# CROSS INVOLVING ELEVEN WINTER WHEAT 

## ALIEN-TRANSLOCATION LINES

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Submitted to the Faculty of the Graduate College of the Oklahoma State University in partial fulfillment of the requirements

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            DOCTOR OF PHILOSOPHY
    
        December, 1980
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AN ANALYSIS OF YIELD AND OTHER TRAITS IN A DIALLEL CROSS INVOLVING ELEVEN WINTER WHEAT

ALIEN -TRANSLOCATION LINES

## Thesis Approved:



## ACKNOWLEDGMENTS

Special thanks are extended to Dr. E. L. Smith, major advisor, for his guidance and counsel throughout the course of this study. Grateful acknowledgments are also extended to the other members of my graduate committee, i.e., Dr. C. M. Taliaferro, Dr. B. B. Tucker, Dr. L. M. Verhalen, and Dr. H. C. Young, Jr., for their help and constructive criticism in the preparation of this manuscript.
A special note of thanks is given to Dr. R. W. McNew for his aid in conducting the statistical analyses herein.
The author is also grateful to the Agronomy Department of Oklahoma State University for the use of its facilities and for the financial assistance which made this study possible.

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## CHAPTER I

## INTRODUCTION

Rusts have long been major disease problems in wheat (Triticum aestivum L. em. Thell.) and have caused substantial losses in yield and quality. Many relatives of wheat contain genes which impart resistance to those pests. To incorporate the resistance into wheat, attempts have been made to transfer specific genes from those relatives; and through the aid of irradiation and meiotic control, several translocations between wheat and a segment of alien chromatin carrying the desired gene(s) have been successfully induced in wheat.
Because the transfer of alien genes is not a precise procedure, it is rare that transfer of a single gene takes place. More often, the portion of alien chromatin is sufficiently large to contain genes conditioning other effects as well. Many of those additional effects may be eliminated by successive generations of backcrossing to wheat. In other instances, valuable portions of wheat chromatin may have been lost during the transfer process, also leading to undesirable effects. However, the alien chromatin may compensate, at least in part, for such loss.
At present, a number of alien-translocation lines conditioning resistance to several pests are available for use in wheat breeding programs. Thus the possibility exists for the combination of two or
more alien translocations into the same genotype, thereby imparting multiple-pest resistance.

Due to additional genes in certain combinations, two translocations may result in an $F_{1}$ exhibiting hybrid vigor. On the other hand, if two translocations are united and either or both are missing portions of valuable wheat chromatin, a decline in the vigor of the $F_{1}$ may result. This increase or reduction in vigor may be noted by measuring agronomically and economically important characters.

The objectives of this research were to study the $F_{1}$ 's and parents of a diallel cross involving 11 alien-translocation lines of winter wheat to measure the response of eight performance traits and to determine the potential usefulness of these lines in a breeding program. Standard analyses of variance, analyses of general and specific combining ability, and examination of hybrid vs. parental contrasts were used to attain the above objectives.

## CHAPTER II

## LITERATURE REVIEW

Several relatives of wheat contain genes conditioning desirable features such as disease and insect resistance. Resistance to the rusts, powdery mildew, and several insects may be found in Secale cereale L. and in several of the Aegilops species. Species of Agropyron are noted for resistance to all three wheat rusts, wheat streak mosaic virus, barley yellow dwarf mosaic virus, and bunt (15). The gene pool available to wheat breeders could be greatly enlarged by exploiting the genetic variability of these wheat relatives (23). The incorporation of such desirable characters into wheat would be of great potential value in breeding programs.

Wheat is an allohexaploid composed of the $A, B$, and $D$ genomes each containing seven pairs of chromosomes. Due to its hexaploid nature, unique opportunities exist for the introduction of desirable alien variation. According to Kimber (11), the relationship of the alien chromosome carrying the desired gene(s) to the chromosome(s) of wheat must be the basic consideration in any attempt to introduce such variation into cultivated wheat. If a close relationship exists between the two chromosomes, the alien character may be transferred by normal hybridization. Knott (15) further adds that a diploid species carrying one of the $A, B$, or $D$ genomes can be crossed directly with common wheat or first to a tetraploid and later to a hexaploid. After a wide
cross has been made, the doubled or non-doubled $F_{1}$ can be backcrossed to wheat and eventually "wheat-like" lines containing one or more characters from the alien species can be recovered. The desired character, however, is almost always accompanied by undesirable traits. These undesirable traits are present because alien chromosomes do not ordinarily pair and cross over with wheat chromosomes, so at least one entire chromosome must be added to the wheat complement or substituted for a wheat chromosome in order to retain the desired gene (22).

Besides this normal introgression of alien genes into the wheat genome through hybridization, there are two other means of introducing alien variation, i.e., by irradiation or by meiotic contro1 (15). Sears (19) was the first to describe an irradiation technique used to transfer leaf rust resistance from Triticum umbellulatum (Aegilops umbellulata) to $T$. aestivum. Leaf rust resistant plants carrying an added T. umbellulatum isochromosome were $X$-rayed just prior to meiosis. The $X$-rayed pollen was then used to pollinate emasculated spikes of normal 'Chinese Spring' wheat. Resistant $F_{1}$ plants were examined for the presence of a translocation between the T. umbellulatum and a wheat chromosome. Seventeen translocations were obtained, but only one was transmitted normally through pollen. This translocation was later designated as "Transfer."

Since the pioneering work of Sears, Knott (12) and several others have used similar techniques to successfully transfer alien genes to wheat. The success of Knott's program of transferring stem rust resistance from Agropyron elongatum was attributed to two reasons: first, because wheat is an allohexaploid, it can stand the loss of up to a whole chromosome without seriqus.deleterious effects; and second,
the added Agropyron chromatin had no serious effects except on the gametes. He also found that in families segregating for one of the translocations, resistant and susceptible sibs showed no observable differences. Both X-rays and thermal neutrons were used by Sharma and Knott (29) to translocate a segment of the Agropyron chromosome contained in a wheat-1ike Agropyron derivative, "Agrus," to wheat. One translocation involving chromosome 7D appeared to have no detrimental effects and was normally transmitted through the gametes. This translocation is now known as "Agatha."

Not only have Aegilops and Agropyron species been used as donors of alien material, but Secale (rye) has also been used as a source of resistance. Driscoll and Jensen (7) reported the translocation of a gene for wheat leaf rust resistance from a rye chromosome to wheat. It was later shown that the translocated alien segment present in this line, "Transec," occupied a terminal position and was of considerable length. The wheat chromosome involved was identified as chromosome 4A (5) and was found to involve the loss of genetically unimportant wheat chromatin (6). More recently, Sebesta and Wood (26) have successfully transferred greenbug resistance from rye to wheat through the use of X-rays.

The major advantage of induced translocations through the use of $X$-rays is that the procedure will work regardless of the pairing affinity of the alien chromosome with its wheat homoeologues (15). Knott (14) considered the Agropyrons the easiest to handle and the most valuable source of potentially useful genes. Usually, lines can be produced in which the translocated segment is transmitted as though it were a dominant gene. He further added that the only real problem is
that a segment of wheat chromosome will have been lost, and this could also be deleterious. According to Sears (20), a more desirable approach to this problem would be to produce a reciprocal translocation in which the desired alien chromosome segment replaces a segment of homoeologous wheat chromosome. Because the majority of transfers thus obtained are reciprocal in nature, the alien segment may then, at least partially compensate for the missing wheat segment, as well as introducing the desired alien gene. Sears (23) indicated that the attachment of a terminal segment of the alien chromosome to a nonhomoeologous chromosome would likely result in deficiencies for critical wheat genes in every case.

Because transfers of this type usually involve overall phenotypic modifications or are too arbitrary, alternative procedures have been sought. According to Knott (15), Okamoto, as well as Riley and Chapman, was among the first to report that pairing of homoeologous chromosomes in wheat is genetically controlled. Several methods utilizing homoeologous crossing over to transfer alien genes to wheat chromosomes have since been reported (15).

Riley et al. (18) were the first to report a successful technique using induced homoeologous pairing to transfer yellow rust resistance from Triticum comosum (Aegilops comosa) to wheat. An alien addition line which contained the $T$. comosum chromosome conditioning resistance to the disease was first produced. This was then crossed with $T$. speltoides (Aegilops speltoides) in order to suppress the activity of chromosome 5 Which inhibits homoeologous pairing. The suppression of $5 B$ activity permitted the $T$. comosum chromosome to recombine with its wheat homoeologue. A backcrossing program, combined with selection of
rust-resistant plants, was undertaken to develop a 42-chromosome plant which was also resistant to yellow rust.

Through the use of a deficiency for chromosome 5B, Sears (20) was able to transfer resistance to leaf rust from two different Agropyron elongatum chromosomes to wheat chromosomes, i.e., 3D in "Tap 67" and 7D in "Agrus." For these two chromosomes, the induction of homoeologous pairing appeared to be a superior method for effecting the transfer of the desired gene when compared to those methods utilizing ionizing radiation (21).

This method of transferring desirable characters from the relatives to wheat appears to be more precise than irradiation techniques. Such transfers may have no deleterious effects, especially if the desired alien gene is near the end of one arm, permitting transfer of the gene through exchange of only a short terminal segment (20).

According to Driscoll (3), this method of induced homoeologous pairing does have limitations. The wheat chromosome that is to be involved in the translocation would presumably have to be one of the three in the pertinent homoeologous groups. For irradiation-induced transfers, any chromosome is capable of being involved in any translocation. The position of the alien gene on each of the three wheat chromosomes is also predestined, as crossing over would presumably take place between homoeologous segments of the chromosomes. If such positions were to coincide with wheat genes of significant value, the loss may not be tolerable. Some alien chromosomes with very limited possibilities for crossing over with their wheat homoeologues may never give rise to fully satisfactory transfers through induced pairing. Radiation-induced transfers from such chromosomes would have a better
chance for success (18).
Sears (20) described another technique to transfer part of an alien chromosome to wheat. This transfer would be accomplished by allowing a telocentric for one arm of the alien chromosome to combine with a wheat telocentric following misdivision of two monosomes. This transfer chromosome may be useful if the alien arm is not deleterious and replaces a nonessential wheat arm.

Besides being induced through irradiation or meiotic control, alien translocations occasionally arise spontaneously. Some alien translocations have been produced in this manner, including the "Agent" (32) and "Neuzucht" (36) translocations.

Prior to the work of Sebesta (27) and of Sebesta and Smith (28), the 1 iterature did not contain information regarding the combining of: translocation lines. Knott (13) , however, did test lines which had a pair of Agropyron chromosomes either added to the normal chromosome complement or substituted for wheat chromosome 6A. None of the lines equaled the check cultivar for all characters, but there was also no conclusive evidence for a detrimental effect of the Agropyron chromosome on any character studied. This work did not involve translocation lines containing segments of alien chromatin. Extensive reviews of the literature regarding translocation lines have been published by Knott (15) and Sears (20) in the last 10 years.

If alien-gene transfers are to be useful (regardless of the method used to induce them), they should be relatively free from deleterious effects associated with the exchange of alien chromatin. Driscoll (4) pointed out that an alien translocation may be acceptable in some cultivars but not in others. Thus, alien translocations should be tested
in a wide variety of genetic backgrounds.

Several alien-translocation lines have been used successfully in the development of improved cultivars. 'Arthur $71, '$ grown commercially in the soft red winter wheat region of the U.S., contains the "Transfer" translocation (17). The "Neuzucht" translocation is present in the cultivars 'Aurora' and 'Kavkaz' grown in Eastern Europe (36). 'Sage' and 'Osage,' grown commercially in the hard red winter wheat region of the U.S., contain the "Agent" translocation (31). The "Teewon" translocation is present in the cultivar 'Payne' (30), also grown in the hard red winter wheat region of the U.S. The germplasm lines 'Amigo' and CI 15322 contain the "Amigo" and "Peethree" translocations, respectively (26, 25).

Sebesta (27) and Sebesta and Smith (28) studied the $\mathrm{F}_{1}$ 's and parents of a diallel cross among six alien translocation lines of winter wheat to determine the effects on eight performance traits when two different alien gene complexes were combined two at a time. Significant GCA and SCA variances were detected for all eight characters. The GCA components were larger than those for SCA for tiller number, kernel weight, plant height, and heading date while the reverse was true for kernels/spike, grain yield, percent sterility, and percent protein. Four hybrids showed high positive heterosis for grain yield, while four other hybrids exhibited unexpectedly high levels of sterility, low values for kernels/spike, and low grain yield. The results of their study suggested that an increase in vigor may occur when certain alien translocations are combined; however, other combinations may result in detrimental effects on performance traits. Only by studying
a possible combination is one likely to know whether the translocations
involved are complementary or antagonistic to one another.

CHAPTER III

MATERIALS AND METHODS

## Materials

Eleven winter wheat cultivars or selections were used in this study. Nine of the 11 are known to be alien-translocation lines, and two are presumed to be. The 11 lines are listed in Table $I$ along with the designation and origin of the alien segment (known for nine lines) and the chromosome arms involved (known for six).

Arthur 71 and 'Purdue $6615 D^{\prime}$ ( $6 *$ Arthur/Agatha) are soft red winter wheats developed at Purdue University, Lafayette, Indiana. Arthur 71 is a midtall, apically awnletted cultivar which is moderately early while Purdue 6615 D is an Arthur-type wheat. Arthur 71 and Purdue 6615D contain the "Transfer" and "Agatha"' translocations, respectively. Aurora is a semi-hard red winter wheat developed at Krasnodar, USSR, and contains the "Neuzucht" translocation. 'Winter Transec' contains the "Transec" translocation and is a white winter wheat developed at Cornell University, Ithaca, New York. Osage, Payne, Amigo, and CI 15322 are all hard red winter wheats developed at Oklahoma State University, Stillwater, Oklahoma. Osage is midtall in height, awned, medium late in maturity, and contains the "Agent" translocation. Payne is a medium early maturing, short-strawed, awned cultivar containing the "Teewon" translocation. Amigo and CI 15322 are both germplasm

TABLE I

## ALIEN-TRANSLOCATION LINES STUDIED

| Cultivar or Selection | Translocation Designation | Origin of Alien Gene Complex | $\frac{\text { Arms I }}{\text { Alien }}$ | $\frac{\text { volved }}{\text { Wheat }}$ | Reference |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Osage | Agent | Agropyron elongatum | 3 Ag | 3D | (32) |
| Arthur 71 | Transfer | Triticum umbellulatum | $6 c^{\text {u }}$ | 6B1 | (19) |
| Aurora | Neuzucht | Secale cereale | 1R | 1 Bs | (36) |
| Winter Transec | c Transec | Secale cereale | 2R | 4A | (7) |
| Purdue 6615D | Agatha | Agropyron elongatum | 7 e 1 | 7D1 | (29) |
| Payne | Teewon | Agropyron elongatum | Unknown |  | (24) |
| Salmon | Salmon | Secale cereale | 1R | 1B | (36) |
| Amigo | Amigo | Secale cereale | Unknown |  | (26) |
| CI 15322 | Peethree | Agropyron elongatum | Unknown |  | (25) |
| Flex | --- | ? | Unknown |  | (35) |
| Plainsman V | --- | ? | Unknown |  | (1) |

lines containing the "Amigo" and "Peethree" translocations, respectively. 'Sa1mon' is a white winter wheat obtained from the progeny of a hybrid between two strains of octaploid Triticale, as reported by Tsunewaki (34). Salmon contains the "Salmon" translocation. 'Flex' is a tall, hard red winter wheat with high grain protein developed at South Dakota State University, Brookings, South Dakota. 'Plainsman V,' also a high protein cultivar was developed by Seed Research, Inc., Scott City, Kansas. It is presumed but as yet uncertain that Flex and Plainsman V contain alien translocations. Descriptive information concerning these lines can be found elsewhere (1, 7, 8, 17, 25, 26, 29, $30,31,34,35,36)$.

The 55 different $F_{1}$ 's in this study resulted from a diallel crossing scheme involving the 11 lines listed in Table I (reciprocal crosses were not kept separate). Crosses were made in the greenhouse during the 1977-78 crossing season. Crossed and parental seed were planted in flats on October 6 and 7,1978 . The seedlings were vernalized at outside temperatures for approximately 6 weeks before transplanting.

Field Layout and Nursery Management

The seedlings were transplanted to the field on December 12, 1978. The experiment was conducted in a randomized, complete-block design with four replications. Single-row plots were spaced 30 cm apart, and the 10 plants in each plot were spaced 30 cm apart. The study was conducted during the 1978-1979 growing season at the Agronomy Research Station, Stillwater, Oklahoma, on a Bethany silt loam soil type.

A preplant application of ammonium nitrate was applied on September 5,1978 , at the rate of $30 \mathrm{~kg} / \mathrm{ha}$ actual N . On March 8,1979 , a top dressing of ammonium nitrate was applied at the rate of $40 \mathrm{~kg} / \mathrm{ha}$ actual N. Supplemental water was applied by sprinkler on two occasions. Insecticides were applied twice to control armyworms and greenbugs while weeds were controlled manually.

The study was harvested on June $18-25,1979$, by pulling and bagging individual plants. Eight vigorous, bordered plants from each plot were evaluated.

## Characters Evaluated

The number of tillers/plant, kernels/spike, kernel weight, grain yield, percent spike fertility, plant height, heading date; and percent grain protein were evaluated in this experiment. A11 measurements, with the exception of percent protein and heading date, were taken on an individual plant basis. The measurements were made as follows:

## Tiller Number

Tiller number/plant was recorded as the number of fertile (seedbearing) spikes/plant at the time of threshing and was expressed on a per plant basis.

## Kernels/Spike

The number of kernels/spike was determined by selecting the three best heads from each plant. These were threshed in bulk and the seed were counted and the total divided by three to determine the mean number of kernels/spike.

## Kerne1 Weight

The weight of the seeds of the three selected heads measured to the nearest $1 / 10 \mathrm{~g}$ was divided by the number of seeds produced on the same three heads. This character was expressed as grams per 1000 kernels (per plant).

## Grain Yield

Grain yield was taken as the weight of threshed grain from each individual plant including the three-spike sample and was expressed as grams per plant.

## Percent Fertility

Percent fertility was determined as an average of the same three main heads used to obtain the other yield components and was expressed on a per plant basis. The number of florets/spike was counted before the three heads were threshed. This character was determined as follows: [number of seed per spike $\div$ (number of spikelets per spike X 2$)] \times 100 \%$. Basal and apical spikelets were excluded, as were tertiary and quaternary florets. Therefore, it is possible for fertility percentages to exceed values above $100 \%$ if any of these other florets set seed.

## Plant Height

The measurement of this character was taken as the distance in centimeters from the crown (soil line) to the tip of the tallest spike, excluding awns, and was recorded on a per plant basis.

## Heading Date

This character was recorded as a visual estimation of the date when $50 \%$ of the plants in a row were fully headed. This trait was determined on a per row basis and was expressed as the number of days after March 31.

## Percent Protein

Percent protein of the grain was determined at the wheat quality laboratory at Oklahoma State University using an infrared analyzer. This character was determined on a per row basis.

## Statistical Analyses

Standard analyses of variance were conducted on all data in order to detect the presence or absence of significant differences among the entries for the eight previously mentioned characters. The analyses for percent protein and heading date were conducted on a row basis, the other characters on a plant basis.

Heterosis was determined for all characters with respect to both the mid-parent (MP) and high-parent (HP) values. Least significant differences (LSD) were used to test each hybrid-parent contrast. The standard error (SE) of the difference between treatment means for a hybrid vs. its high-parent value was defined as $\sqrt{\frac{2 \text { EMS }}{r}}$. Adjusted LSD values were used to test the hybrid vs. its mid-parent value because hybrid means were based on only half as many observations as midparent values. The $S E$ for such comparisons was defined as $\sqrt{\frac{3 E M S}{2 r}}$ where EMS is the experimental error mean square and represents the
number of observations comprising the treatment mean.

## Combining Ability Analyses

Combining ability analyses were conducted using $\mathrm{F}_{1}$ and parental data for all eight characters according to the procedure outlined by Griffing (9) as method 2 , model I. Under this model, the genotypes and blocks are regarded as fixed effects. Inferences, therefore, are restricted to the genotypes entering the diallel cross, because the experimental material was not a random sample of a definable larger population. Griffing's analysis provides for the partitioning of the sum of squares due to genotypes into general combining ability (GCA) and specific combining ability (SCA) sums of squares associated with $p-1$ and $[p(p-1)] / 2$ degrees of freedom, respectively, where $p$ represents the number of parents involved in the diallel cross. Under this model, the tests of significance for $G C A$ and $S C A$ are: $M S_{G C A} / M S_{E}$ and $M S_{S C A} / M S_{E}$, respectively where $M S_{G C A}$ and $M S_{S C A}$ are the mean squares associated with GCA and SCA, respectively, and $M S_{E}$ is the error mean square. The variance components for GCA and SCA were calculated according to the procedures outlined by Griffing.

All of the plants of one of the $\mathrm{F}_{1}$ 's died in the seedling state. Therefore, no data could be recorded nor could means or SCA effects be calculated for this hybrid. Because of this, three separate analyses for GCA and SCA were calculated, i.e., all parents and hybrids were included except for the hybrid that died; one of the parents that resulted in the lethal combination was excluded along with all its hybrids; and the other parent and all its hybrids were excluded. Each of the last two methods excluded one parent and 10 hybrids from the
GCA and SCA analyses, respectively, while the first method excluded only one hybrid from the SCA analysis. Correlation coefficients. between the first method and the other two were no lower than 0.98 for any character measured. Therefore, the data reported herein with respect to GCA and SCA are the result of the first method of analysis.

CHAPTER IV

## EXPERIMENTAL RESULTS

## General Considerations

No serious problems with diseases or insects were encountered during the course of this study; and in general, the yields were good. However, all plants of the Osage/Winter Transec hybrid died as seedlings prior to transplanting to the field. Therefore, no data was recorded for that entry nor could means or SCA effects be calculated for it. Several plants of the Osage/Plainsman V hybrid exhibited some degree of stunting and leaf injury at various times during the growing season. The damage to both these hybrids was apparently due to hybrid necrosis, a genetic defect as described by Hermsen (10). In the case of Osage/Winter Transec, the symptoms were severe, resulting in death of the plants. In the case of Osage/Plainsman $V$, the symptoms were less severe, but affecting grain yield and other traits.

After heading, an abnormal amount of spike sterility was observed in several hybrids particularly Arthur 71/Aurora, Arthur 71/Winter Transec, Aurora/Purdue 6615D, Winter Transec/Flex, Aurora/F1ex, Winter Transec/Purdue 6615D, and Purdue 6615D/CI 15322. In these hybrids, however, no indication of necrosis or chlorosis was observed; the plants appeared normal throughout the season.

## Analyses of Variance

Mean squares from analyses of variance for each of the eight traits are presented in Tables II and III. Because heading date and percent protein were measured on a row basis, their mean squares are presented separately (Table III) from the others. In both analyses (Tables II and III), there were only 64 degrees of freedom for entries instead of 65 because all plants of the Osage/Winter Transec hybrid died as seedlings. Highly significant entry (i.e., genotype) mean squares were obtained for all eight characters. Mean squares for replications were highly significant for six of the eight traits studied indicating that blocking of replications was effective for those traits in removing significant amounts of variation from experimental error, thus increasing the precision of the analysis. All characters measured on a plant basis (Table II) exhibited highly significant differences between replication by entry mean squares (experimental error) and residual mean squares (sampling error). This indicates that experimental error for those traits contained variation in addition to that among subsamples.

## Comparisons Among Means

The means for each trait are presented in Table IV. In this and subsequent tables, the following abbreviations were used: Arthur $71=$ Atr71, Aurora = Aura, Winter Transec $=W$.Trans, Purdue $6615 \mathrm{D}=6615 \mathrm{D}$, Salmon $=$ Salm, CI $15322=15322$, and Plainsman $V=P m V$. No abbreviations were necessary for Osage, Payne, Amigo, or Flex. All means, with the exception of those for heading date and percent protein, were based on 32 observations. The means for describing heading date and percent

TABLE II
MEAN SQUARES FOR SIX TRAITS FROM ANALYSES OF VARIANCE OF $\mathrm{F}_{1}$ HYBRIDS AND PARENTS

| Source of Variation | df | Tiller <br> Number | $\begin{gathered} \text { Kernels/ } \\ \text { Spike } \\ \hline \end{gathered}$ | Kernel <br> Weight | Grain <br> Yield | Percent Fertility | Plant <br> Height |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Replication | 3 | 296.6** | 476.1** | 232.4** | $826.7 \div$ * | 782.9 | 2294.8** |
| Entry | 64 | 121.1** | $4133.6 * *$ | 672.1** | 457.1 - | 29908. $5 * *$ | 2096.7 ネぇ |
| Rep X Entry | 192 | 21.7** | 90.5** | 25.3** | 47.9** | 593.5** | 78.6** |
| Residual | 1820 | 9.8 | 27.7 | 6.8 | 17.0 | 197.4 | 36.2 |

*,** Significant at the 0.05 and 0.01 levels of probability, respectively.

## TABLE III

## MEAN SQUARES FOR HEADING DATE AND PERCENT PROTEIN FROM THE ANALYSES OF VARIANCE OF $\mathrm{F}_{1}$ HYBRIDS AND PARENTS

| Source of <br> Variatinn | df | Heading <br> Date | Percent <br> Protein |
| :--- | :---: | :---: | :---: |
| Replication | 3 | 14.6 | $125.7 * *$ |
| Entry | 64 | $599.8 * *$ | $82.0 \% *$ |
| Error | 192 | 13.7 | 4.0 |

[^0]MEANS FOR EIGHT CHARACTERS FROM AN 11-PARENT DIALLEL CROSS OF WINTER WHEAT ALIEN-TRANSLOCATION LINES

| Entry | Tiller Number (No./Plant) | $\begin{aligned} & \hline \text { Kernels/ } \\ & \text { Spike } \\ & \text { (No.) } \\ & \hline \end{aligned}$ | $\begin{gathered} \text { Kernel } \\ \text { Weight } \\ (g / 1000) \\ \hline \end{gathered}$ | Grain Yield (g/Plant) | Percent <br> Fertility | Plant Height (cm) | $\begin{gathered} \text { Heading } \\ \text { Date }{ }^{1} \\ \hline \end{gathered}$ | Percent Protein |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| F1 Hybrids |  |  |  |  |  |  |  |  |
| Osage/Atr 71 | 15.3 | 30.2 | 44.0 | 16.5 | 107.6 | 70.4 | 39.8 | 17.4 |
| /Aura | 13.4 | 51.3 | 37.5 | 20.0 | 145.9 | 79.2 | 39.5 | 15.2 |
| /W.Trans | --- | --- | --- | --- | --- | --- | --- | --- |
| /6615D | 16.8 | 32.6 | 40.8 | 17.2 | 105.2 | 71.4 | 38.0 | 16.0 |
| /Payne | 14.6 | 52.4 | 34.2 | 21.2 | 153.7 | 76.4 | 40.1 | 15.7 |
| /Salm | 12.5 | 41.4 | 41.6 | 17.7 | 130.7 | 85.2 | 45.3 | 18.5 |
| /Amigo | 14.4 | 40.4 | 42.8 | 19.6 | 130.8 | 76.0 | 38.5 | 15.3 |
| /15322 | 17.2 | 34.2 | 36.7 | 17.8 | 108.3 | 79.5 | 41.0 | 17.9 |
| /Flex | 13.8 | 39.3 | 37.1 | 17.2 | 126.9 | 76.8 | 45.5 | 17.9 |
| $/ \mathrm{PmV}$ | 10.0 | 32.3 | 30.3 | 8.3 | 121.7 | 64.9 | -38.3 | 16.2 |
| Atr $71 /$ Aura | 11.8 | 14.1 | 45.2 | 6.5 | 41.6 | 72.0 | 37.2 | 21.2 |
| /W.Trans | 14.2 | 13.9 | 36.6 | 6.3 | 41.3 | 76.3 | 41.5 | 20.2 |
| /6615D | 11.2 | 29.1 | 38.7 | 10.8 | 100.4 | 64.0 | 37.3 | 17.1 |
| /Payne | 14.3 | 26.7 | 43.1 | 12.9 | 89.0 | 66.9 | 39.0 | 17.6 |
| /Salm | 13.8 | 22.8 | 45.0 | 12.4 | 70.1 | 78.0 | 40.0 | 19.6 |
| /Amigo | 13.9 | 26.2 | 45.2 | 13.9 | --. 87.7 | 70.8 | 37.0 | 17.3 |
| /15322 | 15.1 | 23.9 | 41.6 | 12.5 | 83.2 | 73.7 | 38.8 | 16.9 |
| /Flex | 14.9 | 29.6 | 38.3 | 13.8 | 101.6 | 71.7 | 39.8 | 16.8 |
| /Prov | 13.9 | 25.8 | 40.2 | 10.1 | 92.2 | 67.9 | 36.0 | 18.9 |
| Aura/W.Trans | 11.1 | 55.1 | 39.9 | 20.1 | 140.6 | 81.4 | 41.0 | 15.2 |
| /6615D | 10.4 | 12.4 | 45.2 | 4.9 | 36.9 | 66.0 | 37.8 | 20.9 |
| /Payne | 10.0 | 56.0 | 36.6 | 16.6 | 153.8 | 71.7 | 37.3 | 15.9 |
| /Salm | 10.1 | 45.6 | 43.6 | 17.7 | 134.2 | 80.8 | 40.0 | 18.0 |
| /Amigo | 10.3 | 47.5 | 44.0 | 17.1 | 132.6 | 77.3 | 36.5 | 15.8 |
| /15322 | 11.6 | 43.6 | 41.7 | 16.7 | 122.9 | 82.8 | 38.0 | 16.5 |
| /Flex | 13.6 | 21.3 | 45.2 | 11.6 | 58.3 | 76.8 | 39.8 | 20.8 |
| /PmV | 10.4 | 49.8 | 39.9 | 14.6 | 147.2 | 70.7 | 36.8 | 16.1 |
| W. Trans/6615D | 11.8 | 21.6 | 40.4 | 8.5 | 68.8 | - 73.6 | 39.5 | 18.6 |
| /Payne | 12.9 | 60.1 | 33.8 | 22.1 | 164.1 | 83.2 | 43.8 | 15.0 |
| /Salm | 12.5 | 34.0 | 38.4 | 13.5 | 88.7 | 94.2 | 49.0 | 19.6 |
| /Amigo | 12.5 | 34.5 | 39.0 | 14.9 | 104.4 | 80.3 | 43.0 | 16.2 |
| / 15322 | 11.6 | 44.4 | 33.8 | 14.6 | 126.2 | ... | 49.3 | 17.7 |
| /Flex | 14.4 | 22.2 | 35.1 | 9.1 | 58.7 | - 88.8 | 50.3 | 20.1 |
| /PmV | 14.2 | 49.8 | 29.5 | 16.4 | 152.4 | 75.3 | 38.3 | 16.3 |

TABLE IV (Continued)

| Encry | $\begin{gathered} \text { Tiller } \\ \text { Number } \\ \text { Go.'Plant } \\ \hline \end{gathered}$ | $\begin{gathered} \text { Kernels/ } \\ \text { Spike } \\ \text { (.3o.) } \\ \hline \end{gathered}$ | $\begin{gathered} \text { Kernel } \\ \text { keight } \\ (\mathrm{g} / 1000) \\ \hline \end{gathered}$ | $\begin{gathered} \text { Grain } \\ \text { Yield } \\ \text { (g/Plant) } \\ \hline \end{gathered}$ | Percent Fertility | Plant Height (cm) | Heading Date ${ }^{1}$ | Percent <br> Protein |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 661 晹 Payne | 15.2 | 29.2 | 42.8 | 15.7 | 88.8 | 70.7 | 38.0 | 18.2 |
| Salm | 14.8 | 28.0 | 48.5 | 17.4 | 83.7 | 80.0 | 38.3 | 19.1 |
| 'sizigo | 13.7 | 29.0 | 41.0 | 13.6 | 94.1 | 67.4 | 37.5 | 17.8 |
| -15322 | 15.0 | 20.2 | 42.6 | 10.1 | 66.5 | 74.0 | 38.0 | 19.6 |
| ;Elex | 15.7 | 34.4 | 37.9 | 15.0 | 109.8 | 73.8 | 40.3 | 17.1 |
| /Pmot | 12.9 | 24.4 | 40.0 | 9.2 | 83.4 | 66.5 | 32.8 | 19.9 |
| पayne sale | 12.8 | 51.1 | 38.9 | 19.6 | 149.6 | 80.6 | 42.0 | 16.4 |
| - Amiz | 12.5 | 42.6 | 39.0 | 15.0 | 137.6 | 68.9 | 37.8 | 16.6 |
| 15322 | 13.8 | 39.4 | 35.0 | 16.1 | :32.1 | 74.2 | 40.3 | 16.6 |
| Elex | 14.9 | 34.3 | 37.7 | 16.3 | 109.0 | 71.4 | 39.5 | 18.5 |
| Priv | 12.5 | 38.4 | 35.1 | 14.0 | 127.5 | 63.1 | 36.0 | 17.3 |
| Salmamigo | 13.4 | 40.1 | 43.1 | 19.1 | 126.9 | 80.0 | 38.8 | 16.1 |
| '15322 | $1 \mathrm{1i} .3$ | 35.4 | 35.8 | 12.4 | 107.8 | 90.0 | 46.5 | 19.1 |
| Fiex | 12.3 | 30.3 | 41.2 | 13.5 | 87.1 | 86.7 | 48.8 | 20.6 |
| /Pmil | 11.8 | 47.6 | 34.3 | 15.3 | 153.6 | 72.6 | 37.8 | 16.6 |
| Amizoil5322 | 13.2 | 33.2 | 38.3 | 14.3 | 113.2 | 76.0 | 38.5 | 17.8 |
| flex | 15.0 | 27.6 | 42.1 | 14.5 | 90.7 | 72.1 | 39.3 | 19.1 |
| / Pai | 13.3 | 33.6 | 40.1 | 14.8 | 116.2 | 65.4 | 34.8 | 17.7 |
| 15322/Flex | 15.2 | 33.4 | 37.9 | 16.2 | 103.9 | 83.3 | 43.3 | 19.5 |
| Pmav | 13.3 | 36.8 | 33.1 | 13.4 | 130.6 | 69.5 | 37.0 | 17.4 |
| clexipev | 15.3 | 27.4 | 30.8 | 12.9 | 95.1 | 57.0 | 37.0 | 20.7 |
| Parencs |  |  |  |  |  |  |  |  |
| Osage | 12.3 | 42.1 | 36.1 | 15.0 | 142.6 | 73.0 | 45.5 | 16.3 |
| Atr71 | 10.8 | 24.1 | 37.6 | 8.2 | 96.9 | 61.1 | 38.0 | 17.7 |
| Aura | 7.9 | 52.3 | 41.2 | 13.8 | 131.4 | 69.7 | 40.3 | 17.5 |
| k. Trans | 8.5 | 60.2 | 27.0 | 11.4 | 154.8 | 93.2 | 53.8 | 16.9 |
| 66150 | 11.6 | 32.5 | 34.4 | 9.8 | 111.7 | 62.7 | 37.0 | 16.5 |
| Payne | 11.3 | 51.2 | 32.8 | 16.5 | 160.8 | 69.5 | 43.3 | 26.5 |
| Salm | 9.6 | 47.8 | 35.9 | 13.7 | 135.7 | 81.6 | 49.0 | 18.6 |
| Amigo | 11.0 | 37.1 | 38.9 | 13.2 | 131.5 | 67.1 | 39.0 | 15.7 |
| 15322 | 11.2 | 32.7 | 28.9 | 9.0 | 116.7 | 73.0 | 40.3 | 17.1 |
| flex | 12.4 | 39.3 | 30.8 | 13.2 | 124.3 | 77.2 | 49.5 | 18.8 |
| Pmi | 10.8 | 34.2 | 28.7 | 8.3 | 132.3 | 56.4 | 32.2 | 18.3 |
| $\mathrm{F}_{1}$ Mean ( $\bar{F}_{1}$ ) | 13.3 | 34.9 | 39.4 | 14.5 | 108.0 | 75.4 | - 39\%\% | 17.8 |
| Parental Mean ( $\overline{\mathrm{P}}$ ) | 10.7 | 41.2 | 33.9 | 12.0 | 130.8 | 71.3 | 42.5 | 17.3 |
| $\bar{F}_{1}{ }^{\prime} \mathrm{P}$, \% | 124.3 | 84.7 | 116.2 | 120.8 | 82.6 | 105.8 | 93.9 | 102.9 |
| Experimental Mean | 12.8 | 35.9 | 38.5 | 14.1 | 111.9 | 74.7 | 40.3 | 17.7 |
| LsD ¢0.c5] | 2.3 | 4.7 | 2.5 | 3.4 | 11.9 | 4.4 | 1.8 | 1.0 |
| c. ¢ $^{\text {d }}$ | 36.3 | 26.4 | 13.1 | 49.1 | 21.7 | 11.9 | 9.2 | 11.2 |

${ }^{1}$ Number of days after Marcl: 31.
protein were based on four observations (one observation/entry/ replication). The performance of the hybrids expressed as a percent relative to their respective mid-parent and high-parent values are shown in Table V. Parental and hybrid means, ranked in order, along with hybrid-parent deviations may be found in the Appendix (Tables XIVXXI) for each character separately.

As shown in Table IV, the means for tiller number ranged from 7.9 for Aurora to 17.2 for Osage/CI 15322. The parent with the highest number of tillers was Flex with 12.4 . The overall means for $F_{1}$ 's and parents were 13.3 and 10.7 , respectively, with the $F_{1}$ 's displaying $24.3 \%$ more tillers overall than their parents. Thirty-seven of the 54 $F_{1}$ 's studied exhibited significant positive mid-parent heterosis while $27 \mathrm{~F}_{1}$ 's had significantly more tillers than their respective highparents. One $F_{1}$, Osage/Plainsman $V$, had significantly fewer tillers than its high-parent (Table V).

The number of kernels/spike ranged from 60.2 for Winter Transec to 12.4 for Aurora/Purdue 6615D (Table IV). Winter Transec/Payne, with a value of 60.1 , was the $F_{1}$ with the highest number of kernels. Arthur 71 was the lowest parent with a mean of 24.1 kernels/spike. The overall $\mathrm{F}_{1}$ mean was 34.9 while the overall parental mean was 41.2 . The mean of all $F_{1}$ 's expressed as a percentage of the overall parental mean was $84.7 \%$. As shown in Table $V$, five hybrids were significantly higher in kernels/spike than their mid-parent while none was significantly higher than its respective high-parent. Twenty-six hybrids were significantly lower than their mid-parents while 41 were significantly 1ower than their high-parents.

TABLE $V$

MEAN PERFORMANCE OF HYBRIDS EXPRESSED AS A PERCENT OF MID－PARENT（MP）AND HIGH－PARENT（HP）VALUES

| Fi Hubrids | Tiller Number |  | KernelsiSpike |  | Kernel Weight |  | Grain Yield |  | Percent Fertility |  | Plant Height |  | Heading Date |  | Percent Protein |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 3 MP | ${ }^{3} \mathrm{HP}$ | 2 YP | ${ }_{2} \mathrm{H}^{2}$ | YMP | IHP | TMP | 2HP | ${ }_{3} 78$ | 3 HP | ZMP | \％ H P | Z．MP | \％HP | 2MP | 7 HP |
| Osage／atr 71 | 132＊＊ | 124＊＊ | 91 | 72＊＊ | 119＊＊ | 117＊＊ | 142＊＊＊ | 110 | 90＊ | 75＊＊ | 105 | 96 | 95＊ | 87＊＊ | 102 | 98 |
| ＇Aura | 133＊＊ | 109＊＊ | 109＊ | 98 | 97 | 91＊＊ | 139＊＊ | 133＊＊ | 106 | 102 | 111 | 108 | 92＊＊ | 87＊＊ | 90＊＊ | 87＊＊ |
| ik．Trans | －－－ | －－－ | －－ | －－－ | －－－ | －－－ | －－－ | －－－ | －－－ | －－－ | －－－ | －－－ | －－－ | －－－ | －－－ | －－－ |
| ／6615D | 141＊＊ | 137＊＊ | 87＊ | 77＊＊ | 116＊＊ | 113＊＊ | 139＊＊ | 115 | 83＊＊ | 74＊＊ | 105 | 98 | 92＊＊ | 84＊＊ | 98 | 97 |
| ／Payne | 124＊＊ | 119＊ | 112＊＊ | 102 | 99 | 95 | 135＊＊ | 128＊＊ | 89＊＊ | 84＊＊ | 107＊＊ | 105 | 90＊＊ | 88＊＊ | 96 | 95 |
| ／Salm | 114 | 102 | 92 | 87＊＊ | 116＊＊ | 115＊＊ | 123＊ | 118 | 94 | 92＊ | 110 | 104 | 96＊＊ | 92＊＊ | 106＊＊ | 99＊＊ |
| Amigo | 12 | 117 | 102 | 96 | 114＊＊ | 110＊＊ | 139＊＊ | ．131＊＊ | 95 | 92 | 108＊＊ | 104 | 91＊＊ | 85＊＊ | 96 | 94 |
| ／15322 | 146＊＊＊ | $140 \div$ | 91 | 81＊＊ | 113＊＊ | 102 | 148＊＊ | 119 | 84 | 76＊＊ | 109 | 109 | 96＊ | 90＊＊ | 107＊＊ | 105＊＊ |
| frlex | 112 | 111 | 97 | 93 | 111＊＊ | 103 | 122＊ | 115 | 95 | 89＊＊ | 102 | 99 | 96＊ | 92＊＊ | 102＊＊ | 95＊＊ |
| Pmi | 87 | 81＊ | 85＊＊ | 77ヶ＊ | 94＊ | 84＊＊ | 71＊ | 55＊＊ | 89＊＊ | 85\％＊ | 100 | 89＊＊ | 99 | 84＊＊ | 94 | 89＊＊ |
| Atril／Aura | 126＊＊ | 109＊ | 37＊＊ | 27＊＊ | 115＊＊ | 110\％＊ | 59＊＊ | 47＊＊ | 36＊＊ | 32＊＊ | 110＊＊ | 103 | 95＊＊ | 92＊＊ | 120＊＊ | 120＊＊ |
| ／h．Trans | 146＊＊ | 131＊＊ | 33＊＊ | 23＊＊ | 113＊＊ | 97 | 64＊ | 55＊＊ | 33＊＊ | 27＊＊ | 99 | 82＊＊ | 90＊＊ | 77＊＊ | 117＊＊ | 114＊＊ |
| ／0615D | 100 | 97 | 103 | 90 | 107＊ | 103 | 120 | 110 | 96 | 90 | 103 | 102 | 99 | 98 | 100 | 97 |
| ／Payne | 129シ＊ | 127＊＊ | 71＊＊ | 52＊＊ | 122＊＊ | 115＊＊ | 104 | 78＊ | 69＊＊ | 55＊＊ | 102 | 96 | 96 | 90＊＊ | 103 | 99 |
| ／Salr | 135＊＊ | 128＊＊ | 63＊＊ | 48＊＊ | 122＊＊ | 120＊＊ | 113 | 91 | 60＊＊ | 52＊＊ | 109＊＊ | 96 | 92＊＊ | 82＊＊ | 108＊＊ | 105＊＊ |
| ／Amigo | 128＊＊ | 126＊ | 86＊ | 71＊＊ | 118＊＊ | 116＊＊ | 130＊ | 105 | 77＊＊ | 67＊＊ | 110＊： | 106 | 96 | 95＊ | 104 | 98 |
| 115322 | 137＊＊ | 135＊＊ | 84＊＊ | 73＊＊ | 125＊＊ | 111＊＊ | 145＊＊ | 139＊＊ | 78＊＊ | 71ヶ＊ | 110＊＊ | 101 | 99 | 96 | 97 | \％＊95 |
| ／Flex | 128＊＊ | $120 *$ | 93 | 75＊＊ | 112＊＊ | 102 | 129＊ | 105 | 92 | 82＊＊ | 104 | 93＊ | 91＊＊ | 80＊＊ | 92＊＊ | 89＊＊ |
| $/ \mathrm{PmV}$ | 129＊＊： | 129＊＊ | 89. | 75＊＊ | 121＊＊ | 107＊ | 122 | 122 | 80＊＊ | 70＊＊ | 116＊＊ | 120＊＊ | 103 | 95＊ | 105＊ | 103 |
| Aura／w．Trans | 135＊＊ | 129＊ | 98 | 92＊ | 117\％＊ | 97 | 160＊＊ | 146＊＊ | 98 | 91＊＊ | 100 | 87＊＊ | 87＊＊ | 76＊＊ | 88＊＊ | 87＊＊ |
| ／6615D | 107 | 90 | 29\％＊＊ | 24＊＊ | 120＊＊ | 110＊＊ | 42\％＊ | 36＊＊ | 30\％＊＊ | 28＊＊ | 100 | 95 | 98 | 94＊＊ | 123＊＊ | 119＊＊ |
| i＇Payne | 104 | 88 | 108 | 107 | 99 | 89＊＊ | 110 | 101 | 105 | 96 | 103 | 103 | 89＊＊ | 86＊＊ | 94＊＊ | 91＊＊ |
| ／Salm | 115 | 105 | 91＊ | 87＊＊ | 113＊＊ | 106 | 129＊＊ | 128＊ | 100 | 99 | 107＊＊ | 99 | 90＊＊ | 82＊＊ | 100 | 97 |
| ／Amigo | 109 | 94 | 106 | $91 *$ | 110～\％ | 107＊ | 127＊ | 124 | 101 | 101 | 113＊＊ | 111＊＊ | 92＊＊ | 91＊＊ | 95＊ | 90＊＊ |
| ／15322 | 121＊＊ | 104 | 103 | 83＊＊ | 119＊＊ | 101 | 146＊＊＊ | 121 | 99 | 94 | 116＊＊ | 113＊＊ | 94＊ | 94＊ | 95＊ | 94＊ |
| ／Flex | 134＊＊ | 110 | 47＊＊ | 41＊＊ | 126＊＊ | 110＊＊ | 86 | 84 | 46＊＊ | 44＊＊ | 105 | 99 | 89＊＊ | 80＊＊ | 115＊＊ | 111＊＊ |
| ／ Pnv | 111 | 96 | 115\％\％ | 95 | 114＊＊＊ | 97 | 132＊ | 106 | 112＊＊ | 111＊ | 112＊＊ | 101 | 102 | 91＊＊ | 90＊＊ | 88＊＊ |

TABLE V（Continued）

| E．Avbrids | Tiller Sumber |  | KernelsiSpike |  | Karnel meight |  | Grain Yield |  | Percent Fertility |  | Plant Height |  | Heading Date |  | Percent Protein |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 2？ | \％ B ？ | $\square$ | SHP | UP | H？ | MP | SHP | \％ MP | 沮 | ZMP | 沮 | S／4P | \％HP | PMP | 3 HP |
| A．Erans 66150 | 127 | 102 | $47 \%$ | 36＊＊ | 132＊＊ | 117＊＊ | 80 | 75 | j2＊＊ | 44＊＊ | 94＊ | 79＊＊ | 87＊＊ | 73＊＊ | 111＊＊ | 110＊＊ |
| ＇Payne | ：30\％ | 114 | 108＊ | 100 | 113＊＊ | 103 | 158＊＊ | 134＊＊ | 104 | 102 | 102 | 89＊＊ | 90＊＊ | 81＊＊ | 90＊＊ | 89＊＊ |
| ＇Sale | 137＊＊ | 130＊ | 63＊＊ | 56＊＊ | 122＊＊ | 107＊ | 108 | 99 | 61\％ | 57＊＊ | 108＊＊＊ | 101 | 95\％ | 91＊＊ | 110＊＊ | 105＊ |
| Amigo | 128＊＊ | 114 | 71＊＊ | 57ニ＊ | 118\％＊ | 100 | 121 | 113 | 73＊＊ | 67＊＊ | 100 | 86＊＊ | 93＊＊ | $80 * *$ | 99＊＊＊ | 103＊＊ |
| ／15322 | 117 | 104 | 96 | 74＊＊ | 121＊＊ | 117＊＊ | 143＊＊ | 128 | 93 | 82＊＊ | 116＊＊ | 104 | 105＊ | 92＊＊ | 104＊＊ | 104＊＊ |
| flex | 137\＃＊ | 116 | 45＊＊ | 37＊ | 121＊＊ | 114＊＊ | 74＊ | 69＊ | 42＊ | 38＊＊ | 104 | 95＊ | 97 | 93＊＊ | 113＊＊ | 107＊＊ |
| ／PmV | 146＊＊＊ | 131＊＊ | 106 | 83＊＊ | 106 | 103 | 166＊＊ | 144＊＊ | 106 | 98 | 101 | 81＊＊． | 89＊＊ | 71＊ | 93＊＊ | 89＊＊ |
| 66：50＇Payne | 133＊： | 131＊＊ | 70＊＊ | 57\％ | 127＊＊ | 124＊＊ | 119 | 95 | 65＊＊ | 55＊＊ | 107＊ | 102 | 95＊ | 88 ＊＊ | 110＊＊ | 110＊＊ |
| ／Salm | 140＊＊ | 128＊＊ | 70＊＊ | 59＊＊ | 138＊＊ | 135＊＊ | 148＊＊ | 127＊ | 68＊： | 62＊＊ | 111＊＊ | 98 | 89＊＊ | 78＊＊ | 109＊： | 103 |
| －Amigo | 121＊ | 118 | 83＊＊ | 78＊＊ | 112＊＊ | 105 | 118 | 103 | 77＊＊ | 72＊＊ | 104 | 100 | 99 | 96 | 111 | 108＊＊ |
| ：15322 | 132＊＊： | 129＊＊ | 62＊＊ | 62＊＊＊ | 135＊＊ | 124＊＊ | 107 | 103 | 58＊＊ | 57＊＊ | 109＊＊ | 101 | 98 | 94＊ | 117＊＊， | 115＊＊ |
| Flex | 131＊＊ | 127＊＊ | 96 | 88＊ | 116＊＊ | 110＊＊ | 130\％ | 114 | 93 | 88＊ | 106＊ | 96 | 93＊＊ | 81＊＊ | 97 | 91＊＊ |
| Prov | 115 | 111 | 73＊＊ | 71＊＊ | 127＊＊ | 116＊＊ | 102 | 94 | 68＊＊ | 63＊＊ | 112＊＊ | 106 | 95＊ | 102＊＊ | 114＊＊． | 109＊＊ |
| Payneisalm | 122＊ | 113 | 103 | 100 | 113＊＊ | 108＊． | 130＊＊ | 119 | 101 | 93 | 107＊＊ | 99 | 91＊＊ | 86＊＊ | 93＊＊＊ | 88＊＊ |
| －Amigo | 113 | 112 | 96 | 83＊＊ | 109＊＊ | 100 | 121＊＊ | 109 | 94 | 86＊＊ | 101 | 99 | 92＊＊ | 97＊＊ | 103 | 101 |
| ／15322 | 123＊＊＊ | 122＊＊ | 94 | $77 \%$ | 113＊＊ | 107 | 126＊ | 98 | 95 | 82＊＊＊ | 104 | 102 | 96 | 93＊ぇ | 99 | 97 |
| ／flex | 126＊＊ | 120＊＊ | 75＊＊ | 67＊＊ | 119＊＊＊ | 115＊＊ | 110 | 99 | $76 * *$ | 68＊＊ | 97 | 92＊＊ | 85＊＊ | $80 \div$＊ | 105＊ | 98 |
| ／ mav | 113 | 111 | 90＊ | 75＊＊ | 114＊＊ | 107 | 113 | 85 | 87＊＊ | 79＊＊ | 100 | 91＊＊ | 95＊ | 83＊＊ | 99 | 95＊ |
| Salm／Amigo | 130＊＊ | 122＊ | 94 | 84\％＊ | 115\％＊ | 111＊＊ | 142＊＊ | 139＊＊ | 95 | 94 | 108＊＊ | 98 | 88＊＊ | 79：＊ | 94＊＊＊ | 87＊＊ |
| ／15322 | 113 | 105 | 88＊ | 74＊＊ | 110＊＊ | 100 | 109 | 91 | 85＊＊ | 79ッ＊ | 116＊＊ | 110＊＊ | 104＊ | $95 \div *$ | 107＊＊ | 103 |
| ／Flex | 112 | 99 | 70＊＊ | 63＊＊ | 124＊＊＊ | 115＊＊ | 100 | 99 | 67ヶ＊ | 64＊＊ | 109＊＊ | 106＊ | 99 | 99 | 110＊＊ | 110＊＊ |
| $/ \mathrm{Pay}$ | 116 | 109 | 116＊＊ | 100 | 106 | 96 | 139＊＊ | 112 | 115\％： | 113＊＊ | 105 | 89＊＊ | 93＊＊ | 77＊＊ | 90＊＊ | 89＊＊ |
| Amigo／15322 | 119＊＊ | 118 | 95 | 89 | 113＊＊ | 98 | 129＊＊ | 108 | 91＊ | 86＊＊ | 108＊＊ | 104 | 97 | 96＊ | 109＊＊ | 104 |
| ／Flex | 128\％＊ | 121＊ | 72＊＊＊ | 70＊＊ | 121＊＊ | 108＊＊ | 110 | 110 | 71＊＊ | 69＊＊＊＊＊＊＊＊＊ | 100 | 93＊ | $98 * *$ | 79＊＊ | 111＊＊＊＊＊＊＊＊＊ | 102 |
| 1 PmV | 122＊ | 121＊ | 94 | 91 | 119\％ | 103 | 138＊＊ | 112 | 88＊＊ | $88 * *$ | 106 | 97 | 98 | 89＊＊ | 104 | 97 |
| 15322／Flex | 129＊＊ | 123＊ | 93 | 85＊ | 127＊＊ | 123＊＊ | 146＊＊ | 123 | 86＊＊ | 84＊＊ | 111＊＊ | 108＊＊ | 96 | 87＊＊ | 109＊＊ | 104 |
| ／Puv | 121＊ | 119 | 110 | 108 | 115＊＊ | 115＊＊ | 155\％＊ | 149＊＊ | 105 | 99 | 107＊ | 95 | 102 | 92＊＊ | 98 | 95 |
| Flex／PmV | 132\％＊ | 123＊＊ | 75＊＊ | 70\％ | 128「＊ | 124＊＊ | 120 | 98 | 74＊＊ | 72＊＊ | 100 | 87＊＊ | 91 | 75＊＊ | 112＊＊ | 110＊＊ |

＊，$=*$ Significantly different from the mid－parent（or nigh－parent）at the 0.05 and 0.01 levels of probability，respectively．

The overall $\mathrm{F}_{1}$ mean for kernel weight was $39.4 \mathrm{~g} / 1000$ seeds and the overall parental mean was $33.9 \mathrm{~g} / 1000$ (Table IV). On the average, $F_{1}$ 's had $16.2 \%$ higher kernel weight than their parents. Means for kernel weight ranged from 48.5 g for Purdue $6615 \mathrm{D} / \mathrm{Salmon}$ to 27.0 g for Winter Transec. Aurora was the highest parent with a kernel weight of 41.2 g . Forth-eight hybrids exceeded their mid-parent by significant margins while 30 hybrids were significantly greater than their respective high-parents (Table V). Three hybrids had significantly lower kernel weight than their high-parents, and one hybrid was significantly lower than its mid-parent.

Winter Transec/Payne was the highest yielding entry with a mean of $22.1 \mathrm{~g} / \mathrm{plant}$ while Aurora/Purdue 6615D was the lowest with 4.9 g (Table IV). The overall parental mean was 12.0 g compared to the overall $\mathrm{F}_{\mathrm{i}}$ mean of 14.5 g , i.e., a $20.8 \%$ average yield advantage of $\mathrm{F}_{1}$ 's over their parents. Thirty hybrids yielded significantly greater than their respective mid-parents while 11 hybrids yie1ded significantly more than their high-parents. Five and six hybrids yielded significantly less than their mid-parents and high-parents, respectively (Table V).

Percent fertility ranged from $164.1 \%$ for Winter Transec/Payne to 36.9\% for Aurora/Purdue 6615D (Table IV). The parent with the highest percent fertility was Payne with $160.8 \%$ while the lowest parent was Arthur 71 with $96.9 \%$. Eight hybrids displayed less than $80 \%$ spike fertility. Those hybrids were Arthur 71/Salmon, Winter Transec/ Purdue 6615D, Purdue 6615D/CI 15322, Winter Transec/Flex, Aurora/F1ex, Arthur 71/Aurora, Arthur 71/Winter Transec, and Aurora/Purdue 6615D. The possible reasons for their depressed fertility and its implications
in breeding programs will be considered in the following chapter. The overall means for fertility were $108.0 \%$ and $130.8 \%$ for $F_{1}{ }^{\prime} s$ and parents, respectively, which is a $17.4 \%$ decline in fertility in the $F_{1}{ }^{\prime}$ s overall. Forty hybrids had significantly lower values for spike fertility than their respective high-parents and 31 were significantly lower than their mid-parent values for this trait. Only two hybrids, Salmon/ Plainsman V and Aurora/Plainsman V exhibited significantly greater fertility values than either their mid-parent or high-parent values (Table V).

The highest parental mean for plant height was Winter Transec with 93.2 cm (Table IV) while Winter Transec/CI 15322 was the tallest hybrid with a mean plant height of 96.6 cm . Plainsman $V$ was the entry with the shortest stature having a mean of 56.4 cm . The shortest hybrid was Payne/Plainsman $V$ with a mean of 63.1 cm . As shown in Table $V, 14$ hybrids were significantly shorter than their tallest parent while only one hybrid, Winter Transec/Purdue 6615 D was significantly shorter than its mid-parent. On the other hand, 25 were significantly taller than their mid-parents; and six were significantly taller than their tallest parent.

Means for heading date ranged from the earliest value of 32.2 days for Plainsman $V$ to the latest value of 53.8 days for Winter Transec. The earliest hybrid was Purdue $6615 \mathrm{D} / \mathrm{Plainsman} V$ with a value of 32.8 days while the latest hybrid was Winter Transec/Flex with a value of 50.3 days (Table IV). The $\mathrm{F}_{1}$ 's averaged $6.1 \%$ earlier than the parents with means of 39.9 and 42.5 days for the $F_{1}$ 's and parents, respectively, one hybrid was later than its latest parent (high-parent), and two hybrids were significantly later than their respective mid-parents
(Table V). Thirty-four of 55 hybrids were significantly earlier than their mid-parent values while 49 of 55 were significantly earlier than their respective later parent.

Percent protein means ranged from $21.3 \%$ for Arthur 71/Aurora to $15.0 \%$ for Winter Transec/Payne. The highest parent was Flex with $18.8 \%$ and the lowest was Amigo with $15.7 \%$ (Table IV). The overall means for $\mathrm{F}_{1}$ 's and parents were 17.8 and $17.3 \%$, respectively, which results in a $2.9 \%$ advantage for the $F_{1}$ 's. Twenty-four hybrids were significantly higher than their mid-parents while 17 hybrids were significantly higher than their high-parents. Thirteen hybrids were significantly lower than their mid-parents for percent protein while 17 were significantly lower than their high-parents (Table V).

## Diallel Analyses for General and Specific Combining Ability

All available data were subjected to diallel analyses (9) for general combining ability (GCA) and specific combining ability (SCA) for each of the eight characters. As stated previously, all the plants of the Osage/Winter Transec hybrid died as seedlings; therefore, no SCA effects could be calculated for that hybrid. The mean squares from the combining ability analyses of variance are presented in Table VI. Highly significant GCA and SCA variances were observed for all characters.

The variance components for GCA (associated with additive genetic effects) and SCA (nonadditive effects) were calculated to obtain estimates of additive and nonadditive effects. Although estimates of variance components for GCA and SCA are valid in the strict sense only

## TABLE VI

MEAN SQUARES FOR GENERAL COMBINING ABILITY, SPECIFIC COMBINING ABILITY, AND ERROR FOR EIGHT CHARACTERS

| Trait | GCA ${ }^{1}$ | SCA | Error |
| :---: | :---: | :---: | :---: |
| Tiller Number | $41.47 \times *$ | $10.25 * *$ | 2.70 |
| Kernels/Spike | 1795.95** | $265.30 \% *$ | 11.31 |
| Kernel Weight | 299.90\% | 44.02** | 3.16 |
| Grain Yield | 173.33** | 35.62\% | 5.93 |
| Percent Fertility | 11140.11** | $2367.88 * *$ | 74.19 |
| Plant Height | $1421.63 * *$ | 47.36** | 9.78 |
| Heading Date | 350.38\% | 15.43\% | 1.71 |
| Percent Protein | 25.98** | 7.34** | 0.94 |
| *,** Significant at the 0.05 and 0.01 levels of probability, respectively. |  |  |  |
| $1_{\text {The degrees of }} \mathrm{fr}$ 54 , and 192 , resp | GCA, SCA, | mean sq | 10, |

with random models, estimates of those components were made in this study even though the genotypes included herein were considered fixed rather than random. This was done to provide information regarding the relative magnitude of the components while recognizing at the same time that restrictions are placed on the interpretations of those estimates. The variance components, along with the GCA/SCA ratios for each trait, are presented in Table VII. The GCA components were 2.82 and 4.17 times greater than the SCA components for heading date and plant height, respectively, indicating the importance of additive genetic effects for those two traits. On the other hand, the ratios of GCA to SCA were less than unity for tiller number, kernels/spike, kernel weight, grain yield, percent fertility, and percent protein which indicates that those six traits are influenced to a greater degree by dominance, epistasis, or both.

## GCA Effects

Because the GCA mean squares were significant for all traits (Table VI), GCA effects for the individual parents could be estimated for each of the eight characters studied. Those estimates and their standard errors are shown in Table VIII. The GCA effects, as expected, agreed roughly in rank with the parental means for all characters (comparisons between Tables IV and VIII). This suggests that selection of parents for use in a plant breeding program to improve these traits may be based largely on phenotypic performance of the parents themselves and also suggests the presence of considerable additive effects. The parents Osage and Payne displayed significant positive GCA effects for grain yield, kernels/spike, and percent fertility. Those

TABLE VII

## ESTIMATES OF VARIANCE COMPONENTS FOR GENERAL AND SPECIFIC COMBINING ABILITY FOR EIGHT CHARACTERS

| Trait | GCA <br> Component | SCA <br> Component | GCA/SCA |
| :--- | :---: | :---: | :---: |
| Tiller Number | 4.37 | 7.54 | 0.78 |
| Kernels/Spike | 198.29 | 253.99 | 0.78 |
| Kernel Weight | 32.97 | 40.86 | 0.81 |
| Grain Yield | 18.60 | 29.69 | 0.64 |
| Percent Fertility | 1229.50 | 2293.69 | 0.54 |
| Plant Height | 156.87 | 13.72 | 4.17 |
| Heading Date | 38.74 | 6.85 | 2.82 |
| Percent Protein | 2.83 |  | 0.41 |

TABLE VIII
ESTIMATES OF GENERAL COMBINING ABILITY EFFECTS AMONG ELEVEN PARENTS FOR EIGHT CHARACTERS

| Parent | Tiller <br> Number | $\begin{gathered} \text { Kernels/ } \\ \text { Spike } \\ \hline \end{gathered}$ | Kernel <br> Weight | Grain <br> Yield | Percent Fertility | $\begin{array}{r} \text { Plant } \\ \text { Height } \end{array}$ | $\begin{gathered} \text { Heading } \\ \text { Date } \\ \hline \end{gathered}$ | Percent Protein |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Osage | 0.89 | 3.85* | -0.73 | 2.46* | 15.41* | 0.94 | 1.45* | -1.00* |
| Atr 71 | 0.46 | -10.99* | 2.49* | -2.89* | -25.97* | -4.98* | -1.77* | 0.48 |
| Aura | -1.98* | 5.53* | 3.11* | 0.29 | 2.35 | -0.06 | -1.62* | -0.14 |
| W.Trans | -0.67 | 5.22* | -3.56* | -0.39 | 3.00 | 9.41* | 4.97* | -0.22 |
| 6615D | 0.51 | -8.26* | 1.99* | -2.11* | -21.90* | -5.09* | -2.62* | 0.39 |
| Payne | 0.16 | 7.64* | -1.45* | $2.76 *$ | 21.59* | -2.50* | -0.37 | -0.87* |
| Salm | -0.70 | 3.29* | 1.65* | 1.25 | 4.44 | 7.11* | 2.99* | 0.68 |
| Amigo | 0.01 | -0.34 | 2.44* | 1.27 | 3.92 | -2.33 | -1.98* | -0.85* |
| 15322 | 0.47 | -1.80 | -2.03* | -0.57 | -1.40 | 3.60* | 0.44 | 0.10 |
| F1ex | 1.22 | -4.22* | -0.65 | -0.24 | -12.03* | 1.83 | 2.84* | 1.29* |
| PmV | -0.37 | 0.08 | -3.26* | -1.83 | 10.59* | -7.93* | -4.33* | 0.14 |
| S.E. | 0.64 | 1.32 | 0.70 | 0.96 | 3.38 | 1.23 | 0.51 | 0.38 |

two parents also exhibited the largest negative GCA effects for percent protein. Osage displayed a tendency toward later heading date in its hybrids while Payne showed tendencies toward lighter kernel weight and shorter plant height. Arthur 71 showed significant negative GCA effects for kernels/spike, grain yield, percent fertility, plant height (shorter plants) and heading date (earlier maturing) as well as a tendency toward increased kernel weight. The greatest amount of positive effects for plant height were recorded for Winter Transec, Salmon, and CI 15322 indicating that they contribute toward taller stature. in their hybrids. Winter Transec and Salmon also showed trends in their offspring toward later heading dates, as did Osage and Flex. Flex showed the greatest positive effects for percent protein and tiller number (though the latter was not significant at the 0.05 probability level). The parent Aurora showed significant negative GCA effects for tiller number; however, it displayed positive effects for kernels/spike and kernel weight. Winter Transec and Plainsman V had the largest negative GCA effects for kernel weight. Arthur 71 and Purdue 6615D had significant negative effects for yield, percent fertility, and kernels/spike.

SCA Effects

The SCA effects for tiller number and kernels/spike are given in Table IX along with their respective standard errors. None of the hybrids exhibited significant positive SCA effects for tiller number. One hybrid, Osage/Plainsman $V$, exhibited significant negative SCA effects for this trait while Payne/Plainsman V displayed substantial, but not significant, negative effects. All hybrids with Plainsman V

TABLE IX
ESTIMATES OF SPECIFIC COMBINING ABILITY EFFECTS FOR TILLER NUMBER AND KERNELS/SPIKE

| Parent | Tiller Number (Upper Diagonal) |  |  |  | 6615D | Payne | Salm | Kernels/Spike (Lower Diagonal) |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Osage | Atr 71 | Aura | W.Trans |  |  |  | Amigo | 15322 | Flex | PmV |
| Osage |  | 1.13 | 1.66 | --- | 2.55 | 0.68 | -0.52 | 0.67 | 3.02 | -1.10 | -5.73 |
| Atr 71 | 1.21 |  | 0.43 | 1.56 | -2.58 | 0.86 | 1.22 | 0.57 | 1.29 | 0.36 | -1.93 |
| Aura | 5.77 | 16.52 |  | 0.90 | -0.96 | -1.04 | -0.03 | -0.62 | -0.29 | 1.48 | -1.11 |
| W.Trans | --- | -16.40 | 8.19 |  | 0.87 | 0.58 | 0.99 | 0.32 | -1.05 | 1.01 | -0.50 |
| 6615D | 0.91 | 12.91 | -21.03 | -11.47 |  | 1.69 | 2.10 | 0.37 | 1.15 | 1.09 | -2.28 |
| Payne | 4.83 | -6.09 | 6.76 | 11.10 | -6.31 |  | 0.55 | -0.44 | 0.29 | 0.66 | -4.22 |
| Salm | -1.80 | -5.63 | 4.65 | -10.60 | -3.10 | 4.09 |  | 1.23 | -0.76 | -1.04 | -1.87 |
| Amigo | 0.74 | 1.44 | 6.17 | -6.46 | 1.51 | -0.79 | 1.01 |  | -0.10 | 0.90 | -1.08 |
| 15322 | -3.93 | 0.58 | 3.72 | 4.82 | -5.81 | -2.57 | -2.18 | 0.77 |  | 0.68 | -2.22 |
| Flex | 3.55 | 8.70 | -16.15 | -14.91 | 10.81 | -5.24 | -4.91 | -3.92 | 3.35 |  | -1.20 |
| PmV | -9.56 | -22.50 | -19.76 | 22.04 | 9.39 | -5.58 | -13.32 | -0.60 | 2.60 | 9.66 |  |
|  | S.E. | $=4.57$ | S.E. ${ }^{(2)}$ | $=4.37$ |  |  | S.E. ${ }^{(1)}=2.23$ S.E. ${ }^{(2)}=2.14$ |  |  |  |  |

(1) Standard error between two crosses having one common parent.
(2) Standard error between two crosses having no parents in common.
as a parent showed negative SCA effects for tiller number.
Seven hybrids, i.e., Arthur 71/Aurora, Arthur 71/Purdue 6615D, Winter Transec/Payne, Winter Transec/P1ainsman V, Purdue 6615D/F1ex, Purdue 6615D/Plainsman V, and Flex/Plainsman V, displayed significant positive SCA effects for kernels/spike. Winter Transec/Plainsman V exhibited the largest positive SCA effect. Ten hybrids, on the other hand, displayed significant negative SCA effects for this trait. These were Osage/P1ainsman V, Arthur 71/Winter Transec, Aurora/Purdue 6615D, Aurora/Flex, Aurora/Plainsman V, Winter Transec/Purdue 6615D, Winter Transec/Salmon, 'Winter Transec/Flex, Salmon/Plainsman V, and Arthur 71/ Plainsman V. The largest negative SCA effect was exhibited by Arthur 71/P1ainsman V.

Table $X$ contains SCA effects for kernel weight and grain yield. Purdue 6615D/Salmon had the highest positive SCA effect for kernel weight with a value of 6.43 . Two other hybrids, Purdue 6615D/CI 15322 and Aurora/Flex, had considerable, though non-significant at the 0.05 level, positive.effects for this trait. A11 the hybrids having Plainsman $V$ as a parent displayed negative SCA effects for kernel weight with three of these hybrids, i.e., Winter Transec/P1ainsman $V$, Salmon/Plainsman $V$, and CI 15322/Plainsman $V$, displaying negative effects of statistically significant magnitude. Three other hybrids, Osage/Aurora, Arthur 71/Purdue 6615D, and Aurora/Payne, showed considerable negative effects for kernel weight though none was significantly different from zero. These results are in good agreement with the ranked means of the $\mathrm{F}_{1}$ hybrids for kernel weight (Table XVI).

None of the hybrids showed significant positive SCA effects for grain yield. Two hybrids, Osage/Plainsman V and Aurora/Purdue 6615D,

TABLE X
ESTIMATES OF SPECIFIC COMBINING ABILITY EFFECTS FOR KERNEL WEIGHT AND GRAIN YIELD

(1) Standard error between two crosses having one common parent.
(2) Standard error between two crosses having no parents in common.
displayed significant negative SCA effects for this trait. These results are in good agreement with the rank of the $\mathrm{F}_{1}$ means. SCA effects for percent fertility and plant height are given in Table XI. Ten hybrids, Arthur 71/Purdue 6615D, Winter Transec/Payne, Arthur 71/Flex, Purdue 6615D/Flex, Arthur 71/P1ainsman V, Winter Transec/Plainsman V, Purdue 6615D/P1ainsman V, Salmon/P1ainsman V, Aurora/Flex, and Flex/Plainsman V exhibited significant positive SCA effects for percent fertility. The Flex/Plainsman $V$ hybrid showed the highest positive effect. Arthur 71/Aurora, Arthur 71/Winter Transec, Aurora/Purdue 6615D, Winter Transec/Purdue 6615D, Winter Transec/ Salmon, Winter Transec/Flex, and Aurora/Plainsman V displayed significant negative SCA effects for this trait with Aurora/Purdue 6615D showing the greatest negative effect.

Only Winter Transec/CI 15322 displayed significant positive SCA effects for plant height. Two hybrids, Salmon/Plainsman V and CI 15322/Plainsman $V$, showed significant negative SCA effects.

The SCA effects for heading date and percent protein are presented in Table XII. All the hybrids having Plainsman $V$ as one of their parents exhibited positive SCA effects for heading date (later heading), and three of them (i.e., Aurora/Plainsman V, Payne/Plainsman V, and CI 15322/P1ainsman V) showed significant positive SCA effects for this trait. Two hybrids, Winter Transec/Purdue 6615D and Payne/Flex, showed substantial but non-significant negative SCA effects.

Arthur 71/Aurora, Arthur 71/Winter Transec, Aurora/Purdue 6615D, and Aurora/Flex exhibited significant positive SCA effects for percent protein. Five hybrids, Arthur 71/Flex, Aurora/Winter Transec,

## TABLE XI

ESTIMATES OF SPECIFIC COMBINING ABILITY EFFECTS FOR PERCENT FERTILITY AND PLANT HEIGHT

(1) Standard error between two crosses having one common parent.
(2) Standard error between two crosses having no parents in common.

TABLE XII

ESTIMATES OF SPECIFIC COMBINING ABILITY EFFECTS FOR hEADING DATE AND PERCENT PROTEIN

| Parent | Heading Date (Upper Diagonal) |  |  |  | 6615D | Payne | Salm | Percent Protein (Lower Diagonal) |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Osage | Atr 71 | Aura | W.Trans |  |  |  | Amigo | 15322 | F1ex | PmV |
| Osage |  | -0.37 | -0.77 | --- | $-1.27$ | -1.02 | 0.37 | -1.40 | -1.33 | 0.77 | 2.86 |
| Atr 71 | 0.22 |  | 0.20 | $-2.13$ | 1.20 | 0.70 | $-1.67$ | 0.32 | -0.36 | $-1.76$ | 2.77 |
| Aura | $-1.35$ | 3.14 |  | -2.79 | 1.55 | -1. 20 | -1.82 | -0.34 | -1.26 | -1.92 | 5.31 |
| W.Trans | --- | 2.29 | -2.10 |  | -3.29 | -1.29 | 0.59 | -0.42 | 3.40 | 1.99 | 0.56 |
| 6615D | -1.08 | -1.43 | 2.94 | 0.79 |  | 0.55 | -2.57 | 1.66 | -0.26 | -0.42 | 1.06 |
| Payne | -0.11 | 0.30 | -0.85 | -1.62 | 0.99 |  | -1.07 | -0.33 | -0.26 | -3.42 | 3.84 |
| Sa1m | 1.18 | 0.79 | -0.25 | 1.47 | 0.32 | -1.05 |  | -2.70 | 2.62 | 2.46 | 1.22 |
| Amigo | -0.56 | -0.04 | -0.92 | -0.44 | 0.55 | 0.61 | -1.42 |  | -0.39 | -2.05 | 3.12 |
| 15322 | 1.08 | -1.37 | -1.09 | 0.18 | 1.45 | -0.27 | 0.68 | 0.91 |  | -0.47 | 4.62 |
| F1ex | -0.05 | -2.64 | 1.94 | 1.39 | -2.24 | 0.39 | 0.94 | 1.02 | 0.47 |  | 1.45 |
| PmV | 0.06 | -0.33 | -1. 52 | $-1.62$ | -1.92 | 1.00 | -2.26 | 0.51 | 1.27 | 0.20 |  |
|  | S.E. | $=0.95$ | S.E. | $=0.89$ |  |  | S.E. ${ }^{(1)}=1.78$ S.E. ${ }^{(2)}=1.67$ |  |  |  |  |

(1) Standard error between two crosses having one common parent.
(2) Standatd error between two crosses having no parents in common.
Purdue 6615D/Flex, Purdue 6615D/P1ainsman V, and Salmon/Plainsman $V$, showed significant negative effects.

## CHAPTER V

## DISCUSSION

According to Sprague and Tatum (33) GCA provides a measure of additive gene action. Brown et al (2) and Kronstad and Foote (16) reported that most of the genetic variability of performance traits studied in wheat was due to additive effects of genes. The ratios of variance components obtained in the present study (Table VII) indicated that most of the genetic variation for $p l a n t$ height and heading date could be accounted for by variation due to GCA. However, SCA components were more important for the other six traits in this study.

Tiller number, kernels/spike, kernel weight, grain yield, percent fertility, and percent protein showed larger components for SCA than for GCA. The number of kernels/spike is related to percent fertility, and some of the hybrids did show unusually high levels of spike sterility. This could account, at least in part, for the larger SCA component for kernels/spike. Grain yield in wheat is a complex trait; the major components of which are tiller number, kernels/spike, and kernel weight. Interactions among these traits could also account in part for the high SCA estimate for grain yield. Yield is also associated with spike fertility; and the high SCA component for this trait could account in part for the high SCA estimate for grain yield. The occurrence of significant $S C A$ effects for all traits in this study suggests that the genetic variation for each trait was also partly due
to dominance, epistasis, or both.
The GCA effects, considering all traits, indicated Osage and Payne were the better parents of the 11 lines studied in this test, while Aurora, Salmon, Amigo, CI 15322, Flex, and Plainsman V appeared to be next in potential value. On the other hand, Arthur 71, Purdue 6615D, and Winter Transec seemed to be the least favorable in this set. Some combinations of these translocation lines resulted in heterosis for grain yield while others resulted in low fertility accompanied byea reduction in kernels/spike, low yield, and high protein content.

As shown in Appendix Tables XIV to XXI, the best hybrid exceeded the best parent for tiller number, kernel weight, grain yield, percent fertility, plant height, and percent protein. Eighteen hybrids exceeded or were equal to the highest yielding parent, Payne. Deviations of those high-yielding hybrids from their high-parents ranged from 0.1 to $6.3 \mathrm{~g} / \mathrm{plant}$ while average heterosis (compared to the highparent) of those hybrids was 123\%. Three hybrids, Arthur 71/Aurora, Arthur 71/Winter Transec, and Aurora/Purdue 6615D, had lower grain yield than Arthur 71, the lowest yielding parent. Deviations of the low-yielding hybrids from their high-parents ranged from -5.1 to -8.9 while the average high-parent heterosis of the three low-yielding hybrids was only $46 \%$. These three hybrids also exhibited the lowest values for percent fertility, ranging from 41.6 to $36.9 \%$, and the lowest values for kernels/spike.

The relationships among the characters of grain yield, percent fertility, kernels/spike, and percent protein as observed for the 10 highest and 10 lowest yielding hybrids are illustrated in Table XIII. The highest yielding hybrids generally displayed increased levels of

## TABLE XIII

## HIGH-YIELDING VS. LOW-YIELDING HYBRIDS

|  | Grain |  | Kernels/ |  |
| :---: | :---: | :---: | :---: | :---: |
|  | Yield | Percent | Spike | Percent |
| F1 Hybrid | $(g /$ Plant $)$ | Fertility | (No.) | Protein |

## High-Yielding Group

| W.Trans/Payne | 22.1 | 164.1 | 60.1 | 15.0 |
| :--- | ---: | ---: | ---: | ---: |
| Osage/Payne | 21.2 | 135.7 | 52.4 | 15.7 |
| Aura/W.Trans | 20.1 | 140.6 | 55.1 | 15.2 |
| Osage/Aura | 20.0 | 145.9 | 51.3 | 15.2 |
| Osage/Amigo | 19.6 | 130.8 | 40.4 | 15.3 |
| Payne/Salm | 19,6 | 149.6 | 51.1 | 16.4 |
| Salm/Amigo | 19.1 | 126.9 | 40.1 | 16.1 |
| Payne/Amigo | 18.0 | 137.6 | 42.6 | 16.1 |
| Osage/15322 | 17.8 | 108.3 | 34.2 | 17.9 |
| Osage/Salm | 17.7 | 130.7 | 41.4 | 18.5 |

Low-Yielding Group

| Atr71/6615D | 10.8 | 100.4 | 29.1 | 17.1 |
| :--- | ---: | ---: | ---: | ---: |
| Atr71/PmV | 10.1 | 92.2 | 25.8 | 18.9 |
| $6615 \mathrm{D} / 15322$ | 10.1 | 66.5 | 20.2 | 19.6 |
| $6615 \mathrm{D} / \mathrm{PmV}$ | 9.2 | 83.4 | 24.4 | 19.9 |
| W.Trans/Flex | 9.1 | 58.7 | 22.2 | 20.1 |
| W.Trans/6615D | 8.5 | 68.8 | 21.6 | 18.6 |
| Osage/PmV | 8.3 | 121.7 | 32.3 | 16.2 |
| Atr71/Aura | 6.5 | 41.6 | 14.1 | 21.2 |
| Atr71/W.Trans | 6.3 | 41.3 | 13.9 | 20.2 |
| Aura/6615D | 4.9 | 36.9 | 12.4 | 20.9 |

fertility in relation to the overall $\mathrm{F}_{1}$ mean for that character, with the exception of the Osage/CI 15322 hybrid. This particular hybrid had the highest tiller number (which also contributes to yield) of all parents and $F_{1}$ 's. With the exception of Osage/CI 15322 , these top 10 yielding hybrids also had increased numbers of kernels/spike and were among the lowest in percent protein.

The 10 lowest yielding hybrids generally had depressed levels of spike fertility, fewer kernels/spike, and increased levels of protein with two exceptions, Arthur $71 /$ Purdue 6615 D and Osage/P1ainsman $V$. These two hybrids were also among the lowest hybrids for tiller number and kernel weight, respectively. Both of those traits are components of yield. The reduction in kernel weight exhibited by Osage/Plains man $V$ was probably due to the hybrid necrosis of this hybrid. Four of the parents associated with the high-yielding hybrids were also present in combinations which resulted in low yield and low fertility, i.e., Aurora, Winter Transec, Osage, and CI 15322. Either Payne, Osage, or both were parents in eight of the 10 high-yielding $\mathrm{F}_{1}$ 's while Osage was present in only one of the lowest yielding hybrids.

If the chromosomes involved in the translocations are considered (Table I), it should be noted that those $\mathrm{F}_{1}{ }^{\prime}$ s which exhibited positive heterosis for yield contained either the $2 R-4 \mathrm{~A}$ of Winter Transec, the $3 A g-3 D$ of Osage, the $1 R-1 B s$ of Aurora, the $1 R-1 B$ of Salmon, or the unknown translocations of Payne, Amigo, or CI 15322. The $\mathrm{F}_{1}$ 's having reduced yield contained either the $6 C^{\mathrm{u}}-6 \mathrm{~B} 1$ translocation of Arthur 71 or the $7 e 1-7 D 1$ translocation of Purdue 6615D. There are two exceptions, the Winter Transec/Flex and the Osage/Plainsman V F ${ }_{1}$ 's. These contain the $2 R-4 A$ and $3 A g-3 D$ translocations of Winter Transec and

Osage, respectively, along with the unknown translocations of Flex and Plainsman V.

The extra vigor for yield apparently imparted by Osage and Payne may or may not have been due to the particular alien translocation involved in these two lines. It should be noted that these two were developed specifically for the Southern Great Plains region within which this study was conducted. Their potency for yield may be due to their background genotypes rather than the particular alien-gene complex involved. To make a completely definitive study of the effects of combining alien translocation lines, each translocation should be in a common genetic background which, in this case, could be approached by backcrossing each line onto a single adapted cultivar.

In a previous study $(27,28), F_{1}$ 's and parents of a diallel cross among six alien translocation lines were examined. The results of that study were that GCA components were larger than those for SCA for tiller number, kernel weight, plant height, and heading date, while SCA components were larger for kernels/spike, grain yield, percent sterility, and percent protein. Four hybrids were found to have high positive heterosis for grain yield while four others exhibited high levels of sterility, low values for kernels/spike, and low grain yields. It was also found that two parents, Aurora and Purdue 6615D, were involved as parents in the four highest, as well as the four lowest yielding hybrids.

In the present study GCA components were larger than those for SCA for plant height and heading date while SCA components were larger for kernels/spike, tiller number, kernel weight, grain yield, percent fertility, and percent protein. Several hybrids were found to have
high positive heterosis for grain yield. Several others displayed low levels of spike fertility accompanied by low grain yield, decreased kernels/spike, and increased levels of protein. It is noteworthy that four of the five lowest yielding hybrids in this present study were also the four lowest yielding hybrids in the previous study. It is of further interest to note that one of the four parents present in combinations common to both high-yielding and low-yielding groups in the present study, was also common to both groups in the previous study. The results of this study are in good general agreement with those of the previous study.

The number of kernels/spike in the hybrids in this, as well as in the previous study (27), were lower than the parental mean. In the previous study, only one hybrid had a significantly greater number of kernels/spike than its high-parent while two hybrids had significantly more kernels/spike than their respective mid-parents. In the present study, no hybrid had significantly more kernels/spike than its highparent while six hybrids had a significantly greater number of kernels/ spike than their mid-parents. On the other hand, in both studies there was a preponderance of hybrids with significantly fewer kernels/ spike than both their high- and mid-parents.

The low values for kernels/spike, as well as the low levels of spike fertility exhibited by some of these hybrids, may have been due to the loss of portions of wheat chromatin when two of the alien translocation lines were brought together. The loss of wheat chromatin may result in reduction of chromosome pairing which could result in chromosome imbalance. This would cause a decrease in spike fertility, which in turn would result in a decrease in kernels/spike.
Wheat breeding programs in the Hard Red Winter Wheat Area have progressed to the point that more attention is now being given to the development of cultivars having multiple-pest resistance. Because many of the pest-resistant sources now available are alien translocation lines, it is important to know something about possible beneficial vs. detrimental effects when those translocations are combined into a single genotype.
The results of this study indicate that while certain combinations of these alien translocation lines give an increase in fertility accompanied by increased yield, other combinations should probably be avoided in breeding programs on the Southern Great Plains, e.g. Arthur 71/Aurora, Arthur 71/Winter Transec, Aurora/Purdue 6615D, Osage/Plainsman V, and Winter Transec/Purdue 6615D.

## CHAPTER VI

## SUMMARY AND CONCLUSIONS

The $F_{1}{ }^{\prime} s$ and parents of a diallel cross among 11 alientranslocation 1 ines of winter wheat were studied to determine the response of eight performance traits when alien gene complexes were combined in all possible two-way combinations. The 11 parents and 54 $F_{1}$ 's (one $F_{1}$ died in the seedling stage) were grown in a space-planted nursery on the Agronomy Research Station, Stillwater, Oklahoma, during the 1978-79 season. The experiment was conducted in a randomized, complete-block design with four replications. Eight vigorous, bordered plants from each single-row plot were evaluated. Measurements were taken on tiller number, kernels/spike, kernel weight, grain yield, percent fertility, plant height, heading date, and percent protein. A11 data were subjected to standard analyses of variance, as well as Griffing's analyses for GCA and SCA. Hybrid-parent contrasts were also examined for each trait.

The GCA components were larger than those for SCA for plant height and heading date, indicating that additive gene effects were more important for those two traits than were nonadditive effects. On the other hand, SCA components were larger for tiller number, kernels/ spike, kernel weight, grain yield, percent fertility, and percent protein, indicating that in this particular set of genotypes nonadditive gene effects for those traits were more important than additive
effects.

The combining ability data for yield and yield components indicated that Osage and Payne were probably the better parents in this set while Arthur 71 , Winter Transec, and Purdue 6615 D were the least so. Eleven hybrids, i.e., Winter Transec/Payne, Osage/Payne, Aurora/ Winter Transec, Osage/Aurora, Osage/Amigo, Salmon/Amigo, Aurora/Salmon, Purdue 6615D/Salmon, Winter Transec/P1ainsman V, CI 15322/P1ainsman V, and Arthur $71 / C I 15322$ showed significant positive high-parent heterosis for yield. On the other hand, several exhibited low levels of fertility, low numbers of kernels/spike, and low grain yields.

Aurora, Winter Transec, Osage, and CI 15322 were involved as parents in some of the highest yielding hybrids as well as some of the lowest yielding. This indicates that specific combinations of alientranslocation lines could do very well; whereas, others would do very poorly. Certain combinations of alien-translocation lines might present some problems in a wheat breeding program on the Southern Great Plains, e.g., Aurora/Purdue 6615D, Arthur 71/Winter Transec, Arthur 71/Aurora, Osage/Plainsman V, and Winter Transec/Purdue 6615D. This situation would have important implications in wheat breeding programs for multiple-pest resistance because many of the present sources of pest resistance exist in alien-translocation lines.

## LITERATURE CITED

1. Anonymous. 1976. Hard winter wheat varieties of the $70^{\prime}$ s. Hard Winter Wheat Quality Advisory Counci1, Manhattan, Kan.
2. Brown, C. M., R. O. Weibel, and R. D. Seif. 1966. Heterosis and combining ability in common winter wheat. Crop Sci. 6:382383.
3. Drisco11, C. J. 1965. Induced intergeneric transfers of chromosome segments. In The use of induced mutations in plant breeding, Pergamon Press, New York. (Supp1. to Radiation Bot. 5:727-739).
4. $\qquad$ . 1968. Alien transfer by irradiation and meiotic control. p.196-203. In K. W. Finlay and K. W. Shépherd (ed.) Proc. 3rd Int. Wheat Genet. Symp., Canberra, Aust.
5. $\qquad$ , and L. M. Anderson. 1967. Cytogenetic studies of Transec-A wheat-rye translocation line. Can. J. Genet. Cytol. 9:375-380.
6. $\qquad$ , and L. M. Bielig. 1968. Mapping of the Transec wheat-rye translocation. Can. J. Genet. Cytol. 10:421-425.
7. $\qquad$ , and N. F. Jensen. 1964. Characteristics of leaf rust resistance transferred from rye to wheat. Crop Sci. 4:372374.
8. $\qquad$ , and $\qquad$ . 1969. Registration of Transec wheat germplasm (Reg. No. GP1). Crop Sci. 9:682.
9. Griffing, B. 1956. Concept of general and specific combining ability in relation to diallel crossing systems. Aust. J. Biol. Sci. 9:463-493.
10. Hermsen, J. G. Th. 1963. Hybrid necrosis as a problem for the wheat breeder. Euphytica 12:1-16.
11. Kimber, G. 1971. The design of a method, using ionising radiation, for the introduction of alien variation into wheat. Indian J. Genet. Plant Breed. 31:580-584.
12. Knott, D. R. 1961. The inheritance of rust resistance. VI. The transfer of stem rust resistance from Agropyron elongatum to common wheat. Can. J. Plant Sci. 41:109-123.
$\qquad$ . 1964. The effect on wheat of an Agropyron chromosome carrying rust resistance. Can. J. Genet. Cytol. 6:500507.
13. $\qquad$ . 1968. Agropyrons as a source of rust resistance in wheat breeding. p.204-212. In K. W. Finlay and K. W. Shepherd (ed.) Proc. 3rd Int. Wheat Genet. Symp., Canberra, Aust.
14. $\qquad$ . 1971. The transfer of genes for disease resistance from alien species to wheat by induced translocations. p. 67-77. In Mutation breeding for disease resistance, Int. Atomic Energy Agency, Vienna.
15. Kronstad, W. E., and W. H. Foote. 1964. General and specific combining ability estimates in winter wheat (Triticum aestivum Vill., Host). Crop Sci. 4:616-619.
16. Patterson, F. L., R. L. Gallun, J. J. Roberts, R. E. Finney, and G. E. Shaner. 1975. Registration of Arthur 71 and Abe wheat (Reg. Nos. 560 and 562). Crop Sci. 15:736.
17. Riley, R., V. Chapman, and R. Johnson. 1968. The incorporation of alien disease resistance in wheat by genetic interference with the regulation of meiotic chromosome synapsis. Genet. Res., Camb. 12:199-219.
18. Sears, E. R. 1956. The transfer of leaf-rust resistance from Aegilops umbellulata to wheat. Brookhaven Symp. Biol. 9:1-22.
19. $\qquad$ . 1972. Chromosome engineering in wheat. p. 23-28. Proc. 4th Stadler Symp., Columbia, Mo.
20. $\qquad$ . 1973. Agropyron-wheat transfers induced by homoeologous pairing. p.191-199. In E. R. Sears and L. M. S. Sears (ed.) Proc. 4 th Int. Wheat Genet. Symp., Columbia, Mo.
$\qquad$ . 1975. The wheats and their relatives. p. 59-91. In R. C. King (ed.) Handbook of Genetics, Vol. 2. Plenum Press.
21. $\qquad$ . 1977. Analysis of wheat-Agropyron recombinant chromosomes. p. 63-72. Proc. 8th Eucarpia Cong., Madrid, Spain.
22. Sebesta, E. E. 1976. Personal communication.
23. $\qquad$ - 1980. Personal communication.
24. $\qquad$ , and E. A. Wood, Jr. 1978. Transfer of greenbug resistance from rye to wheat with X-rays. Agron. Abstrs. p. 61-62.
25. Sebesta, P. G. 1977. Combining ability analysis of yield and yield components involving six winter wheat alien-translocation lines. M.S. Thesis. Oklahoma State Univ.
26. $\qquad$ , and E. L. Smith. 1977. Combining ability of six wheat alien translocation lines. Agron. Abstrs. p. 71.
27. Sharma, D., and D. R. Knott. 1966. The transfer of leaf-rust resistance from Agropyron to Triticum by irradiation. Can. J. Genet. Cytol. 8:137-143.
28. Smith, E. L. 1980. Personal communication.
29. $\qquad$ , L. H. Edwards, H. Pass, H. C. Young, Jr., and D. C. Abbott. 1976. Registration of Osage wheat (Reg. No. 570). Crop Sci. 16:445-446.
30. $\qquad$ , A. M. Schlehuber, H. C. Young, Jr., and L. H. Edwards. 1968. Registration of Agent wheat (Reg. No. 471). Crop Sci. 8:511-512.
31. Sprague, G. F., and L. A. Tatum. 1942. General vs. specific combining ability in single crosses of corn. J. Am. Soc. Agron. 34:923-932.
32. Tsunewaki, K. 1964. Genetic studies of a $6 x$-derivative from an 8x Triticale. Can. J. Genet. Cytol. 6:1-11.
33. Wells, D. G., and C. R. Cowley. 1976. Registration of SD69103, Hand, and Flex winter wheat germplasm (Reg. Nos. GP70, GP71, and GP72). Crop Sci. 16:888.
34. Zeller, F. J. 1973. 1B/1R wheat-rye chromosome substitutions and translocations. p. 209-221. In E. R. Sears and L. M. S. Sears (ed.) Proc. 4 th Int. Wheat Genet. Symp., Columbia, Mo.

APPENDIX

PARENTAL AND $\mathrm{F}_{1}$ RANKED MEANS AND HYBRID-PARENT DEVIATIONS FOR TILLER NUMBER

| Parent or F1 Hybrid | Kank | TillerNumber(No. /Plant) | Hybrid-Parent Deviation |  |
| :---: | :---: | :---: | :---: | :---: |
|  |  |  | Mid-Parent | High-Parent |
| Osage/15322 | 1 | 17.2 | 5.4** | 4.9*** |
| Osage/661.5D | 2 | 16.8 | 4.9** | 4.5** |
| $0615 \mathrm{D} / \mathrm{Flex}$ | 3 | 1.5 .7 | 3.6** | 3.4** |
| Osage/Atr 71 | 4 | 15.3 | 3.8** | 3.0** |
| Flex/PmV | 4 | 15.3 | 3.7** | 2.9* |
| 15322/F1ex | 6 | 15.2 | 3.4** | 2.8* |
| 6615D/Payne | 6 | 15.2 | 3.8** | 3.6** |
| Atr 71/1.5322 | 8 | 15.1 | 4.1** | 3.9 \%* |
| Amigo/flex | 9 | 15.0 | 3.3** | 2.6* |
| 66151/1.5322 | 9 | 15.0 | 3.6** | 3.4** |
| Atr7l/Flex | 11 | 14.9 | 3.3** | 2.5* |
| Payne/Flex | 11 | 14.9 | 3.1** | 2.5* |
| $6615 \mathrm{p} / \mathrm{Salm}$ | 13 | 14.8 | 4.2** | 3.2** |
| Osage/Payne | 14 | 14.6 | 2.8** | 2.3* |
| W. Trans/Flex | 15 | 14.4 | 3.9** | 2.0 |
| Ogage/Amigo | 15 | 14.4 | 2.8** | 2.1 |
| Atr 71/Payne | 17 | 14.3 | 3.3** | 3.0\%* |
| Atr71/w.Trans | 18 | 14.2 | 4.5** | 3.4** |
| W.Tramb/PmV | 18 | 14.2 | 4.5** | 3.4** |
| Atr $71 / \mathrm{PmV}$ | 20 | 13.9 | 3.1** | 3.1** |
| Acr 71/Amigo | 20 | 13.9 | 3.0** | 2.9* |
| Atr71/Salm | 22 | 13.8 | 3.6** | 3.0** |
| Osage/Flex. | 22 | 13.8 | 1.5 | 1.4 |
| Payne/15322 | 22 | 13.8 | 2.6** | 2.5* |
| 6615d/Amigo | 25 | 13.7 | 2.4* | 2.1 |
| Aura/Flex | 26 | 13.6 | 3.5** | 1.2 |
| Osage/Aura | 27 | 13.4 | 5.6** | 5.0** |
| Salm/Amigo | 27 | 13.4 | 3.1** | 2.4* |
| Amigo/PmV | 29 | 13.3 | 2.4* | 2.3* |
| $15322 / \mathrm{PmV}$ | 29 | 13.3 | 2.3* | 2.1 |
| Amigo/ 15322 | 31 | 13.2 | 2.1* | 2.0 |
| W.Trans/Payne | 32 | 12.9 | 3.0** | 1.6 |
| 6615D/PmV | 32 | 12.9 | 1.7 | 1.3 |
| Payne/Salm | 34 | 12.8 | 2.4* | 1.5 |
| Payne/Amigo | 35 | 12.6 | 1.5 | 1.3 |
| W.Trans/Anigo | 36 | 12.5 | 2.7** | 1.5 |
| W.Trans/Salm | 36 | 12.5 | 3.4** | 2.9* |
| Payne/PmV | 36. | 12.5 | 1.5 | 1.2 |
| Osage/Salm | 36 | 12.5 | 1.6 | 0.2 |
| Flex | 40 | 12.4 | -- | - |
| Salm/Flex | 41 | 12.3 | 1.3 | -0.1 |
| Osage | 41 | 12.3 | --- | --- |
| Atr71/Aura | 43 | 11.8 | 6.6** | 2. ${ }^{*}$ |
| Salm/L;322 | 43 | 1.1 .8 | 1.4 | 0.6 |
| Salin/ PmV | 43 | 11.8 | 1.6 | 1.0 |
| W.Trama/66150 | 43 | 11.8 | 1.7 | 0.2 |
| W.Trana/15322 | 47 | 11.6 | 1.7 | 0.4 |
| Aura/15322 | 47 | 11.6 | 2.1** | 0.4 |
| 66150 | 47 | 11.6 | --- | --- |
| Payne | 50 | 11.3 | --- | --- |
| Atr71/66150 | 51 | 11.2 | 0.0 | -0.4 |
| 1532.2 | 51 | 11.2 | --- | --- |
| Aura/W.Trans | 53 | 11.1 | 2.9** | 2.5* |
| Amigo | 54 | 11.0 | --- | --- |
| Atr 71 | 55 | 10.8 | --- | --- |
| PmV | 55 | 10.8 | --- | --- |
| Aura/PmV | 57 | 10.4 | 1.1 | -0.4 |
| Aura/6615i | 57 | 10.4 | 0.6 | -1.2 |
| Aura/Amigo | 59 | 10.3 | 0.9 | -0.7 |
| Aura/Salm | 60 | 10.1 | 1.4 | 0.5 |
| Aura/Payne | 61 | 10.0 | 0.4 | -1.3 |
| Osage / PmV | 61 | 10.0 | -1.5 | -2.3* |
| Salm | 63 | 9.6 | --- | --- |
| W.Trans | 64 | 8.6 | --- | --- |
| Aura | 65 | 7.9 | --- | --- |
| Osage/W.trans | 66 | --- | Seedli | ng Lethal |

*,** Significant at the 0.05 and 0.01 levels of probability, respectively.

TABLE XV

PARENTAL AND $\mathrm{F}_{1}$ RANKED MEANS AND HYBRID－PARENT DEVIATIONS FOR KERNELS／SPIKE

| Parent or IL Hybrid | Kornels／ |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  | Rank | spike <br> （ NO. ） | Hybrid－Parent Deviation |  |
|  |  |  | Mid－parent | ligh－Paront |
| W．Trans | 1 | 60.2 | －－－ | －－－ |
| W．Trans／Pityme | 2 | 60.1 | 4．4＊ | －0．1 |
| Aura／Payme | 3 | 56.0 | － 2.8 | 3.7 |
| Aura／W ．＇Tralu： | 1 | 「「． | －1．1 | $-5.1 \%$ |
| Us：as：／Paynu | 5 | 52.4 | ＇，． $8 \% \%$ | 1.2 |
| Aura | 6 | ＇）． 3 | ．$-\cdots$ | －－－ |
|  | 7 | ＇1． 3 | 4．0） | －1．0 |
| P＇ayme | 8 | 4.2 | －－－ | －－－ |
| l＇ayme／sialm | 9 | 51.1 | 1.6 | －0． 1 |
| Sura／Pmi | 10 | 49.8 | 6．6＊＊ | －2．5 |
| W．Trans／binV | 10 | 49.8 | 2.6 | －10．4＊＊ |
| ：$a^{1} 1 \mathrm{~m}$ | 12. | $4 \% .8$ | －－． | －－－ |
| ： $\mathrm{a} 1 \mathrm{~m} / \mathrm{PmV}$ | 13 | 47.6 | $6.6 * *$ | －0．2 |
| Auria／Amigo | 1／1 | 47．＇5 | 2.8 | －4．8： |
| Aura／Salm | 1＇； | 45.6 | －4．4＊ | －6． $7 * *$ |
| W．Tran：i／1，3\％2 | 16 | 14.1 | －2．1 | －15．8\％ |
| Aura／1，32\％ | 17 | 43.6 | 1.1 | －8．7＊＊ |
| l＇ayme／Amis： | 18 | 42.6 | －1．5 | －8．6＊＊ |
| 1）stas： | 19 | 42.1 | －－－ | －－－ |
| 1satge／stlo | 20 | 11．1 | －3．6 | －6．4＊＊ |
| （1）：age／Amisu | 21 | 40.1 | 0.8 | －1．7 |
| $\because a / m / A m i s$, | 22 | 40.1 | －2．3 | －7．7＊＊ |
| Payne／15322 | 23 | 39.4 | －2．6 | －11．8＊＊ |
| （1：age／Flex | 24 | 39.3 | －1．4 | －2．8 |
| Flax | 24 | 39.3 | －－－ | －－－ |
| Payne／PmV | 26 | 38.4 | $-1+3 *$ | $-12.8 * *$ |
| Amipoo | 27 | 37.1 | －－－ | －－－ |
| $15322 / \mathrm{PmV}$ | 28 | 36.8 | 3.4 | 2.6 |
| Salm／1532\％ | 29 | 35.4 | －4．9\％ | －12．4＊＊ |
| W．Trans／Amipol | 30 | 34.5 | －14．1＊＊ | －25．7＊＊ |
| 66150／Flex | 31 | 34.4 | －1．5 | －4．9\％ |
| Payne／Flex | 3. | 34.3 | －10．9＊＊ | －16．9＊＊ |
|  | 33 | 34.2 | －3．2 | －7．9＊＊ |
| I＇mV | 33 | 34.2 | －－－ | －－－ |
| W．Trans／salm | 35 | 34.0 | －20．0\％＊ | $-26.2 * *$ |
| Amigo／prov | 16 | 33.6 | －2．0 | －3．5 |
| 1＇322／F1ex | 37 | 33.4 | －2．6 | －5．9＊ |
| Sulyol 1 ＇32\％ | 18 | 33.2 | －1．7 | －3．9 |
| 1，3\％2 | 39 | 32.7 | －－－ | －－－ |
|  | 40 | 32.6 | －4．7\％ | －9．5\％\％ |
| 60150 | 1.1 | 32.5 | －－－ | －－－ |
| （）：aige／$/ \mathrm{mV}$ | 1.2 | 32.3 | －5．8\％\％ | －9．8＊＊ |
| Salm／flex | 1.3 | 30.3 | －13．3＊＊ | －17．5\％＊ |
| Osamb／Atr71 | 14 | 30.2 | －2．9 | －11．9＊＊ |
| Strolflex | 15 | 29.6 | －2．1 | －9．7＊＊ |
| 601 5 ／／Payme | 1.6 | 29.2 | $-12.6 \% \%$ | －22．0％＊＊ |
| 人tr71／601＇j） | 1.7 | 29.1 | 0.8 | －3．4 |
| 6ol Sb／Amigen | 118 | 29.0 | －5．8\％＊ | －8．1＊＊ |
| 66151）／salm | 19 | 28.0 | －12．1＊＊ | －19．8＊＊ |
| Mul so／Flex | ， 0 | 27.6 | －10．6＊＊ | －11．7火＊ |
| Flex／PmV | ＇，1． | 27.4 | －9．4\％\％ | －11．9＊＊ |
| Atrll／Payne | ＇， 2 | 26.7 | －10．9＊＊ | －24． $5 \% \%$ |
| Atr7L／Amig＇ | ＇，3 | 26.2 | －4．4＊ | －10．9＊＊ |
| Atr／t／PmV | \％ 1 | 25.8 | －3．3 | －8．4＊＊ |
| 661 ${ }^{\circ} \mathrm{O}$／PmV | י3 | 24.4 | －8．9\％\％ | －9．8＊＊ |
| Atril | ＇6 | 24.1 | －－－ | －－－ |
| Atr71／1＇322． | ＇，7 | 23.9 | ．$-4.5 * *$ | －8．8＊＊ |
| Atr71／Salm | ＇8 | 22.8 | －13．1＊＊ | －25．0\％\％ |
| W．Prans／Fle\％ | \％9 | 22．？ | －27．6＊＊ | －38．0\％ |
| W．Irans／66l＇， | 60 | 21.6 | $-24.7 \% \%$ | －38．6\％＊ |
| Mura／flex | 61 | 2L． 3 | －24． 5 号\％ | －31．0\％＊ |
| （6） $1513 / 15322$ | 62 | 20.2 | －12．4＊＊＊ | －12．5＊＊ |
| Atr7l／Aura | 1,3 | 14.1 | －24．1\％ | －38．2＊＊ |
| Atrol／W．Iranm | 64 | 13.9 | －2．8．2＊＊ | －46．3 $\%$ \％ |
|  | 65 | 1.2 .1 | －30．0\％\％ | －39．9\％\％ |
| Osabe／W I＇rans | 66 | － | Scedl | ng lothal |

[^1] respectively．

TABLE XVI

PARENTAL AND $F_{1}$ RANKED MEANS AND HYBRID－PARENT DEVIATIONS FOR KERNEL WEIGHT

| Piarent or Fh llyhrld | Kank | $\begin{aligned} & \text { Kernel } \\ & \text { We ight } \end{aligned}$ | Ilybrid－Parent Deviation |  |
| :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |
|  |  | （g／1000） | Mid－parcol | igh－Par |
| 66150／Salm | 1 | 48.5 | 13．1\％＊ | 12．6\％ |
| Aura／bfl＇， | 2 | 45.2 | $7.4 * *$ | 4．0）\％ |
| Mura／flex | 2 | 4 4，．？ | 9． $2 \times *$ | 4．0） $1 \times$＊ |
| A1rol／Aut： | 2 | $4{ }^{\circ} \mathrm{F}$ ．${ }^{\text {a }}$ | ＇）． $8 * *$ | 4．0） |
| 心tro／大miso | 2 | 45.2 | 6．9＊＊ | 6． $3 \% \%$ |
| Atrl／Salm | 6 | 45.0 | 8． $3 \% \%$ | 7．14\％ |
| W：ay，／／（r）l | 7 | 44.0 | 7．1\％＊ | 6． $4 \times \%$ |
| Nuta／Amis：＂ | 7 | 111．0 | $4.0 \% *$ | 2．8＊ |
| Aura／Salin | 9 | 13.6 | ＇j． $1 * *$ | 2.4 |
| Salm／Amisu | 10 | 13.1 | $5.7 \% \%$ | 4．2\％ |
| Atrol／fayme | 10 | 13.1 | 7．9\％\％ | 5．5＊＊ |
| Onage／Amisu | 12 | 42.8 | 5． $3 \% \%$ | 3．9＊＊ |
| 661＇sl）／Paym． | 12 | 42.8 | 9． $2 \cdots *$ | 8． $4 * \%$ |
| （6，15）／1＇3\％2 | 14 | 42.6 | $11.0 \%$ | 8． $2 \%$ \％ |
| Amiss／Flax | 1＇1 | 4）． 1 | $7.32 \%$ | 3． 2 * \％ |
| Aura／1，32\％ | 16 | 11.7 | $6.7 * *$ | 0 －＇ |
| Ntr／l／lsov | 17 | 11.6 | 8．1＊＊ | 4．0\％： |
| Ostapre／fitlin | 17 | 11.6 | 6． $1 * \%$ | 5． $5: 4$ |
| Aura | 19 | $41 . ?$ | －－． | －－－ |
| Salm／Flex | 19 | 41．2 | $7.9 * \%$ | 5． $3 \% \%$ |
| 661速／Amis | 21 | 41.0 | 4． $4.4 \%$ | 2.1 |
| 9：sami／66150 | 22 | 40.8 | 5． $6 \% \%$ | 4．7\％\％ |
| Watrans／66！5l） | 23 | 40.4 | 9． $7 \% \%$ | b． $0 \times \%$ |
| Ner71／PmV | 34 | 10.2 | 7．1＊＊ | 2．6\％ |
| Amigolinv | 25 | 40.1 | 6． 3 ＊＊ | 1.2 |
| $66 \mathrm{LSO} / \mathrm{PmV}$ | 26 | 40.0 | 8．${ }^{\text {＇，} 2 \times \%}$ | 5．6\％＊ |
| Aura／W．Trams | 27 | 39.9 | 5．8＊＊ | －1．3 |
| Aura／Prav | 27 | 39.9 | 4．9＊＊ | －1．3 |
|  | 29 | 39.0 | 6．1＊＊ | 0.1 |
| Payne／Amisio | 29 | 39.0 | 3．2\％＊ | 0.1 |
| Payne／Salm | 31 | 38.9 | 4． $6 \times \%$ | $3.0 \%$ |
| Amiso | 31 | 38.9 | －．． | －－－ |
| Atr71／66150 | 33 | 38.7 | 2．7＊ | 1.1 |
| W．Trans／ialm | 34 | 38．1 | 6．9＊＊ | 2．5\％ |
| Atr7lflox | 35 | 38.3 | 4．3＊＊ | 0.7 |
| Amisoldr，32\％ | 35 | 38.3 | 1．1ヶ＊） | －0．6 |
| Flex／PmV | 37 | 38.2 | 8．4\％ | 7．4＊＊ |
| 1；322／flex | 38 | 37.9 | 8．1＊＊ | 7．1．4． |
| 66150／Flex | 38 | 37.9 | 5．3＊＊ | 3． $5 * *$ |
| Pityme／flex | 40 | 37.7 | 5．9＊＊ | 4．9\％＊＊ |
| Atril． | 11 | 37.6 | －－． |  |
| Osape／Aura | 42 | 37.5 | －1．1 | $-3.7 \% \%$ |
| 1）Satge／Flex | 43 | 37.1 ． | 3．7＊2＊ | 1.0 |
| Unage／1532\％ | $1 / 4$ | 36.7 | 4．2＊＊ | 0.6 |
| Aura／Payuc | 45 | 36.6 | －0．4 | $-4.6 * *$ |
| Atrol／w．Irams | 45 | 36.6 | 4．3＊＊ | －1．0 |
| Osamge | 47 | 36.1 | ， | ． |
| Salm | 48 | 35.9 | －－－ |  |
| Salm／L5322 | 49 | 3 3．8 | 3．4＊\％ | －0．1 |
| W．Trans／Flex | 50 | 35.1 | 6．2\％ | 4．3＊\％ |
| Paync／PmV | 50 | 35.1 | 4.4 \％ | 2.3 |
| Payne／15322 | ＇2 | 35.0 | 4．2＊＊ | 2.2 |
| 66150 | 53 | 34.4 | －－－ |  |
| Salm／ Priv | 54 | 34.3 | 2.0 | －1．6 |
| ＂sage／Paync | ＇，5 | 34.2 | －0．3 | －1．9 |
| W．Trans／Paync | ＇36 | 33.8 | 3．9\％＊ | 1.0 |
| W．Trans／15322 | 56 | 33.8 | 5．9\％\％ | 4．9＊＊ |
| 15322／PmV | \％8 | 33.1 | 4．3＊＊ | 4． $2 \%$ |
| payne | 59 | 32.8 |  | －－－ |
| Flux | 60 | 30.8 | － | －－ |
| Osage／PmV | 61 | 30.3 | －2．1\％ | －5．8\％＊ |
| W．Trans／PmV | 62 | 29.5 | 1.7 | 0.8 |
| 1 1322 | 63 | 28.9 | －．． | 0.8 |
| PmV | 64 | 28.7 | －－－ | －－－ |
| W．Tratis | 65 | 27.0 | －－－ | －－ |
| Osage／W．Irans | 66 | －－－ | Scedl | Lethal |

[^2] respectively．
＇I＇ヘBIE XVII

PAREN＇IAL AND F1 RANKED MEANS AND HYBRID－PARENT DEVIATIONS FOR GRAIN YIELAD

| Parent or1．llybrid | （rain |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  | Rank | Yicld | Hybrid－Parent Deviat ion |  |
|  |  | （\％／P1ant） | Mid－1are | 11ish－raron！ |
| W．Trans／＇aym． | 1 | $2 ? .1$ | 3．2＊＊ | 「．6\％\％ |
| 0 sater／Payme | \％ | 21.7 | r，． $4 \times \%$ | 4． $7 \%=$ |
| Aura／W．Tram： | 3 | 20.1 | \％． $4 \times \%$ | $6.3 * *$ |
|  | 1 | 20．0 | r） $6 \times$＊ | － 5 ，0） |
|  | ＇， | 19.6 | ＇，＇，\％\％ | 4． $62 \% \%$ |
| P＇Mrı／S：1曲 | ＇） | 19．6 | 1． $12 \%$ | 3.1 |
| ：$\because$／m／Amis！ | 7 | 19.1 | ＇， $1 \%$ \％ | ＇，．4：\％ |
|  | 3 | ． 18.0 | 3．2\％ | 1．＇ |
|  | 9 | 1／8．8 | ， 8.8 ＊ | ： 8 |
|  | 11 | 17：1 | 3．3＊ | $? .1$ |
|  | 10 | 11．1 | 3． $9 \% \%$ | 3．9＊＊ |
| わい1吅／：alm | 1 ！ | 1／1．4 | 5．6．： | 3． $1 \%$ |
| 6－．．ts：／／6，1＇61） | 13 | 11．？ | 1．8\％\％ | $?$ ？ |
| （1：ats：／1／1．x | 13 | 1／．？ | 3．1\％ | ？．2 |
| 人⿻上丨a／Amisas | $1{ }^{\prime}$ | $1 / .1$ | 3．6＊ | 3.3 |
| Ansa／1湲 | 16 | 16：1 | 5． $3 * \%$ | 2.9 |
| Amra／bitur | 17 | －16．6， | 1.5 | 0.1 |
| 1：ats：e／Atr／1 | 18 | 16．＇） | 4.9 ＊＊ | 1．＇s |
| Pityme | 18 | 16．． | －－ | －－． |
| W．Trans／Puv | ？ 0 | 16.4 | 6．${ }^{\text {2 }}$ \％＊ | ＇） 0 \％\％ |
| layme／fley． | 91 | 16．3 | 1． 5 | －0．2 |
| 1）322／F10x | 22 | 16．？ | 5． $1 * *$ | 3.0 |
| Payme／1＇32： | 23 | 16.1 | 3．4＊ | －0．4 |
|  | 24 | 15.1 | 2.6 | －0．8 |
| Salm／PmV | 25 | 15.3 | 4．3＊＊＊ | 1.6 |
| Us：asd | 26 | 15.0 | －－ | －－－ |
| 6，15D／Flex | 26 | 13.0 | 3． $5 \%$ | 1.8 |
| W．Trams／Amisu | 28 | 14.9 | 2.6 | 1.7 |
| Amipo／PmV | 29 | 14.8 | 4．1\％\％ | 1． 6 |
| W．Irans／／i32？ | 30 | 14.6 | $4.4 * *$ | 3.2 |
| Mura／PmV | 30 | 14.6 | 3．6\％ | 0.8 |
| Amisp／rlex | 32 | 1.4 .5 | 1.3 | 1.3 |
| Amiso／l＇32\％ | 33 | 14.3 | 3．2\％ | 1.1 |
| Payne／Pmv | 34 | 14．0 | 1.6 | $-2.5$ |
| Atrll／Amigu | $3 \%$ | 13.9 | 3．2＊ | 0.7 |
| A1r7l／fla\％ | 36 | 13.8 | 3．1＊ | 0.6 |
| 八ura | 36 | 13.8 | －－－ | －－－ |
| ＂alm | 38 | 13.7 | －－－ | －－． |
|  | 39 | 13.6 | 2.1 | 0.4 |
| W． 1 rams／salm | 10 | 13.5 | 1.0 | －0．2 |
| $\cdots \mathrm{alm} / \mathrm{Fl} \cdot \mathrm{x}$ | 10 | 13.5 | 0.1 | －0．2 |
| 1，3\％2／$/ 1 \mathrm{mV}$ | 4.2 | 13.4 | 4．8\％\％ | 4．4\％ |
| flar | 43 | 13.2 | －．． | －．． |
| Amfser | 43 | 13.2 | －．－ | －－－ |
| $1 \cdot 1 \times x /$ Prov | 41 | 12.9 | 2.1 | －0． 3 |
| Aerll／payme | 4, | 12.9 | 0.6 | －3．6\％ |
| 1tr／1／15322． | 41 | 12．5 | 3． $9 * \%$ | 3． $5 \%$ |
| ｜1r71／salm | 188 | 12.4 | 1.5 | －1．3 |
| ：alm／15322 | 48 | 12.4 | 1.1 | －1．3 |
| Sura／flex | ＇） | 11.6 | $-1.9$ | －2．2 |
| W．Trans | 51 | 11.4 | －－ | －－－ |
| Atr71／6，61＇， | ＇3 | 10.8 | 1.8 | 1.0 |
| Atrolld＇mv | ＇3 | 10.1 | 1.9 | 1.8 |
| 66．1＇0）／1＇3） | 53 | 10.1 | 0.7 | 0.3 |
| $\left.6,11^{\circ} \mathrm{j}\right)$ | 5 | 9.8 | －－－ | ． |
| 661 51）／PmV | \％ 6 | 9.2 | 0.2 | －0．6 |
| W．Trames／Flax | 57 | 9.1 | -3.2 \％ | －4．1\％ |
| 15322 | 58 | 9.0 | －－－ | ． |
| W．＇Trans／661＇1） | 59 | 8.5 | －2．1 | －2．9 |
| Usager／PmV | 60 | －8．3 | －3．3\％ | －6．7＊＊ |
| PuV | 60 | 8.3 | －－－ | － |
| Alril | 62 | 8.2 | －－－ | －－ |
| Str7l／Aura | 63 | 6.5 | －4．5＊＊ | －7．3＊＊ |
| Atr71／W．Trans | 64 | 6.3 | －3．5＊ | －5．1＊＊ |
| Aura／66150 | 65 | 4.9 | $-6.9 \% *$ | －8．9\％＊ |
| Us：ase／W．Trans | 66 | －－－ | Seedl | Lethal |

[^3]TABLE,XVIII
PARENTAL AND $\mathrm{F}_{1}$ RANKED MEANS AND HYBRID-PARENT DEVIATIONS FOR PERCENT FERTILITY

| parent or Fi Hybrid | Rank | percont Hercility | Hybrid-Parent Deviation |  |
| :---: | :---: | :---: | :---: | :---: |
|  |  |  | Mid-Parent | igh-Parent |
| W.'Trans/Payor | 1 | 16\%1. 1 | 4 6.3 | 3.3 |
| Payne | 2 | 16, (). 8 | , --2. | -- |
| W.'Trains | 3 | 154.8 | , --- | ~- |
| Aura/Payne | 4 | 1.33 .8 | 7.7 | -7.0 |
| Siam/ PmV | ' | 1.83.6 | - $19.6 \%$ \% | 17.9.9\% |
| W. Trans / Pruv | 6 | $1.52: 4$ | - 8.9 | -2.4 |
| l'aynes/Salm | 7 | 149.6 | $=1.4$ | -11.2 |
| Aura/P'mV | 8 | 147.2 | $\therefore 15.4 \% \%$ | 14.9\% |
| Osage/Aura | 9 | $14{ }^{\text {t }} 5.9$ | - -8.9 | 3.3 |
| Osage | 10 | 142.6 | $\vdots$ | --- |
| Aura/W.Trans | 11. | 140.6 | -2.5 | -14.2** |
| Payne/Amigio | 12 | 137.6 | 1-8.5 | $-23.2 * *$ |
| osage/Paync | 13 | 135.7 | ! $-16.0 \times *$ | -25.1** |
| salm | 13 | 135.7 | - | --- |
| Aura/Salm | 1.5 | 134.2 | : 0.7 | -1.5 |
| Nura/Amigo | 16 | 132.6 | ; 1.2 | 1.1 |
| l'uv | 17 | 132.3 | ; --- | --- |
| l'ayne/ 15322 | 18 | 132.1 | - -6.6 | -28.7 ** |
| Amigo | 19 | 17.', | : | - --- |
| Nura | 20 | 131.4 | $i$ | -- |
| Osage/Amigo | 21 | 130.8 | - -6.2 | -11.8 |
| Osage/salm | 22 | 130.7 | - -8.4 | -11.9* |
| $15322 / \mathrm{PmV}$ | 23 | 130.6 | + 6.1 | -1.7 |
| Payne/PmV | 24 | 127.5 | -19.0** | -33.3** |
| Osage/Flex | 25 | 126.9 | - -6.5 | -15.7\% \% |
| Silm/Amigo | 25 | 126.9 | - -6.7 | -8.8 |
| W.Trans/15322 | 27 | $\because \quad 126.2$ | : -9.6 | $-28.6 * *$ |
| Flex | 28. | 124.3 | $\cdots$ | --- |
| Aura / 15322 | 29 | 122.9 | -1.1 | -8.5 |
| Osage / PmV | 30 | 121.7 | -15.7** | -20.9** |
| 15,322 | 31 | 1116.7 | --- | --- |
| Ainf go/ PmV | 32 | $\therefore 140.2$ | -15.7\% | -16.1\% |
| Amigo/ 15322 | 33 | $\cdots 113.2$ | -10.9* | -18.3** |
| 06150 | 34 | 111.7 | $\therefore$--- | --- |
| 60150/Flex | 35 | 109.8 | -8 2 | -14.5** |
| Payne/Flex | 36 | 109.0 | -33.5** | -51.8** |
| Ogage/L5322 | 37 | 108.3 | -8.4 | -34.3** |
| Salm/15322 | 38 | -107: | -18.4** | -27.9** |
| Osage/Atr71 | 39 | 107.6 | -12.1* | -35.0\%* |
| Osage/66151) | $41)$ | 195.2 | -21.9** | -37.4** |
| W. Trans/Amigo | 41 | 104.4 | -38.7** | -50.4** |
| 15322/Flex | 42 | 103.9 | -1.6.6** | -20.4\% \% |
| Atr71/Flex | 43 | 101. 6 | -9.0. | -22.7** |
| Atr71/66150 | 44 | 100.4 | -3.9 | $-11.3$ |
| Acr71 | 4.5 | 96.9 | --- | --. |
| Plex/PmV | 46 | 95.1 | -33.2\% $\%$ | -37.2** |
| : $66150 /$ mixo | 47 | .94 .1 | -27.5\%\%. | -37.4\%* |
| Atrel / PmV | 48 | 92.2 | -22.4** | -40 1\% |
| \| Amigo/Flex | 49 | 90.7 | -37.2** | -40.8** |
| Ntr7!/Payne | 150 | 89.0 | -39.8** | -71.8** |
| 66150/Payme | 51 | 88.8 | -47.4** | -72.0x\% |
| W.Trans/sillm | 52 | - 88.7 | -56.6** | -66.1** |
| Atr71/Amigo | 53. | -87.7 | -26.5** | -43.8\%* |
| Sa/m/Flex | 14. | 87.1 | -42.9** | -48.6** |
| 66,50/Salm | 55 | 83.7 | -40.0**. | $-52.0 * * *$ |
| 661.50/PmV | 56 | 83.4 | -38.6*\% | -48.9:\% |
| Atr71/15322 | 57 | 83.2 | -23.6** | -33.5\%* |
| Atr71/Salm | 58 | 70.1 . | -46.2** | -65.6\%\% |
| W.'Trans/66150 | 59 | 68.8 | -64.4** | -86.0\%* |
| 66150/15322 | 60 | 66.5 | $-47.7 * *$ | -50.2\% \% |
| W.Trans/Flex | 61 | 58.7 | -80.8** | -96.1\%\% |
| Aura/flex | 62 | 58.3 | -69.5** | -73.1** |
| Atr71/Aura | 63 | 41.6 | -72.5** | -89.8** |
| Atr71/W.Trans | 64 | 41.3 | -84.5\%\% | -113. $5 \% \times$ |
| Aura/66150 | 65 | 36.9 | -84.6\%\% | -94.5*\% |
| Osage/W.Trans | 66 | , | Soedli | Lethal |

TABLE XIX
PARENTAL AND F1 RANKED MEANS AND HYBRID-PARENT DEVIATIONS FOR PLANT HEIGHT

*,** Significant at the 0.05 and 0.01 levels of probability, respectively.

TABLA XX

## PARENTAL AND $\mathrm{F}_{1}$ RANKED MEANS AND HYBRID-PARENT DEVIATIONS FOR HEADING DATE


${ }^{1}$ Number of day aftar March 31.
*, ** Significant at the 0.05 and 0.01 levels of probability, respectively.

TABLE XXI
PARENTAL AND $\mathrm{F}_{1}$ RANKED MEANS AND HYBRID-PARENT DEVIATIONS FOR PERCENT PROTEIN


[^4]
## VITA $^{2}$

Paul Gregory Sebesta<br>Candidate for the Degree of<br>Doctor of Philosophy

## Thesis: AN ANALYSIS OF YIELD AND OTHER TRAITS IN A DIALLEL CROSS INVOLVING ELEVEN WINTER WHEAT ALIEN-TRANSLOCATION LINES

Major Field: Crop Science

Biographical:
Persona1 Data: Born in Ithaca, New York, October 13, 1953, the son of Dr. and Mrs. Emil E. Sebesta. Married Deborah Nan King on July 19, 1980.

Education: Graduated from C. E. Donart High School, Stillwater, Oklahoma, in 1971; received the Bachelor of Science degree from Oklahoma State University in May, 1975, with a major in Agronomy; received the Master of Science degree in Agronomy from Oklahoma State University in May, 1977; and completed the requirements for the Doctor of Philosophy degree in Crop Science at Oklahoma State University in December, 1980.

Professional Experience: Student Assistant, Department of Agronomy, Oklahoma State University from September, 1971 to May, 1975; Graduate Research Assistant, Department of Agronomy, Oklahoma State University from May, 1975 to September, 1980.

Professiona1 Organizations: American Society of Agronomy, Crop Science Society of America, Sigma Xi, and Genetics Society of Canada.


[^0]:    *,** Significant at the 0.05 and 0.01 levels of probability, respectively.

[^1]:    ＊，＊＊Significant at the 0．0＇s and 0.01 levels of probability，

[^2]:    $\%$ ，Significant at the 0.05 and 0.01 levels of probability，

[^3]:    $\therefore$ ， $\operatorname{six}$ Sinificant at the 0．0r and 0．01 levols of probability，
    ruspertively．

[^4]:    *,** Signifleant at the 0.05 and 0.01 levels of probability, respectively.

