137	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-24	649	364	401	645	248	126	179	155	84	119	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-25	135	100	165	296	140	91	133	138	59	75	ND	ND	ND	ND	ND	ND	ND	ND
EFMK-26	265	175	263	476	195	136	187	315	90	108	ND	ND	ND	ND	ND	ND	ND	ND

Bouldin Creek outcrop

Sample Name	C13 AI	C14 AI	C15 AI	C16 AI	C17 AI	C18 AI	C19 AI	C20 AI	C21 AI	C22 AI	C23 AI	C24 AI	C25 AI	C26 AI	C27 AI	C28 AI	C30 AI	C31 AI
EFBC-5	0	1	0	1	0	0	1	0	0	1	ND	1	ND	ND	ND	ND	ND	ND
EFBC-7	ND	0	0	1	0	0	1	0	0	0	0	1	ND	ND	ND	ND	ND	ND
EFBC-9	0	1	1	1	0	0	1	0	0	1	ND	1	ND	ND	ND	ND	ND	ND
EFBC-11	0	0	0	1	0	0	1	0	0	0	ND	1	ND	ND	ND	ND	ND	ND
EFBC-15	0	0	0	0	0	0	0	0	0	0	0	0	ND	ND	ND	ND	ND	ND
EFBC-19	0	2	1	2	1	1	3	1	1	1	0	2	ND	ND	ND	ND	ND	ND
EFBC-20	0	1	1	3	1	2	4	2	1	2	0	2	ND	ND	ND	ND	ND	ND
EFBC-23	0	1	1	3	1	2	3	1	1	2	0	2	ND	ND	ND	ND	ND	ND
EFBC-25	3	6	3	4	1	2	2	1	1	1	0	1	ND	ND	ND	ND	ND	ND
EFBC-27	3	8	4	4	1	2	2	1	1	1	0	2	ND	ND	ND	ND	ND	ND
EFBC-30	3	12	6	11	1	6	10	3	2	3	0	2	ND	ND	ND	ND	ND	ND
EFBC-32	10	34	12	24	2	10	23	6	6	6	ND	4	ND	ND	ND	ND	ND	ND
EFBC-34	8	26	10	19	2	9	20	6	5	4	ND	3	ND	ND	ND	ND	ND	ND

ACC #1 core

Sample Name	C13 AI	C14 AI	C15 AI	C16 AI	C17 AI	C18 AI	C19 AI	C20 AI	C21 AI	C22 AI	C23 AI	C24 AI	C25 AI	C26 AI	C27 AI	C28 AI	C30 AI	C31 AI
EFAC-1	ND	1	4	22	9	29	43	15	10	14	ND	ND	ND	ND	ND	ND	ND	ND
EFAC-3	ND	13	22	50	14	25	60	15	9	15	ND	ND	ND	ND	ND	ND	ND	ND
EFAC-6	ND	1	6	24	8	16	36	15	5	8	ND	ND	ND	ND	ND	ND	ND	ND
EFAC-7	ND	1	7	34	8	30	70	18	18	15	ND	ND	ND	ND	ND	ND	ND	ND
EFAC-8	ND	4	18	67	8	42	132	26	32	22	ND	ND	ND	ND	ND	ND	ND	ND
EFAC-10	ND	3	43	149	35	142	172	63	59	45	ND	ND	ND	ND	ND	ND	ND	ND

W. Brechtel #1 core

Sample Name	C₁₃ AI	C₁₄ AI	C₁₅ AI	C₁₆ AI	C₁₇ AI	C₁₈ AI	C₁₉ AI	C₂₀ AI	C₂₁ AI	C₂₂ AI	C₂₃ AI	C₂₄ AI	C₂₅ AI	C₂₆ AI	C₂₇ AI	C₂₈ AI	C₃₀ AI	C₃₁ AI
EFWB-1	17	91	151	214	90	149	171	89	46	66	8	46	9	ND	49	142	ND	ND
EFWB-3	53	186	175	213	44	108	112	44	33	21	64	22	ND	ND	35	96	ND	ND
EFWB-4	63	179	132	144	30	86	77	31	23	26	3	18	ND	ND	27	69	ND	ND
EFWB-7	195	351	422	462	79	230	235	131	83	75	12	26	ND	ND	60	103	ND	ND
EFWB-8	142	206	145	150	26	74	79	39	28	25	5	12	ND	ND	29	54	ND	ND
EFWB-9	990	762	746	702	173	424	341	311	108	98	18	29	11	ND	22	36	ND	ND

C.J. Hendershot #1 core

Sample Name	C13 AI	C14 AI	C15 AI	C16 AI	C17 AI	C18 AI	C19 AI	C20 AI	C21 AI	C22 AI	C23 AI	C24 AI	C25 AI	C26 AI	C27 AI	C28 AI	C30 AI	C31 AI
EFHE-1	152	286	357	410	141	314	257	143	103	121	23	64	ND	ND	25	108	19	34
EFHE-2	128	246	303	360	117	281	236	126	84	88	58	64	ND	ND	13	108	22	38
EFHE-3	312	485	434	441	133	296	226	92	60	95	56	54	ND	ND	23	115	28	49
EFHE-5	315	507	426	490	97	321	309	150	99	103	43	53	ND	ND	31	151	23	40
EFHE-6	252	422	369	401	85	246	235	112	78	67	27	39	ND	ND	19	82	19	36
EFHE-7	338	541	462	495	106	340	302	154	97	86	97	44	ND	ND	31	121	23	37
EFHE-9	558	969	806	869	155	634	610	318	190	158	76	91	ND	ND	10	144	20	19
EFHE-10	27	68	150	288	78	319	288	167	106	101	23	31	ND	ND	27	28	10	11

Lily Hoppess #1 core

Sample Name	C13 AI	C14 AI	C15 AI	C16 AI	C17 AI	C18 AI	C19 AI	C20 AI	C21 AI	C22 AI	C23 AI	C24 AI	C25 AI	C26 AI	C27 AI	C28 AI	C30 AI	C31 AI
EFLH-1	1430	1844	1362	1302	398	924	747	412	260	265	76	285	ND	ND	48	181	65	65
EFLH-5	1870	2741	2472	2510	715	1841	1656	848	595	495	149	556	ND	ND	74	190	159	200
EFLH-9	1695	2056	1827	1742	544	1235	1028	580	337	412	99	373	ND	ND	51	187	103	140
EFLH-13	181	239	274	278	105	147	119	70	37	34	15	52	ND	ND	12	20	16	11
EFLH-15	1076	1187	1076	1015	339	553	503	251	146	138	74	236	ND	ND	43	124	61	32
EFLH-20	464	669	680	758	270	467	454	219	125	165	50	149	ND	ND	30	80	38	43

Oils and condensates (Concentrations are expressed as µg biomarkers/g whole oil; ND = not determined)

Sample Name	C₁₃ AI	C₁₄ AI	C₁₅ AI	C₁₆ AI	C₁₇ AI	C₁₈ AI	C₁₉ AI	C₂₀ AI	C₂₁ AI	C₂₂ AI	C₂₃ AI	C₂₄ AI	C₂₅ AI	C₂₆ AI	C₂₇ AI	C₂₈ AI	C₃₀ AI	C₃₁ AI
NWMK-533.2	2716	1316	785	919	577	433	517	739	263	300	ND	ND	ND	ND	ND	ND	ND	ND
NFMK-526-1H	2720	980	554	777	564	248	333	1530	245	367	ND	ND	ND	ND	ND	ND	ND	ND
MJOG-UEF	845	133	79	137	120	ND	62	406	50	62	ND	ND	ND	ND	ND	ND	ND	ND
MJOG-LEF	658	97	95	171	195	ND	61	365	48	77	ND	21	28	48	19	58	31	28
GRES-1-110	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND

O. Quantitative biomarker analysis results for monoaromatic steroids (Concentrations are expressed as µg biomarkers/g TOC; ND = not determined)

Lozier Canyon outcrop

Sample Name	Preg	20-Mpreg	C ₂₇ 5 -Chol (20S)	C ₂₇ Diachol (20S)	C ₂₇ MAs	C ₂₈ 5 -Ergost (20S) + C ₂₈ Diaergost (20S)	C ₂₇ 5 -Chol (20R)	C ₂₈ 5 -Ergost (20S)	C ₂₈ 5 -Ergost (20R) + C ₂₈ Diaergost (20R)	C ₂₉ 5 -Stigmast (20S)+C ₂₉ Diastig (20R)	C ₂₉ 5 -Stigmast (20S)	C ₂₈ 5 -Ergost (20R)	C ₂₉ 5 -Stigmast (20R) + C ₂₉ Diastigmast (20R)	C ₂₉ 5 -Stigmast (20R)
EFLC-28	3	3	5	2	5	25	4	35	1	1	7	10	4	3
EFLC-26	7	5	6	3	5	28	4	38	1	1	8	11	4	3
EFLC-25	1	1	1	1	1	7	1	11	0	0	2	3	1	1
EFLC-20	3	4	8	3	7	36	4	47	2	2	8	12	4	3
EFLC-18	0	0	0	0	0	2	0	2	0	0	0	1	0	0
EFLC-14	2	1	1	0	1	5	0	5	0	0	1	1	1	0
EFLC-12	8	5	6	2	5	25	2	28	1	1	4	6	3	2
EFLC-10	16	4	7	2	6	26	3	29	1	1	6	6	4	1
EFLC-8	18	6	7	2	6	26	3	29	1	1	6	6	4	1
EFLC-6	2	2	2	1	2	7	1	8	0	0	1	2	1	1
EFLC-4	1	0	0	0	0	1	0	2	0	0	0	0	0	0
EFLC-1	234	368	770	360	802	2825	ND	3665	100	91	489	856	337	155

Ferguson McKnight #526-1H core

Sample Name	Preg	20-Mpreg	C ₂₇ 5 -Chol (20S)	C ₂₇ Diachol (20S)	C ₂₇ MAS	C ₂₈ 5 -Ergost (20S) + C ₂₈ Diaergost (20S)	C ₂₇ 5 -Chol (20R)	C ₂₈ 5 -Ergost (20S)	C ₂₈ 5 -Ergost (20R) + C ₂₈ Diaergost (20R)	C ₂₉ 5 -Stigmast (20S)+C ₂₉ Diastig (20R)	C ₂₉ 5 -Stigmast (20S)	C ₂₈ 5 -Ergost (20R)	C ₂₉ 5 -Stigmast (20R) + C ₂₉ Diastigmast (20R)	C ₂₉ 5 -Stigmast (20R)
EFMK-1	135	62	17	13	26	51	13	52	4	11	14	13	9	5
EFMK-4	75	36	8	5	11	21	6	21	2	5	6	5	3	2
EFMK-8	158	70	15	8	19	37	9	38	2	8	11	10	7	6
EFMK-12	72	35	7	4	9	20	5	20	2	4	6	7	2	2
EFMK-15	51	29	7	4	10	24	6	24	2	5	8	9	2	3
EFMK-17	166	92	30	16	44	102	24	104	7	19	34	23	18	12
EFMK-20	107	63	17	9	23	54	10	52	6	12	19	21	4	5
EFMK-23	109	62	18	8	26	50	11	51	6	14	21	21	5	6
EFMK-24	94	53	15	6	24	45	8	45	4	12	17	18	3	5
EFMK-25	56	30	7	3	13	27	6	21	3	6	9	6	4	2
EFMK-26	60	22	6	5	10	22	5	18	2	3	5	5	4	3

Bouldin Creek outcrop

Sample Name	Preg	20-Mpreg	C ₂₇ 5 -Chol (20S)	C ₂₇ Diachol (20S)	C ₂₇ MAS	C ₂₈ 5 -Ergost (20S) + C ₂₈ Diaergost (20S)	C ₂₇ 5 -Chol (20R)	C ₂₈ 5 -Ergost (20S)	C ₂₈ 5 -Ergost (20R) + C ₂₈ Diaergost (20R)	C ₂₉ 5 -Stigmast (20S)+C ₂₉ Diastig (20R)	C ₂₉ 5 -Stigmast (20S)	C ₂₈ 5 -Ergost (20R)	C ₂₉ 5 -Stigmast (20R) + C ₂₉ Diastigmast (20R)	C ₂₉ 5 -Stigmast (20R)
EFBC-5	2	5	2	3	5	47	7	85	3	4	15	8	41	12
EFBC-7	2	4	2	3	4	43	6	81	2	3	13	7	39	10
EFBC-9	3	6	3	4	6	66	9	124	4	5	21	11	60	17
EFBC-11	2	5	3	3	5	62	7	115	4	5	19	10	53	15
EFBC-15	1	3	1	1	2	20	4	40	1	2	12	5	19	9
EFBC-19	7	17	7	9	14	132	22	255	9	11	78	37	114	59
EFBC-20	14	32	12	14	24	208	36	379	13	18	94	51	173	72
EFBC-23	12	22	10	11	19	162	25	265	9	14	54	32	113	40
EFBC-25	11	21	10	12	19	164	25	269	12	16	49	33	114	37
EFBC-27	12	23	11	11	20	140	25	228	9	13	49	35	94	38
EFBC-30	7	17	5	6	11	58	16	103	6	5	29	15	50	23
EFBC-32	11	28	11	11	21	125	38	239	15	10	76	35	131	63
EFBC-34	7	18	7	7	13	78	21	156	6	7	53	23	78	43

ACC #1 core

Sample Name	Preg	20-Mpreg	C ₂₇ 5 -Chol (20S)	C ₂₇ Diachol (20S)	C ₂₇ MAS	C ₂₈ 5 -Ergost (20S) + C ₂₈ Diaergost (20S)	C ₂₇ 5 -Chol (20R)	C ₂₈ 5 -Ergost (20S)	C ₂₈ 5 -Ergost (20R) + C ₂₈ Diaergost (20R)	C ₂₉ 5 -Stigmast (20S)+C ₂₉ Diastig (20R)	C ₂₉ 5 -Stigmast (20S)	C ₂₈ 5 -Ergost (20R)	C ₂₉ 5 -Stigmast (20R) + C ₂₉ Diastigmast (20R)	C ₂₉ 5 -Stigmast (20R)
EFAC-1	40	183	81	91	143	1382	237	3077	108	100	572	184	1471	443
EFAC-3	72	216	72	91	139	1328	216	3071	106	99	668	226	1493	514
EFAC-6	48	200	76	104	147	1151	156	1950	80	88	289	16	785	209
EFAC-7	28	73	23	28	46	267	103	513	24	20	152	46	261	116
EFAC-8	16	81	25	31	48	312	132	668	27	23	187	58	371	147
EFAC-10	46	206	81	100	155	869	321	1734	86	71	555	234	885	431

W. Brechtel #1 core

Sample Name	Preg	20-Mpreg	C ₂₇ 5 -Chol (20S)	C ₂₇ Diachol (20S)	C ₂₇ MAS	C ₂₈ 5 -Ergost (20S) + C ₂₈ Diaergost (20S)	C ₂₇ 5 -Chol (20R)	C ₂₈ 5 -Ergost (20S)	C ₂₈ 5 -Ergost (20R) + C ₂₈ Diaergost (20R)	C ₂₉ 5 -Stigmast (20S)+C ₂₉ Diastig (20R)	C ₂₉ 5 -Stigmast (20S)	C ₂₈ 5 -Ergost (20R)	C ₂₉ 5 -Stigmast (20R) + C ₂₉ Diastigmast (20R)	C ₂₉ 5 -Stigmast (20R)
EFWB-1	941	1575	2672	302	2341	7283	2052	8464	117	131	1272	1503	1537	580
EFWB-3	1041	2116	4070	432	3218	16606	3033	1925	331	228	2929	3202	6149	900
EFWB-4	717	1466	2929	422	2438	11878	2449	13723	227	162	1471	2278	2313	719
EFWB-7	864	1700	3315	456	2695	10670	2640	12025	188	126	1476	2063	1954	646
EFWB-8	464	1027	2064	390	1774	7955	1708	8697	141	117	972	1414	1447	410
EFWB-9	443	848	1267	163	1004	3070	876	3403	38	35	505	654	498	195

C.J. Hendershot #1 core

Sample Name	Preg	20-Mpreg	C ₂₇ 5 -Chol (20S)	C ₂₇ Diachol (20S)	C ₂₇ MAS	C ₂₈ 5 -Ergost (20S) + C ₂₈ Diaergost (20S)	C ₂₇ 5 -Chol (20R)	C ₂₈ 5 -Ergost (20S)	C ₂₈ 5 -Ergost (20R) + C ₂₈ Diaergost (20R)	C ₂₉ 5 -Stigmast (20S)+C ₂₉ Diastig (20R)	C ₂₉ 5 -Stigmast (20S)	C ₂₈ 5 -Ergost (20R)	C ₂₉ 5 -Stigmast (20R) + C ₂₉ Diastigmast (20R)	C ₂₉ 5 -Stigmast (20R)
EFHE-1	751	764	1125	263	1163	3573	1284	891	119	117	913	1147	769	432
EFHE-2	777	802	1113	234	1161	3688	1387	4587	164	89	963	1173	836	472
EFHE-3	471	796	938	298	1106	3881	6202	4792	230	109	924	1195	825	445
EFHE-5	507	1112	1683	537	1719	8516	2478	9624	495	329	1407	2157	1478	597
EFHE-6	424	1028	1340	451	1318	5683	1752	6011	317	222	898	1387	891	371
EFHE-7	465	1160	1617	604	1650	7107	2121	7958	406	304	1099	1734	1079	463
EFHE-9	363	853	1222	355	1168	4300	6046	4683	217	143	753	1069	711	283
EFHE-10	210	464	591	208	596	1902	653	2160	116	74	378	583	287	153

Fasken "A" #1 core

Sample Name	Preg	20-Mpreg	C ₂₇ 5 -Chol (20S)	C ₂₇ Diachol (20S)	C ₂₇ MAS	C ₂₈ 5 -Ergost (20S) + C ₂₈ Diaergost (20S)	C ₂₇ 5 -Chol (20R)	C ₂₈ 5 -Ergost (20S)	C ₂₈ 5 -Ergost (20R) + C ₂₈ Diaergost (20R)	C ₂₉ 5 -Stigmast (20S)+C ₂₉ Diastig (20R)	C ₂₉ 5 -Stigmast (20S)	C ₂₈ 5 -Ergost (20R)	C ₂₉ 5 -Stigmast (20R) + C ₂₉ Diastigmast (20R)	C ₂₉ 5 -Stigmast (20R)
EFFA-1	4	1	0	0	0	1	1	0	0	0	0	0	0	0
EFFA-3	0	0	0	0	0	0	0	0	0	0	0	0	0	0
EFFA-5	14	5	1	1	2	5	1	5	0	1	1	1	1	0
EFFA-11	18	7	2	1	2	7	2	7	1	1	2	2	1	1
EFFA-17	14	4	1	1	2	5	1	4	0	1	1	1	1	0
EFFA-18	15	6	1	1	2	5	1	5	0	1	1	1	1	1
EFFA-20	3	1	0	0	0	1	0	1	0	0	0	0	0	0
EFFA-21	6	1	0	0	0	1	0	1	0	0	0	0	0	0
EFFA-23	10	2	1	0	1	2	0	2	0	0	0	1	0	0
EFFA-24	49	13	3	2	4	12	2	11	1	1	2	3	1	1
EFFA-28	34	9	2	2	2	8	1	8	1	1	2	2	1	1
EFFA-32	57	19	5	4	5	17	4	18	1	2	3	5	2	1
EFFA-37	9	2	0	0	0	2	0	2	0	0	0	0	0	0
EFFA-38	26	9	2	2	3	8	2	9	1	1	1	2	1	1
EFFA-40	7	2	1	1	1	2	1	2	0	0	0	1	0	0

Lily Hoppess #1 core

Sample Name	Preg	20-Mpreg	C ₂₇ 5 -Chol (20S)	C ₂₇ Diachol (20S)	C ₂₇ MAS	C ₂₈ 5 -Ergost (20S) + C ₂₈ Diaergost (20S)	C ₂₇ 5 -Chol (20R)	C ₂₈ 5 -Ergost (20S)	C ₂₈ 5 -Ergost (20R) + C ₂₈ Diaergost (20R)	C ₂₉ 5 -Stigmast (20S)+C ₂₉ Diastig (20R)	C ₂₉ 5 -Stigmast (20S)	C ₂₈ 5 -Ergost (20R)	C ₂₉ 5 -Stigmast (20R) + C ₂₉ Diastigmast (20R)	C ₂₉ 5 -Stigmast (20R)
EFLH-1	424	214	111	217	205	607	168	710	63	105	152	297	111	48
EFLH-5	556	267	149	277	271	920	232	1007	89	144	205	257	149	69
EFLH-9	446	208	111	212	202	600	150	708	62	102	159	207	98	62
EFLH-13	67	26	10	39	33	77	120	120	14	25	19	39	5	5
EFLH-15	221	126	59	170	138	431	121	543	66	108	127	172	65	44
EFLH-20	211	119	47	130	112	317	87	393	48	78	87	130	56	31

Oils and condensates (Concentrations are expressed as µg biomarkers/g whole oil; ND = not determined)

Sample Name	Preg	20-Mpreg	C ₂₇ 5 -Chol (20S)	C ₂₇ Diachol (20S)	C ₂₇ MAS	C ₂₈ 5 -Ergost (20S) + C ₂₈ Diaergost (20S)	C ₂₇ 5 -Chol (20R)	C ₂₈ 5 -Ergost (20S)	C ₂₈ 5 -Ergost (20R) + C ₂₈ Diaergost (20R)	C ₂₉ 5 -Stigmast (20S)+C ₂₉ Diastig (20R)	C ₂₉ 5 -Stigmast (20S)	C ₂₈ 5 -Ergost (20R)	C ₂₉ 5 -Stigmast (20R) + C ₂₉ Diastigmast (20R)	C ₂₉ 5 -Stigmast (20R)
NWMK-533.2	572	207	70	57	92	240	45	234	19	47	72	74	21	28
NFMK-526-1H	398	110	33	9	51	98	24	94	8	12	48	18	28	14
MJOG-UEF	91	24	3	5	6	13	6	17	5	4	14	4	8	4
MJOG-LEF	163	33	5	5	9	17	6	20	4	11	16	4	10	4
GREF-1-110	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND

P. Quantitative biomarker analysis results for phenanthrenes and triaromatic steroids (Concentrations are expressed as µg biomarkers/g TOC; ND = not determined)

Lozier Canyon outcrop

Sample Name	P	3-MP	2-MP	9+4-MP	1-MP	C ₂₀ Preg	C ₂₁ 20-Mpreg	C ₂₆ Chol (20S)	C ₂₆ Chol (20R) + C ₂₇ Ergost (20S)	C ₂₈ Stig (20S)	C ₂₇ Ergost (20R)	C ₂₈ Stig (20R)
EFLC-28	4004	262	472	505	597	32	33	41	194	115	117	117
EFLC-26	6639	195	448	415	518	24	30	32	159	94	96	86
EFLC-25	3184	158	331	369	439	30	37	50	249	165	155	155
EFLC-20	5217	233	521	538	652	34	45	83	441	201	283	190
EFLC-18	2940	61	166	123	189	18	15	12	68	28	38	22
EFLC-14	1751	71	148	128	164	34	14	9	48	18	28	13
EFLC-12	11659	3573	4252	6133	4792	801	390	568	3081	948	2024	799
EFLC-10	5357	1597	2100	2544	2036	120	95	108	525	179	335	146
EFLC-8	6408	975	1556	1673	1580	228	46	59	263	97	163	74
EFLC-6	11033	2094	3149	3792	3298	274	161	192	819	279	525	238
EFLC-4	1683	47	107	84	105	55	10	10	50	18	30	15
EFLC-1	1591	1064	989	2465	1153	1375	1312	2423	8984	2652	5384	2566

276

Ferguson McKnight #526-1H core

Sample Name	P	3-MP	2-MP	9+4-MP	1-MP	C₂₀ Preg	C₂₁ 20-Mpreg	C₂₆ Chol (20S)	C₂₆ Chol (20R) + C₂₇ Ergost (20S)	C₂₈ Stig (20S)	C₂₇ Ergost (20R)	C₂₈ Stig (20R)
EFMK-1	6280	3977	4734	6788	5027	462	622	82	311	182	206	175
EFMK-4	7852	4257	5607	8187	5809	494	616	48	196	119	143	117
EFMK-8	15290	10216	12489	19042	12212	1063	960	66	268	194	195	177
EFMK-12	9036	5134	6768	9851	7226	476	527	35	140	97	108	93
EFMK-15	3974	2692	3413	5070	3623	316	394	28	136	87	99	83
EFMK-17	16140	10525	13369	19619	14079	1083	1330	116	489	311	343	272
EFMK-20	11271	7357	9545	13832	9990	781	966	65	315	218	231	208
EFMK-23	11624	7908	10406	15083	10664	853	1037	85	363	272	254	235
EFMK-24	10489	7015	9423	14465	9922	768	901	66	294	226	210	196
EFMK-25	8998	6077	8574	12678	9015	695	767	56	242	142	182	131
EFMK-26	115578	80223	110884	161085	117384	6826	6771	432	1861	1092	1325	1322

277

Fasken "A" #1 core

278

Sample Name	P	3-MP	2-MP	9+4-MP	1-MP	C ₂₀ Preg	C ₂₁ 20-Mpreg	C ₂₆ Chol (20S)	C ₂₆ Chol (20R) + C ₂₇ Ergost (20S)	C ₂₈ Stig (20S)	C ₂₇ Ergost (20R)	C ₂₈ Stig (20R)
EFFA-1	1598	659	805	419	338	2	1	0	0	0	0	0
EFFA-3	2364	1013	1123	734	506	7	4	0	2	1	1	1
EFFA-5	2872	1348	1524	1042	832	11	4	1	2	1	1	1
EFFA-11	2748	1435	1678	1161	951	15	7	1	3	2	2	1
EFFA-17	1945	909	1023	759	538	7	3	0	1	1	1	1
EFFA-18	2864	1358	1569	1132	845	12	6	1	2	1	1	1
EFFA-20	453	203	239	163	122	1	1	0	0	0	0	0
EFFA-21	909	347	385	254	175	2	1	0	0	0	0	0
EFFA-23	1178	530	605	430	305	3	1	0	1	0	0	0
EFFA-24	3833	1864	1910	1616	1058	15	5	1	3	2	2	2
EFFA-28	3107	1485	1540	1241	851	12	4	1	2	2	1	1
EFFA-32	4203	2179	2310	1912	1271	18	8	1	4	3	2	2
EFFA-37	1642	552	627	389	261	2	1	0	0	0	0	0
EFFA-38	2828	1542	1632	1347	954	10	4	1	3	2	1	1
EFFA-40	1043	449	493	361	245	3	2	0	1	0	0	0

Bouldin Creek outcrop

Sample Name	P	3-MP	2-MP	9+4-MP	1-MP	C ₂₀ Preg	C ₂₁ 20-Mpreg	C ₂₆ Chol (20S)	C ₂₆ Chol (20R) + C ₂₇ Ergost (20S)	C ₂₈ Stig (20S)	C ₂₇ Ergost (20R)	C ₂₈ Stig (20R)
EFBC-5	15	7	5	16	12	3	1	2	13	4	7	3
EFBC-7	11	4	4	8	7	2	1	1	9	3	5	2
EFBC-9	15	6	5	12	10	2	1	1	13	4	7	3
EFBC-11	7	4	3	8	7	2	1	1	11	3	6	2
EFBC-15	19	5	6	10	10	2	2	2	12	5	7	3
EFBC-19	53	6	6	15	13	2	1	1	10	5	6	4
EFBC-20	19	4	4	11	12	2	1	1	9	4	5	4
EFBC-23	27	4	5	10	9	2	1	1	9	3	5	3
EFBC-25	17	4	5	11	9	2	1	1	10	4	6	3
EFBC-27	11	6	7	18	13	3	1	2	11	5	6	3
EFBC-30	6	3	4	9	6	1	1	1	3	1	2	2
EFBC-32	7	5	6	13	9	2	1	1	7	3	4	3
EFBC-34	11	8	9	17	10	2	1	1	7	3	4	2

ACC #1 core

Sample Name	P	3-MP	2-MP	9+4-MP	1-MP	C ₂₀ Preg	C ₂₁ 20-Mpreg	C ₂₆ Chol (20S)	C ₂₆ Chol (20R) + C ₂₇ Ergost (20S)	C ₂₈ Stig (20S)	C ₂₇ Ergost (20R)	C ₂₈ Stig (20R)
EFAC-1	90	15	16	25	31	3	2	2	14	5	8	4
EFAC-3	65	11	12	17	23	2	3	1	12	5	7	3
EFAC-6	35	8	8	14	12	4	2	2	11	4	7	3
EFAC-7	56	10	11	16	17	2	3	1	7	3	4	2
EFAC-8	25	7	8	12	16	2	2	1	6	3	4	2
EFAC-10	155	39	38	41	59	4	3	1	9	4	5	3

W. Brechtel #1 core

Sample Name	P	3-MP	2-MP	9+4-MP	1-MP	C₂₀ Preg	C₂₁ 20-Mpreg	C₂₆ Chol (20S)	C₂₆ Chol (20R) + C₂₇ Ergost (20S)	C₂₈ Stig (20S)	C₂₇ Ergost (20R)	C₂₈ Stig (20R)
EFWB-1	572	200	272	215	247	45	70	106	387	119	242	285
EFWB-3	699	215	306	242	248	77	110	207	1054	244	715	451
EFWB-4	530	169	227	197	191	67	100	194	916	232	611	348
EFWB-7	9181	3431	4616	6977	4599	135	154	258	1061	289	695	555
EFWB-8	4264	1092	1526	2211	1630	82	105	205	887	232	579	372
EFWB-9	6754	2980	4187	6302	4020	297	217	90	356	120	228	186

C.J. Hendershot #1 core

Sample Name	P	3-MP	2-MP	9+4-MP	1-MP	C₂₀ Preg	C₂₁ 20-Mpreg	C₂₆ Chol (20S)	C₂₆ Chol (20R) + C₂₇ Ergost (20S)	C₂₈ Stig (20S)	C₂₇ Ergost (20R)	C₂₈ Stig (20R)
EFHE-1	219	89	79	121	84	29	24	41	150	48	82	46
EFHE-2	172	78	62	110	77	26	20	35	132	41	74	40
EFHE-3	186	80	70	114	80	29	21	34	148	40	88	50
EFHE-5	162	71	57	110	79	45	40	91	452	108	263	104
EFHE-6	115	54	41	84	60	42	39	84	348	85	206	79
EFHE-7	157	72	56	115	80	53	48	106	450	111	257	105
EFHE-9	223	93	69	131	93	3	45	105	391	92	225	91
EFHE-10	588	231	232	427	289	36	24	54	162	48	84	47

Lily Hoppess #1 core

Sample Name	P	3-MP	2-MP	9+4-MP	1-MP	C₂₀ Preg	C₂₁ 20-Mpreg	C₂₆ Chol (20S)	C₂₆ Chol (20R) + C₂₇ Ergost (20S)	C₂₈ Stig (20S)	C₂₇ Ergost (20R)	C₂₈ Stig (20R)
EFLH-1	1905	883	780	1519	1087	186	152	127	468	304	244	250
EFLH-5	1722	788	707	1366	1025	143	115	89	378	217	204	174
EFLH-9	3044	1257	1147	2181	1685	59	140	91	371	247	197	203
EFLH-13	906	382	353	628	529	42	28	12	41	30	23	24
EFLH-15	2158	883	848	1441	1195	100	77	41	168	130	91	101
EFLH-20	2170	894	826	1524	1174	127	87	42	183	151	99	114

Oils and condensates (Concentrations are expressed as µg biomarkers/g whole oil; ND = not determined)

Sample Name	P	3-MP	2-MP	9+4-MP	1-MP	C₂₀ Preg	C₂₁ 20-Mpreg	C₂₆ Chol (20S)	C₂₆ Chol (20R) + C₂₇ Ergost (20S)	C₂₈ Stig (20S)	C₂₇ Ergost (20R)	C₂₈ Stig (20R)
NWMK-533.2	8110	4994	5447	10290	6419	686	628	38	198	119	111	87
NFMK-526-1H	4123	3144	3984	7703	4513	335	292	12	59	33	35	24
MJOG-UEF	5738	4506	5535	8830	5914	180	110	4	6	7	9	3
MJOG-LEF	8507	6263	7283	9639	6863	234	166	4	4	6	6	3
GREF-1-110	13775	7404	8028	7589	5555	50	62	ND	ND	ND	ND	ND

Q. ^{13}C values for *n*-alkanes of Eagle Ford Shale bitumens and oils analyzed in this study. ^{13}C values are expressed in per mil (‰) relative to the Vienna Pee Dee belemnite (VPDB) standard

Lozier Canyon outcrop

	C ₁₂	C ₁₃	C ₁₄	C ₁₅	C ₁₆	C ₁₇	Pr	C ₁₈	Ph	C ₁₉	C ₂₀	C ₂₁	C ₂₂	C ₂₃	C ₂₄	C ₂₅	C ₂₆	C ₂₇	C ₂₈	C ₂₉	C ₃₀	C ₃₁		
EFLC-28					-29.3	-29.4	-29.6	-29.3	-30.5	-29.4	-28.7	-29.7	-30.1	-30.1	-28.8	-29.2	-29.5	-29.1	-27.5					
EFLC-26				-23.3	-26.7	-27	-27.5	-26.7	-28.7	-26.8	-28.2	-27.8	-27.3	-28.3	-27.9	-28	-27.7	-27.3	-25.6					
EFLC-20																	-32.7		-34					
EFLC-14			-31.2	-30.4	-30.6	-29.9	-30.2	-30.1	-30.7	-30.1	-30	-30.3	-30.6	-30.7	-30.6	-31.3	-30.3	-30.1	-30.8	-31.4				
EFLC-10		-31.5	-32.4	-30.7	-31	-31	-31.1	-30.6	-31.9	-32.5	-30.5	-31.5	-31.6	-31.7	-31.2	-32	-31.6	-32	-31.3	-31.3				
EFLC-8		-31.5	-31.7	-30.8	-31	-31.1	-30.7	-31.2	-32	-31.6	-31.1	-31.5	-31.1	-31.3	-31.2	-30.5	-31.7	-31	-30.8	-28.1				

Ferguson McKnight #526-1H core

	C ₁₂	C ₁₃	C ₁₄	C ₁₅	C ₁₆	C ₁₇	Pr	C ₁₈	Ph	C ₁₉	C ₂₀	C ₂₁	C ₂₂	C ₂₃	C ₂₄	C ₂₅	C ₂₆	C ₂₇	C ₂₈	C ₂₉	C ₃₀	C ₃₁		
EFMK-1				-29.2	-29.0	-29.4	-31.0	-30.1	-30.0	-29.6	-30.2	-30.0	-29.7	-30.7										
EFMK-24		-31.0	-30.5	-30.1	-29.8	-29.6	-30.3	-29.8	-30.4	-30.2	-30.1	-30.3	-29.3	-30.8	-30.5	-29.3	-30.6	-31.6						
EFMK-25			-29.9	-29.8	-30.2	-29.7	-30.2	-29.8	-30.2	-30.0	-29.7	-30.1	-30.3	-30.6	-31.4	-27.8	-31.1							

Fasken "A" #1 core

	C ₁₂	C ₁₃	C ₁₄	C ₁₅	C ₁₆	C ₁₇	Pr	C ₁₈	Ph	C ₁₉	C ₂₀	C ₂₁	C ₂₂	C ₂₃	C ₂₄	C ₂₅	C ₂₆	C ₂₇	C ₂₈	C ₂₉	C ₃₀	C ₃₁	
EFFA-3	-31.2	-30.0	-30.9	-29.3	-29.3	-29.5	-30.5	-29.4	-31.3	-30.2	-30.1	-30.6	-30.7	-30.1	-29.9	-30.2							
EFFA-18		-30.2	-30.7	-29.3	-29.2	-29.3	-30.2	-29.3	-31.3	-30.6	-30.4	-30.1	-29.8	-30.2	-30.5	-29.9							
EFFA-21		-29.8	-29.3	-29.2	-29.9	-29.3	-29.9	-30.2	-30.2	-29.6	-30.4	-30.2	-29.6	-30.3	-30.0	-32.3							
EFFA-24	-28.9	-30.2	-30.6	-30.1	-30.2	-29.7	-30.6	-30.0	-30.8	-30.6	-29.7	-30.2	-30.1	-30.3	-29.8	-30.6	-29.7	-30.2	-30.7	-30.0	-29.9	-30.4	
EFFA-40			-29.9	-29.6	-29.6	-29.3	-29.8	-29.4	-30.6	-30.4	-29.4	-29.7	-29.5	-29.9									

Bouldin Creek outcrop

	C ₁₂	C ₁₃	C ₁₄	C ₁₅	C ₁₆	C ₁₇	Pr	C ₁₈	Ph	C ₁₉	C ₂₀	C ₂₁	C ₂₂	C ₂₃	C ₂₄	C ₂₅	C ₂₆	C ₂₇	C ₂₈	C ₂₉	C ₃₀	C ₃₁	
EFBC-5						-33.7	-31.8	-33.1	-30.7	-34.5	-32.2	-32.3	-31.3	-29.7	-32.5	-32.1							
EFBC-20						-32.2	-32.3	-33.2	-31.3	-33.0	-32.1	-32.0	-30.8	-30.3	-31.2	-30.6							
EFBC-25				-34.4	-33.0	-33.5	-32.2	-33.5	-31.3	-34.2	-31.6	-32.8	-31.1	-29.9	-32.0	-32.0							

ACC #1 core

	C ₁₂	C ₁₃	C ₁₄	C ₁₅	C ₁₆	C ₁₇	Pr	C ₁₈	Ph	C ₁₉	C ₂₀	C ₂₁	C ₂₂	C ₂₃	C ₂₄	C ₂₅	C ₂₆	C ₂₇	C ₂₈	C ₂₉	C ₃₀	C ₃₁	
EFAC-1						-31.2	-32.4	-31.5	-31.4	-32.2	-31.4	-30.1	-29.9	-30.2	-30.3	-30.8	-30.4						
EFAC-7					-30.6	-31.7	-32.7	-31.9	-31.6	-32.9	-31.8	-31.7	-30.4	-30.9	-30.3	-32.3							
EFAC-10					-31.2	-32.0	-32.1	-32.4	-31.6	-34.0	-33.3	-32.5	-31.5	-31.7	-32.3	-32.2							

W. Brechtel #1 core

	C ₁₂	C ₁₃	C ₁₄	C ₁₅	C ₁₆	C ₁₇	Pr	C ₁₈	Ph	C ₁₉	C ₂₀	C ₂₁	C ₂₂	C ₂₃	C ₂₄	C ₂₅	C ₂₆	C ₂₇	C ₂₈	C ₂₉	C ₃₀	C ₃₁	
EFWB-1					-31.3	-30.2	-30.7	-30.8	-31.3	-31.8	-30.4	-30.0	-29.9	-31.3	-31.2	-31.8	-30.5						
EFWB-3					-31.9	-31.1	-32.4	-32.0	-32.7	-34.0	-30.7	-31.2	-30.7	-32.9	-30.8	-31.4	-30.7						
EFWB-7					-33.2	-31.7	-32.1	-32.3	-33.1	-34.0	-31.1	-31.1	-30.2	-30.4	-30.9	-31.9	-32.6						
EFWB-8					-31.4	-30.4	-31.8	-31.0	-32.8	-31.9	-31.0	-30.3	-29.9	-30.7	-29.8	-30.5	-29.6						

C.J. Hendershot #1 core

	C ₁₂	C ₁₃	C ₁₄	C ₁₅	C ₁₆	C ₁₇	Pr	C ₁₈	Ph	C ₁₉	C ₂₀	C ₂₁	C ₂₂	C ₂₃	C ₂₄	C ₂₅	C ₂₆	C ₂₇	C ₂₈	C ₂₉	C ₃₀	C ₃₁	
EFHE-1			-	30.6	30.5	29.7	30.3	30.7	31.8	-29.9	-30.7	-31.6	-33.0	-30.2	-30.1	-30.0	-30.1						
EFHE-5			-	32.2	30.7	31.9	31.4	32.2	31.5	-32.3	-32.6	-30.8	-32.1	-31.6	-32.6	-31.8	-31.0	-31.8					
EFHE-7			-	31.7	31.4	33.4	31.1	32.0	31.6	-32.3	-34.2	-31.8	-32.1	-31.8	-33.7	-31.5	-31.9	-32.3					
EFHE-9			-	32.2	32.5	33.6	32.5	31.8	32.0	-32.1	-33.3	-32.0	-33.1	-32.2	-34.4	-32.1	-33.1	-32.8					
EFHE-10			-	32.4	31.9	31.7	32.0	32.1	32.1	-32.3	-33.1	-32.5	-33.2	-33.1	-34.4	-32.6	-32.5	-32.3	-32.4	-32.0	-32.0	-34.3	

Lily Hoppess #1 core

	C ₁₂	C ₁₃	C ₁₄	C ₁₅	C ₁₆	C ₁₇	Pr	C ₁₈	Ph	C ₁₉	C ₂₀	C ₂₁	C ₂₂	C ₂₃	C ₂₄	C ₂₅	C ₂₆	C ₂₇	C ₂₈	C ₂₉	C ₃₀	C ₃₁	
EFLH-5			-	31.9	31.4	30.1	31.2	29.9	31.3	-29.7	-33.6	-32.0	-32.3	-32.4	-31.5	-30.9	-30.9	-31.1	-27.1	-30.0			
EFLH-13			-		32.0	32.2	32.0	31.9	32.1	-31.8	-31.9	-32.3	-32.7	-32.4	-32.6	-32.2	-31.6	-31.7	-32.8	-31.1	-31.8	-30.6	
EFLH-20		-31.3	-	32.7	31.7	32.0	32.2	32.0	32.2	-31.6	-32.0	-32.5	-33.4	-32.8	-33.3	-32.5	-32.3	-32.7	-34.3	-32.3	-32.1	-32.4	-35.0

Oils and condensates

	C₁₁	C₁₂	C₁₃	C₁₄	C₁₅	C₁₆	C₁₇	Pr	C₁₈	Ph	C₁₉	C₂₀	C₂₁	C₂₂	C₂₃	C₂₄	C₂₅	C₂₆	C₂₇	C₂₈	C₂₉
NWMK-533.2		-29.6	-31.7	-32.3	-31.6	-30.1	-32.5	-31.1	-32.2	-31.0	-31.0	-30.7	-31.3	-30.6	-31.2	-31.7	-31.8	-31.8	-29.7	-32.0	
NFMK-526-1H		-28.6	-31.8	-32.7	-30.7	-30.0	-31.3	-29.6	-31.6	-30.2	-29.7	-30.2	-31.2	-30.0	-31.3	-31.4	-28.7				
MJOG-UEF	-28.8	-28.1	-31.3	-31.1	-30.0	-30.4	-30.5	-30.6	-30.2	-31.2	-31.1	-30.2	-30.3	-30.1	-30.5	-30.0	-30.5	-29.8	-28.9	-30.1	
MJOG-LEF	-30.4	-27.7	-29.4	-31.2	-29.9	-30.4	-31.6	-32.0	-31.2	-31.5	-30.7	-31.3	-30.6	-30.0	-30.8	-30.7	-30.7	-30.8	-30.0	-31.3	-29.4
GRAF-1-110	-28.5	-26.6	-28.4	-30.4	-28.9	-29.6	-29.4	-30.1	-30.5	-29.3	-28.3	-29.7	-29.9	-29.6	-29.9	-30.2	-29.6	-30.5	-28.2		

