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Abstract:

ATRT (Atypical Teratoid Rhabdoid Tumor) is an aggressive pediatric brain tumor. Scientists believe that INI-1's loss from the SWI/SNF complex is what causes the ATRT tumor to develop. However, I hypothesized that rather than the loss of INI-1 from the SWI/SNF complex, it is the loss of INI-1 from some other unidentified INI-1 associated protein complex that causes the tumor to develop. To begin to answer this question, I set out to identify proteins that associated with INI-1. Multiple techniques were used over the duration of this project. Cell culture techniques were employed to grow two HeLa cell lines, one line had a FLAG tag version of INI-1 and the other HeLa cell line was used a control. These cells were grown and then lysed. Then half of the protein sample was run on a SDS-PAGE gel and the other half of the sample was sent to mass spectrum for analysis. Protein lists were then compiled and compared for different samples to determine the significance of the proteins attached to INI-1.

Project Title:

INI-1 and All of Its Friends: The Analysis of the INI-1 Protein and Associated Proteins That Contribute to Cancer Formation.

Goal:

Identify INI-1 associated proteins and determine their role in tumor formation.

Introduction:

Cancer is a devastating, progressive disease that affects millions of people each year. One type of cancer is a pediatric atypical teratoid rhabdoid tumor (ATRT). This cancer's victims are usually less than 3 years old and undergo a variety of treatments, such as chemotherapy, radiation, and surgery, to combat the very progressive malignant tumor [2]. ATRT is very aggressive, and, unfortunately, extremely difficult to diagnosis. Some of these symptoms are early morning headaches, vomiting, and lethargy [2]. Unfortunately, these symptoms can be associated with a variety of medical issues. Consequently, this increases the difficulty of diagnosis since multiple variables need to be ruled out in order to come to the conclusion of the possibility of a tumor. Not much is known about the direct cause of the ATRT tumor, but there is a known correlation between the INI-1 protein and the tumor's development. The INI-1 protein is a very elusive protein that scientists know exists, but they do not know the function of it. Consequently, they cannot determine its precise role in the development of the pediatric tumor.

Although it is known that INI-1 is part of the SWI/SNF complex, which is comprised of about ten proteins that unravel DNA, INI-1's specific role in the SWI/SNF complex has not been found to directly correlate to tumor formation. The current theory states that it is the loss of INI-1 from the SWI/SNF complex leads to the cancer development. However, what if it is not the loss

of INI-1 from the SWI/SNF complex that is causing the cancer, but rather what if the cancer is caused by the unknown proteins that INI-1 is also occasionally attached to?

Figure 1

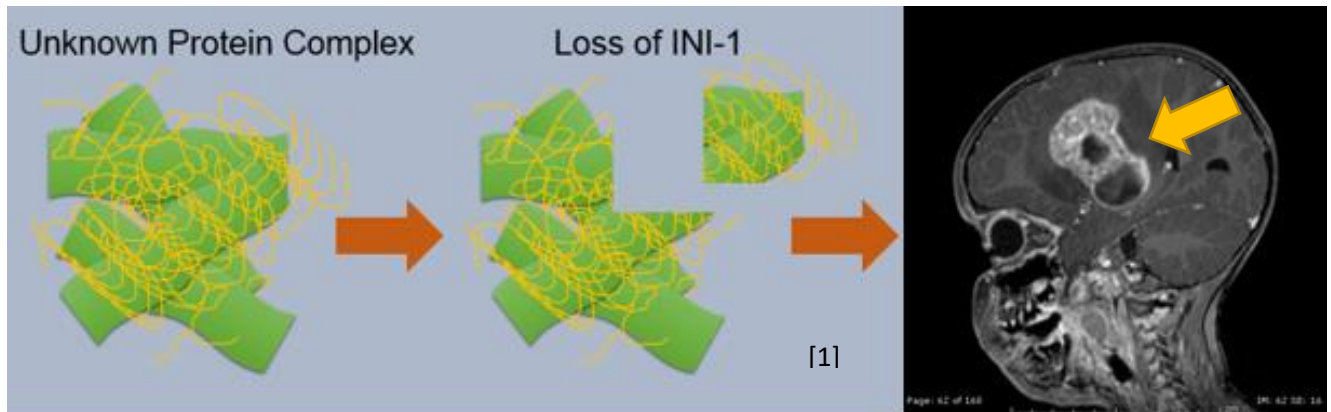


Figure 1: This is a visual that illustrates my hypothesis. The loss of the INI-1 from some unknown protein complex is what causes ATRT to develop.

Experimentation:

My hypothesis is that the loss of INI-1 leads to the malfunction of an unknown protein complex, thus leading to tumor formation. To test the hypothesis, I will identify interacting INI-1 proteins.

Cell Culture

Experimentation began with the use of sterile techniques and methods for growing a HeLa culture, which is a cell culture of a line of cancer cells. Both a control cell line of HeLa cells and a HeLa cell line containing a FLAG tagged version of INI-1 that allows INI-1 to be isolated by column chromatography were grown. HeLa cells were used because they are easily grown and are

not extremely prone to contamination. I learned a wide variety of sterile cell culture techniques in this stage of the experiment.

The cells grew quite well most of the time. The incubator did malfunction over the course of my production and carbon dioxide levels fell below that necessary for HeLa cell growth. Also, one instance of contamination did occur. This contamination was isolated to one plate. This plate was immediately disposed of and sterilized. After the incubator failed, it took a month to grow the cells back up to the confluency needed to begin to split them and save them at a rapid rate. However, by the end of the year, I had developed quite a stock of large cell plates for both the tagged and control versions of the HeLa cells. The picture below is one of a plate of HeLa cells that contained the FLAG tag version of INI-1.

Figure 2

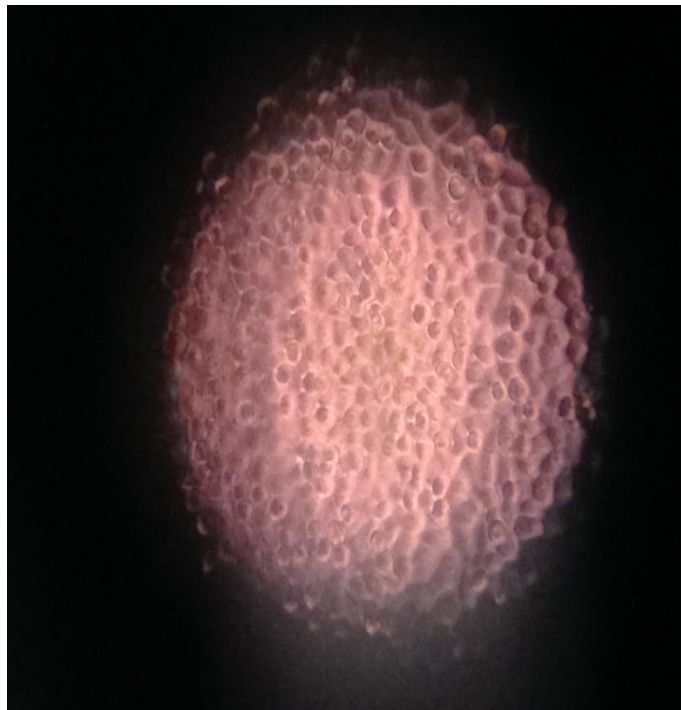


Figure 2: This is a picture of the FLAG tag INI-1 HeLa cells growing on a plate. This was viewed using a light microscope.

My original lines were grown from a cell sample that had been placed in a cryovial and frozen at -80°C . The exact date of freezing of the frozen samples was unknown; however, it is safe to conclude that they had been frozen for many years. Consequently, it was difficult to get them to grow effectively after being frozen for so long. However, eventually the cells began to grow.

In order to properly grow the cells, one must be sure to make everything as sterile as possible. All work must be done in a cell culture room in a hood with proper air circulation as well as proper UV light to sterilize the room when finished. Cellular media was made 3 days prior to attempting to grow the cells. All of the components of the media were filtered in order to prevent contamination from occurring. This media also contained Fetal Bovine Serum, which was not filtered to make sure the nutrients necessary for cellular growth did not get removed.

The pH of the media was at a physiological pH of 7.4. This was necessary because the cells used were human cells and the human body is at a pH of 7.4. Thus, this pH optimizes growth of the cells. After the media was created, it was plated and left in the incubator for 3 days, after which the plate was removed from the incubator and examined under a microscope to look for contamination. If the media was contaminated, it was not used on the cells. Fortunately, there was never a case of contaminated media.

In order to actually begin growing cells, the media needs to be room temperature. Then the cells start out growing on a small plate. Ten milliliters of media was plated and then the cryovial was warmed until the contents were completely thawed. After this, the cryovial was dumped onto the small media containing plate and the lid was put on top. A figure-eight motion was made with the plate in order to make sure that the cells spread out. Then the plate was properly labeled and placed in an incubator at a temperature of 37°C . The plates were checked every other day to make sure that growth was normal and no contamination had occurred.

After the cells had reached 85-100% confluency, the cells were removed from the small plates and placed on larger plates to grow. One can tell when the cells are ready to be split by the fact that they begin to stack up on top of each other. The stacking prevents cells from truly being able to grow to their fullest potential, because the cells on top of the stacking blob essentially smother the cells underneath. Cells were transferred from the plate by adding Trypsin. Trypsin cleaves the cells from the plates so that they can be pipetted up and placed onto a new plate. For the larger plate 30 milliliters of media was used to grow the cells. Multiple figure eights were used to make sure that the cells were evenly spread around so that they are able to grow most efficiently.

Unfortunately, my supply of INI-1 Flag tagged and control cell cryovials were limited. To combat the total loss of all stored cryovials, I froze some cryovials of the HeLa cell lines I grew for later use. I did this because it is easier to grow samples that have not been frozen for a long time rather than ones that have been frozen for years. I wanted to create a security that I could look to in case my cell plates that were growing had spontaneous death or if I wanted to resume cell culture again after stopping for any reason in the future. The cryovials that I froze are going to be used by another member of the laboratory who will continue this project.

After the cultures had grown to approximately 85-100% confluency, the cells were scraped off of the plate. This was done by first washing the cells with PBS. After the washing was completed, 1mL of PBS was added to the plate to make sure that there was enough liquid to scrap the cells off the plate. The actual scraping of the cells was done with a Rubber Policeman. When scrapping cells, it is very important to not be too rough with them. The cells could possibly burst open if they are pressed too hard. After the cells were scrapped off of the plate, the cells were centrifuged and stored in the -80°C freezer until I had enough built up pelleted cells that I could begin experimentation.

After I had an ample supply of frozen cell pellets, the cells were then lysed or broken open and the cell extract from the cells were collected. Column chromatography techniques were employed with a resin that is only capable of removing INI-1 and its associated proteins (anti-FLAG resin) to isolate the INI-1 protein from the HeLa cell extract. The resin was washed with the same buffer that was used in lysing open the cells. This was done to make sure that the resin was compatible with the proteins and able to induce protein-resin binding. A HeLa cell line that did not contain the tagged INI-1 was used as a control. The control line was grown using the same methods as the tagged version of INI-1.

Figure 3

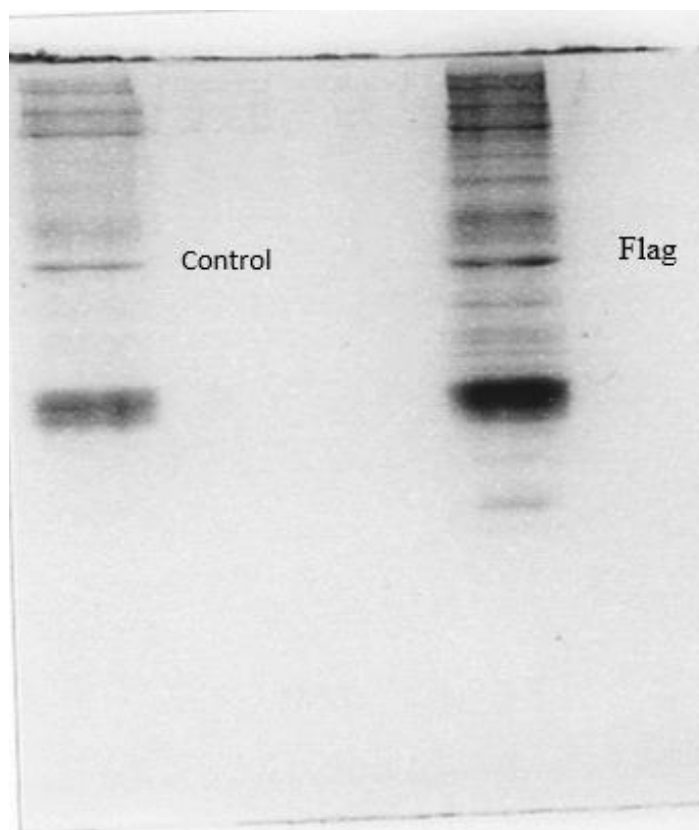


Figure 3: This is an SDS-PAGE gel that indicates the different protein banding patterns of the two lanes. The banding pattern for both the FLAG tag INI-1 Lane and the Control Lane was the same. This is a 12% gel.

An SDS-PAGE gel was used to analyze the proteins. This is because SDS-PAGE gels can separate proteins based on size. A high molecular weight ladder was chosen to be the comparison for protein weights from a high molecular ladder. When analyzing the gel above, it is evident that the protein banding pattern is the same in the control lane and the FLAG lane. This protein banding pattern was not expected to be the same for the flagged and control lines of HeLa cells. This was not expected because one of the lines has the FLAG tag version of INI-1 that would attach to the anti-FLAG resin. Therefore there should have been more proteins in the FLAG lane than the Control lane. This unexpected result was quite puzzling.

Control Tests

Consequently, different scenarios were proposed to explain the phenomenon. Were the gel techniques poor? Were the cells actually lysed open? Many more questions were asked. In order to answer these questions, control experiments were run to determine the cause of the unexpected banding patterns in the cell lines. For instance, in one control experiment, the resin was washed more than the previous gel to see if the resin was able to bind INI-1 more efficiently if the resin was immersed in the buffer prior to introduction to the cellular extracts. This method, however, only caused a slight protein binding increase. Therefore, it was concluded that the resin was not the reason why the two cell line proteins looked the same on the gels.

Another control test was conducted in order to see to see if the SDS-PAGE gel was being correctly run with good technique. (An SDS-PAGE gel is simply a gel that separate proteins to distinguish between different cell lines.) The control sample was a sample of INI-1 protein and those associated with it. This sample was from a frozen immunoprecipitation experiment

conducted by Dr. Ruhl several years before. The exact date of the creation of this nuclear extract sample was not known. The buffer was also run on the SDS-PAGE gel to determine if the protein bands on the gel were simply from the buffer being used or if they were actually from the cell extracts. The results from this control run indicated that the buffer did not contain any of the proteins that were associated with INI-1. The only proteins evidenced in the buffer lane were the antibody chains. This was expected due to the fact that resin was added to the buffer. The resin is the part of the combination of resin and sample that actually contains the antibody.

Figure 4

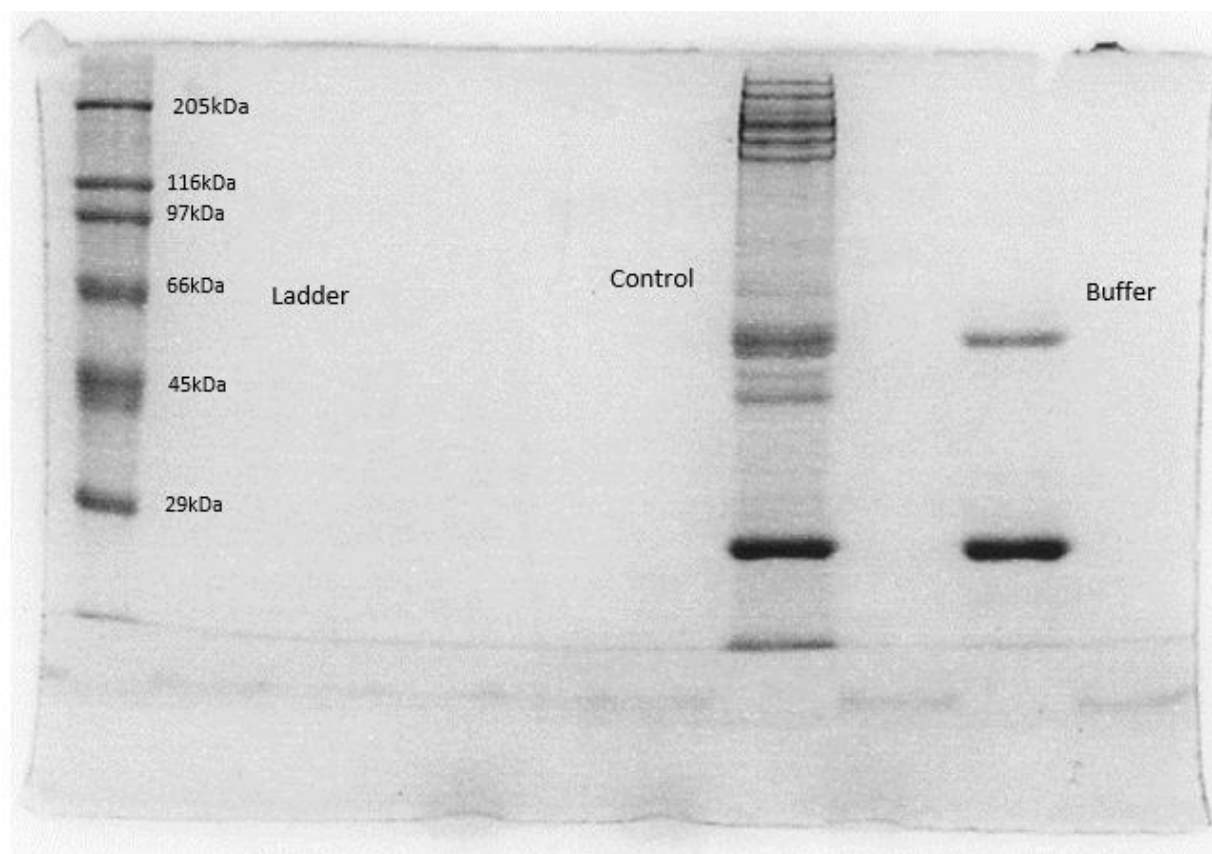


Figure 4: This is a SDS-PAGE gel that indicates that the gel techniques used were effective. This is an example of a type of control tests conducted to see if cell lysis was effective. This is an 8% gel.

This gel also proved that I did have good technique in my gel electrophoresis and that the two darker bands in the lanes were actually from the buffer not from the cell extract. This still did not answer the question of why the banding patterns were the same in the original SDS-PAGE gel. Eventually, the conclusion was made that the cell was not being lysed completely. Since INI-1 is found in the SWI/SNF complex and that is a complex that helps cells read DNA, INI-1 is expected to be inside of the nucleus of the cell. This is because DNA is actually found inside of the nucleus of the cell and is replicated inside the nucleus as well. Since INI-1 is not being removed from the cell, the conclusion was made that the nucleus of the cell was not being lysed, only the cell wall was being broken open. This explains why the gel (Figure 3) at the beginning of this page had similar protein banding patterns. The only difference between the two different cell lines is really the flagging of the INI-1 protein inside of the nucleus.

Consequently, a majority of project was trying to figure out a way to break the nucleus open without damaging the protein. This was a puzzling problem in that the proteins are quite sensitive and destroying them is very possible if too harsh of techniques are being employed. Mechanical, chemical, and freeze-thaw lysis techniques were all utilized; however, none of the techniques were very effective. Some would consider conducting a nuclear extract of the cell since a nuclear extract is only all of the proteins from inside the nucleus and nothing else. However, it is not known whether or not INI-1 interacts with proteins that are located outside of the nucleus. Consequently, this eliminates the nuclear extracts as a viable option since it could eliminate possible INI-1 to other protein interaction.

The Analytical Biochemistry Laboratory class graciously served as a method testing unit for testing different methods of breaking open the cells to extract protein. All of the cell pellets they used for their method testing were pellets previously saved over the course of my experiment.

The pellet sizes were all relatively the same. This was done to make sure that there was not a lane that had a significantly darker banding pattern for one group. The laboratory class used a mechanical lysis with a dounce. This method yielded excellent results that were able to be sent for proteomics. Each laboratory group conducted experiments on two sets of cells. One sample was the proteins from the cell with the FLAG tag version of INI-1 and the other sample was the control cells. The lanes in the graph are in pairs. The first lane is the FLAG tag and the second lane is the control. As an example, lane 2 and 3 is from one laboratory group. Lane 2 is the FLAG tag sample and lane 3 is the control sample. Lane 4 and 5 are from the next laboratory group. This pattern continues all the way to the final group with the lanes 22 and 23. Lane 1 is the High Molecular Weight Ladder.

Figure 5

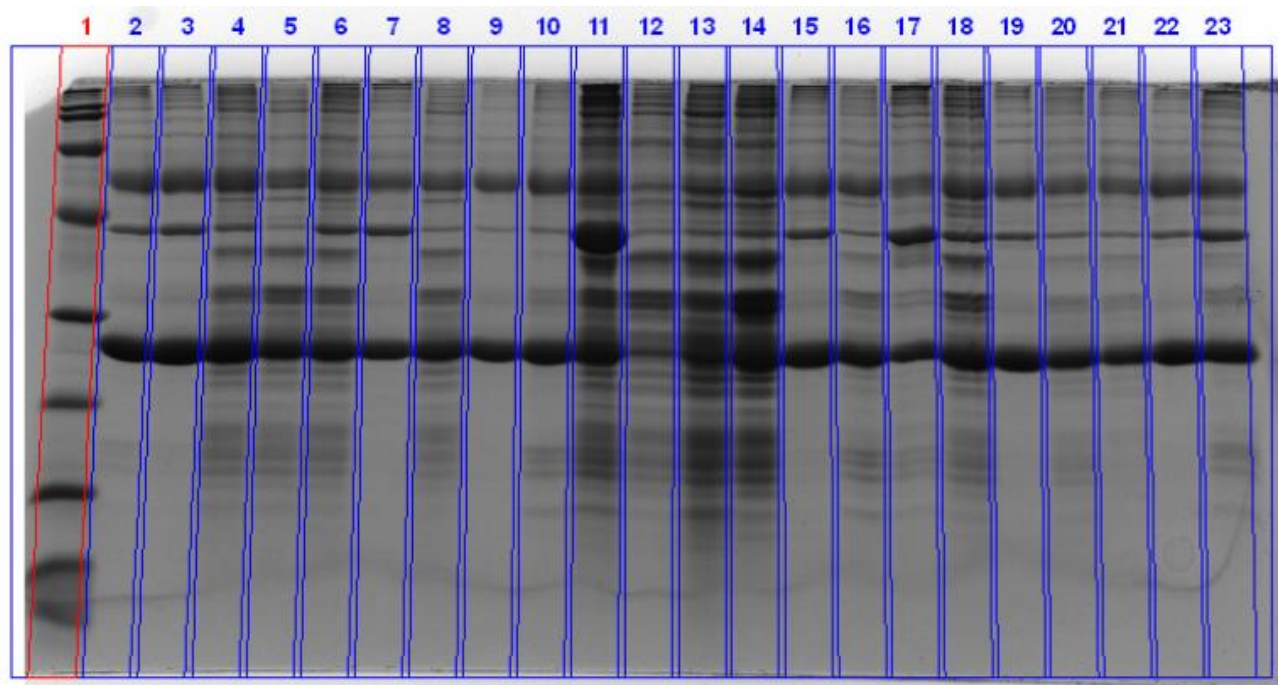


Figure 5: SDS-PAGE gel created by the Analytical Biochemistry Lab course. Lane 1 is the High Molecular Ladder. The lanes chosen to continue onward to Mass Spectrum Analysis were Lanes 2 (FLAG tag INI-1) and Lane 3 (Control). Lanes 6 (FLAG tag INI-1) and Lane 7 (Control) were also chosen. This is a 12% gel.

After the chromatography was completed, both the tagged and non-tagged samples were sent off for mass spectrum analysis. The tagged and non-tagged samples were the samples from lanes 2,3 and lanes 6,7. These two groups of lanes were chosen due to the fact that they seemed to have the most expected banding pattern. By this it is meant that that the INI-1 Flag tag lane visually looks different than the Control Lane. There are different banding patterns. However, these lanes are not so different that it indicates suspicion in the group's methods.

Proteomics

This statistical analysis provided a set of other proteins associated with INI-1. This statistical analysis was completed by sending the mass spectrum samples saved from the laboratory's experiment to mass spectrum for analysis. After the analysis was complete a list of proteins was sent back. This protein list was read in the computer program Scaffold 4. Scaffold 4 can organize proteomic data and look for significance. After comparing both the tagged and non-tagged lines using the "Total Spectrum Count" setting, it was determined which proteins are only associated with INI-1. (The "Total Spectrum Count setting was used to distinguish all of the differences between the proteins when analyzing the proteomic data for significance.) This was done by evaluating the p-value of the proteins. If the p-value of the protein was less than 0.05, then the protein was deemed to significantly associate with INI-1. Since INI-1 attaches to the SWI/SNF complex, it was expected that some of the identified proteins would be members of the SWI/SNF complex. (There were 5 proteins that are identified with the SWI/SNF complex that were significantly associated with INI-1 in both the 2,3 and 6,7 trials.) It would have been interesting to see the other Analytical Biochemistry Laboratory group's proteomic data results. However,

running the mass spectrum requires money and time. The two items limited the number of samples that could be sent at once.

The submission of only two trials of this experiment, however, are not enough trials for me to conclusively determine whether or not the proteins are actually associated with INI-1. More trials must be implemented to conclusively determine which proteins are associated with INI-1. This is important due to the fact that human error is always present. More experimental trials need to be conducted so that this human error would be less significant. For instance, the protein keratin was found in both cell line protein lists. This keratin is not from the cells. Rather, the keratin is from human skin cells. Granted this protein was marked off significance list due to the lack of a significant p-value. There is still the possibility for other human error like possible protein contamination. It is very unlikely that this happened; however, more trials would really strengthen the findings of the experiment. Consequently, more trials were conducted in order to determine the significance of the experimental finding.

More Trials Conducted

More cells were broken open over the course of the semester; however, there, once again, were difficulties in breaking open the cells. This problem in the methodology was unexpected. The Analytical Biochemistry Laboratory used the protocol and it worked well. For reasons unknown, their protocol was not achieving the results desired when replicated. Multiple different steps were taken to try to replicate the results obtained by the Analytical Biochemistry Laboratory. Unfortunately, this proved to be challenging.

Figure 6

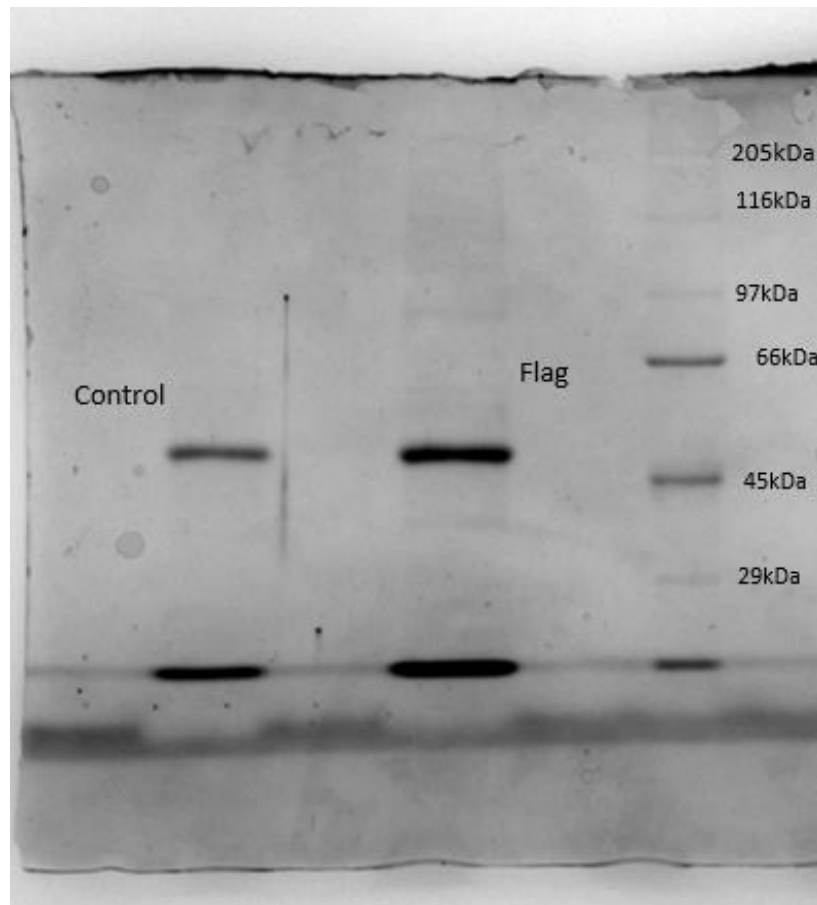


Figure 6: This is an SDS-PAGE gel Kristy Johnson and I prepared that illustrates the poor results obtained when we used the same cell lysis methods utilized by the Analytical Biochemistry Lab course. This is an 8% gel.

The previous douncing method was not as effective as it was for the laboratory. There also were more washing methods implemented in the new trails, and the addition of more washing could have possibly caused protein to become lost. This is evidenced by the gel above. The banding pattern was the same between the lane containing the Flag tag version of INI-1 and the control HeLa cells. The antibody resin does show up on the gel. The light chain is the darker band on the gel below the 29 kDa mark on the ladder. The light chain should be at about the 27 kDa mark. The heavy chain is at about the 50 kDa mark on the gel. This band, however, is about where it should be on the gel. This indicates that the lighter proteins moved through the gel faster than the ladder.

After multiple trials, it was determined that the method of breaking open cells should be the freeze/thaw method. This method was very effective and 2 more sample pairs were run on a gel.

Figure 7

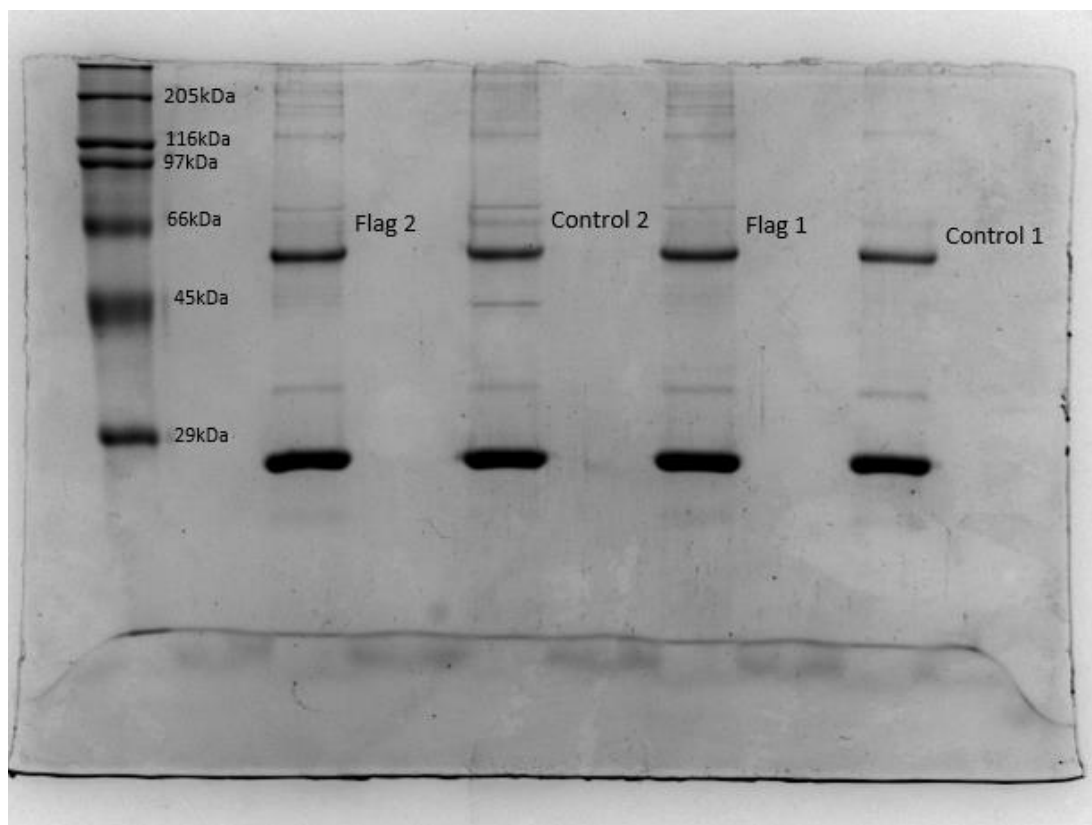


Figure 7: This SDS-PAGE gel illustrates that the cells were effectively lysed with the freeze/thaw method. This is a 12 % gel.

This gel contains all the markers expected for a good gel. Both the light chain and the heavy chain from the antibody are present on the gel. The INI-1 band is also located on the gel at about the 47kDa mark. It is very light in the picture; however, when viewing the gel directly with light, the band was much more evident. There are obvious differences between the lanes of the FLAG tag version of the INI-1 protein and the control lane. These different bands can be viewed between the flag and the control between the 116kDa mark and the 205 kDa mark. This indicates that there are differences in protein for the two samples. These samples have not been sent to mass spectrum

for analysis, yet. However, the samples are saved at -80°C with little to no buffer in the vials. They are ready to be sent off to mass spectrum for analysis whenever the laboratory is ready to send more samples.

Future Studies:

More immunoprecipitation experiments will take place to provide more quality samples to be sent off to mass spectrum for analysis. This will provide more N-values so that it can be better confirmed what proteins associate with INI-1 consistently. Also, the provision of more experimental runs will better enable a consistent experimental technique to be obtained for breaking open the HeLa cells. The mass spectrum results will then be confirmed by using coimmunoprecipitation to pull INI-1 out of the HeLa cell and using an IP-Western blot. Using this technique, specific proteins like INI-1 can be identified. This process will allow me to reconfirm the exact protein that attaches to INI-1.

Results and Conclusion:

The results from the experiment were that I identified proteins associated with INI-1 from the mass spectrum and proteomic analysis of the lanes 2,3 and 6,7 from the Analytical Biochemistry Laboratory course. (A list of proteins that associated with INI-1 protein exclusively is provided in Appendix 1. The descriptions of the proteins found significant in both 2,3 and 6,7 can be found in Appendix 2.) In results of the 6,7 lanes, 312 proteins were identified. However, only 212 of the proteins significantly associated with INI-1. Significance was determined by those proteins whose p-value was under 0.05. In the results of the 2,3 lanes, 162 proteins were identified. However, only 93 of these proteins were significantly associated with INI-1. The fact that not all proteins that were found were significant is an important reminder of why the control sample is

important. Without the control, it would be easy to assume that all the proteins that were pulled out of the cell were attached to INI-1. This is simply not the case. Sometimes non-specific proteins attach to resins. The use of the control allows for the elimination of these non-specific proteins and helps strengthen the conclusions of this experiment.

Fifty-four proteins were found to have a significant p-value in both the 2,3 lanes and the 6,7 lanes. The common theme between a lot of the proteins is that they are involved in some form with cytokinesis. This is an interesting observation that needs to be explored in more depth in order to fully understand the INI-1 proteins role in the development of ATRT. Some other proteins were found to be crucial in maintaining a proper cytoskeletal structure. This is also an interesting observation in that INI-1 may be contributing to the structural integrity of the cell, and, possibly, when that integrity falters ATRT develops. One of the significant proteins identified in both of the trials that went to mass spectrum for analysis was Myosin. This protein plays a major role in cellular development, specifically cytokinesis. Myosin is also important in maintaining a proper cellular shape and structure [3]. Another protein identified was Isoform 2 of Filamin A. This protein is also involved in cytoskeleton maintenance. It binds transmembrane protein to the cytoskeleton and is crucial to the signaling mechanisms of the cell [4]. These are just two proteins from the list of 54 proteins that were significant in both the proteomics results from the 6,7 lanes as well as the 2,3 lanes. However, these are the two proteins I am most interested in exploring their role with INI-1 in the development of ATRT.

The finding of all of these non-SWI/SNF proteins associating with INI-1 was initially expected because INI-1's role in tumor formation is not directly known to be associated with the SWI/SNF complex. It is simply the current theory. INI-1 may have a variety of different roles in the cell that are not currently known. The identification of these new proteins could help scientists

understand the correlation and causation INI-1 has with the development of the pediatric tumor ATRT. Could this newly discovered protein be the key to unlocking the cause of the devastating pediatric cancer? Could the analyses of this new protein have the potential to save lives? These are questions that must be answered, and they will be answered in future studies. More research time and money is necessary to determine the extent to which these newly associated proteins could be significant in finding a treatment or even a cure for ATRT.

Bibliography:

1. Dixon, A., MD. (n.d.). Atypical teratoid rhabdoid tumour (ATRT) | Radiology Case | Radiopaedia.org. Retrieved April 28, 2016, from <http://radiopaedia.org/cases/atypical-teratoid-rhabdoid-tumour-atrt>
2. National Cancer Institute. Childhood Atypical Teratoid/Rhabdoid Tumor Treatment. (n.d.). Retrieved April 28, 2016, from <http://www.cancer.gov/types/brain/hp/child-cns-atrt-treatment-pdq>
3. Uniprot. (n.d.). Myosin-9. Retrieved May 05, 2016, from <http://www.uniprot.org/uniprot/P35579>
4. Uniprot. (n.d.). Filamin-A. Retrieved May 05, 2016, from <http://www.uniprot.org/uniprot/P21333>
5. National Center for Biotechnology Information. (n.d.). Retrieved May 08, 2016, from <http://www.ncbi.nlm.nih.gov/>
6. UniProt. (n.d.). Retrieved May 08, 2016, from <http://www.uniprot.org/>
7. Human Gene GSN (uc004blf.1) Description and Page Index. (n.d.). Retrieved May 08, 2016, from <http://moma.ki.au.dk/genome-mirror/cgi-bin/hgGene?org=Human>

Appendix 1:List of proteins for 6,7 and 2,3.

A significant p-value is highlighted in green.

#	Identified Proteins (312)	Accession Number	Molecular Weight	Protein Grouping Ambiguity	Lanes 6,7 P-values	Lanes 2,3 P-values
1	Myosin-9 OS=Homo sapiens GN=MYH9 PE=1 SV=4 Actin, cytoplasmic 2 OS=Homo sapiens	sp P35579 MYH9_HUMAN	227 kDa	TRUE	<0.00010	<0.00010
2	GN=ACTG1 PE=1 SV=1 Isoform 2 of Filamin-A OS=Homo sapiens	sp P63261 ACTG_HUMAN	42 kDa	TRUE	0.0024	0.09
3	GN=FLNA Spectrin alpha chain, non-erythrocytic 1 OS=Homo sapiens	sp P21333-2 FLNA_HUMAN (+2)	280 kDa	TRUE	0.00014	0.02
4	GN=SPTAN1 PE=1 SV=3 Spectrin beta chain, non-erythrocytic 1 OS=Homo sapiens	sp Q13813 SPTN1_HUMAN	285 kDa		0.00027	0.00012
5	GN=SPTBN1 PE=1 SV=2 RecName: Full=Keratin, type II cytoskeletal 1; AltName: Full=67 kDa cytokeratin; AltName: Full=Cytokeratin-1; Short=CK-1; AltName: Full=Hair alpha protein; AltName: Full=Keratin-1; Short=K1; AltName: Full=Type-II keratin Kb1	sp Q01082 SPTB2_HUMAN	275 kDa	TRUE	<0.00010	<0.00010
6	RecName: Full=Keratin, type II cytoskeletal 2 epidermal; AltName: Full=Cytokeratin-2e; Short=CK-2e; AltName: Full=Epithelial keratin-2e; AltName: Full=Keratin-2 epidermis; AltName: Full=Keratin-2e;	gi 238054406 sp P04264.6 K2C1_HUMAN (+1)	66 kDa	TRUE	0.00019	0.017
7		gi 239938650 sp P35908.2 K22E_HUMAN (+1)	65 kDa	TRUE	0.00095	0.0032

Short=K2e; AltName:
Full=Type-II keratin Kb2

	LIM domain only protein 7 OS=Homo sapiens					
8	GN=LMO7 PE=2 SV=1 RecName: Full=Keratin, type I cytoskeletal 10; AltName: Full=Cytokeratin-10; Short=CK-10; AltName: Full=Keratin-10;	tr E9PMS6 E9PMS6_HUMAN	145 kDa	TRUE	0.0013	0.00016
9	Short=K10 Keratin, type I cytoskeletal 17 OS=Homo sapiens	gi 269849769 sp P13645.6 K1C10_HUMAN (+1)	59 kDa	TRUE	0.018	0.076
10	GN=KRT17 PE=1 SV=2 Isoform 8 of Filamin-B OS=Homo sapiens	sp Q04695 K1C17_HUMAN	48 kDa	TRUE	0.058	0.34
11	GN=FLNB Keratin, type I cytoskeletal 18 OS=Homo sapiens	sp O75369-8 FLNB_HUMAN	282 kDa	TRUE	<0.00010	
12	GN=KRT18 PE=1 SV=2 Histone H1.2 OS=Homo sapiens GN=HIST1H1C	sp P05783 K1C18_HUMAN	48 kDa	TRUE	0.0043	0.011
13	PE=1 SV=2 RecName: Full=Keratin, type I cytoskeletal 9; AltName: Full=Cytokeratin-9; Short=CK-9; AltName: Full=Keratin-9; Short=K9	sp P16403 H12_HUMAN	21 kDa	TRUE	0.0012	0.49
14	Antigen KI-67 OS=Homo sapiens GN=MKI67 PE=1 SV=2	gi 239938886 sp P35527.3 K1C9_HUMAN (+1)	62 kDa	TRUE	0.043	0.19
15	Isoform 2 of Plectin OS=Homo sapiens	sp P46013 KI67_HUMAN	359 kDa		0.00044	
16	GN=PLEC 40S ribosomal protein S4, X isoform OS=Homo sapiens GN=RPS4X PE=1 SV=2	sp Q15149-2 PLEC_HUMAN	518 kDa		0.069	
17	Uncharacterized protein C19orf21 OS=Homo sapiens GN=C19orf21	sp P62701 RS4X_HUMAN	30 kDa		0.00019	0.0059
18	PE=1 SV=1	sp Q8IVT2 CS021_HUMAN	75 kDa		0.0024	<0.00010

19	40S ribosomal protein S3a OS=Homo sapiens GN=RPS3A PE=1 SV=2 Isoform 2 of Keratin, type II cytoskeletal 8 OS=Homo sapiens	sp P61247 RS3A_HUMAN	30 kDa		0.0014	
20	GN=KRT8 Alpha-actinin-4 OS=Homo sapiens	sp P05787-2 K2C8_HUMAN (+1)	57 kDa	TRUE	<0.00010	0.12
21	GN=ACTN4 PE=1 SV=2 Nucleolar RNA helicase 2 OS=Homo sapiens	sp O43707 ACTN4_HUMAN	105 kDa	TRUE	<0.00010	0.0029
22	GN=DDX21 PE=1 SV=5 60S ribosomal protein L6 OS=Homo sapiens	sp Q9NR30 DDX21_HUMAN	87 kDa	TRUE	0.00012	
23	GN=RPL6 PE=1 SV=3 Protein RRP5 homolog OS=Homo sapiens	sp Q02878 RL6_HUMAN	33 kDa		<0.00010	0.049
24	GN=PDCD11 PE=1 SV=3 Thyroid hormone receptor-associated protein 3 OS=Homo sapiens GN=THRAP3	sp Q14690 RRP5_HUMAN	209 kDa		<0.00010	
25	PE=1 SV=2 Isoform 2 of Heterogeneous nuclear ribonucleoprotein M OS=Homo sapiens	sp Q9Y2W1 TR150_HUMAN	109 kDa	TRUE	<0.00010	0.00068
26	GN=HNRNPM 60S ribosomal protein L7 OS=Homo sapiens	sp P52272-2 HNRPM_HUMAN (+1)	74 kDa	TRUE	<0.00010	
27	GN=RPL7 PE=1 SV=1 Protein arginine N-methyltransferase 5 OS=Homo sapiens	sp P18124 RL7_HUMAN	29 kDa		0.00017	0.026
28	GN=PRMT5 PE=1 SV=4 Nucleophosmin OS=Homo sapiens	sp O14744 ANM5_HUMAN	73 kDa		0.06	0.027
29	GN=NPM1 PE=1 SV=2 Tubulin alpha-1B chain OS=Homo sapiens	sp P06748 NPM_HUMAN	33 kDa	TRUE(6,7)	0.0026	0.44
30	GN=TUBA1B PE=1 SV=1 60S ribosomal protein L7a OS=Homo sapiens	sp P68363 TBA1B_HUMAN	50 kDa	TRUE	0.12	
31	GN=RPL7A PE=1 SV=2 60S ribosomal protein L3 OS=Homo sapiens	sp P62424 RL7A_HUMAN	30 kDa		0.00024	0.0059
32	GN=RPL3 PE=1 SV=2 Isoform 3 of Unconventional myosin-XVIIIa OS=Homo sapiens	sp P39023 RL3_HUMAN	46 kDa		0.00034	0.15
33	GN=MYO18A	sp Q92614-3 MY18A_HUMAN (+3)	227 kDa	TRUE (2,3)	<0.00010	0.00017

	DNA topoisomerase 1 OS=Homo sapiens					
34	GN=TOP1 PE=1 SV=2	sp P11387 TOP1_HUMAN	91 kDa		<0.00010	
	60S ribosomal protein L8 OS=Homo sapiens					
35	GN=RPL8 PE=1 SV=2	sp P62917 RL8_HUMAN	28 kDa	TRUE (2,3)	0.007	0.00094
	Ribosomal L1 domain- containing protein 1 OS=Homo sapiens					
36	GN=RSL1D1 PE=1 SV=3	sp O76021 RL1D1_HUMAN	55 kDa		<0.00010	
	Histone H2B type 1- C/E/F/G/I OS=Homo sapiens GN=HIST1H2BC					
37	PE=1 SV=4	sp P62807 H2B1C_HUMAN	14 kDa	TRUE	0.0034	0.0017
	Isoform 3 of Bcl-2- associated transcription factor 1 OS=Homo					
38	sapiens GN=BCLAF1	sp Q9NYF8-3 BCLF1_HUMAN (+1)	100 kDa	TRUE	0.00044	
	60S ribosomal protein L28 OS=Homo sapiens					
39	GN=RPL28 PE=1 SV=3	sp P46779 RL28_HUMAN	16 kDa		0.008	
	SWI/SNF complex subunit SMARCC2 OS=Homo sapiens					
40	GN=SMARCC2 PE=1 SV=1	sp Q8TAQ2 SMRC2_HUMAN	133 kDa	TRUE	<0.00010	<0.00010
	LIM domain and actin- binding protein 1 OS=Homo sapiens					
41	GN=LIMA1 PE=1 SV=1	sp Q9UHB6 LIMA1_HUMAN	85 kDa		0.0015	0.0023
	Nucleolin OS=Homo sapiens GN=NCL PE=1					
42	SV=3	sp P19338 NUCL_HUMAN	77 kDa	TRUE	<0.00010	
	60S ribosomal protein L13 OS=Homo sapiens					
43	GN=RPL13 PE=1 SV=4	sp P26373 RL13_HUMAN	24 kDa		0.00057	0.00054
	Leucine zipper protein 1 OS=Homo sapiens					
44	GN=LUZP1 PE=1 SV=2	sp Q86V48 LUZP1_HUMAN	120 kDa		<0.00010	0.0035
	Unconventional myosin- Ib OS=Homo sapiens					
45	GN=MYO1G PE=1 SV=2	sp B0I1T2 MYO1G_HUMAN	116 kDa	TRUE	0.00017	<0.00010
	60S ribosomal protein L21 OS=Homo sapiens					
46	GN=RPL21 PE=1 SV=2	sp P46778 RL21_HUMAN	19 kDa		0.001	0.15
	Isoform Beta of Nucleolar and coiled- body phosphoprotein 1 OS=Homo sapiens					
47	GN=NOLC1	sp Q14978-2 NOLC1_HUMAN	75 kDa		0.00017	
	H/ACA ribonucleoprotein complex subunit 4					
48		sp O60832 DKC1_HUMAN	58 kDa		<0.00010	

	OS=Homo sapiens GN=DKC1 PE=1 SV=3					
	40S ribosomal protein S6 OS=Homo sapiens					
49	GN=RPS6 PE=1 SV=1 Isoform 4 of Putative ribosomal RNA methyltransferase NOP2 OS=Homo sapiens	sp P62753 RS6_HUMAN	29 kDa		0.00013	0.12
50	GN=NOP2 60S ribosomal protein L18a OS=Homo sapiens	sp P46087-4 NOP2_HUMAN (+1)	93 kDa		<0.00010	
51	GN=RPL18A PE=1 SV=2 40S ribosomal protein S8 OS=Homo sapiens	sp Q02543 RL18A_HUMAN	21 kDa		0.00028	
52	GN=RPS8 PE=1 SV=2 60S ribosomal protein L18 OS=Homo sapiens	sp P62241 RS8_HUMAN	24 kDa		<0.00010	
53	GN=RPL18 PE=1 SV=2 40S ribosomal protein S2 OS=Homo sapiens	sp Q07020 RL18_HUMAN	22 kDa		0.028	
54	GN=RPS2 PE=1 SV=2 Tubulin beta chain OS=Homo sapiens	sp P15880 RS2_HUMAN	31 kDa		0.0042	
55	GN=TUBB PE=1 SV=2 N-acetyltransferase 10 OS=Homo sapiens	sp P07437 TBB5_HUMAN (+4)	50 kDa	TRUE	0.0042	
56	GN=NAT10 PE=1 SV=2 Keratin, type I cytoskeletal 14 OS=Homo sapiens	sp Q9H0A0 NAT10_HUMAN	116 kDa		<0.00010	
57	GN=KRT14 PE=1 SV=4 Eukaryotic translation initiation factor 4B OS=Homo sapiens	sp P02533 K1C14_HUMAN	52 kDa	TRUE	0.0012	0.0051
58	GN=EIF4B PE=1 SV=2 Keratin, type II cytoskeletal 5 OS=Homo sapiens GN=KRT5 PE=1 SV=3	sp P23588 IF4B_HUMAN (+1)	69 kDa		0.00086	0.69
59	Nucleolar protein 56 OS=Homo sapiens	sp P13647 K2C5_HUMAN	62 kDa	TRUE	0.00043	0.31
60	GN=NOP56 PE=1 SV=4 ATP-dependent RNA helicase A OS=Homo sapiens GN=DHX9 PE=1 SV=4	sp O00567 NOP56_HUMAN	66 kDa		<0.00010	
61	Isoform 2 of Unconventional myosin- Ic OS=Homo sapiens	sp Q08211 DHX9_HUMAN	141 kDa		0.00057	
62	GN=MYO1C	sp O00159-2 MYO1C_HUMAN (+3)	118 kDa		0.12	0.0026

63	rRNA 2'-O-methyltransferase fibrillar OS=Homo sapiens GN=FBL PE=1 SV=2	sp P22087 FBRL_HUMAN	34 kDa		0.00022	
64	Histone H4 OS=Homo sapiens GN=HIST1H4A PE=1 SV=2	sp P62805 H4_HUMAN	11 kDa		0.055	0.0059
65	40S ribosomal protein S13 OS=Homo sapiens GN=RPS13 PE=1 SV=2	sp P62277 RS13_HUMAN	17 kDa		0.0091	<0.00010
66	SWI/SNF complex subunit SMARCC1 OS=Homo sapiens GN=SMARCC1 PE=1 SV=3	sp Q92922 SMRC1_HUMAN	123 kDa	TRUE	0.0019	<0.00010
67	Isoform 2 of Annexin A2 OS=Homo sapiens GN=ANXA2	sp P07355-2 ANXA2_HUMAN (+1)	40 kDa		0.0002	0.027
68	60S ribosomal protein L17 OS=Homo sapiens GN=RPL17 PE=1 SV=3	sp P18621 RL17_HUMAN	21 kDa		0.00019	
69	Isoform 2 of Transcription activator BRG1 OS=Homo sapiens GN=SMARCA4	sp P51532-2 SMCA4_HUMAN (+5)	181 kDa	TRUE	0.0019	<0.0010
70	Keratin, type II cytoskeletal 7 OS=Homo sapiens GN=KRT7 PE=1 SV=5	sp P08729 K2C7_HUMAN	51 kDa	TRUE	0.023	0.025
71	60S ribosomal protein L26 OS=Homo sapiens GN=RPL26 PE=1 SV=1	sp P61254 RL26_HUMAN (+3)	17 kDa		0.0039	
72	60S ribosomal protein L14 OS=Homo sapiens GN=RPL14 PE=1 SV=4	sp P50914 RL14_HUMAN	23 kDa	TRUE	0.00096	0.0086
73	60S ribosomal protein L4 OS=Homo sapiens GN=RPL4 PE=1 SV=5	sp P36578 RL4_HUMAN	48 kDa		0.002	
74	40S ribosomal protein S11 OS=Homo sapiens GN=RPS11 PE=1 SV=3	sp P62280 RS11_HUMAN	18 kDa		0.0097	
75	Isoform 2 of Synaptopodin OS=Homo sapiens GN=SYNPO	sp Q8N3V7-2 SYNPO_HUMAN	96 kDa		0.26	0.011
76	40S ribosomal protein S16 OS=Homo sapiens GN=RPS16 PE=1 SV=2	sp P62249 RS16_HUMAN	16 kDa		0.00031	
77	60S ribosomal protein L10a OS=Homo sapiens GN=RPL10A PE=1 SV=2	sp P62906 RL10A_HUMAN	25 kDa		0.0079	

78	60S ribosomal protein L19 OS=Homo sapiens GN=RPL19 PE=1 SV=1	sp P84098 RL19_HUMAN (+2)	23 kDa		0.011	0.12
79	Tropomodulin-3 OS=Homo sapiens GN=TMOD3 PE=1 SV=1	sp Q9NYL9 TMOD3_HUMAN	40 kDa		0.0036	0.043
80	40S ribosomal protein S23 OS=Homo sapiens GN=RPS23 PE=1 SV=3	sp P62266 RS23_HUMAN	16 kDa		0.0044	0.0084
81	RNA-binding motif protein, X chromosome OS=Homo sapiens GN=RBMX PE=1 SV=3	sp P38159 RBMX_HUMAN	42 kDa		0.0015	
82	Unconventional myosin-le OS=Homo sapiens GN=MYO1E PE=1 SV=2	sp Q12965 MYO1E_HUMAN	127 kDa		0.06	
83	Histone H1x OS=Homo sapiens GN=H1FX PE=1 SV=1	sp Q92522 H1X_HUMAN	22 kDa		<0.00010	0.12
84	60S ribosomal protein L32 OS=Homo sapiens GN=RPL32 PE=1 SV=2	sp P62910 RL32_HUMAN (+1)	16 kDa		0.0079	0.0051
85	40S ribosomal protein S15a OS=Homo sapiens GN=RPS15A PE=1 SV=2	sp P62244 RS15A_HUMAN	15 kDa	TRUE	0.0042	
86	Isoform 3 of Myosin-10 OS=Homo sapiens GN=MYH10	sp P35580-3 MYH10_HUMAN (+1)	231 kDa	TRUE	0.00082	
87	40S ribosomal protein S25 OS=Homo sapiens GN=RPS25 PE=1 SV=1	sp P62851 RS25_HUMAN	14 kDa		0.0024	0.98
88	Histone H3.1 OS=Homo sapiens GN=HIST1H3A PE=1 SV=2	sp P68431 H31_HUMAN (+1)	15 kDa		0.0031	
89	60S ribosomal protein L24 OS=Homo sapiens GN=RPL24 PE=1 SV=1	sp P83731 RL24_HUMAN	18 kDa		<0.00010	
90	Heterochromatin protein 1-binding protein 3 OS=Homo sapiens GN=HP1BP3 PE=1 SV=1	sp Q5SSJ5 HP1B3_HUMAN	61 kDa		<0.00010	
91	Isoform 2 of F-actin-capping protein subunit beta OS=Homo sapiens GN=CAPZB	sp P47756-2 CAPZB_HUMAN (+2)	31 kDa		0.1	0.13
92	60S ribosomal protein L23a OS=Homo sapiens GN=RPL23A PE=1 SV=1	sp P62750 RL23A_HUMAN	18 kDa		0.0092	
93	Isoform 2 of Alpha-actinin-1 OS=Homo sapiens GN=ACTN1	sp P12814-2 ACTN1_HUMAN (+3)	103 kDa	TRUE	0.15	0.055

94	Heat shock cognate 71 kDa protein OS=Homo sapiens GN=HSPA8 PE=1 SV=1 RecName: Full=Gelsolin; AltName: Full=AGEL; AltName: Full=Actin-depolymerizing factor; Short=ADF; AltName: Full=Brevin; Flags:	sp P11142 HSP7C_HUMAN	71 kDa	TRUE	0.013	
95	Precursor Putative pre-mRNA-splicing factor ATP-dependent RNA helicase DHX15 OS=Homo sapiens GN=DHX15 PE=1 SV=2	gi 121116 sp P06396.1 GELS_HUMAN (+4)	86 kDa		<0.00010	0.00013
96	CCAAT/enhancer-binding protein zeta OS=Homo sapiens	sp O43143 DHX15_HUMAN	91 kDa		0.0015	
97	GN=CEBPZ PE=1 SV=3 F-actin-capping protein subunit alpha-1 OS=Homo sapiens	sp Q03701 CEBPZ_HUMAN	121 kDa		0.00019	
98	GN=CAPZA1 PE=1 SV=3 60S acidic ribosomal protein P0 OS=Homo sapiens GN=RPLP0 PE=1 SV=1	sp P52907 CAZA1_HUMAN	33 kDa	TRUE	0.023	0.087
99	40S ribosomal protein S14 OS=Homo sapiens	sp P05388 RLA0_HUMAN (+2)	34 kDa		0.00031	
100	GN=RPS14 PE=1 SV=3 60S ribosomal protein L34 OS=Homo sapiens	sp P62263 RS14_HUMAN	16 kDa		0.0022	0.14
101	GN=RPL34 PE=1 SV=3 Probable ATP-dependent RNA helicase DDX5 OS=Homo sapiens	sp P49207 RL34_HUMAN	13 kDa		0.0024	0.079
102	GN=DDX5 PE=1 SV=1 40S ribosomal protein S3 OS=Homo sapiens	sp P17844 DDX5_HUMAN (+1)	69 kDa	TRUE	0.00056	
103	GN=RPS3 PE=1 SV=2 Isoform 2 of Nucleolar protein 6 OS=Homo sapiens GN=NOL6	sp P23396 RS3_HUMAN (+2)	27 kDa		0.00028	
104	AT-rich interactive domain-containing protein 1A OS=Homo sapiens GN=ARID1A PE=1 SV=3	sp Q9H6R4-2 NOL6_HUMAN (+2)	112 kDa		0.00084	
105	Nucleolar complex protein 2 homolog	sp O14497 ARI1A_HUMAN	242 kDa	True (2,3)	<0.00010	0.00018
106		sp Q9Y3T9 NOC2L_HUMAN	85 kDa		<0.00010	

	OS=Homo sapiens GN=NOC2L PE=1 SV=4					
	60S ribosomal protein L27 OS=Homo sapiens					
107	GN=RPL27 PE=1 SV=2 Isoform 2 of 40S ribosomal protein S24 OS=Homo sapiens	sp P61353 RL27_HUMAN	16 kDa		0.0039	0.47
108	GN=RPS24 60S ribosomal protein L27a OS=Homo sapiens	sp P62847-2 RS24_HUMAN (+4)	15 kDa		<0.00010	
109	GN=RPL27A PE=1 SV=2 POTE ankyrin domain family member E OS=Homo sapiens	sp P46776 RL27A_HUMAN	17 kDa		0.00073	
110	GN=POTEE PE=1 SV=3 40S ribosomal protein S18 OS=Homo sapiens	sp Q6S8J3 POTEE_HUMAN	121 kDa	TRUE	0.37	0.37
111	GN=RPS18 PE=1 SV=3 40S ribosomal protein S19 OS=Homo sapiens	sp P62269 RS18_HUMAN	18 kDa		0.00072	0.035
112	GN=RPS19 PE=1 SV=2 60S ribosomal protein L29 OS=Homo sapiens	sp P39019 RS19_HUMAN	16 kDa		0.00017	
113	GN=RPL29 PE=1 SV=2 Drebrin OS=Homo sapiens GN=DBN1 PE=1 SV=4	sp P47914 RL29_HUMAN	18 kDa		0.03	
114	60S ribosomal protein L36 OS=Homo sapiens	sp Q16643 DREB_HUMAN (+1)	71 kDa		0.013	0.0058
115	GN=RPL36 PE=1 SV=3 Methylosome protein 50 OS=Homo sapiens	sp Q9Y3U8 RL36_HUMAN	12 kDa	TRUE	0.0054	
116	GN=WDR77 PE=1 SV=1 Nucleolar protein 58 OS=Homo sapiens	sp Q9BQA1 MEP50_HUMAN	37 kDa		0.025	0.65
117	GN=NOP58 PE=1 SV=1 Isoform 2 of Ribosomal RNA processing protein 1 homolog B OS=Homo sapiens GN=RRP1B	sp Q9Y2X3 NOP58_HUMAN	60 kDa		<0.00010	
118	RecName: Full=Trypsin; Flags: Precursor ATP-dependent RNA helicase DDX24 OS=Homo sapiens	sp Q14684-2 RRP1B_HUMAN (+1)	82 kDa		<0.00010	
119	GN=DDX24 PE=1 SV=1 Enhancer of rudimentary homolog OS=Homo sapiens	gi 136429 sp P00761.1 TRYP_PIG	24 kDa		0.33	0.18
120	GN=ERH PE=1 SV=1	sp Q9GZR7 DDX24_HUMAN (+1)	96 kDa		0.0025	
121		sp P84090 ERH_HUMAN	12 kDa		1	0.091

122	60S ribosomal protein L35a OS=Homo sapiens GN=RPL35A PE=1 SV=2 RecName: Full=Serum albumin; Flags:	sp P18077 RL35A_HUMAN	13 kDa	TRUE (6,7)	0.038	0.24
123	Precursor Isoform 2 of SWI/SNF-related matrix-associated actin-dependent regulator of chromatin subfamily E member 1 OS=Homo sapiens GN=SMARCE1 RNA-binding protein 28 OS=Homo sapiens	gi 113576 sp P02768.2 ALBU_HUMAN (+1)	69 kDa	True (2,3)	0.00017	0.37
124	sapiens GN=SMARCE1 RNA-binding protein 28 OS=Homo sapiens	sp Q969G3-2 SMCE1_HUMAN (+2)	42 kDa		0.0023	0.0003
125	GN=RBM28 PE=1 SV=3 Histone H1.0 OS=Homo sapiens GN=H1FO PE=1 SV=3	sp Q9NW13 RBM28_HUMAN	86 kDa	TRUE	<0.00010	
126	Isoform 2 of Fragile X mental retardation syndrome-related protein 1 OS=Homo sapiens GN=FXR1	sp P07305 H10_HUMAN	21 kDa		0.034	0.13
127	Isoform 2 of Serine/threonine-protein phosphatase PP1-alpha catalytic subunit OS=Homo sapiens GN=PPP1CA	sp P51114-2 FXR1_HUMAN (+1)	61 kDa	TRUE	0.0031	
128	Isoform 2 of RNA-binding protein 10 OS=Homo sapiens GN=RBM10	sp P62136-2 PP1A_HUMAN (+1)	39 kDa	TRUE	0.48	0.0034
129	Histone H1.5 OS=Homo sapiens GN=HIST1H1B PE=1 SV=3	sp P98175-2 RBM10_HUMAN (+3)	103 kDa		0.023	0.0078
130	60S ribosomal protein L31 OS=Homo sapiens GN=RPL31 PE=1 SV=1	sp P16401 H15_HUMAN	23 kDa	TRUE	0.00014	
131	60S ribosomal protein L23 OS=Homo sapiens GN=RPL23 PE=1 SV=1	sp P62899 RL31_HUMAN	14 kDa		0.027	0.073
132	Actin, alpha cardiac muscle 1 OS=Homo sapiens GN=ACTC1 PE=1 SV=1	sp P62829 RL23_HUMAN	15 kDa		0.16	0.35
133	60S ribosomal protein L30 OS=Homo sapiens GN=RPL30 PE=1 SV=2	sp P68032 ACTC_HUMAN (+1)	42 kDa	TRUE	0.37	0.45
134	Histone H1.4 OS=Homo sapiens GN=HIST1H1E PE=1 SV=2	sp P62888 RL30_HUMAN	13 kDa		<0.00010	
135		sp P10412 H14_HUMAN	22 kDa	TRUE	<0.00010	0.1

136	60S ribosomal protein L11 OS=Homo sapiens GN=RPL11 PE=1 SV=2 Isoform 2 of Tropomyosin alpha-3 chain OS=Homo sapiens	sp P62913 RL11_HUMAN	20 kDa		0.0062	
137	GN=TPM3 Ras GTPase-activating-like protein IQGAP1 OS=Homo sapiens	sp P06753-2 TPM3_HUMAN (+2)	29 kDa	TRUE	0.0014	
138	GN=IQGAP1 PE=1 SV=1 Isoform 2 of SWI/SNF-related matrix-associated actin-dependent regulator of chromatin subfamily D member 2 OS=Homo sapiens	sp P46940 IQGA1_HUMAN	189 kDa		<0.00010	
139	GN=SMARCD2 RNA-binding protein 14 OS=Homo sapiens	sp Q92925-2 SMRD2_HUMAN (+3)	57 kDa	TRUE	0.0034	
140	GN=RBM14 PE=1 SV=2 EF-hand domain-containing protein D2 OS=Homo sapiens	sp Q96PK6 RBM14_HUMAN	69 kDa		0.0025	
141	GN=EFHD2 PE=1 SV=1 Isoform 2 of ATP-dependent RNA helicase DDX54 OS=Homo sapiens	sp Q96C19 EFHD2_HUMAN	27 kDa		0.00018	0.0048
142	GN=DDX54 Serine/arginine-rich splicing factor 1 OS=Homo sapiens	sp Q8TDD1-2 DDX54_HUMAN (+1)	99 kDa		0.00046	
143	GN=SRSF1 PE=1 SV=2 60S ribosomal protein L35 OS=Homo sapiens	sp Q07955 SRSF1_HUMAN (+1)	28 kDa		0.0075	
144	GN=RPL35 PE=1 SV=2 60S ribosomal protein L37 OS=Homo sapiens	sp P42766 RL35_HUMAN	15 kDa		0.0078	0.12
145	GN=RPL37 PE=1 SV=2 Keratin, type I cytoskeletal 16 OS=Homo sapiens	sp P61927 RL37_HUMAN	11 kDa		0.0011	
146	GN=KRT16 PE=1 SV=4 Neurabin-2 OS=Homo sapiens	sp P08779 K1C16_HUMAN	51 kDa	TRUE	0.057	0.041
147	GN=PPP1R9B PE=1 SV=2 Flotillin-2 OS=Homo sapiens	sp Q96SB3 NEB2_HUMAN	89 kDa	TRUE	0.0059	0.013
148	GN=FLOT2 PE=1 SV=2 Isoform 2 of Myb-binding protein 1A OS=Homo sapiens	sp Q14254 FLOT2_HUMAN (+2)	47 kDa		0.00028	0.001
149	GN=MYBBP1A	sp Q9BQG0-2 MBB1A_HUMAN (+1)	149 kDa		<0.00010	

150	Isoform Smooth muscle of Myosin light polypeptide 6 OS=Homo sapiens GN=MYL6	sp P60660-2 MYL6_HUMAN (+8)	17 kDa		0.0012	
151	Keratin, type II cytoskeletal 6A OS=Homo sapiens GN=KRT6A PE=1 SV=3	sp P02538 K2C6A_HUMAN	60 kDa	TRUE	0.023	0.026
152	Probable ATP-dependent RNA helicase DDX27 OS=Homo sapiens GN=DDX27 PE=1 SV=2	sp Q96GQ7 DDX27_HUMAN	90 kDa		0.00039	
153	60S ribosomal protein L12 OS=Homo sapiens GN=RPL12 PE=1 SV=1	sp P30050 RL12_HUMAN	18 kDa		0.00039	
154	Heterogeneous nuclear ribonucleoprotein U OS=Homo sapiens GN=HNRNPU PE=1 SV=6	sp Q00839 HNRPU_HUMAN	91 kDa		<0.00010	
155	Isoform UBF2 of Nucleolar transcription factor 1 OS=Homo sapiens GN=UBTF	sp P17480-2 UBF1_HUMAN (+2)	85 kDa		0.01	
156	60S ribosomal protein L22 OS=Homo sapiens GN=RPL22 PE=1 SV=2	sp P35268 RL22_HUMAN	15 kDa	TRUE	0.012	
157	60S ribosomal protein L5 OS=Homo sapiens GN=RPL5 PE=1 SV=3	sp P46777 RL5_HUMAN	34 kDa		0.002	
158	Ribosome biogenesis protein BRX1 homolog OS=Homo sapiens GN=BRX1 PE=1 SV=2	sp Q8TDN6 BRX1_HUMAN	41 kDa		0.00068	
159	X-ray repair cross-complementing protein 5 OS=Homo sapiens GN=XRCC5 PE=1 SV=3	sp P13010 XRCC5_HUMAN	83 kDa		0.00027	<0.00010
160	Probable rRNA-processing protein EBP2 OS=Homo sapiens GN=EBNA1BP2 PE=1 SV=2	sp Q99848 EBP2_HUMAN (+1)	35 kDa		0.00089	
161	Elongation factor 1-alpha 1 OS=Homo sapiens GN=EEF1A1 PE=1 SV=1	sp P68104 EF1A1_HUMAN (+1)	50 kDa		0.026	
162	40S ribosomal protein S9 OS=Homo sapiens GN=RPS9 PE=1 SV=3	sp P46781 RS9_HUMAN	23 kDa		0.00043	
163	Isoform 2 of Unconventional myosin-	sp O43795-2 MYO1B_HUMAN (+2)	125 kDa		0.00056	0.18

	Ib OS=Homo sapiens GN=MYO1B					
	X-ray repair cross-complementing protein					
	6 OS=Homo sapiens					
164	GN=XRCC6 PE=1 SV=2 Isoform B of SWI/SNF-related matrix-associated actin-dependent regulator of chromatin subfamily B member 1 OS=Homo sapiens	sp P12956 XRCC6_HUMAN (+1)	70 kDa		0.00013	0.0005
165	GN=SMARCB1 Isoform 2 of Guanine nucleotide-binding protein-like 3 OS=Homo sapiens	sp Q12824-2 SNF5_HUMAN (+2)	43 kDa		0.00039	0.00011
166	GN=GNL3 Probable ATP-dependent RNA helicase DDX47 OS=Homo sapiens	sp Q9BVP2-2 GNL3_HUMAN (+1)	61 kDa		<0.00010	
167	GN=DDX47 PE=1 SV=1 Nucleolar GTP-binding protein 1 OS=Homo sapiens	sp Q9H0S4 DDX47_HUMAN	51 kDa			1
168	GN=GTPBP4 PE=1 SV=3 Isoform 2 of Pleckstrin homology-like domain family B member 2 OS=Homo sapiens	sp Q9BZE4 NOG1_HUMAN	74 kDa		0.011	
169	GN=PHLDB2 Isoform 3 of Neurabin-1 OS=Homo sapiens	sp Q86SQ0-2 PHLB2_HUMAN (+2)	137 kDa		0.067	
170	GN=PPP1R9A Nucleolar protein 10 OS=Homo sapiens	sp Q9ULJ8-3 NEB1_HUMAN (+3)	154 kDa	TRUE	<0.00010	
171	GN=NOL10 PE=1 SV=1 Isoform 2 of RRP12-like protein OS=Homo sapiens	sp Q9BSC4 NOL10_HUMAN (+1)	80 kDa		0.00013	
172	GN=RRP12 40S ribosomal protein S15 OS=Homo sapiens	sp Q5JTH9-2 RRP12_HUMAN (+2)	133 kDa		0.00015	
173	GN=RPS15 PE=1 SV=2 Hornerin OS=Homo sapiens	sp P62841 RS15_HUMAN (+1)	17 kDa		0.051	0.14
174	GN=HRNR PE=1 SV=2 pre-rRNA processing protein FTSJ3 OS=Homo sapiens	sp Q86YZ3 HORN_HUMAN	282 kDa		0.0037	0.21
175	GN=FTSJ3 PE=1 SV=2 Isoform 2 of Probable ATP-dependent RNA	sp Q8IY81 SPB1_HUMAN	97 kDa		0.00084	
176		sp Q92841-1 DDX17_HUMAN (+3)	72 kDa	TRUE	0.00026	

	helicase DDX17 OS=Homo sapiens GN=DDX17					
	Histone H2A type 1 OS=Homo sapiens GN=HIST1H2AG PE=1					
177	SV=2	sp POC0S8 H2A1_HUMAN (+3)	14 kDa	True (2,3)	0.87	0.61
	Isoform 2 of Nuclear mitotic apparatus protein 1 OS=Homo sapiens GN=NUMA1	sp Q14980-2 NUMA1_HUMAN (+1)	237 kDa		<0.00010	
178	40S ribosomal protein S17 OS=Homo sapiens GN=RPS17 PE=1 SV=2	sp P08708 RS17_HUMAN (+3)	16 kDa		0.052	
179	Ig alpha-1 chain C region OS=Homo sapiens GN=IGHA1 PE=1 SV=2	sp P01876 IGHA1_HUMAN	38 kDa		0.00033	
180	Isoform 2 of Actin- related protein 2 OS=Homo sapiens GN=ACTR2	sp P61160-2 ARP2_HUMAN (+1)	45 kDa		0.052	
181	RRP15-like protein OS=Homo sapiens GN=RRP15 PE=1 SV=2	sp Q9Y3B9 RRP15_HUMAN	31 kDa		0.01	
182	Isoform 2 of Probable ATP-dependent RNA helicase DDX31 OS=Homo sapiens GN=DDX31	sp Q9H8H2-2 DDX31_HUMAN (+2)	86 kDa		0.00027	
183	78 kDa glucose- regulated protein OS=Homo sapiens GN=HSPA5 PE=1 SV=2	sp P11021 GRP78_HUMAN	72 kDa	TRUE	0.58	
184	Folate receptor alpha OS=Homo sapiens GN=FOLR1 PE=1 SV=3	sp P15328 FOLR1_HUMAN	30 kDa		0.37	
185	Myosin regulatory light chain 12B OS=Homo sapiens GN=MYL12B PE=1 SV=2	sp O14950 ML12B_HUMAN (+2)	20 kDa		0.003	0.0011
186	Isoform 2 of Heat shock 70 kDa protein 1A/1B OS=Homo sapiens GN=HSPA1A	sp P08107-2 HSP71_HUMAN (+1)	64 kDa	TRUE	0.64	
187	Beta-actin-like protein 2 OS=Homo sapiens GN=ACTBL2 PE=1 SV=2	sp Q562R1 ACTBL_HUMAN	42 kDa	TRUE	0.33	0.12
188	Fragile X mental retardation syndrome- related protein 2 OS=Homo sapiens GN=FXR2 PE=1 SV=2	sp P51116 FXR2_HUMAN	74 kDa	TRUE	<0.00010	
189						

190	Zinc finger protein ubi- d4 OS=Homo sapiens GN=DPF2 PE=1 SV=2	sp Q92785 REQU_HUMAN	44 kDa	0.00027	
191	Ribosomal RNA- processing protein 7 homolog A OS=Homo sapiens GN=RRP7A PE=1 SV=2	sp Q9Y3A4 RRP7A_HUMAN	32 kDa	0.00027	
192	60S ribosomal protein L37a OS=Homo sapiens GN=RPL37A PE=1 SV=2	sp P61513 RL37A_HUMAN	10 kDa	0.079	
193	Isoform A1-A of Heterogeneous nuclear ribonucleoprotein A1 OS=Homo sapiens GN=HNRNPA1	sp P09651-2 ROA1_HUMAN (+5)	34 kDa	0.0039	
194	Isoform 2 of Actin-like protein 6A OS=Homo sapiens GN=ACTL6A	sp O96019-2 ACL6A_HUMAN (+1)	43 kDa	0.023	<0.00010
195	Cell growth-regulating nucleolar protein OS=Homo sapiens GN=LYAR PE=1 SV=2	sp Q9NX58 LYAR_HUMAN	44 kDa	0.00069	
196	Transferrin receptor protein 1 OS=Homo sapiens GN=TFRC PE=1 SV=2	sp P02786 TFR1_HUMAN	85 kDa	0.0039	
197	THO complex subunit 4 OS=Homo sapiens GN=ALYREF PE=1 SV=3	sp Q86V81 THOC4_HUMAN	27 kDa	0.002	
198	Phostensin OS=Homo sapiens GN=PPP1R18 PE=1 SV=1	sp Q6NYC8 PPR18_HUMAN (+1)	68 kDa	0.009	0.12
199	Isoform Short of TATA- binding protein- associated factor 2N OS=Homo sapiens GN=TAF15	sp Q92804-2 RBP56_HUMAN (+1)	62 kDa	0.00069	
200	40S ribosomal protein S26 OS=Homo sapiens GN=RPS26 PE=1 SV=3	sp P62854 RS26_HUMAN (+1)	13 kDa	0.012	
201	p21-activated protein kinase-interacting protein 1 OS=Homo sapiens GN=PAK1IP1 PE=1 SV=2	sp Q9NWT1 PK1IP_HUMAN	44 kDa	0.00048	
202	Nucleolar complex protein 3 homolog OS=Homo sapiens GN=NOC3L PE=1 SV=1	sp Q8WTT2 NOC3L_HUMAN (+1)	93 kDa	1	
203	Isoform 2 of Probable ATP-dependent RNA helicase DDX56	sp Q9NY93-2 DDX56_HUMAN (+1)	57 kDa	0.0039	

	OS=Homo sapiens GN=DDX56						
	Actin, cytoplasmic 1 OS=Homo sapiens						
204	GN=ACTB PE=1 SV=1 Ubiquitin-40S ribosomal protein S27a OS=Homo sapiens GN=RPS27A	sp P60709 ACTB_HUMAN	42 kDa	TRUE	0.37	0.37	
205	PE=1 SV=2 Protein KRI1 homolog OS=Homo sapiens	sp P62979 RS27A_HUMAN	18 kDa		0.3	0.37	
206	GN=KRI1 PE=1 SV=2 40S ribosomal protein S30 OS=Homo sapiens	sp Q8N9T8 KRI1_HUMAN	83 kDa		0.0011		
207	GN=FAU PE=1 SV=1 Isoform 2 of ATP- dependent RNA helicase DDX3X OS=Homo	sp P62861 RS30_HUMAN	7 kDa		0.12		
208	sapiens GN=DDX3X 60S ribosomal protein L39 OS=Homo sapiens	sp O00571-2 DDX3X_HUMAN (+1)	71 kDa		0.033		
209	GN=RPL39 PE=2 SV=2 Keratin, type I cytoskeletal 19 OS=Homo sapiens	sp P62891 RL39_HUMAN	6 kDa		0.37		
210	GN=KRT19 PE=1 SV=4 Isoform 1 of Complement decay- accelerating factor OS=Homo sapiens	sp P08727 K1C19_HUMAN	44 kDa	TRUE	0.021		
211	GN=CD55 Isoform 2 of Supervillin OS=Homo sapiens	sp P08174-2 DAF_HUMAN (+7)	49 kDa		0.025		
212	GN=SVIL NHP2-like protein 1 OS=Homo sapiens	sp O95425-2 SVIL_HUMAN (+1)	201 kDa		0.0038	0.022	
213	GN=NHP2L1 PE=1 SV=3 Transformer-2 protein homolog beta (Fragment) OS=Homo sapiens GN=TRA2B PE=4	sp P55769 NH2L1_HUMAN	14 kDa		0.011		
214	SV=1 60S ribosomal protein L36a OS=Homo sapiens	tr H7C2L4 H7C2L4_HUMAN	13 kDa	TRUE	<0.00010		
215	GN=RPL36A PE=1 SV=2 40S ribosomal protein S7 OS=Homo sapiens	sp P83881 RL36A_HUMAN (+1)	12 kDa		<0.00010		
216	GN=RPS7 PE=1 SV=1 40S ribosomal protein S29 OS=Homo sapiens	sp P62081 RS7_HUMAN (+1)	22 kDa		0.12		
217	GN=RPS29 PE=1 SV=2	sp P62273 RS29_HUMAN	7 kDa		0.091	0.00099	

218	Src substrate cortactin OS=Homo sapiens GN=CTTN PE=1 SV=2	sp Q14247 SRC8_HUMAN	62 kDa		1	
219	Unhealthy ribosome biogenesis protein 2 homolog OS=Homo sapiens GN=URB2 PE=1 SV=2	sp Q14146 URB2_HUMAN	171 kDa		0.021	
220	Periodic tryptophan protein 2 homolog OS=Homo sapiens GN=PWP2 PE=1 SV=2	sp Q15269 PWP2_HUMAN	102 kDa		0.00098	
221	H/ACA ribonucleoprotein complex subunit 3 OS=Homo sapiens GN=NOP10 PE=1 SV=1	sp Q9NPE3 NOP10_HUMAN	8 kDa		0.00098	
222	Isoform 2 of Protein polybromo-1 OS=Homo sapiens GN=PBRM1	sp Q86U86-2 PB1_HUMAN (+6)	187 kDa		0.12	
223	Serine/threonine- protein phosphatase PP1-beta catalytic subunit OS=Homo sapiens GN=PPP1CB PE=1 SV=3	sp P62140 PP1B_HUMAN	37 kDa	TRUE	0.12	
224	Isoform 2 of Protein phosphatase 1 regulatory subunit 12A OS=Homo sapiens GN=PPP1R12A	sp O14974-2 MYPT1_HUMAN (+5)	111 kDa		0.016	0.29
225	60S ribosomal protein L15 OS=Homo sapiens GN=RPL15 PE=1 SV=2	sp P61313 RL15_HUMAN	24 kDa		0.031	
226	Transducin beta-like protein 3 OS=Homo sapiens GN=TBL3 PE=1 SV=2	sp Q12788 TBL3_HUMAN (+1)	89 kDa		0.0022	
227	RNA-binding protein 34 OS=Homo sapiens GN=RBM34 PE=1 SV=2	sp P42696 RBM34_HUMAN	49 kDa		0.0061	
228	Isoform 3 of G patch domain-containing protein 4 OS=Homo sapiens GN=GPATCH4	sp Q5T3I0-3 GPTC4_HUMAN (+2)	51 kDa		0.0061	
229	Isoform 2 of Double- stranded RNA-specific adenosine deaminase OS=Homo sapiens GN=ADAR	sp P55265-2 DSRAD_HUMAN (+6)	133 kDa		0.0061	
230	Calmodulin OS=Homo sapiens GN=CALM1 PE=1 SV=2	sp P62158 CALM_HUMAN (+3)	17 kDa		0.12	0.37

	F-actin-capping protein subunit alpha-2 OS=Homo sapiens						
231	GN=CAPZA2 PE=1 SV=3	sp P47755 CAZA2_HUMAN	33 kDa	TRUE	0.00028	0.12	
	Reticulocalbin-1 OS=Homo sapiens						
232	GN=RCN1 PE=1 SV=1	sp Q15293 RCN1_HUMAN	39 kDa		0.0023		
	Probable ATP-dependent RNA helicase DDX10 OS=Homo sapiens GN=DDX10 PE=1						
233	SV=2	sp Q13206 DDX10_HUMAN (+1)	101 kDa		0.0023		
	Brain acid soluble protein 1 OS=Homo sapiens GN=BASP1 PE=1						
234	SV=2	sp P80723 BASP1_HUMAN	23 kDa		0.0002		
	Isoform SRP55-3 of Serine/arginine-rich splicing factor 6 OS=Homo sapiens						
235	GN=SRSF6	sp Q13247-3 SRSF6_HUMAN (+1)	38 kDa		0.008		
	mRNA turnover protein 4 homolog OS=Homo sapiens GN=MRT04						
236	PE=1 SV=2	sp Q9UKD2 MRT4_HUMAN	28 kDa		0.008		
	Isoform 3 of Death-associated protein kinase 1 OS=Homo sapiens GN=DAPK1						
237	ATP-dependent RNA helicase DDX50 OS=Homo sapiens	sp P53355-3 DAPK1_HUMAN (+3)	161 kDa		0.0029		
238	GN=DDX50 PE=1 SV=1	sp Q9BQ39 DDX50_HUMAN	83 kDa	TRUE	0.002		
	Flotillin-1 OS=Homo sapiens GN=FLOT1 PE=1						
239	SV=3	sp O75955 FLOT1_HUMAN (+1)	47 kDa		0.0002		
	Histone H2B type 2-E OS=Homo sapiens GN=HIST2H2BE PE=1						
240	SV=3	sp Q16778 H2B2E_HUMAN	14 kDa	TRUE	0.58	0.0081	
	Heterogeneous nuclear ribonucleoprotein F OS=Homo sapiens						
241	GN=HNRNPF PE=1 SV=3	sp P52597 HNRNPF_HUMAN	46 kDa		0.00039		
	WD repeat-containing protein 5 OS=Homo sapiens GN=WDR5 PE=1						
242	SV=1	sp P61964 WDR5_HUMAN	37 kDa		0.026		
	Isoform Short of Probable global transcription activator SNF2L2 OS=Homo sapiens GN=SMARCA2						
243		sp P51531-2 SMCA2_HUMAN (+1)	179 kDa	TRUE	0.00022	<0.00010	

	Serine/arginine-rich splicing factor 9 OS=Homo sapiens						
244	GN=SRSF9 PE=1 SV=1 WD repeat-containing protein 36 OS=Homo sapiens GN=WDR36	sp Q13242 SRSF9_HUMAN	26 kDa			0.0053	
245	PE=1 SV=1 Isoform 2 of RNA- binding protein 39 OS=Homo sapiens	sp Q8NI36 WDR36_HUMAN (+1)	105 kDa			0.0053	
246	GN=RBM39 Isoform C1 of Heterogeneous nuclear ribonucleoproteins C1/C2 OS=Homo sapiens	sp Q14498-2 RBM39_HUMAN (+4)	59 kDa			0.0053	
247	GN=HNRNPC ATP-dependent RNA helicase DDX18 OS=Homo sapiens	sp P07910-2 HNRPC_HUMAN (+14)	32 kDa			0.038	
248	GN=DDX18 PE=1 SV=2 Isoform 1 of Fragile X mental retardation protein 1 OS=Homo	sp Q9NVP1 DDX18_HUMAN	75 kDa			0.00039	
249	sapiens GN=FMR1 Isoform 2 of KRR1 small subunit processome component homolog OS=Homo sapiens	sp Q06787-2 FMR1_HUMAN (+2)	67 kDa	TRUE		0.16	
250	GN=KRR1 Tubulin beta-4B chain OS=Homo sapiens	sp Q13601-2 KRR1_HUMAN (+1)	37 kDa			0.16	
251	GN=TUBB4B PE=1 SV=1 Actin-related protein 2/3 complex subunit 1B OS=Homo sapiens	sp P68371 TBB4B_HUMAN	50 kDa	TRUE		0.12	
252	GN=ARPC1B PE=1 SV=3 DnaJ homolog subfamily C member 9 OS=Homo sapiens GN=DNAJC9	sp O15143 ARC1B_HUMAN	41 kDa			0.37	
253	PE=1 SV=1 Serine/threonine- protein kinase 38 OS=Homo sapiens	sp Q8WXX5 DNJC9_HUMAN	30 kDa			0.0075	
254	GN=STK38 PE=1 SV=1 Probable ATP- dependent RNA helicase DDX52 OS=Homo sapiens GN=DDX52 PE=1	sp Q15208 STK38_HUMAN	54 kDa			0.00056	0.37
255	SV=3 PIN2/TERF1-interacting	sp Q9Y2R4 DDX52_HUMAN	68 kDa			0.0075	
256	telomerase inhibitor 1	sp Q96BK5 PINX1_HUMAN	37 kDa			0.12	

	OS=Homo sapiens GN=PINX1 PE=1 SV=2				
	Nucleolar pre-ribosomal-associated protein 1 OS=Homo sapiens GN=URB1 PE=1 SV=4	sp O60287 NPA1P_HUMAN	254 kDa	0.14	
257	Isoform 2 of Basigin OS=Homo sapiens GN=BSG	sp P35613-2 BASI_HUMAN (+3)	29 kDa	0.12	
	Isoform 4 of Serine/arginine-rich splicing factor 7 OS=Homo sapiens GN=SRSF7	sp Q16629-4 SRSF7_HUMAN (+1)	26 kDa	0.13	
259	Isoform 2 of Actin-related protein 2/3 complex subunit 4 OS=Homo sapiens GN=ARPC4	sp P59998-2 ARPC4_HUMAN (+3)	72 kDa	0.37	0.0016
260	Protein flightless-1 homolog OS=Homo sapiens GN=FLII PE=1 SV=2	sp Q13045 FLII_HUMAN	145 kDa	0.0065	
261	Small nuclear ribonucleoprotein Sm D3 OS=Homo sapiens GN=SNRPD3 PE=1 SV=1	sp P62318 SMD3_HUMAN (+1)	14 kDa	0.00039	0.37
262	Isoform 5 of Putative oxidoreductase GLYR1 OS=Homo sapiens GN=GLYR1	sp Q49A26-5 GLYR1_HUMAN	52 kDa	1	
263	Signal recognition particle 9 kDa protein OS=Homo sapiens GN=SRP9 PE=1 SV=2	sp P49458 SRP09_HUMAN	10 kDa	1	
264	RNA 3'-terminal phosphate cyclase-like protein OS=Homo sapiens GN=RCL1 PE=1 SV=3	sp Q9Y2P8 RCL1_HUMAN	41 kDa	0.12	
265	Isoform 2 of Protein SSXT OS=Homo sapiens GN=SS18	sp Q15532-2 SSXT_HUMAN (+7)	42 kDa	0.12	0.16
266	Isoform 2 of Signal-induced proliferation-associated 1-like protein 1 OS=Homo sapiens GN=SIPA1L1	sp O43166-2 SI1L1_HUMAN (+2)	197 kDa	0.12	0.0025
267	Keratin, type II cytoskeletal 73	sp Q86Y46 K2C73_HUMAN	59 kDa	TRUE	0.12
268					

	OS=Homo sapiens GN=KRT73 PE=1 SV=1					
	Isoform 2 of SWI/SNF-related matrix-associated actin-dependent regulator of chromatin subfamily D member 1 OS=Homo sapiens GN=SMARCD1 Splicing factor 3B subunit 3 OS=Homo sapiens GN=SF3B3 PE=1 SV=4	sp Q96GM5-2 SMRD1_HUMAN (+1)	53 kDa	TRUE	0.019	0.00049
269						
270	Protein C16orf88 OS=Homo sapiens GN=C16orf88 PE=1 SV=1	sp Q15393 SF3B3_HUMAN	136 kDa		0.016	
271	Matrin-3 OS=Homo sapiens GN=MATR3 PE=1 SV=2	sp Q1ED39 CP088_HUMAN	52 kDa		0.16	
272	Protein AATF OS=Homo sapiens GN=AATF PE=1 SV=1	sp P43243 MATR3_HUMAN (+3)	95 kDa		0.12	
273	Isoform 2 of Spectrin beta chain, non-erythrocytic 1 OS=Homo sapiens GN=SPTBN1	sp Q9NY61 AATF_HUMAN	63 kDa		0.12	
274	Isoform 2 of Spectrin beta chain, non-erythrocytic 2 OS=Homo sapiens GN=SPTBN2	sp Q01082-3 SPTB2_HUMAN	251 kDa	TRUE	0.37	
275	Periodic tryptophan protein 1 homolog OS=Homo sapiens GN=PWP1 PE=1 SV=1	sp O15020-2 SPTN2_HUMAN (+1)	268 kDa	TRUE	0.37	
276	Isoform 2 of DNA topoisomerase 2-alpha OS=Homo sapiens GN=TOP2A	sp Q13610 PWP1_HUMAN (+1)	56 kDa		0.12	
277	Isoform 2 of Suppressor of SWI4 1 homolog OS=Homo sapiens GN=PPAN	sp P11388-2 TOP2A_HUMAN (+3)	178 kDa	TRUE	0.12	
278	Tropomyosin alpha-4 chain OS=Homo sapiens GN=TPM4 PE=1 SV=3	sp Q9NQ55-2 SSF1_HUMAN (+2)	52 kDa	TRUE	0.12	
279	Isoform 2 of Neuroguidin OS=Homo sapiens GN=NGDN	sp P67936 TPM4_HUMAN	29 kDa	TRUE	0.12	
280	Q8TD57 DYH3_HUMAN-DECOY	sp Q8NEJ9-2 NGDN_HUMAN (+2)	35 kDa		0.16	
281		Q8TD57 DYH3_HUMAN-DECOY	?		0.37	

282	60S ribosomal protein L10 OS=Homo sapiens GN=RPL10 PE=1 SV=4	sp P27635 RL10_HUMAN (+1)	25 kDa		0.37
	Sodium/potassium- transporting ATPase subunit alpha-1 OS=Homo sapiens				
283	GN=ATP1A1 PE=1 SV=1	sp P05023 AT1A1_HUMAN	113 kDa		0.13
	H/ACA ribonucleoprotein complex subunit 2 OS=Homo sapiens				
284	GN=NHP2 PE=1 SV=1	sp Q9NX24 NHP2_HUMAN	17 kDa		0.37
	Isoform 6 of Cyclin- dependent kinase inhibitor 2A, isoform 4 OS=Homo sapiens				
285	GN=CDKN2A	sp Q8N726-2 CD2A2_HUMAN (+1)	9 kDa		0.13
	Isoform 2 of Nucleolar protein 8 OS=Homo				
286	sapiens GN=NOL8	sp Q76FK4-2 NOL8_HUMAN (+5)	124 kDa		0.37
	Isoform 2 of 40S ribosomal protein S20 OS=Homo sapiens				
287	GN=RPS20	sp P60866-2 RS20_HUMAN (+1)	16 kDa		0.37
	ATP-dependent RNA helicase DDX51 OS=Homo sapiens				
288	GN=DDX51 PE=1 SV=3	sp Q8N8A6 DDX51_HUMAN	72 kDa		0.37
	Isoform 2 of Heterogeneous nuclear ribonucleoprotein K OS=Homo sapiens				
289	GN=HNRNPK	sp P61978-2 HNRPK_HUMAN (+4)	51 kDa		0.12
	60S ribosomal protein L9 OS=Homo sapiens				
290	GN=RPL9 PE=1 SV=1	sp P32969 RL9_HUMAN (+1)	22 kDa		0.12
	SAFB-like transcription modulator OS=Homo sapiens GN=SLTM PE=1				
291	SV=2	sp Q9NWH9 SLTM_HUMAN	117 kDa		0.37
	Ribosome biogenesis protein NSA2 homolog OS=Homo sapiens				
292	GN=NSA2 PE=1 SV=1	sp O95478 NSA2_HUMAN	30 kDa	TRUE	0.12
	U3 small nucleolar RNA- associated protein 18 homolog OS=Homo sapiens GN=UTP18 PE=1				
293	SV=3	sp Q9Y5J1 UTP18_HUMAN	62 kDa		0.37
	Isoform 2 of Protein ELYS OS=Homo sapiens				
294	GN=AHCTF1	sp Q8WYP5-2 ELYS_HUMAN (+2)	256 kDa		0.37

	Isoform 2 of DNA-dependent protein kinase catalytic subunit OS=Homo sapiens			
295	GN=PRKDC	sp P78527-2 PRKDC_HUMAN (+2)	466 kDa	0.37
	Q9C0G0-2 ZN407_HUMAN-DECOY			
296	DECOY	Q9C0G0-2 ZN407_HUMAN-DECOY	?	0.37
	Heat shock protein HSP90-beta OS=Homo sapiens GN=HSP90AB1			
297	PE=1 SV=4	sp P08238 HS90B_HUMAN	83 kDa	0.37
	Isoform 2 of Myosin phosphatase Rho-interacting protein OS=Homo sapiens			
298	GN=MPRIP	sp Q6WCQ1-2 MPRIP_HUMAN (+3)	118 kDa	0.37
	U3 small nucleolar ribonucleoprotein protein IMP4 OS=Homo sapiens GN=IMP4 PE=1			
299	SV=1	sp Q96G21 IMP4_HUMAN (+3)	34 kDa	0.37
	FACT complex subunit SPT16 OS=Homo sapiens			
300	GN=SUPT16H PE=1 SV=1	sp Q9Y5B9 SP16H_HUMAN	120 kDa	0.37
	Isoform A2 of Heterogeneous nuclear ribonucleoproteins A2/B1 OS=Homo sapiens			
301	GN=HNRNPA2B1	sp P22626-2 ROA2_HUMAN (+1)	36 kDa	0.37
	Isoform 2 of AT-rich interactive domain-containing protein 2 OS=Homo sapiens			
302	GN=ARID2	sp Q68CP9-3 ARID2_HUMAN (+1)	191 kDa	0.37
	Zinc finger protein 234 OS=Homo sapiens			
303	GN=ZNF234 PE=2 SV=3	sp Q14588 ZN234_HUMAN	81 kDa	0.37
	Putative ATP-dependent RNA helicase DHX33 OS=Homo sapiens			
304	GN=DHX33 PE=1 SV=2	sp Q9H6R0 DHX33_HUMAN	79 kDa	0.37
	Isoform 2 of Interleukin enhancer-binding factor 3 OS=Homo sapiens			
305	GN=ILF3	sp Q12906-2 ILF3_HUMAN (+6)	76 kDa	0.37
	NF-kappa-B-repressing factor OS=Homo sapiens			
306	GN=NKRF PE=1 SV=2	sp O15226 NKRF_HUMAN (+1)	78 kDa	0.37
	Isoform 4 of Apoptotic chromatin condensation inducer in the nucleus			
307		sp Q9UKV3-5 ACINU_HUMAN (+2)	151 kDa	0.37

0.0057

	OS=Homo sapiens GN=ACIN1				
	RecName: Full=Lactotransferrin; Short=Lactoferrin; AltName: Full=Talalactoferrin; Contains: RecName: Full=Kalioicin-1; Contains: RecName: Full=Lactoferroxin-A; Contains: RecName: Full=Lactoferroxin-B; Contains: RecName: Full=Lactoferroxin-C;				
308	Flags: Precursor Isoform 2 of Poly(rC)- binding protein 2 OS=Homo sapiens	gi 85700158 sp P02788.6 TRFL_HUMAN (+3)	78 kDa		0.37
309	GN=PCBP2 Ribosome biogenesis protein BMS1 homolog OS=Homo sapiens	sp Q15366-2 PCBP2_HUMAN (+9)	39 kDa	TRUE	0.37
310	GN=BMS1 PE=1 SV=1 Interferon-stimulated 20 kDa exonuclease-like 2 OS=Homo sapiens	sp Q14692 BMS1_HUMAN	146 kDa		0.37
311	GN=ISG20L2 PE=1 SV=1 Q7Z7A1- 5 CNTRL_HUMAN-	sp Q9H9L3 I20L2_HUMAN	39 kDa		0.37
312	DECOY	Q7Z7A1-5 CNTRL_HUMAN-DECOY	?		0.37

List of Proteins Only in Lanes 2,3

#	Identified Proteins (162) RecName: Full=Serum albumin; AltName: Full=BSA; AltName: Allergen=Bos d 6; Flags: Precursor	Accession Number	Molecular Weight	Protein Grouping Ambiguity	Lanes 2,3 P-values	What protein from 6,7 is it similar to?
16	Isoform 2 of Filamin-B OS=Homo sapiens	gi 1351907 sp P02769.4 ALBU_BOVIN	69 kDa	TRUE	<0.00010	Similar to 123
19	GN=FLNB Glycogen phosphorylase, muscle form OS=Homo sapiens GN=PYGM PE=1 SV=6	sp O75369-2 FLNB_HUMAN (+3)	276 kDa	TRUE	0.0011	
22	RecName: Full=Carbonic anhydrase 2; AltName: Full=Carbonate dehydratase II; AltName: Full=Carbonic anhydrase II; Short=CA-II	sp P11217 PYGM_HUMAN	97 kDa		<0.00010	
32	Histone H3.1 OS=Homo sapiens GN=HIST1H3A PE=1 SV=2	gi 41019480 sp P00921.3 CAH2_BOVIN	29 kDa		<0.00010	Extremely similar to 88
33	Tubulin alpha-1B chain OS=Homo sapiens GN=TUBA1B PE=1 SV=1	sp P68431 H31_HUMAN (+2)	15 kDa		0.00053	Extremely similar to 30
34	Isoform 2 of Bcl-2- associated transcription factor 1 OS=Homo sapiens GN=BCLAF1	sp P68363 TBA1B_HUMAN (+5)	50 kDa		0.11	
39	RecName: Full=Beta- galactosidase; Short=Beta-gal; AltName: Full=Lactase	sp Q9NYF8-2 BCLF1_HUMAN (+1)	106 kDa	TRUE	0.0011	
40	RecName: Full=Ovalbumin; AltName: Full=Allergen Gal d II; AltName: Full=Egg albumin; AltName: Full=Plakalbumin; AltName: Allergen=Gal d 2	gi 114939 sp P00722.2 BGAL_ECOLI	116 kDa		0.00031	
42		gi 129293 sp P01012.2 OVAL_CHICK	43 kDa		0.00067	

44	Isoform 2 of Plectin OS=Homo sapiens GN=PLEC	sp Q15149-2 PLEC_HUMAN (+7)	518 kDa		0.016	Extremely similar to 16
47	Isoform 8 of Protein polybromo-1 OS=Homo sapiens GN=PBRM1	sp Q86U86-8 PB1_HUMAN (+1)	190 kDa		<0.00010	
48	Isoform 3 of SWI/SNF-related matrix-associated actin-dependent regulator of chromatin subfamily D member 2 OS=Homo sapiens GN=SMARCD2	sp Q92925-3 SMRD2_HUMAN (+2)	55 kDa	TRUE	<0.00010	
62	Isoform 2 of AT-rich interactive domain-containing protein 1B OS=Homo sapiens GN=ARID1B	sp Q8NFD5-2 ARI1B_HUMAN (+2)	238 kDa	TRUE	<0.00010	
66	Isoform Smooth muscle of Myosin light polypeptide 6 OS=Homo sapiens GN=MYL6	sp P60660-2 MYL6_HUMAN (+1)	17 kDa		0.028	Extremely similar to 150
70	Isoform 3 of Neurabin-1 OS=Homo sapiens GN=PPP1R9A	sp Q9ULJ8-3 NEB1_HUMAN (+2)	154 kDa	TRUE	0.18	Extremely similar to 170
72	40S ribosomal protein S11 OS=Homo sapiens GN=RPS11 PE=1 SV=3	sp P62280 RS11_HUMAN (+1)	18 kDa		0.045	Extremely similar to 74
75	Isoform 2 of 60S ribosomal protein L28 OS=Homo sapiens GN=RPL28	sp P46779-2 RL28_HUMAN (+3)	18 kDa		0.00051	
78	Isoform 2 of Myosin-10 OS=Homo sapiens GN=MYH10	sp P35580-2 MYH10_HUMAN (+4)	231 kDa	TRUE	<0.00010	
79	Isoform 2 of Pleckstrin homology-like domain family B member 2 OS=Homo sapiens GN=PHLDB2	sp Q86SQ0-2 PHLB2_HUMAN (+3)	137 kDa		0.00067	Extremely similar to 169
85	40S ribosomal protein S16 OS=Homo sapiens GN=RPS16 PE=1 SV=2 RecName: Full=Lysozyme C; AltName: Full=1,4-beta-N-acetylmuramidase C; AltName: Full=Allergen	sp P62249 RS16_HUMAN (+1)	16 kDa		0.037	Extremely similar to 76
86	Gal d IV; AltName:	gi 126608 sp P00698.1 LYSC_CHICK	16 kDa		0.0099	

Allergen=Gal d 4; Flags: Precursor							
96	40S ribosomal protein S2 OS=Homo sapiens GN=RPS2 PE=1 SV=2	sp P15880 RS2_HUMAN (+1)	31 kDa		0.018	Extremely similar to 54	
100	Zinc finger protein ubi-d4 OS=Homo sapiens GN=DPF2 PE=1 SV=2	sp Q92785 REQU_HUMAN (+2)	44 kDa		0.00084	Extremely similar to 190	
101	Elongation factor 1-alpha 1 OS=Homo sapiens GN=EEF1A1 PE=1 SV=1	sp P68104 EF1A1_HUMAN (+2)	50 kDa		0.45	Extremely similar to 161	
104	40S ribosomal protein S8 OS=Homo sapiens GN=RPS8 PE=1 SV=2	sp P62241 RS8_HUMAN (+1)	24 kDa		0.0011	Extremely similar to 52	
109	40S ribosomal protein S3a OS=Homo sapiens GN=RPS3A PE=1 SV=2	sp P61247 RS3A_HUMAN (+1)	30 kDa		0.1	Extremely similar to 19	
110	Heat shock cognate 71 kDa protein OS=Homo sapiens GN=HSPA8 PE=1 SV=1	sp P11142 HSP7C_HUMAN (+1)	71 kDa	TRUE	0.76	Extremely similar to 94	
114	40S ribosomal protein S15a OS=Homo sapiens GN=RPS15A PE=1 SV=2	sp P62244 RS15A_HUMAN (+1)	15 kDa		0.0091	Extremely similar to 85	
117	60S ribosomal protein L26 OS=Homo sapiens GN=RPL26 PE=1 SV=1	sp P61254 RL26_HUMAN (+5)	17 kDa		0.11	Extremely similar to 71	
118	60S ribosomal protein L22 OS=Homo sapiens GN=RPL22 PE=1 SV=2	sp P35268 RL22_HUMAN (+3)	15 kDa		0.00055	Extremely similar to 156	
119	60S ribosomal protein L27a OS=Homo sapiens GN=RPL27A PE=1 SV=2	sp P46776 RL27A_HUMAN (+1)	17 kDa		0.14	Extremely similar to 109	
120	OTU domain-containing protein 4 OS=Homo sapiens GN=OTUD4 PE=1 SV=3	sp Q01804 OTUD4_HUMAN (+1)	124 kDa		0.31		
127	Isoform 3 of Keratin, type I cytoskeletal 13 OS=Homo sapiens GN=KRT13	sp P13646-3 K1C13_HUMAN (+2)	46 kDa	TRUE	0.13		
128	60S ribosomal protein L37a OS=Homo sapiens GN=RPL37A PE=1 SV=2	sp P61513 RL37A_HUMAN (+3)	10 kDa		0.0034	Extremely similar to 192	

136	Isoform Gamma-2 of Serine/threonine-protein phosphatase PP1-gamma catalytic subunit OS=Homo sapiens GN=PPP1CC	sp P36873-2 PP1G_HUMAN (+2)	39 kDa	TRUE	0.37	
137	40S ribosomal protein S17 OS=Homo sapiens GN=RPS17 PE=1 SV=2	sp P08708 RS17_HUMAN (+5)	16 kDa		0.37	Extremely similar to 179
138	60S ribosomal protein L23a OS=Homo sapiens GN=RPL23A PE=1 SV=1	sp P62750 RL23A_HUMAN (+4)	18 kDa		<0.00010	Extremely similar to 92
140	Isoform 2 of Actin-related protein 2 OS=Homo sapiens GN=ACTR2	sp P61160-2 ARP2_HUMAN (+2)	45 kDa		0.37	Extremely similar to 181
141	60S ribosomal protein L38 OS=Homo sapiens GN=RPL38 PE=1 SV=2	sp P63173 RL38_HUMAN	8 kDa		0.36	
147	60S ribosomal protein L11 (Fragment) OS=Homo sapiens GN=RPL11 PE=2 SV=1	tr Q5VVC8 Q5VVC8_HUMAN	20 kDa		0.37	
148	40S ribosomal protein S19 OS=Homo sapiens GN=RPS19 PE=1 SV=2	sp P39019 RS19_HUMAN (+4)	16 kDa		0.00039	Extremely similar to 112
149	Isoform 3 of B-cell CLL/lymphoma 7 protein family member B OS=Homo sapiens GN=BCL7B	sp Q9BQE9-3 BCL7B_HUMAN (+4)	18 kDa		0.37	
152	Histone H2B type 1-K OS=Homo sapiens GN=HIST1H2BK PE=1 SV=3	sp O60814 H2B1K_HUMAN	14 kDa	TRUE	0.37	
154	60S ribosomal protein L18a OS=Homo sapiens GN=RPL18A PE=1 SV=2	sp Q02543 RL18A_HUMAN (+3)	21 kDa		0.37	Extremely similar to 51
155	Myosin-1 OS=Homo sapiens GN=MYH1 PE=1 SV=3	sp P12882 MYH1_HUMAN	223 kDa	TRUE	0.13	
156	Actin-related protein 2/3 complex subunit 2 OS=Homo sapiens GN=ARPC2 PE=1 SV=1	sp O15144 ARPC2_HUMAN	34 kDa		0.12	
157	60S ribosomal protein L18 (Fragment) OS=Homo sapiens GN=RPL18 PE=3 SV=1	tr HOYHA7 HOYHA7_HUMAN	19 kDa		0.37	

158	60S ribosomal protein L24 OS=Homo sapiens GN=RPL24 PE=1 SV=1	sp P83731 RL24_HUMAN (+2)	18 kDa	0.37	Extremely similar to 89
159	Ras GTPase-activating- like protein IQGAP1 OS=Homo sapiens GN=IQGAP1 PE=1 SV=1	sp P46940 IQGA1_HUMAN (+1)	189 kDa	0.12	Extremely similar to 138
160	Isoform 2 of PHD finger protein 10 OS=Homo sapiens GN=PHF10	sp Q8WUB8-2 PHF10_HUMAN (+2)	56 kDa	0.37	
161	Isoform 2 of Src substrate cortactin OS=Homo sapiens GN=CTTN	sp Q14247-2 SRC8_HUMAN (+2)	71 kDa	0.37	
162	60S ribosomal protein L17 OS=Homo sapiens GN=RPL17 PE=1 SV=3	sp P18621 RL17_HUMAN (+5)	21 kDa	0.37	Extremely similar to 68

Appendix 2: List of Protein Descriptions for Proteins with a Significant P-value in both the 6,7 and 2,3 Proteomics Results

(Some of the description were unable to be determined.)

References 5,6, and 7 from the Bibliography were used to find the protein descriptions.

Protein Name	Description
Myosin-9 OS=Homo sapiens GN=MYH9 PE=1 SV=4	Cellular myosin that appears to play a role in cytokinesis, cell shape, and specialized functions such as secretion and capping. During cell spreading, plays an important role in cytoskeleton reorganization, focal contacts formation (in the margins but not the central part of spreading cells), and lamellipodial retraction; this function is mechanically antagonized by MYH10. http://www.uniprot.org/uniprot/P35579
Isoform 2 of Filamin-A OS=Homo sapiens GN=FLNA	Promotes orthogonal branching of actin filaments and links actin filaments to membrane glycoproteins. Anchors various transmembrane proteins to the actin cytoskeleton and serves as a scaffold for a wide range of cytoplasmic signaling proteins. Interaction with FLNA may allow neuroblast migration from the ventricular zone into the cortical plate. Tethers cell surface-localized furin, modulates its rate of internalization and directs its intracellular trafficking (By similarity). Involved in ciliogenesis. http://www.uniprot.org/uniprot/P21333
Spectrin alpha chain, non-erythrocytic 1 OS=Homo sapiens GN=SPTAN1 PE=1 SV=3	Fodrin, which seems to be involved in secretion, interacts with calmodulin in a calcium-dependent manner and is thus candidate for the calcium-dependent movement of the cytoskeleton at the membrane. http://www.uniprot.org/uniprot/Q13813
Spectrin beta chain, non-erythrocytic 1 OS=Homo sapiens GN=SPTBN1 PE=1 SV=2	Fodrin, which seems to be involved in secretion, interacts with calmodulin in a calcium-dependent manner and is thus candidate for the calcium-dependent movement of the cytoskeleton at the membrane. http://www.uniprot.org/uniprot/Q01082
LIM domain only protein 7 OS=Homo sapiens GN=LMO7 PE=2 SV=1	<ul style="list-style-type: none"> • ubiquitin-protein transferase activity • zinc ion binding • protein ubiquitination • regulation of cell adhesion • regulation of signaling http://www.uniprot.org/uniprot/Q8WW11
RecName: Full=Keratin, type II cytoskeletal 1; AltName: Full=67 kDa cyokeratin; AltName:	<ul style="list-style-type: none"> • May regulate the activity of kinases such as PKC and SRC via binding to integrin beta-1 (ITB1) and the receptor of activated protein C kinase 1

<p>Full=Cytokeratin-1; Short=CK-1; AltName: Full=Hair alpha protein; AltName: Full=Keratin-1; Short=K1; AltName: Full=Type-II keratin Kb1</p>	<p>(RACK1). In complex with C1QBP is a high affinity receptor for kininogen-1/HMWK.</p> <ul style="list-style-type: none"> • Carbohydrate binding • Receptor activity • Structural molecule activity <p>http://www.uniprot.org/uniprot/P04264</p>
<p>RecName: Full=Keratin, type II cytoskeletal 2 epidermal; AltName: Full=Cytokeratin-2e; Short=CK-2e; AltName: Full=Epithelial keratin-2e; AltName: Full=Keratin-2 epidermis; AltName: Full=Keratin-2e; Short=K2e; AltName: Full=Type-II keratin Kb2</p>	<ul style="list-style-type: none"> • Probably contributes to terminal cornification. Associated with keratinocyte activation, proliferation and keratinization • Structural constituent of cytoskeleton <p>http://www.uniprot.org/uniprot/P35908</p>
<p>RecName: Full=Keratin, type I cytoskeletal 9; AltName: Full=Cytokeratin-9; Short=CK-9; AltName: Full=Keratin-9; Short=K9</p>	<ul style="list-style-type: none"> • May serve an important special function either in the mature palmar and plantar skin tissue or in the morphogenetic program of the formation of these tissues. Plays a role in keratin filament assembly. • Structural constituent of cytoskeleton <p>http://www.uniprot.org/uniprot/P35527</p>
<p>Keratin, type I cytoskeletal 18 OS=Homo sapiens GN=KRT18 PE=1 SV=2</p>	<ul style="list-style-type: none"> • Involved in the uptake of thrombin-antithrombin complexes by hepatic cells (By similarity). When phosphorylated, plays a role in filament reorganization. Involved in the delivery of mutated CFTR to the plasma membrane. Together with KRT8, is involved in interleukin-6 (IL-6)-mediated barrier protection. • poly(A) RNA binding • scaffold protein • structural molecule activity <p>http://www.uniprot.org/uniprot/P05783</p>
<p>40S ribosomal protein S4, X isoform OS=Homo sapiens GN=RPS4X PE=1 SV=2</p>	<ul style="list-style-type: none"> • poly(A) RNA binding • rRNA binding • structural constituent of ribosome • cellular protein metabolic process • gene expression • multicellular organismal development • nuclear-transcribed mRNA catabolic process, nonsense-mediated decay • positive regulation of cell proliferation • positive regulation of translation

	<ul style="list-style-type: none"> • SRP-dependent cotranslational protein targeting to membrane • translation • translational elongation • translational initiation • translational termination • viral life cycle • viral process • viral transcription <p>http://www.uniprot.org/uniprot/P62701</p>
<p>Uncharacterized protein C19orf21 OS=Homo sapiens GN=C19orf21 PE=1 SV=1</p>	<p>Could not find a lot of info about this protein.</p>
<p>Alpha-actinin-4 OS=Homo sapiens GN=ACTN4 PE=1 SV=2</p>	<ul style="list-style-type: none"> • F-actin cross-linking protein which is thought to anchor actin to a variety of intracellular structures. This is a bundling protein (Probable). Probably involved in vesicular trafficking via its association with the CART complex. The CART complex is necessary for efficient transferrin receptor recycling but not for EGFR degradation (PubMed: 15772161). Involved in tight junction assembly in epithelial cells probably through interaction with MICALL2. Links MICALL2 to the actin cytoskeleton and recruits it to the tight junctions (By similarity). May also function as a transcriptional coactivator, stimulating transcription mediated by the nuclear hormone receptors PPARG and RARA • actin binding • actin filament binding • calcium ion binding • chromatin DNA binding • integrin binding • ion channel binding • ligand-dependent nuclear receptor transcription coactivator activity • nuclear hormone receptor binding • nucleoside binding • poly(A) RNA binding • protein homodimerization activity • retinoic acid receptor binding • RNA polymerase II regulatory region sequence-specific DNA binding <p>http://www.uniprot.org/uniprot/O43707</p>

<p>60S ribosomal protein L6 OS=Homo sapiens GN=RPL6 PE=1 SV=3</p>	<ul style="list-style-type: none"> • Specifically binds to domain C of the Tax-responsive enhancer element in the long terminal repeat of HTLV-I. • DNA binding • poly(A) RNA binding • RNA binding • structural constituent of ribosome <p>http://www.uniprot.org/uniprot/Q02878</p>
<p>Thyroid hormone receptor-associated protein 3 OS=Homo sapiens GN=THRAP3 PE=1 SV=2</p>	<p>Involved in pre-mRNA splicing. Remains associated with spliced mRNA after splicing which probably involves interactions with the exon junction complex (EJC). Can trigger mRNA decay which seems to be independent of nonsense-mediated decay involving premature stop codons (PTC) recognition. May be involved in nuclear mRNA decay. Involved in regulation of signal-induced alternative splicing. During splicing of PTPRC/CD45 is proposed to sequester phosphorylated SFPQ from PTPRC/CD45 pre-mRNA in resting T-cells. Involved in cyclin-D1/CCND1 mRNA stability probably by acting as component of the SNARP complex which associates with both the 3'end of the CCND1 gene and its mRNA. Involved in response to DNA damage. Is excluded from DNA damage sites in a manner that parallels transcription inhibition; the function may involve the SNARP complex. Initially thought to play a role in transcriptional coactivation through its association with the TRAP complex; however, it is not regarded as a stable Mediator complex subunit. Cooperatively with HELZ2, enhances the transcriptional activation mediated by PPARG, maybe through the stabilization of the PPARG binding to DNA in presence of ligand. May play a role in the terminal stage of adipocyte differentiation. Plays a role in the positive regulation of the circadian clock. Acts as a coactivator of the CLOCK-ARNTL/BMAL1 heterodimer and promotes its transcriptional activator activity and binding to circadian target genes.</p> <p>http://www.uniprot.org/uniprot/Q9Y2W1</p>
<p>60S ribosomal protein L7 OS=Homo sapiens GN=RPL7 PE=1 SV=1</p>	<p>Part of Ribosome. Can't really find anything</p>
<p>60S ribosomal protein L7a OS=Homo sapiens GN=RPL7A PE=1 SV=2</p>	<ul style="list-style-type: none"> • poly(A) RNA binding • RNA binding • structural constituent of ribosome <p>http://www.uniprot.org/uniprot/P62424</p>

<p>Isoform 3 of Unconventional myosin-XVIIIa OS=Homo sapiens GN=MYO18A</p>	<p>May link Golgi membranes to the cytoskeleton and participate in the tensile force required for vesicle budding from the Golgi. Thereby, may play a role in Golgi membrane trafficking and could indirectly give its flattened shape to the Golgi apparatus. Alternatively, in concert with LURAP1 and CDC42BPA/CDC42BPB, has been involved in modulating lamellar actomyosin retrograde flow that is crucial to cell protrusion and migration. May be involved in the maintenance of the stromal cell architectures required for cell to cell contact. http://www.uniprot.org/uniprot/Q92614 Couldn't really find anything on Isoform 3 specifically</p>
<p>60S ribosomal protein L8 OS=Homo sapiens GN=RPL8 PE=1 SV=2</p>	<ul style="list-style-type: none"> • poly(A) RNA binding • RNA binding • rRNA binding • structural constituent of ribosome <p>http://www.uniprot.org/uniprot/P62917</p>
<p>Histone H2B type 1-C/E/F/G/I OS=Homo sapiens GN=HIST1H2BC PE=1 SV=4</p>	<p>Core component of nucleosome. Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility is regulated via a complex set of post-translational modifications of histones, also called histone code, and nucleosome remodeling.</p> <p>Has broad antibacterial activity. May contribute to the formation of the functional antimicrobial barrier of the colonic epithelium, and to the bactericidal activity of amniotic fluid.</p> <p>http://www.uniprot.org/uniprot/P62807</p>
<p>SWI/SNF complex subunit SMARCC2 OS=Homo sapiens GN=SMARCC2 PE=1 SV=1</p>	<p>SWI/SNF</p>
<p>LIM domain and actin-binding protein 1 OS=Homo sapiens GN=LIMA1 PE=1 SV=1</p>	<ul style="list-style-type: none"> • Binds to actin monomers and filaments. Increases the number and size of actin stress fibers and inhibits membrane ruffling. Inhibits actin filament depolymerization. Bundles actin filaments, delays filament nucleation and reduces formation of branched filaments.1 Publication • actin filament binding • actin monomer binding • zinc ion binding <p>http://www.uniprot.org/uniprot/Q9UHB6</p>

<p>60S ribosomal protein L13 OS=Homo sapiens GN=RPL13 PE=1 SV=4</p>	<ul style="list-style-type: none"> • poly(A) RNA binding • RNA binding • structural constituent of ribosome <p>http://www.uniprot.org/uniprot/P26373#P26373-1</p>
<p>Leucine zipper protein 1 OS=Homo sapiens GN=LUZP1 PE=1 SV=2</p>	<ul style="list-style-type: none"> • artery development • neural fold bending • ventricular septum development <p>http://www.uniprot.org/uniprot/Q86V48</p> <ul style="list-style-type: none"> • This gene encodes a protein that contains a leucine zipper motif. The exact function of the encoded protein is not known. In mice this gene affects neural tube closure. Alternative splicing results in multiple transcript variants. <p>http://www.ncbi.nlm.nih.gov/gene/7798</p>
<p>Unconventional myosin-Ig OS=Homo sapiens GN=MYO1G PE=1 SV=2</p>	<ul style="list-style-type: none"> • Unconventional myosin required during immune response for detection of rare antigen-presenting cells by regulating T-cell migration. Unconventional myosins are actin-based motor molecules with ATPase activity and serve in intracellular movements. Acts as a regulator of T-cell migration by generating membrane tension, enforcing cell-intrinsic meandering search, thereby enhancing detection of rare antigens during lymph-node surveillance, enabling pathogen eradication. Also required in B-cells, where it regulates different membrane/cytoskeleton-dependent processes. Involved in Fc-gamma receptor (Fc-gamma-R) phagocytosis. • Minor histocompatibility antigen HA-2: Constitutes the minor histocompatibility antigen HA-2. More generally, minor histocompatibility antigens (mHags) refer to immunogenic peptide which, when complexed with MHC, can generate an immune response after recognition by specific T-cells. The peptides are derived from polymorphic intracellular proteins, which are cleaved by normal pathways of antigen processing. The binding of these peptides to MHC class I or class II molecules and their expression on the cell surface can stimulate T-cell responses and thereby trigger graft rejection or graft-versus-host disease (GVHD) after

	<p>hematopoietic stem cell transplantation from HLA-identical sibling donor. GVHD is a frequent complication after bone marrow transplantation (BMT), due to mismatch of minor histocompatibility antigen in HLA-matched sibling marrow transplants. HA-2 is restricted to MHC class I HLA-A*0201</p> <ul style="list-style-type: none"> • ATP Binding • Motor Activity • phosphatidylinositol-3,4,5-trisphosphate binding • phosphatidylinositol-3,4-bisphosphate binding • phosphatidylinositol-4,5-bisphosphate binding <p>http://www.uniprot.org/uniprot/B011T2</p>
<p>60S ribosomal protein L21 OS=Homo sapiens GN=RPL21 PE=1 SV=2</p>	<ul style="list-style-type: none"> • poly(A) RNA binding • RNA binding • structural constituent of ribosome <p>http://www.uniprot.org/uniprot/P46778</p>
<p>Keratin, type I cytoskeletal 14 OS=Homo sapiens GN=KRT14 PE=1 SV=4</p>	<ul style="list-style-type: none"> • keratin filament binding • structural constituent of cytoskeleton • The nonhelical tail domain is involved in promoting KRT5-KRT14 filaments to self-organize into large bundles and enhances the mechanical properties involved in resilience of keratin intermediate filaments in vitro. <p>http://www.uniprot.org/uniprot/P02533</p>
<p>40S ribosomal protein S13 OS=Homo sapiens GN=RPS13 PE=1 SV=2</p>	<ul style="list-style-type: none"> • mRNA binding • poly(A) RNA binding • small ribosomal subunit rRNA binding • structural constituent of ribosome <p>http://www.uniprot.org/uniprot/P62277</p>
<p>Keratin, type II cytoskeletal 5 OS=Homo sapiens GN=KRT5 PE=1 SV=3</p>	<ul style="list-style-type: none"> • scaffold protein binding • structural constituent of cytoskeleton <p>http://www.uniprot.org/uniprot/P13647</p>
<p>SWI/SNF complex subunit SMARCC1 OS=Homo sapiens GN=SMARCC1 PE=1 SV=3</p>	<p>SWI/SNF</p>
<p>Isoform 2 of Annexin A2 OS=Homo sapiens GN=ANXA2</p>	<ul style="list-style-type: none"> • calcium-dependent phospholipid binding • calcium-dependent protein binding • calcium ion binding • phosphatidylinositol-4,5-bisphosphate binding • phospholipase A2 inhibitor activity • poly(A) RNA binding • protease binding • S100 protein binding

	http://www.uniprot.org/uniprot/P07355
Isoform 2 of Transcription activator BRG1 OS=Homo sapiens GN=SMARCA4	<ul style="list-style-type: none"> • Transcriptional coactivator cooperating with nuclear hormone receptors to potentiate transcriptional activation. • Component of the CREST-BRG1 complex, a multiprotein complex that regulates promoter activation by orchestrating a calcium-dependent release of a repressor complex and a recruitment of an activator complex. http://www.uniprot.org/uniprot/P51532#P51532-2
Keratin, type II cytoskeletal 7 OS=Homo sapiens GN=KRT7 PE=1 SV=5	<ul style="list-style-type: none"> • Blocks interferon-dependent interphase and stimulates DNA synthesis in cells. Involved in the translational regulation of the human papillomavirus type 16 E7 mRNA (HPV16 E7). • Structural Molecule Activity http://www.uniprot.org/uniprot/P08729
60S ribosomal protein L14 OS=Homo sapiens GN=RPL14 PE=1 SV=4	<ul style="list-style-type: none"> • poly(A) RNA binding • RNA binding • structural constituent of ribosome http://www.uniprot.org/uniprot/P50914
Isoform 2 of Synaptopodin OS=Homo sapiens GN=SYNPO	<p>Actin-associated protein that may play a role in modulating actin-based shape and motility of dendritic spines and renal podocyte foot processes. Seems to be essential for the formation of spine apparatuses in spines of telencephalic neurons, which is involved in synaptic plasticity (By similarity).</p> http://www.uniprot.org/uniprot/Q8N3V7
Tropomodulin-3 OS=Homo sapiens GN=TMOD3 PE=1 SV=1	<ul style="list-style-type: none"> • Blocks the elongation and depolymerization of the actin filaments at the pointed end. The Tmod/TM complex contributes to the formation of the short actin protofilament, which in turn defines the geometry of the membrane skeleton (By similarity). • tropomyosin binding http://www.uniprot.org/uniprot/Q9NYL9
40S ribosomal protein S23 OS=Homo sapiens GN=RPS23 PE=1 SV=3	<ul style="list-style-type: none"> • poly(A) RNA binding • structural constituent of ribosome http://www.uniprot.org/uniprot/P62266
60S ribosomal protein L32 OS=Homo sapiens GN=RPL32 PE=1 SV=2	<ul style="list-style-type: none"> • poly(A) RNA binding • structural constituent of ribosome http://www.uniprot.org/uniprot/P62910
AT-rich interactive domain-containing protein 1A OS=Homo sapiens GN=ARID1A PE=1 SV=3	<ul style="list-style-type: none"> • DNA binding • ligand-dependent nuclear receptor binding • transcription coactivator activity http://www.uniprot.org/uniprot/O14497

<p>40S ribosomal protein S18 OS=Homo sapiens GN=RPS18 PE=1 SV=3</p>	<ul style="list-style-type: none"> • poly(A) RNA binding • rRNA binding • structural constituent of ribosome <p>http://www.uniprot.org/uniprot/P62269</p>
<p>Drebrin OS=Homo sapiens GN=DBN1 PE=1 SV=4</p>	<p>Drebrins might play some role in cell migration, extension of neuronal processes and plasticity of dendrites. Required for actin polymerization at immunological synapses (IS) and for CXCR4 recruitment to IS.</p> <p>http://www.uniprot.org/uniprot/Q16643</p>
<p>RecName: Full=Gelsolin; AltName: Full=AGEL; AltName: Full=Actin-depolymerizing factor; Short=ADF; AltName: Full=Brevin; Flags: Precursor</p>	<p>Calcium-regulated, actin-modulating protein that binds to the plus (or barbed) ends of actin monomers or filaments, preventing monomer exchange (end-blocking or capping). It can promote the assembly of monomers into filaments (nucleation) as well as sever filaments already formed.</p> <p>http://moma.ki.au.dk/genome-mirror/cgi-bin/hgGene?org=Human&hgg_gene=uc004blf.1&hgg_chrom=none&db=hg19</p>
<p>Isoform 2 of SWI/SNF-related matrix-associated actin-dependent regulator of chromatin subfamily E member 1 OS=Homo sapiens GN=SMARCE1</p>	<p>SWI/SNF</p>
<p>EF-hand domain-containing protein D2 OS=Homo sapiens GN=EFHD2 PE=1 SV=1</p>	<ul style="list-style-type: none"> • Calcium ion binding <p>http://www.uniprot.org/uniprot/Q96C19</p>
<p>Neurabin-2 OS=Homo sapiens GN=PPP1R9B PE=1 SV=2</p>	<ul style="list-style-type: none"> • Scaffold Protein in Multiple Signaling Pathways • actin filament binding • protein phosphatase 1 binding • protein phosphatase inhibitor activity <p>http://www.uniprot.org/uniprot/Q96SB3</p>
<p>Flotillin-2 OS=Homo sapiens GN=FLOT2 PE=1 SV=2</p>	<ul style="list-style-type: none"> • May act as a scaffolding protein within caveolar membranes, functionally participating in formation of caveolae or caveolae-like vesicles. May be involved in epidermal cell adhesion and epidermal structure and function. • protein heterodimerization activity <p>http://www.uniprot.org/uniprot/Q14254</p>
<p>Keratin, type II cytoskeletal 6A OS=Homo sapiens GN=KRT6A PE=1 SV=3</p>	<ul style="list-style-type: none"> • Epidermis-specific type I keratin involved in wound healing. Involved in the activation of follicular keratinocytes after wounding, while it does not play a major role in keratinocyte proliferation or migration. Participates in the regulation of epithelial migration by inhibiting the activity of SRC during wound repair.

	<ul style="list-style-type: none"> • Structural Constituent of Cytoskeleton http://www.uniprot.org/uniprot/P02538
X-ray repair cross-complementing protein 5 OS=Homo sapiens GN=XRCC5 PE=1 SV=3	<ul style="list-style-type: none"> • ATP binding • ATP-dependent DNA helicase activity • damaged DNA binding • DNA binding • double-stranded DNA binding • poly(A) RNA binding • protein C-terminus binding • telomeric DNA binding • transcription regulatory region DNA binding • ubiquitin protein ligase binding http://www.uniprot.org/uniprot/P13010
X-ray repair cross-complementing protein 6 OS=Homo sapiens GN=XRCC6 PE=1 SV=2	<ul style="list-style-type: none"> • 5'-deoxyribose-5-phosphate lyase activity • ATP binding • ATP-dependent DNA helicase activity • damaged DNA binding • DNA binding • double-stranded DNA binding • poly(A) RNA binding • protein C-terminus binding • telomeric DNA binding • transcription regulatory region DNA binding http://www.uniprot.org/uniprot/P12956
Isoform B of SWI/SNF-related matrix-associated actin-dependent regulator of chromatin subfamily B member 1 OS=Homo sapiens GN=SMARCB1	SWI/SNF
Myosin regulatory light chain 12B OS=Homo sapiens GN=MYL12B PE=1 SV=2	<ul style="list-style-type: none"> • Myosin regulatory subunit that plays an important role in regulation of both smooth muscle and nonmuscle cell contractile activity via its phosphorylation. Phosphorylation triggers actin polymerization in vascular smooth muscle. Implicated in cytokinesis, receptor cap ping, and cell locomotion (By similarity). • Calcium Ion Binding http://www.uniprot.org/uniprot/O14950
Isoform 2 of Actin-like protein 6A OS=Homo sapiens GN=ACTL6A	<ul style="list-style-type: none"> • chromatin binding • transcription coactivator activity http://www.uniprot.org/uniprot/O96019
Isoform 2 of Supervillin OS=Homo sapiens GN=SVIL	Forms a high-affinity link between the actin cytoskeleton and the membrane. Isoform <u>2</u> may be involved in modulation of focal adhesions. Supervillin-mediated down-regulation of focal adhesions involves binding to

	<p>TRIP6. Plays a role in cytokinesis through KIF14 interaction (By similarity).</p> <p>http://www.uniprot.org/uniprot/O95425</p>
<p>Isoform Short of Probable global transcription activator SNF2L2 OS=Homo sapiens GN=SMARCA2</p>	<ul style="list-style-type: none"> • ATP binding • chromatin binding • DNA-dependent ATPase activity • helicase activity • RNA polymerase II transcription coactivator activity • transcription coactivator activity • transcription regulatory region DNA binding <p>http://www.uniprot.org/uniprot/P51531</p>
<p>Isoform 2 of SWI/SNF-related matrix-associated actin-dependent regulator of chromatin subfamily D member 1 OS=Homo sapiens GN=SMARCD1</p>	<p>Chromatin Remodeling</p> <p>http://www.uniprot.org/uniprot/Q96GM5</p>