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Comparisons of Mass Spectrometry Compatible Surfactants for Global Analysis of the Mammalian Brain Proteome

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Abstract

Methods for the global analysis of protein expression offer an approach to study the molecular basis of disease. Studies of protein expression in tissue, such as brain, are complicated by the need for efficient and unbiased digestion of proteins that permit identification of peptides by shotgun proteomic methods. In particular, identification and characterization of less abundant membrane proteins has been of great interest for studies of brain physiology, but often proteins of interest are of low abundance or exist in multiple isoforms. Parsing protein isoforms as a function of disease will be essential. In this study, we develop a digestion scheme using detergents compatible with mass spectrometry that improves membrane protein identification from brain tissue. We show the modified procedure yields close to 5,000 protein identifications from 1.8 mg of rat brain homogenate with an average of 25% protein sequence coverage. This procedure achieves a remarkable reduction in the amount of starting material required to observe a broad spectrum of membrane proteins. Among the proteins identified from a mammalian brain homogenate, 1897 (35%) proteins are annotated by GeneOntology as membrane proteins, and 1225 (22.6%) proteins are predicted to contain at least one transmembrane domain. Membrane proteins identified included neurotransmitter receptors and ion channels implicated in important physiological functions and disease.

Keywords

brain proteome; mass spectrometry; membrane proteins; proteolysis; shotgun proteomics

INTRODUCTION

The brain is a complex system of thousands of proteins that form a complicated molecular network. Large-scale proteomic analysis can help unravel the complexities of brain function as many of the activities of the brain involve intricate signaling networks and changes in post-translational modifications. Although, bottom up shotgun sequencing is useful for proteome analysis, the complexity of the mammalian brain presents technical and biological challenges. The brain consists of many distinct functional regions, each of which consists of different cell types. The structural morphology between these regions exacerbate the complexity of the biology. For example, the cell body and axon terminal of the same neuron are often located in two different functional regions. In addition, the brain has the highest lipid content of any other tissue (excluding adipose) making it particularly challenging for mass spectrometry-based proteomic analysis¹. Therefore, new approaches to prepare brain tissue are needed to improve the potential of mass spectrometry-based shotgun proteomic technology for the study of disease.

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Gel-based proteomic methods, such as two-dimensional polyacrylamide gel electrophoresis (2D-PAGE), have been used to characterize the brain proteome²⁻⁴. Membrane proteins are typically underrepresented in 2D PAGE analyses due to poor solubility in the starting buffer⁵. A proteomic analysis of whole mouse brain using 2D-PAGE required large amounts of starting material (1–15 g) and yielded less than 500 protein identifications⁶. To increase coverage of the proteome and, in particular, the coverage of membrane proteins, shotgun strategies using multidimensional liquid chromatography separation coupled with tandem mass spectrometry (LC/LC-MS/MS) have been applied to the brain⁷. By using high pH and proteinase K digestion of protein mixtures in combination with LC/LC-MS/MS, Wu et al. reported the identification of 1,610 proteins from 500 µg of rat brain homogenate⁸. Nielsen et al. reported a novel method to enrich membrane proteins from 150 mg of mouse cortex for analysis by LC-MS/MS⁹. 862 proteins were identified from sixteen cortex fractions. They also applied the same technique to analyze proteins extracted from 15 mg of mouse hippocampus and identified 1685 proteins. Recently, Wang et al. reported a method of analyzing both global and cysteinyl-enriched peptides from a whole brain sample¹⁰. A total of 48,328 peptides were identified covering 7792 non-redundant proteins from 200–300 mg of brain homogenate. Among the 7792 proteins, 5636 proteins were identified at a ~1.5% false positive rate estimated by reverse database searching using a minimum of two or more peptides for identification, and 1332 proteins were annotated as membrane proteins. The identification of total and membrane proteins from brain tissue has increased dramatically using bottom up approaches, but a large amount of starting material is usually required for identification of hydrophobic membrane proteins. Although it might be feasible to obtain a large amount of starting material by pooling brain tissue from different animals, it poses a challenge to the number of biological replicates that can be performed for each study and studies of normal versus diseased tissues. A reduction of starting material without improving sample processing and/or analytical procedures usually results in a decrease of protein sequence coverage and thus confidence in identified proteins. Hence, an important issue to reducing the amount of starting material without decreasing protein identifications is to improve the processing of protein mixtures.

Previously, we demonstrated the utility of mass spectrometry (MS) compatible detergents for shotgun analysis of complex protein mixtures¹¹. We showed that a modified trypsin digestion protocol incorporating detergents compatible with mass spectrometry improves proteolytic processing of protein mixtures and results in a dramatic reduction in the amount of material required for proteome analysis. Three commercially available mass spectrometry compatible detergents were compared: RapiGest SF (RPG), Invitrosol (IVS), and PPS Silent Surfactant (PPS). RapiGest SF and PPS Silent Surfactant are acid-labile surfactants. Invitrosol (IVS) is a homogeneous surfactant that elutes in three peaks well separated from the elution times of most peptides. All three surfactants can act as mild denaturants to solubilize and unfold hydrophobic proteins at concentrations that do not inhibit common proteolytic enzymes such as trypsin. In addition to solubilizing and denaturing hydrophobic proteins in solution, we also observed a complementarity to the proteins identified using the different conditions suggesting a strategy to increase coverage of a proteome. In this study, we use this approach to improve global profiling of brain tissue, and more importantly the detection of membrane proteins. As membrane proteins play an important role in brain function, the ability to observe this class of proteins is critical for the study of diseases in brain.

MATERIALS AND METHODS

Materials

InvitrosolTM was purchased from Invitrogen (Carlsbad, CA). RapiGestTM SF acid labile surfactant was purchased from Waters Corp. (Milford, MA). PPSTM Silent surfactant was

provided by Dr. Norris from Protein Discovery Inc. (Knoxville, TN). Trypsin (modified, sequencing grade) was obtained from Roche. Other laboratory reagents were purchased from Sigma-Aldrich (St. Louis, MO) unless noted otherwise.

Sample Preparation

A Sprague-Dawley rat at postnatal day 45 was sacrificed and the brain was quickly removed, and then frozen with liquid nitrogen. One entire rat brain was homogenized in a Teflon dounce grinder in 1–2 ml of 0.32M sucrose, 4mM HEPES, and protease inhibitors (Roche Applied Science). The protein concentration was determined using a BCA protein assay (Pierce).

Digestion protocols

300 µg of cell lysates were precipitated with methanol/chloroform and resuspended in different conditions for trypsin digestion. Each digestion condition (300µg) was run in triplicates (100µg).

Trypsin digestion with the MS detergents—40 µl of invitrosol (5X stock), 20 µl of RapiGest SF (1% stock), and 20 µl of PPS (1% stock) were used to resuspend the protein pellets. The concentration of each detergent used in this study was determined based on the maximum recommended concentration suggested by the manufactures. Then, proteins were incubated at 60°C for 5 minutes and the remaining solvent was added to the samples (200µl final volume). All samples were sonicated for 2 hours in a water bath and digested with trypsin for 16 hours at 37°C. For trypsin digestion in aqueous Tris buffer, samples were reduced in 5mM TCEP for 20 minutes at room temperature, and cysteines were alkylated in the dark in 25mM iodoacetamide (IAM) for 30 minutes at room temperature prior to the addition of trypsin. Trypsin was added to each sample at a ratio of 1:30 enzyme/protein along with 2 mM CaCl₂.

Post-Digestion

Following digestion, all reactions were acidified with 90% formic acid (2% final) to stop proteolysis. Samples with RapiGest SF and PPS were acidified and incubated at 37°C for additional 4 hours to facilitate the hydrolysis of detergents. Then, samples were centrifuged for 30 minutes at 14,000 rpm to remove insoluble material. The soluble peptide mixtures were collected for LC-MS/MS analysis.

Multidimensional chromatography and tandem mass spectrometry

Peptide mixtures were pressure-loaded onto a 250 µm inner diameter (i.d.) fused-silica capillary packed first with 2.5 cm of 5 µm strong cation exchange material (Partisphere SCX, Whatman), followed by 5 cm of 10 µm C12 reverse phase (RP) particles (Jupitor, Phenomenex or Polaris 2000, Metachem Technologies). Loaded and washed microcapillaries were connected *via* a 2 µm filtered union (UpChurch Scientific) to a 100 µm i.d. column, which had been pulled to a 5 µm i.d. tip using a P-2000 CO₂ laser puller (Sutter Instruments), then packed with 13 cm of 4 µm C12 reverse phase (RP) particles (Jupitor, Phenomenex or Polaris 2000, Metachem Technologies) and equilibrated in 5% acetonitrile, 0.1 % formic acid (Buffer A). This split-column was then installed in-line with a Quaternary Agilent 1100 series HPLC pump. Overflow tubing was used to decrease the flow rate from 0.1 ml/min to about 200–300 nl/min. Fully automated 13-step chromatography runs were carried out. 110-minute gradient was used for each chromatographic step. Three different elution buffers were used: 5% acetonitrile, 0.1 % formic acid (Buffer A); 80% acetonitrile, 0.1% formic acid (Buffer B); and 0.5 M ammonium acetate, 5% acetonitrile, 0.1% formic acid (Buffer C). In such sequences of chromatographic events, peptides are sequentially eluted from the SCX resin to the RP resin by increasing salt steps (increase in Buffer C concentration), followed by organic gradients (increase in Buffer

B concentration). The last chromatography step consists in a high salt wash with 100% Buffer C followed by acetonitrile gradient. The application of a 2.5 kV distal voltage electrosprayed the eluting peptides directly into an LTQ ion trap mass spectrometer equipped with a nano-LC electrospray ionization source (Thermo Finnigan). Full MS spectra were recorded on the peptides over a 400 to 2000 m/z range, followed by five tandem mass (MS/MS) events sequentially generated in a data-dependent manner on the first, second, third, and fourth most intense ions selected from the full MS spectrum (at 35% collision energy). Mass spectrometer scan functions and HPLC solvent gradients were controlled by the Xcalibur data system (Thermo Finnigan, San Jose, CA).

Database search and interpretation of MS/MS datasets

Tandem mass spectra were extracted from raw files, and a binary classifier - previously trained on a manually validated data set - was used to remove the low quality MS/MS spectra. The remaining spectra were searched against a *Rat* protein database containing 38,841 protein sequences downloaded as FASTA-formatted sequences from EBI-IPI (database version 3.17, released on May, 18, 2006)¹², and 124 common contaminant proteins, for a total of 38,965 target database sequences. To calculate confidence levels and false positive rates, we used a decoy database containing the reverse sequences of 38,965 proteins appended to the target database 13, and the SEQUEST algorithm 14-15 to find the best matching sequences from the combined database.

SEQUEST searches were done on an Intel Xeon 80-processor cluster running under the Linux operating system. The peptide mass search tolerance was set to 3 Da. No differential modifications were considered. For the aqueous digestion, the mass of the amino acid Cysteine was statically modified by +57 Da, due to carboxyamidomethylation of the sample. No enzymatic cleavage conditions were imposed on the database search, so the search space included all candidate peptides whose theoretical mass fell within the 3 Da mass tolerance window, despite their tryptic status.

The validity of peptide/spectrum matches was assessed in DTASelect2¹⁶ using SEQUEST-defined parameters, the cross-correlation score (XCorr) and normalized difference in cross-correlation scores (DeltaCN). The search results were grouped by charge state (+1, +2, and +3) and tryptic status (fully tryptic, half-tryptic, and non-tryptic), resulting in 9 distinct sub-groups. In each one of the sub-groups, the distribution of XCorr and DeltaCN values for (a) direct and (b) decoy database hits was obtained, and the two subsets were separated by quadratic discriminant analysis. Outlier points in the two distributions (for example, matches with very low Xcorr but very high DeltaCN) were discarded. Full separation of the direct and decoy subsets is not generally possible; therefore, the discriminant score was set such that a false positive rate of 1% was determined based on the number of accepted decoy database peptides. This procedure was independently performed on each data subset, resulting in a false positive rate independent of tryptic status or charge state.

In addition, a minimum sequence length of 7 amino acid residues was required, and each protein on the list was supported by at least two peptide identifications. These additional requirements – especially the latter - resulted in the elimination of most decoy database and false positive hits, as these tended to be overwhelmingly present as proteins identified by single peptide matches. After this last filtering step, the false identification rate was reduced to below 1%. Finally, the Kyte-Doolittle hydropathy score (KD) of all peptides passed the filters were also calculated and reported in the DTASelect2 output.

The identified proteins were categorized based on their cellular localizations and molecular functions accordingly to Gene Ontology (GO) information obtained from the European Bioinformatics Institute at <http://www.ebi.ac.uk/pub/databases/GO/goa/rat>. The TMHMM

(www.cbs.dtu.dk/services/TMHMM/) algorithm was used to predict transmembrane domains (TMDs) from the identified proteins.

Gene Ontology Data Analysis

We assigned cellular components and molecular functions based on the Gene Ontology (GO) consortium using GoMiner. For each GO category, Fisher's exact test was used to calculate the p-value that reflects the significance over-representation of proteins identified in one GO category compared to the proteins identified from the entire protein database in the same GO category. The categories with p-values less than 0.05 were considered significantly over represented or under represented.

RESULTS

The initial step in shotgun proteomic experiments is proteolytic digestion of a complex protein mixture. The challenge is to disrupt existing protein-protein interactions, and denature and solubilize the proteins. Samples of tissue represent a bigger challenge because in addition to a diverse mixture of cell types there are extracellular components present. We showed previously that RPG, IVS, and PPS when used with organic or aqueous solvents can greatly increase protein coverage in complex cellular protein mixtures¹¹. In this study, we show this procedure can increase the identification of membrane proteins from brain tissue. Rat brain tissue was homogenized and the insoluble material was extracted. The material was divided into 6 aliquots of 300 µg each and dissolved in RPG (0.1%), IVS (1X) and PPS (0.2%) mixed with either mixed organic-aqueous (80% acetonitrile) or aqueous buffers (Tris-HCl, pH 8.0) for digestion. A total of 6 different digestion conditions were used for comparison, and each digestion condition was analyzed in triplicate (100 µg per replicate) with an identical experimental setup. A schematic illustration of the sample preparation is shown in Figure 1. The concentration of MS compatible detergents used in this study differs from our previous report¹¹. To increase the solubility of hydrophobic and membrane proteins in the rat brain homogenate in solution, we used the maximum concentration of each detergent recommended by the manufacture to solubilize the protein pellet after the methanol/chloroform precipitation: 5X invitrosol, 1% RapiGest, and 1% PPS. This is a crucial step to ensure the inclusion of membrane proteins known to be important for the normal brain function. After the initial solubilization step, protein samples were diluted further with trypsin digestion buffer to achieve the optimal concentration of the MS compatible detergents for trypsin digestion. Detail protocols for sample preparation are described in the method section.

A total of 18 MudPIT analyses were performed. Tandem mass spectra collected in each experiment were searched using SEQUEST against the rat IPI protein database plus a decoy database (see Methods section). A 1% peptide false discovery rate and a minimum of two-peptide matches to a protein were required for protein identification. Following database search and data filtering, a total of 60,712 peptides that corresponded to 4,797 proteins were identified from 1.8 mg of rat brain homogenate using significantly less starting material compared to similar studies reported previously (Supplementary table 1). By using our method, the number of proteins identified was much improved by combining the different specificities of the detergents used to solubilize the protein mixtures than the technical replicates with the same detergent. The optimized digestion protocol employing different types of detergents led to increased or equivalent protein identifications from less material than reported in the literature.

Increases in Overall Sequence Coverage and Representation of Proteins in the Brain Proteome

The complementary nature of this sample processing strategy increases the protein sequence coverage and thus the confidence of protein identification. In complex protein mixtures such

as tissue homogenates, sequence coverage levels tend to be less overall (e.g. <10%) and even more so for less abundant proteins often having only one or two peptide matches ⁸⁻¹⁰. In our study, a majority (71%) of proteins identified have more than 10% sequence coverage (Figure 2). We found it's an effective strategy to increase protein sequence coverage by combining trypsin digestion with a combination of solvents and detergents. Over 57% of the proteins show at least a 10% increase in sequence coverage by using the three combinations of mixed solvents and detergents compared with triplicates of the same trypsin digestion conditions. Hence, we demonstrate that the modified digestion protocol can be used to improve protein identification in a proteome as complex as the brain compared to methods without detergent (three times less protein identifications, data not shown) and other shotgun-based brain proteome studies ⁸⁻¹⁰.

Improved Identification of Brain Membrane Proteins

Many transmembrane and membrane-associated proteins in the brain form neurotransmitter receptors, ion channels, and G-proteins-couple receptors that are of interest because of their roles in the normal physiology of the brain and the neurological diseases ¹⁷. The hydrophobic nature of membrane proteins makes them difficult to study because of difficulty solubilizing and digesting the proteins. Gel-based proteomic studies using enriched brain membrane fractions identified mostly cytosolic proteins in the brain ^{2, 18}. In other studies using a bottom-up approach, a range from 454 to 1447 (18.6%) transmembrane proteins have been identified from brain tissues ⁸⁻¹⁰.

In our previous study, the modified digestion protocol with MS compatible detergents yielded an increase in total number of proteins identified by improving the efficiency of proteolysis. In this study, we modified the digestion protocol to increase the solubility of membrane proteins. First, our results show good coverage of regions rich in transmembrane proteins. We observe roughly a 2-fold increase in proteins identified with molecular weight greater than 100kDa in our dataset (22%) as compared to the predicted rat proteome (12%). Next, we classified proteins identified from each modified trypsin digestion condition using the TMHMM algorithm ¹⁹ and GeneOntology (GO) ²⁰ to determine the representation of membrane proteins in our data set (Table 1). After merging the data from all six proteolytic conditions, 4,797 proteins were identified from 1.8mg of rat brain homogenate, 1897 (40%) proteins are annotated as membrane proteins, and 1225 (26%) proteins predicted to contain at least one transmembrane domain. The improved digestion strategy increases the identification of transmembrane proteins with 10–100 fold less starting material compared to other shotgun brain proteome studies.

We also observe that the use of different detergents yields complementary information from membrane proteins leading to higher overall sequence coverage. First, the highest number of brain transmembrane proteins was identified in an aqueous protein digestion with RapiGest (Tris RPG, 556) followed by an aqueous protein digestion with Invitrosol (Tris IVS, 423) (Table 1). Interestingly, the different solvent compositions used with RapiGest generated both the highest number of identified transmembrane proteins (Tris RPG, 556 proteins) and lowest (ACN RPG, 450 proteins). This result is consistent with our previous observation that there is a solvent preference for the detergents ¹¹. Trypsin digestion in Tris buffer with the acid labile detergents RPG and PPS resulted in more identified transmembrane proteins than digestion in buffer with 80% acetonitrile. In this study, aqueous Tris RPG digestion increases transmembrane protein identifications by 24% over RPG digestion in 80% acetonitrile. Furthermore, we found interesting differences in peptide hydrophobicity. Aqueous digestions in combination with the MS-compatible detergents yielded more hydrophobic peptides (lower negative KD value) than organic solvent digestions (Figure 3). In particular, Tris PPS digestion yields peptides with the highest hydrophobicity (−2.86) followed by Tris RPG (−3.76) and Tri

INV (−3.78). Therefore, the impact of buffer composition in combination with the detergents provides valuable information to identify the optimal digestion condition of brain membrane proteins.

Improving protein sequence coverage is particularly important for identification of brain transmembrane proteins because they are usually less abundant, hard to digest, and have a variety of isoforms. In table 2, we list twelve identified GABA-A receptors/transporters and sixteen glutamate receptors. These proteins are important for normal brain functions and are multi-span transmembrane proteins. The percentage sequence coverage of each protein identified using each set of digestion conditions (triplicate runs) is listed and compared. We notice that the amino acid sequence coverage of these membrane proteins varies in different digestion conditions after adding together peptides from triplicate runs and may reflect protein solubility in combinations of solvents and MS compatible detergents (Table 2). For example, Gamma-aminobutyric-acid receptor gamma-1 subunit and Gamma-aminobutyric-acid type B receptor subunit 1 (spliced isoform 1E) were only identified in the PRG and PPS digestions in the aqueous solution. On the other hand, Gamma-aminobutyric-acid receptor alpha-5 subunit was identified exclusively in the INV and RPG digestions in the 80% acetonitrile (Table 2).

In addition to providing more confident identification, high levels of sequence coverage are necessary to differentiate isoforms of proteins. For example, an analysis of peptides used to identify three transmembrane proteins, Sodium- and chloride-dependent GABA transporter 1, GPI45-TrkB receptor, and excitatory amino acid transporter 1, shows extensive sequence coverage of the extracellular ligand binding domains and the intracellular signaling domains as a result of the unique and complementary mixture of peptides generated by digestion in combinations of detergents and solvents. Additionally, peptides generated by digesting the excitatory amino acid transporter 1 with Invitrosol in 80% acetonitrile extend into the predicted transmembrane domains of the protein (Supplementary figure 1). In fact, we found 2.8% of total peptides identified contain amino acids extending into the transmembrane region. We find enrichment of peptides containing the transmembrane domains with a significant increase in the number of peptides containing transmembrane regions compared to Wu et al using Proteinase K digestion⁸. Most of these peptides are found in the aqueous digestion with RapiGest (Tris RPG, 88 peptides), followed by the organic digestion with Invitrosol (ACN IVS, 86 peptides), and the aqueous protein digestion with Invitrosol (Tris IVS, 73 peptides) (Table 3). We observe that use of organic solvents with Invitrosol yields the highest percentage of peptides containing transmembrane sequences over the total number of transmembrane proteins identified (20.3%) (Table 3). In other words, trypsin digestion of transmembrane proteins with invitrosol in 80% acetonitrile shows an increasing trend of generating peptides containing transmembrane regions comparing to other digestion conditions. This trend has been previously observed²¹. It suggests that perturbing the lipid bilayer near the transmembrane region in the presence of high percentage organic solvent facilitates the proteolysis of the helical regions of transmembrane proteins. In summary, our results demonstrate that MS compatible detergents provide a valuable tool to increase the identification and sequence coverage of brain membrane proteins, including proteins localized to the synapse (discussed below).

Physiological Relevance of the Identified Brain Proteome

Annotation of the data using GeneOntologies revealed that around 35% (1897 proteins) of the identified proteins are classified as membrane proteins. To explore the physiological relevance of the identified proteins, we clustered proteins identified in our data based on GeneOntology (GO) and identify several GO categories with a significant enrichment ($p < 0.05$) compared to the entire rat proteome. For example, we observed a three to four fold enrichment of proteins localized to the presynaptic membrane, synaptic vesicle membrane, synapse, and synaptosome

(Supplementary figure 2A). This brain-specific compartment receives signals, in the form of neurotransmitters, from the presynaptic terminal of an adjacent neuron. The postsynaptic and presynaptic termini are the signaling center between two neurons, and consist of cell-surface receptors and a large variety of signaling pathways^{22, 23}. In addition to proteins with synaptic functions, we also observed a six and five-fold increase in proteins involved in glutamate transmission and transport activities, which is consistent with the fact that the major excitatory neurotransmitter in the brain is glutamate (Supplementary figure 2B). Glutamate receptors can be broadly divided into ligand-gated ion channels and G-protein coupled receptors. The three major subtypes of ionotropic receptors are AMPA (alpha-amino-3-hydroxy-5-methyl-ioxazole-4-propionic acid), kainate, and NMDA (N-methyl-D-aspartate). In addition, there are two orphan receptors. In total, this family is represented by seventeen genes, and in our analysis, we identified ten of these ionotropic glutamate receptors (Table 2). In addition, we identified six of the eight G-protein coupled glutamate receptors (Table 2). Besides the glutamate receptors, we also identified other classes of membrane proteins essential for maintaining brain functions such as neurotransmitter receptors, ion channels, and transporters (Supplementary table 2). Together, our results demonstrate the utility of our method in identifying membrane proteins essential for the function of the brain.

DISCUSSION

Using total brain homogenate, we identified 5,422 proteins, including 1225 (22.6%) proteins that contained at least one transmembrane domain and 1897 (35%) proteins annotated as membrane proteins. Among these proteins, we identified many proteins currently under intense investigation in neuroscience. For example, we identified numerous proteins involved in synaptic function. The most notable proteins identified in this category are the glutamate and GABA receptors. Direct modulators of GABA receptors include benzodiazepines, barbiturates, neurosteroids and anesthetics, and altered GABA receptor expression has been strongly implicated in insomnia, anxiety, premenstrual syndrome, alcoholism, schizophrenia, depression and epilepsy^{51–54}. The glutamate receptor family plays important roles brain development and in some forms of synaptic plasticity that may underlie higher order processes, such as learning and memory. In terms of neurological disorders, glutamate is crucial to the process of neurodegeneration, and potential therapeutic effects of glutamate neuromodulators have been reported for Alzheimer's disease, Parkinson's disease, ischemic stroke, and neuropathic pain^{55–61}. In addition, we identified numerous membrane proteins, such as ion channels and amino acid transporters, known to play important roles in brain physiology.

In conclusion, we demonstrate that modification of trypsin digestion protocols with MS compatible detergents improves global and membrane protein identification in the mammalian brain. Furthermore, by coupling the trypsin digestion protocol with MS compatible detergents and high sensitivity LC-MS/MS analysis, it is possible to profile and quantify proteins from specific regions of the brain without pooling animal tissues, which often masks the natural variability between individual animals. Lastly, it is also possible to further increase the protein coverage by incorporating other protein digestion strategies such as proteinase K digestion. The combination of simple sample preparation, optimized trypsin digestion technique, and highly sensitive multidimensional LC-MS/MS analysis opens up new possibilities to proteomic-driven biological discoveries in normal brain physiology and neurological abnormality.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 2. Distribution of total sequence coverage in identified proteins. Figure 2 shows that over 60% of proteins identified have more than 10% protein sequence coverage.

	ACN IVS	ACN PPS	ACN RPG	Tris IVS	Tris PPS	Tris RPG
Average KD	-6.59	-3.91	-5.31	-3.78	-2.86	-3.76

Figure 3.

A summary of averaged hydrophobicity (KD) from peptides identified in six proteolytic conditions. The Kyte-Doolittle hydropathy score (KD) of peptides identified in each digestion condition was calculated and plotted in histograms. The average KD of peptides from each digestion condition is summarized in a table. The result shows that aqueous digestions in combination with the MS-compatible detergents yield more hydrophobic peptides (lower negative KD value) than organic solvent digestions.

Categorization of proteins identified in 6 different trypsin digestion conditions with MS compatible detergents based on the cellular localization of Gene Ontology (GO).

Table 1

		ACN IVS	ACN PPS	ACN RPG	Tris IVS	Tris PPS	Tris RPG
Predicted TM (%)		423 (16.0)	466 (16.9)	450 (16.3)	476 (17.8)	431 (15.0)	556 (16.9)
No Predicted TM							
	Integral to membrane	6	4	4	7	8	10
	Plasma membrane	30	27	26	29	27	34
	Mitochondrial membrane	27	30	33	33	32	37
	Nuclear membrane	18	14	17	15	6	8
	Golgi membrane	7	15	9	7	9	9
	ER membrane	6	10	5	6	5	9
	Lysosome membrane	1	1	1	1	1	1
	Membrane	128	140	142	139	151	162
	Cytoskeleton	120	120	114	119	116	128
	Cytoplasmic	344	319	335	318	362	381
	Nucleus	162	168	173	157	286	204
	Mitochondrion	84	95	85	82	84	96
	ER	21	23	23	21	25	26
	Endosome	4	5	5	5	3	4
	Lysosome	4	6	5	4	5	6
	Microsome	2	2	2	2	2	3
	Peroxisome	2	4	5	4	4	6
	Extracellular	29	30	33	20	25	25
No assignment		1227	1278	1302	1229	1295	1586
Total Protein (Spectra) Identified		2645 (23779)	2757 (22902)	2769 (21873)	2674 (22237)	2877 (19642)	3291 (25338)

* TM = transmembrane

Table 2

List of identified GABA-A receptors/transporters and glutamate receptors. Percentage sequence coverage (SeqCov) of each protein is listed in each of digestion condition. Total sequence coverage is calculated based on the collective peptides identified in all six conditions for each protein. x indicates no peptide identification for the listed protein.

Locus	Description	L	MW	pI	ACN INV SeqCov	ACN PPS SeqCov	ACN RPG SeqCov	Tris INV SeqCov	Tris PPS SeqCov	Tris RPG SeqCov
GABA-A receptors and transporters										
IP100192642.1	Gamma-aminobutyric-acid receptor alpha-1 subunit precursor	455	51754	9.3	10.3	13	11.2	x	6.6	11.2
IP100197343.1	Gamma-aminobutyric-acid receptor alpha-3 subunit precursor	493	55433	8.8	4.9	x	11	6.3	x	8.5
IP100325359.3	Gamma-aminobutyric-acid receptor alpha-5 subunit precursor	464	52337	9.1	6.7	x	4.1	x	x	x
IP100209268.1	Gamma-aminobutyric-acid receptor beta-1 subunit precursor	474	54072	8.8	11.2	8.6	6.1	11.2	10.1	x
IP100209269.1	Gamma-aminobutyric-acid receptor beta-2 subunit precursor	474	54634	9.3	11.2	x	x	13.7	10.8	16.7
IP100327083.3	Gamma-aminobutyric-acid receptor beta-3 subunit precursor	473	54166	9.1	18.8	14.2	x	x	x	19.7
IP100211960.1	Gamma-aminobutyric-acid receptor gamma-1 subunit precursor	465	53551	8	x	x	x	x	6.5	8.4
IP100192646.1	Gamma-aminobutyric-acid receptor gamma-2 subunit precursor	466	54077	8.5	9.9	15.9	9	6.4	x	13.1
IP100331966.6	Gamma-aminobutyric acid type B receptor, subunit 2 precursor	940	1E+05	8.7	3	2.9	2.7	3.8	x	3
IP100208182.1	Gamma-aminobutyric acid type B receptor, subunit 1 precursor, Splice Isoform 1E	991	1E+05	8	x	x	x	11.9	8.2	11
IP100187596.1	Sodium- and chloride-dependent GABA transporter 1	599	67002	8	13.2	13.2	12.4	13.4	3.5	8.5
IP100214462.1	Sodium- and chloride-dependent GABA transporter 3	627	69947	7	12.4	7.5	11.8	18	6.1	7.5
Glutamate Receptors										
IP100326054.1	Glutamate [NMDA] receptor subunit epsilon 1 precursor	1464	2E+05	7	x	x	2.3	2.7	4.9	4.6
IP100326061.1	Glutamate [NMDA] receptor subunit epsilon 2 precursor	1482	2E+05	6.9	7.3	5.7	4.5	6.6	3.9	7.6
IP100198625.5	Glutamate [NMDA] receptor subunit zeta 1 precursor, Splice Isoform A	938	1E+05	8.8	3.7	5.5	7.7	3.4	5.4	7.1
IP100207091.1	Glutamate receptor delta-1 subunit precursor	1009	1E+05	6.7	x	x	x	x	5.7	2.7
IP100206854.2	Glutamate receptor delta-2 subunit precursor	1007	1E+05	6.1	x	3.2	x	7.1	x	10
IP100231012.2	Glutamate receptor 1 precursor, Splice Isoform Flip	907	1E+05	7.7	8.6	x	x	10.9	6	8.6
IP100231061.5	Glutamate receptor 2 precursor, Splice Isoform Flip	883	98745	7.6	18.7	20.5	x	32.7	23.8	26
IP100231095.1	Glutamate receptor 3 precursor, Splice Isoform Flip	888	1E+05	8.3	x	5.6	3.8	19.3	x	x
IP100195445.1	Glutamate receptor 4 precursor, Splice Isoform 1	902	1E+05	7.9	x	x	x	9	x	4.9
IP100210260.1	Metabotropic glutamate receptor 1 precursor, Splice Isoform 1A	1199	1E+05	6.9	2.7	5.6	x	3.7	2	2.7

Locus	Description	L	MW	pI	ACN INV SeqCov	ACN PPS SeqCov	ACN RPG SeqCov	Tris INV SeqCov	Tris PPS SeqCov	Tris RPG SeqCov
IP100212618.1	Metabotropic glutamate receptor 2 precursor	872	95774	7.8	2.3	5	2.2	7.3	10.7	4.8
IP100212619.2	Glutamate receptor, metabotropic 3	936	1E+05	8.2	5.8	6.9	x	14	15	11.6
IP100212621.1	Metabotropic glutamate receptor 5 precursor, Splice Isoform 2	1203	1E+05	7.8	3.8	3.3	5.6	7.6	4.1	3.6
IP100198587.1	Metabotropic glutamate receptor 7 precursor	915	1E+05	7.9	x	3	3.9	3.1	5.6	5.7
IP100393591.2	Metabotropic glutamate receptor 8 precursor	910	1E+05	8.2	x	x	x	x	x	5.3
IP100231277.4	Splice Isoform GluR7A of Glutamate receptor, ionotropic kainate 3 precursor	910	1E+05	7.4	X	X	X	X	X	4.4
Neurotransmitter Receptors										
IP100187680.3	5-hydroxytryptamine 2C receptor	460	51917	9.3	x	x	x	5.2	x	x
IP100421955.1	Dopamine and cAMP-regulated phosphoprotein DARPP-32	205	22913	4.6	46.3	37.1	59	44.4	38	61
Ion Channels										
IP100365813.2	Voltage-dependent anion channel 1	234	25729	8.8	27.8	22.6	22.6	22.6	22.6	22.6
IP100421874.3	Voltage-dependent anion-selective channel protein 1	282	30624	8.5	75.9	76.2	70.2	82.3	79.1	77.7
IP100198327.2	Voltage-dependent anion-selective channel protein 2	295	31746	7.5	44.7	40.3	36.6	68.5	68.1	69.8
IP100207891.2	Voltage-dependent anion-selective channel protein 3 (VDAC-3) (rVDAC3)	284	30912	8.7	29.2	36.6	38.7	50.4	43.7	50
IP100190644.1	Potassium voltage-gated channel subfamily A member 1	495	56379	5.1	x	11.3	x	8.3	x	6.1
IP100208365.1	Potassium voltage-gated channel subfamily A member 2	499	56701	4.9	x	11.4	x	16.2	10	11.6
IP100208362.1	Potassium voltage-gated channel subfamily A member 4	655	73390	5.1	x	x	x	x	4.9	5.3
IP100190053.1	Potassium voltage-gated channel subfamily A member 6	530	58884	5.1	x	x	x	7.7	x	x
IP100394218.1	Potassium voltage-gated channel subfamily D member 2	630	70549	8.1	10.6	9.5	12.9	6.7	x	x
IP100389372.1	Potassium voltage-gated channel subfamily D member 3, Splice Isoform 1	655	73513	8.2	x	4.1	x	x	x	x
IP100207012.1	Voltage-gated potassium channel beta-1 subunit	401	44710	9.3	x	x	x	9	x	13.2
IP100211012.1	Voltage-gated potassium channel beta-2 subunit	367	41021	9	16.9	20.2	19.1	29.4	21.3	22.6
IP100197991.5	Kv3.3c voltage-gated potassium channel subunit splice variant C	770	82068	6.9	x	x	x	3.6	x	5.2
IP100391769.1	Voltage-gated calcium channel alpha2/delta-1 subunit	1091	1E+05	5.3	20.7	25.4	22.3	x	x	x
IP100365653.2	Putative voltage-gated calcium channel alpha(2)/delta-4 subunit	1463	2E+05	6	x	x	x	x	3.6	x
IP100211876.1	Voltage-dependent L-type calcium channel beta-3 subunit	484	54564	6.3	x	x	x	6.4	x	x
IP100211870.1	Voltage-dependent P/Q-type calcium channel alpha-1A subunit, Splice Isoform 1	2212	3E+05	8.6	4	3.5	x	x	x	2.8
IP100200639.2	Voltage-dependent N-type calcium channel alpha-1B subunit	2356	3E+05	8.6	2.2	1.1	2.4	x	x	x
IP100199192.3	Voltage-dependent T-type calcium channel alpha-1G subunit	2254	3E+05	7.3	1.4	x	x	x	x	x

Locus	Description	L	MW	pI	ACN INV SeqCov	ACN PPS SeqCov	ACN RPG SeqCov	Tris INV SeqCov	Tris PPS SeqCov	Tris RPG SeqCov
IP100207426.1	Voltage-dependent calcium channel gamma-8 subunit	421	43269	9.2	23	14	23	12.1	12.1	7.8
IP100400699.3	R1IA Voltage-gated sodium channel	1956	2E+05	5.6	6.7	5.1	x	7.4	6	6.5
IP100213193.3	Voltage-gated sodium channel variant rPN4a	1988	2E+05	6.3	x	1.7	x	x	x	2.3
IP100210089.1	sodium channel, voltage-gated, type 10, alpha polypeptide	1957	2E+05	6.2	x	x	x	x	1.5	x
Transporters										
IP100421430.1	Zinc transporter ZnT-3	388	41898	7.3	14.4	8.5	8.5	9.5	9.8	17.5
IP100214787.1	Monocarboxylate transporter 1	494	53238	8.3	4.9	x	5.3	7.9	x	7.1
IP100191391.1	Solute carrier family 2, facilitated glucose transporter member 1	492	53963	8.7	10.8	8.5	6.9	x	6.9	6.9
IP100198723.1	Solute carrier family 2, facilitated glucose transporter member 3	493	53581	5	11.8	8.3	6.5	10.5	x	2.4
IP100395291.1	Neutral amino acid transporter ASCT1	529	55900	6.3	19.1	18.5	9.8	15.9	4.5	13.4
IP100204778.1	Large neutral amino acids transporter small subunit 1	512	55903	7.9	2.9	x	8.2	8.2	x	10
IP100324377.1	Splice Isoform GLAST-1 of Excitatory amino acid transporter 1	543	59698	8.4	25.6	15.3	15.3	19.9	x	x
IP100214057.5	Excitatory amino acid transporter 4	561	60715	9.1	4.6	8.4	4.6	7.5	x	x

Table 3

A summary of identified proteins and peptides containing transmembrane (TM) regions in six digestion conditions. Proteins containing at least one transmembrane domain (TMD) are predicted by the TMHMM algorithm (see method). Peptides containing the sequences of the transmembrane region are identified and summarized in the second row. The last row of data represents the percentage of TMD peptides identified from the identified TMD proteins. Higher ratio of TMD peptide vs. TMD protein indicates higher representation of the transmembrane region in the particular digestion condition.

	Tris RPG	Tris IVS	Tris PPS	ACN IVS	ACN PPS	ACN RPG
TMD Protein	556	476	430	422	324	213
TMD Peptide	88	73	23	86	23	39
TMD peptide/ TMD Protein	15.8%	15.3%	5.3%	20%	4.9%	8.7%