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Quantitative proteomics for identifying biomarkers for tuberculous meningitis

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Abstract

Introduction: Tuberculous meningitis is a frequent extrapulmonary disease caused by *Mycobacterium tuberculosis* and is associated with high mortality rates and severe neurological sequelae. In an earlier study employing DNA microarrays, we had identified genes that were differentially expressed at the transcript level in human brain tissue from cases of tuberculous meningitis. In the current study, we used a quantitative proteomics approach to discover protein biomarkers for tuberculous meningitis.

Methods: To compare brain tissues from confirmed cases of tuberculous meningitis with uninfected brain tissue, we carried out quantitative protein expression profiling using iTRAQ labeling and LC-MS/MS analysis of SCX fractionated peptides on Agilent's accurate mass QTOF mass spectrometer.

Results and conclusions: Through this approach, we identified both known and novel differentially regulated molecules. Those described previously included signal-regulatory protein alpha (SIRPA) and protein disulfide isomerase family A, member 6 (PDIA6), which have been shown to be overexpressed at the mRNA level in tuberculous meningitis. The novel overexpressed proteins identified in our study included amphiphysin (AMPH) and neurofascin (NFASC) while ferritin light chain (FTL) was found to be downregulated in TBM. We validated amphiphysin, neurofascin and ferritin light chain using immunohistochemistry which confirmed their differential expression in tuberculous meningitis. Overall, our data provides insights into the host response in tuberculous meningitis at the molecular level in addition to providing candidate diagnostic biomarkers for tuberculous meningitis.

Keywords: Relative quantitation, Cerebrospinal fluid, Histopathology, Early diagnosis, Tuberculosis

Introduction

Tuberculosis (TB) is a common and sometimes fatal transmissible disease, especially in developing countries. Approximately thirty percent of the global population is exposed to the acid-fast bacilli causing TB. Of those who are infected with tuberculosis, ~10% percent develop a clinical manifestation of the disease during their lifetime. From a global perspective, approximately twenty percent of TB infected population live in India. The World

Health Organization (WHO) has estimated that one million children develop TB annually worldwide which accounts for about 11% of all TB cases [1]. Tuberculous bacilli most commonly infect lungs. *Mycobacterium tuberculosis* (MTB) may also spread to extrapulmonary sites including the meninges, lymph nodes, genitourinary tract, skeletal system and skin [2]. Tuberculous meningitis (TBM) is the infection of meninges caused by MTB, with a mortality rate of ~30%. Further, those who survive TBM are usually left with severe neurological defects [3-5]. There is an increased risk of TBM in HIV-infected patients as compared to non-HIV infected cases although the clinical manifestations of the disease do not differ between the two groups [6,7].

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Culturing mycobacteria and subsequent microbiological examination is considered a gold standard for the diagnosis of TBM. However, this method is time consuming and insensitive, with a positive outcome achieved only in 25–70% of clinically diagnosed cases [8]. Although PCR assays can be an alternative rapid approach for diagnosis, they cannot differentiate between latent or active forms of the disease. Although nucleic acid amplification test (NAAT) has a high specificity when tested in body fluids, it lacks adequate sensitivity in cases of meningitis and pleuritis [9].

Amongst the existing molecular markers, Adenosine deaminase isoenzyme-2 (ADA2) has a sensitivity of 100% and a specificity of 86.4% for the detection of TBM in cerebrospinal fluid (CSF) [10]. Adenosine deaminase (ADA) activity in the CSF of TBM patients has been suggested to be useful for early differential diagnosis of TBM [11]. The ADA activity of CSF and plasma have been evaluated as a diagnostic aid in TBM [12] and ADA activity in CSF was considered to be a simple, useful and rapid diagnostic test for early recognition of TBM in children [13]. However, overexpression of ADA was also often overexpressed in other forms of meningitis including infections with pyogenic bacteria [14]. In the CSF of TBM patients, the presence of 65 kDa heat shock protein antigen might be a marker for early diagnosis of the disease [15]. High levels of CSF lactate and lactate dehydrogenase levels have also been suggested for diagnosing TBM [16].

Early diagnosis of TBM is considered a key to effective treatment and prognosis. Approximately 90% of the patients are diagnosed in stage II or III [17]. Overall, the diagnosis of TBM still remains a major challenge due to inadequate current diagnostic methods and poor sensitivity and/or specificity of existing markers. Although corticosteroids are used extensively to reduce mortality and neurological disability, it may not be the only solution to reduce the mortality and morbidity [18]. In TBM, a number of pathological changes including meningeal adhesion, infarction, tuberculoma and hydrocephalus may occur leading to neurological sequelae [4]. These sequelae are known to correlate with the stage of meningitis at admission. Patients treated at an early stage have a five times higher chance of recovery than those with advanced disease stages [19]. Therefore, patient's clinical condition at admission and delay in starting the treatment are important factors for determining their survival [20]. These findings emphasize the need to focus on identifying candidate molecular markers which can be developed as diagnostic tools in the management of TBM.

Mass spectrometry-based quantitative proteomics has emerged as a powerful approach for identifying and studying disease biomarkers and has become one of the essential tools in biomarker discovery [21,22]. Advances

in quantitative mass spectrometry have led to identification and quantitation of biomarkers which serve as indicators of disease progression, prognosis, drug safety and help to elucidate the mechanism of drug treatment [23]. There are various labeling approaches that one can employ to carry out quantitative proteomic measurements. *In vitro* labeling methods include Isobaric Tags for Relative and Absolute Quantitation (iTRAQ), Isotope-Coded Affinity Tags (ICAT), ^{18}O labeling and *in vivo* methods include Stable Isotope Labeling by Amino acids in Cell culture (SILAC) and ^{15}N labeling [24,25]. iTRAQ labeling is an effective method for studying differential protein expression levels in tissue samples. It has been extensively used for biomarker discovery in various disease contexts [26-34].

In this study, we used an iTRAQ-based quantitative proteomic approach to identify differentially expressed proteins from brain tissues of tuberculous meningitis cases as compared to controls. We identified several proteins which are differentially expressed in TBM. These proteins include both novel and previously reported candidate protein markers. We validated some of these candidate biomarkers using immunohistochemical labeling.

Materials and methods

Sample collection

The study was approved by scientific ethics committee of National Institute of Mental Health and Neuro Sciences (NIMHANS), Bangalore, India. Samples from frontal cortex with overlying meninges from cases of tuberculous meningitis (n=6) and similar, but uninfected, control brain tissues (n=6) from victims of road traffic accidents were collected at the time of autopsy. The autopsy was conducted within 8–16 h postmortem with the body kept at 4°C after death. The samples were obtained from the Human Brain Tissue Repository, Department of Neuropathology, NIMHANS, Bangalore. Sample details including the selection criteria are provided in Additional file 1: Table S1.

Sample preparation and iTRAQ labeling

Brain tissue samples were lysed in 0.5% SDS, sonicated, homogenized and centrifuged at 13,000 rpm for 10 min at 4°C. Supernatant was collected and protein quantitation was carried out by Lowry's assay (Bio-Rad Hercules, CA; USA). For each condition, 160 µg of protein sample was utilized for the experiment. Each sample was treated with 4 µL of reducing agent (tris (2-carboxyethyl) phosphine (TCEP)) at 60°C for 1 h and alkylated with 2 µL of cysteine blocking reagent, methyl methanethiosulfonate (MMTS) for 10 min at room temperature. After alkylation, the samples were subjected to trypsin digestion (Sequencing Grade Modified Trypsin, Promega Cat#:

V511A) using 1:20 (w/w) at 37°C for 16 h. Peptide samples from each condition was split in equal halves (80 µg each) and labeled with iTRAQ 4-plex reagents (catalog # 4352135, Applied Biosystems, Foster City, CA, USA) as per manufacturer's protocol. We used technical replicates for control and TBM samples. Peptides from control samples were labeled with iTRAQ reagents yielding reporter ions 114 and 115 while peptides from TBM were labeled with 116 and 117. The samples were pooled following iTRAQ labeling.

Strong cation exchange chromatography (SCX)

Pooled iTRAQ labeled peptides were fractionated by strong cation exchange chromatography on PolySULFOETHYL A column (200 x 2.1 mm; 5 µm; 200Å PolyLC, Columbia, MD) using Agilent's 1200 series HPLC system. The peptides were reconstituted in SCX solvent A (10 mM potassium phosphate, 25% acetonitrile, pH 2.8) and loaded on SCX column isocratically using 100% solvent A for 20 min at a flow rate of 200 µl per minute. Peptides were eluted using a 30 min gradient from 8% to 35% solvent B (350 mM KCl in solvent A). Fractions were collected every minute using a fraction collector. The fractions were vacuum dried and stored at -80°C until LC-MS/MS analysis.

LC-MS/MS analysis

The samples were analyzed on HPLC chip-cube interfaced with Accurate Mass 6520 quadrupole time of flight mass (QTOF) spectrometer (Agilent Technologies, Santa Clara, CA). The HPLC-Chip contains a 40 nl enrichment column and 43 mm x 75 µm analytical column. These columns were made up of a reversed-phase material Zorbax 300SB-C₁₈, with a particle size of 5 µm. The samples were loaded onto the enrichment column using Agilent's 1200 series capillary liquid chromatography pump at a flow rate of 3 µl/min using 97% solvent A and 3% solvent B. An injection flush volume of 4 µl was applied during enrichment step. The peptides were eluted at a flow rate of 400 nl/min using a gradient of solvent A (0.1% formic acid) and solvent B (0.1% formic acid in 90% acetonitrile). The gradient was started from 3% to 18% of solvent B over 8 min, subsequently changed to 22% B in the next 7 min and finally changed to 45% of solvent B for 25 min. MassHunter workstation data acquisition software (Version B.01.03) was used for data dependent acquisition. MS spectra were acquired for 1 second from m/z 350–1800 followed by three MS/MS spectra in next second comprising the duty cycle of 2.1 second including an interscan delay of 0.1 second. Precursor ions were preferred based on charge state in the order of 2+, 3+ and >3+. The capillary and fragmentor voltage of 1950V and 175 V respectively was applied

with a medium isolation width of 4 m/z and a collision energy slope of 3 V plus at offset of 2 V.

Data analysis

The mass spectrometry raw data was processed to peak list format by using MassHunter Qualitative Analysis software (Agilent Technologies, Version B.03.3). These processed peak list files were then searched against Human RefSeq Protein Database (Release 40) containing 31,811 protein sequences through Proteome Discoverer platform (Thermo Scientific, Version 1.2). The workflow includes spectrum files, spectrum selector, Mascot and Sequest search nodes followed by peptide validator for false discovery analysis whereas a reporter ion quantifier was used for quantitation. Search parameters included trypsin as the enzyme with one missed cleavage allowed, oxidation of methionine, deamidation at asparagine and glutamine were set as a variable modification whereas methyl-thio at cysteine and iTRAQ label at N-terminus of the peptide and lysine were set as a fixed modification. The reporter ion window tolerance was set at 100 ppm. The identified peptides were filtered using 1% false discovery rate derived using decoy database search strategy and top ranked hit based on peptide score, XCorr and IonScore for Sequest and Mascot respectively. The search results from both Mascot and Sequest were merged and unique peptide(s) identified for each protein were used to calculate relative protein quantitation in Proteome Discoverer workflow. The average ratio was used for relative protein quantification for proteins with multiple peptide matches. Bioinformatic analysis was carried out to categorize proteins based on biological processes, cellular component and molecular function classification using annotations in Human Protein Reference Database (HPRD, <http://hprd.org>) [35], which is in compliance with gene ontology (GO) standards.

Immunohistochemical labeling

Formalin fixed and paraffin embedded autopsy tissues were collected and cut into 4 µm thick sections on glass slides. These slides were subjected for deparaffinization and rehydration. Endogenous peroxidase activity was quenched by 3% H₂O₂ for 20 min at room temperature. For antigen retrieval, the tissue sections were microwaved in citrate buffer (pH 6.0) for 30 min. The tissue sections were incubated with 3% skimmed milk in PBS, pH 7.4 at room temperature. The tissue sections were incubated with primary antibodies at following dilutions - anti-amphiphysin (dilution 1:100, catalog # ab52646), anti-neurofascin (dilution 1:250, catalog # ab31457), anti-ferritin light chain (dilution 1:500, catalog # ab69090) which were purchased from Abcam (Cambridge UK).

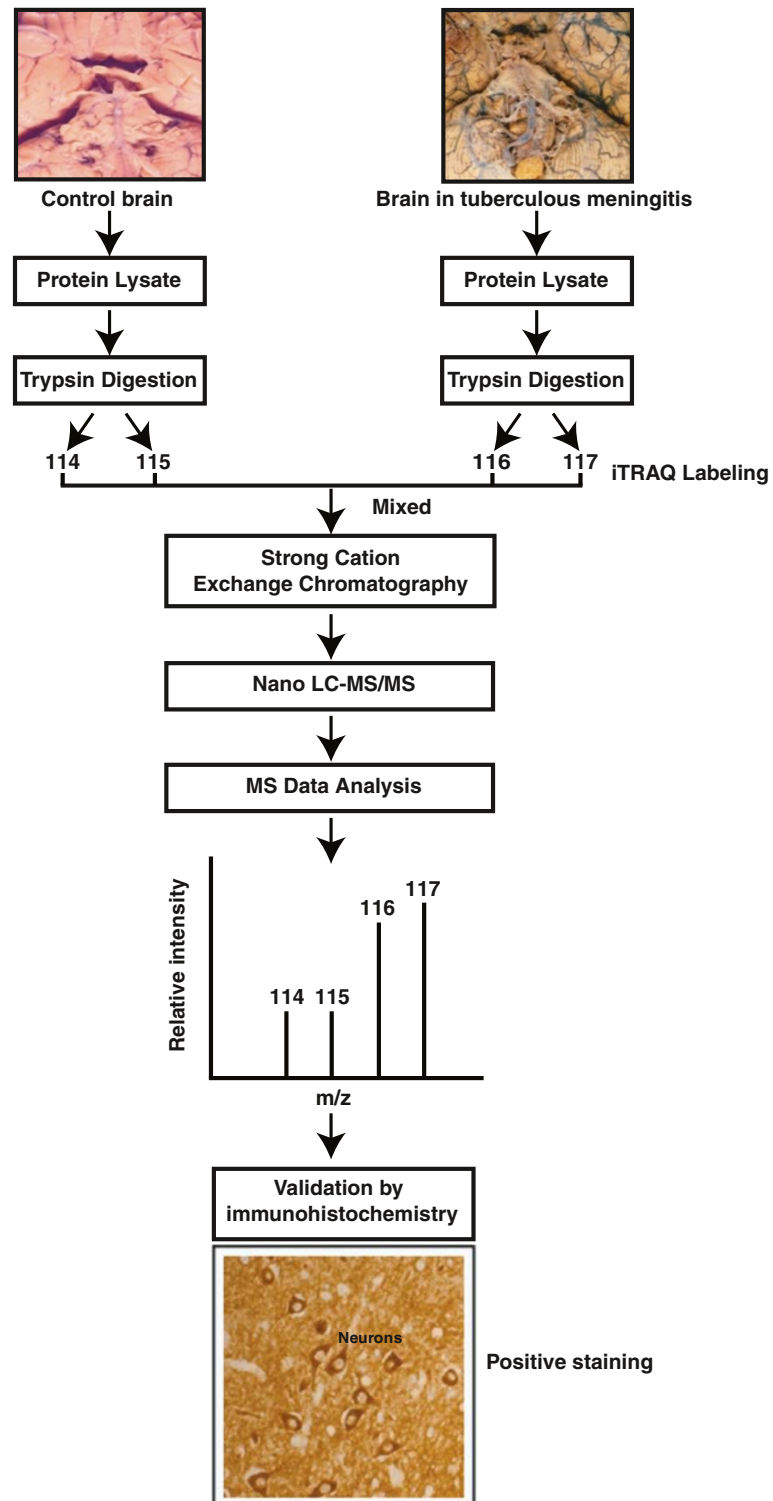


Figure 1 Outline of the quantitative proteomic strategy. TBM-infected and uninfected frontal cortex tissues were used to extract proteins and subjected to digestion with trypsin. Peptides from the two groups were labeled with iTRAQ reagents followed by LC-MS/MS on a QTOF mass spectrometer. A subset of the differentially expressed molecules were validated by immunohistochemistry.

In parallel with test slides, negative and positive controls were also subjected for overnight incubation at room temperature, followed by incubation with prediluted secondary antibody conjugated with poly HRP (catalog # K4011) from Dako. The reaction was visualized with chromogen substrate DAB/H₂O₂ (Dako catalog # K4007) as per manufacturer's instructions. The sections were counterstained with hematoxylin.

Results and discussion

We used six TBM and six uninfected samples for our quantitative proteomic analysis. The overall workflow for the study is depicted in Figure 1. The tandem mass spectra were searched against human NCBI RefSeq release 40 (31,811 protein sequences) using Mascot and Sequest search engines through the Proteome Discoverer software. A total of 81,494 spectra were acquired which resulted in 5,988 peptide-spectrum matches. We identified a total of 434 proteins with 919 unique peptides. The protein and peptide data is provided in Additional file 2: Table S2 and Additional file 3: Table S3, respectively. In all, we identified 110 upregulated and 24

downregulated proteins at 1% FDR. The distribution of cellular localization and biological processes of the differentially expressed proteins is shown in Figure 2A and B. The subcellular localization and biological processes of the differentially expressed proteins were classified based on annotations in Human Protein Reference Database (HPRD, <http://hprd.org>) [35]. Partial list of differentially expressed proteins are provided in Tables 1 and 2.

Proteins previously implicated in TBM and tuberculosis associated studies

Proteins that have previously been implicated in TBM and tuberculosis associated studies included Vimentin (VIM), Peroxiredoxin 5 (PRDX5), Glial fibrillary acidic protein (GFAP), Signal-regulatory protein alpha (SIRPA) and Protein disulfide isomerase family A, member 6 (PDIA6). VIM is an intermediate filament family protein and is also the potential ligand for natural cytotoxicity triggering receptor 1 (NKP46). VIM was shown to be 2.8-fold upregulated in the present study. VIM is involved in the lysis of the MTB-infected cells through NKP46 [36,37]. PRDX5 is a member of the peroxiredoxin family

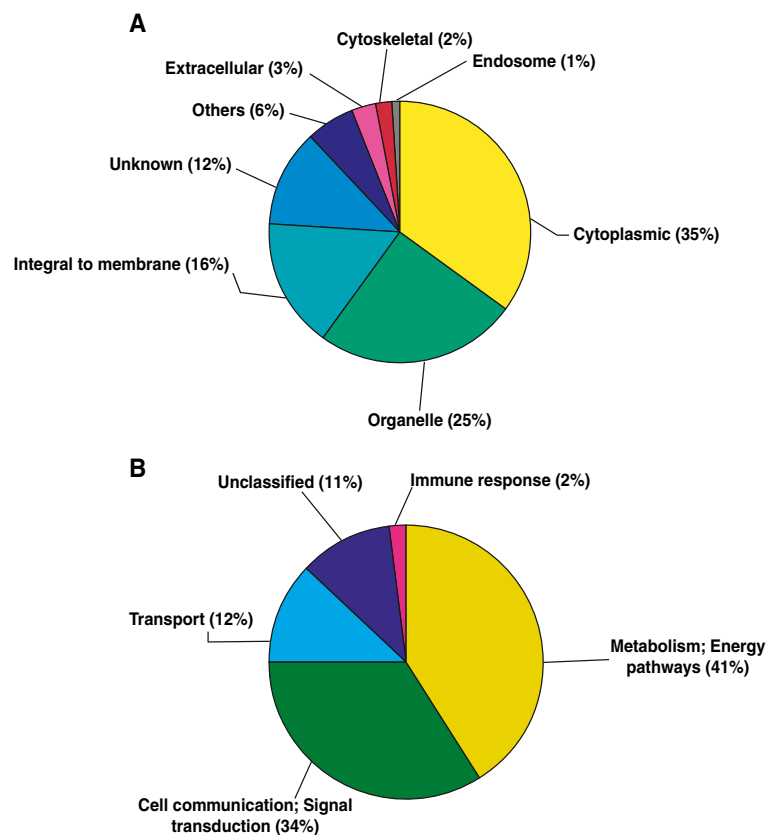


Figure 2 Subcellular localization and biological process of differentially expressed proteins. Subcellular localization (A) and biological process (B) of differential expressed proteins are shown.

Table 1 A partial list of proteins overexpressed in TBM

	Gene Symbol	Protein	Description	Fold-change
1	<i>WARS</i>	Tryptophanyl-tRNA synthetase, cytoplasmic isoform b	It induced by interferon and involved in the catalyzation of the aminoacylation of tRNA (trp) with tryptophan.	7.2
2	<i>SLC4A4</i>	Electrogenic sodium bicarbonate cotransporter 1 isoform 2	It plays a functional role in the regulation of bicarbonate secretion and absorption and intracellular pH.	6.9
3	<i>NPM1</i>	Nucleophosmin isoform 3	It is a phosphoprotein which shuttles between the nucleus and the cytoplasm. It plays a role in ARF/p53 signaling pathway.	4.8
4	<i>DHX9</i>	ATP-dependent RNA helicase A	It localizes to cytoplasm and nucleus. It acts as a transcriptional regulator.	4.9
5	<i>HPCA</i>	Neuron-specific calcium-binding protein hippocalin	It belongs to the neuron-specific calcium-binding proteins family. It may play a role in the neurons of the central nervous system.	4.4
6	<i>GPM6A</i>	Neuronal membrane glycoprotein M6-a isoform 3	It is a transmembrane protein and expressed on neurons in the central nervous system.	3.8
7	<i>GFAP</i>	Glial fibrillary acidic protein isoform 2	It is an intermediate filament protein and a marker for astrocytes. It is overexpressed in astrogliosis.	3.2
8	<i>SLC8A2</i>	Sodium/calcium exchanger 2 precursor	It is a Sodium/calcium exchanger, regulates the intracellular calcium concentrations.	2.8
9	<i>VIM</i>	Vimentin	It belongs to the intermediate filament protein family. It plays a role in cell shape and integrity.	2.8
10	<i>VGF</i>	VGF nerve growth factor inducible precursor	It expressed in neuroendocrine cells and overexpressed by nerve growth factor	2.5

of antioxidant enzymes. *PRDX5* was found to be overexpressed in MTB (H37Rv) infected mice study [38]. In our study, it was found to be 2.2-fold overexpressed in TBM as compared to controls. *PRDX5* plays an antioxidant protective role in tissues. *GFAP* showed 3.2-fold upregulation in TBM as compared to controls. Our finding correlated with a recent study that described a

4.7-fold overexpression of *GFAP* at mRNA level in TBM cases [39]. Representative MS/MS and reporter ion spectra of differentially regulated molecules are provided in Figure 3. MS/MS spectrum of *GFAP* is depicted in Figure 3A. *SIRPA* belongs to the signal regulatory family. The regulation of *SIRPA* expression can change the effectiveness of signaling processes. Growth factor

Table 2 A partial list of downregulated proteins in TBM

	Gene Symbol	Protein	Description	Fold-change
1	<i>PDIA6</i>	Protein disulfide-isomerase A6 precursor	It is endoplasmic reticulum (ER) resident protein and it plays a role in folding of disulfide-bonded proteins	2.0
2	<i>KPNA4</i>	Importin subunit alpha-4	It is cytoplasmic protein, recognizes nuclear localization signals.	2.0
3	<i>PFDN5</i>	Prefoldin subunit 5 isoform alpha	It belongs to the prefoldin alpha subunit family. It is a subunit of the molecular chaperone complex, involved in protein folding.	2.5
4	<i>GARS</i>	Glycyl-tRNA synthetase	It belongs to the class II family of tRNA synthetases and this protein plays a role in autoimmune diseases.	3.0
5	<i>MDH1</i>	Malate dehydrogenase, cytoplasmic	It localized to the cytoplasm and mitochondria. It involved in malate-aspartate shuttle.	3.0
6	<i>AK1</i>	Adenylate kinase isoenzyme 1	It is an enzyme involved in regulating the adenine nucleotide composition in the cell and it is localized in the cytosol.	3.0
7	<i>LANCL1</i>	LanC-like protein 1	It is a loosely associated peripheral membrane protein which belongs to the LanC family of bacterial membrane-associated proteins. It plays a role in antimicrobial peptide synthesis.	5.0
8	<i>SNCG</i>	Gamma-synuclein	It belongs to the member of the synuclein family of proteins and it may play a role in the pathogenesis of neurodegenerative diseases.	5.0
9	<i>PTPLB</i>	Protein-tyrosine phosphatase-like member B	It localizes to endoplasmic reticulum (ER) and involved in the dehydration of very long chain fatty acid synthesis	7.0
10	<i>NCDN</i>	Neurochondrin isoform 2	It is a cytoplasmic protein and may play a role in spatial learning processes.	7.0

receptors and growth hormone receptor signaling is suppressed by the upregulation of *SIRPA* [40-43]. *SIRPA* regulates the NF κ B activity that renders the cells resistant to TNF mediated apoptosis [44]. A recent study reported *SIRPA* to be 2-fold upregulated at the mRNA level in TBM [39]. In our study, also *SIRPA* protein was found to be 2-fold overexpressed in TBM. Protein disulfide isomerase family A, member 6 (PDIA6) is a protein that belongs to the endoplasmic reticulum (ER) and it is involved in the folding of disulphide bonded proteins [45]. This protein was shown to be downregulated (2.6-fold) in TBM at the mRNA level [39] and confirmed to be downregulated (2.5-fold) at the protein level in our study.

In our previous study on TBM using gene expression microarray, we had identified 2,434 differentially regulated transcripts [39]. In current proteomic study, 33 out of 134 differentially regulated proteins were also

identified to be differentially expressed in the microarray dataset. However, only three proteins (*GFAP*, *SIRPA*, *ACTB*) were found to be correlated at both mRNA and protein levels, which is likely due to low correlation frequently observed between transcriptomic and proteomic studies.

Differentially expressed proteins with no previous association with TBM

We identified a large number of differentially expressed proteins, which have not been reported earlier in the literature to be associated with TBM. These novel proteins include Ca⁺⁺-dependent secretion activator 2 (*CADPS2*) which belongs to the calcium-dependent activator of secretion (CAPS) protein family, and was found 3-fold upregulated in the present study. *CADPS2* protein regulates the exocytosis of synaptic and dense-core vesicles in

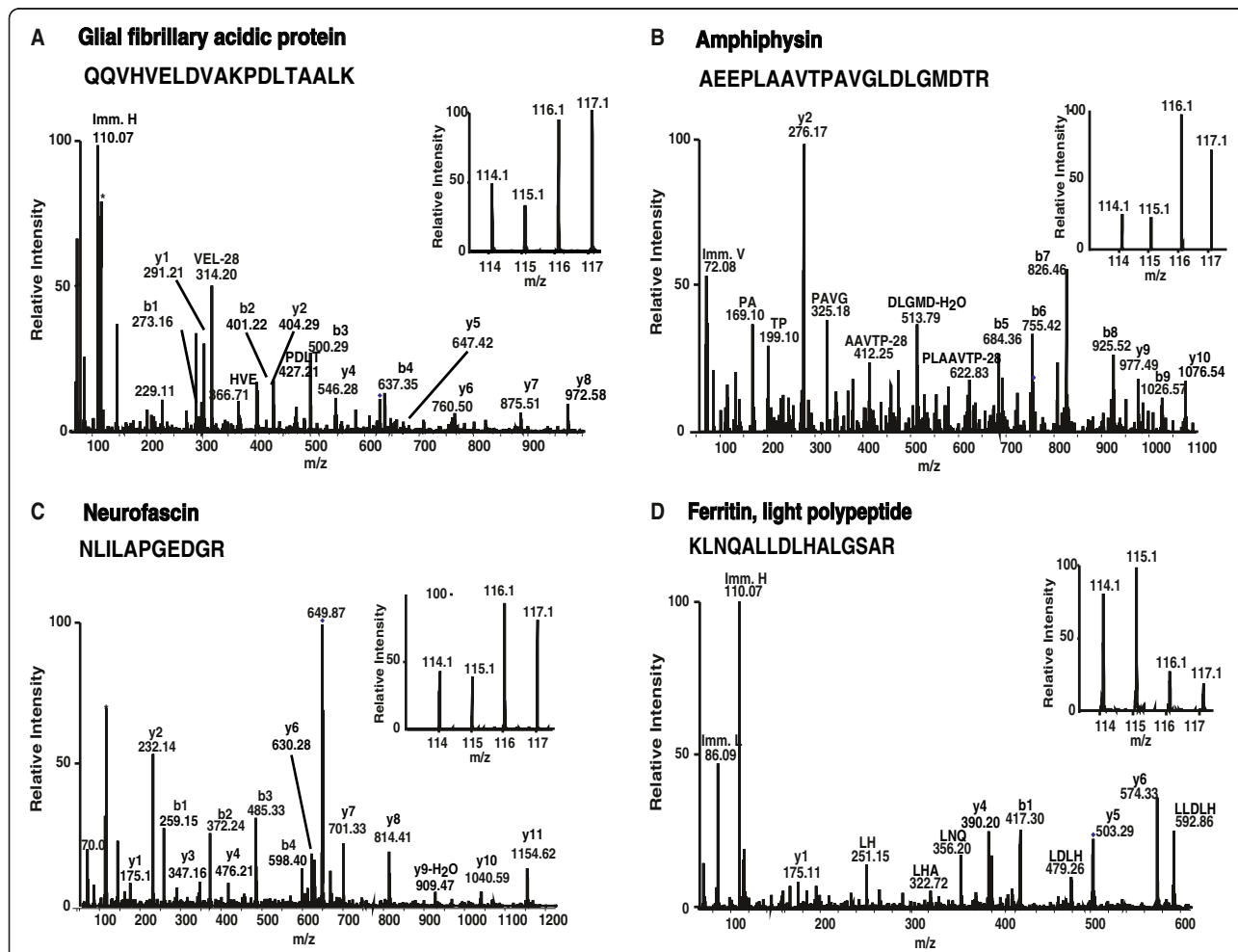


Figure 3 Representative MS and MS/MS spectra of upregulated as well as downregulated proteins. MS and MS/MS spectra of peptides from selected differentially expressed proteins identified in this study. Insets show relative intensities of reporter ions. (A) Glial fibrillary acid protein (GFAP); (B) Amphiphysin (AMPH); (C) Neurofascin (NFASC) (D) Ferritin, light chain (FTL).

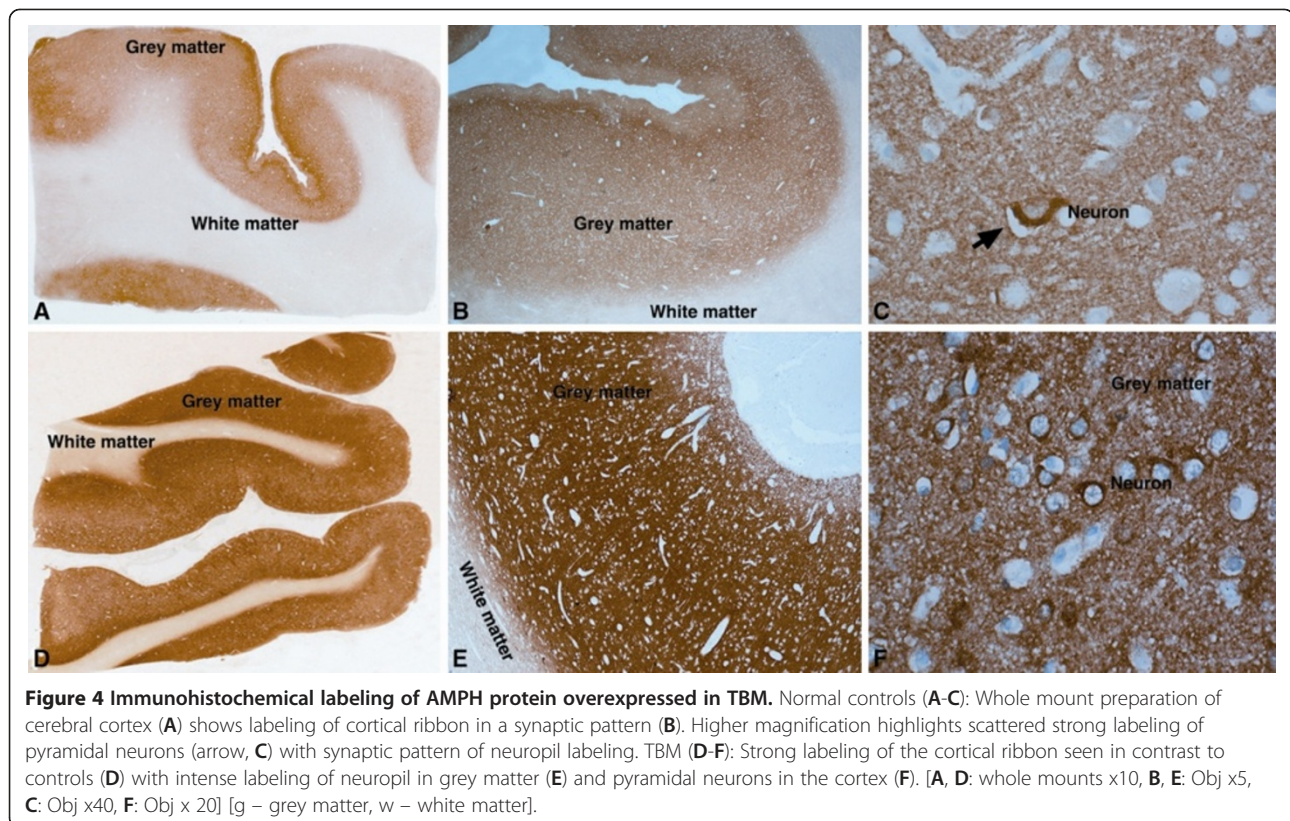
neurons and neuroendocrine cells [46,47]. Amphiphysin (AMPH) (Figure 3B) is an adapter molecule which plays a role in synaptic vesicle endocytosis [48] and it was found to be 3.7-fold overexpressed in TBM compared to control cases. Heat shock protein 90kDa alpha (cytosolic), class A member 1 (HSP90AA1) is a highly conserved molecular chaperone and was found to be 2.1-fold upregulated in the present study. HSP90AA1 is a molecular chaperone, which plays a role in signal transduction, protein degradation, protein folding and is expressed under stress conditions and according to a recent report it plays a role in gene expression in mammalian cells [49]. Neurofascin (NFASC) (Figure 3C) is the L1 family immunoglobulin cell adhesion molecule, found to be 2-fold upregulated in the current study. NFASC plays a role in organization of the axon initial segment (AIS) and nodes of Ranvier in central nervous system, neurite extension and neurite fasciculation [50-52]. Ferritin, light chain (FTL) (Figure 3D) protein is an iron storage protein which, play a role in neurodegeneration [53] and found to be 66-fold downregulated in the present study. Downregulation of FTL causes the depigmentation in metastatic melanoma cells [54] and responds during inflammation especially where oxidative stress and reactive oxygen intermediates are generated [55,56].

Validation of candidate biomarkers by immunohistochemical labeling

We used immunohistochemistry (IHC) to validate a subset of differentially expressed proteins identified in iTRAQ-based study. The results from IHC validation of AMPH, NFASC and FTL from fifteen TBM cases are summarized below.

Amphiphysin (AMPH) is an adapter molecule associated with cytoplasmic surface of the synaptic vesicle, which was confirmed to be overexpressed in all 15 TBM cases tested (Figure 4). AMPH protein expression was seen diffusely distributed in synaptic fashion in the cortical ribbon in control cases (Figure 4A, B). The neurons in the grey matter showed cytoplasmic labeling with AMPH (Figure 4C). In comparison, cases of tuberculous meningitis revealed intense staining of the neuropil of the cortical ribbon reflecting upregulation of protein expression (Figure 4D, E). Several neurons showed cytoplasmic accumulation of protein within the neuronal soma (Figure 4F). Astrocytes and microglia were not labeled and white matter was devoid of labeling. Inflammatory exudates in the meninges in the cases of tuberculous meningitis also did not reveal positivity.

Neurofascin (NFASC) is an adhesion molecule, which plays an important role in cell communication and signal transduction. Overexpression of this protein was validated



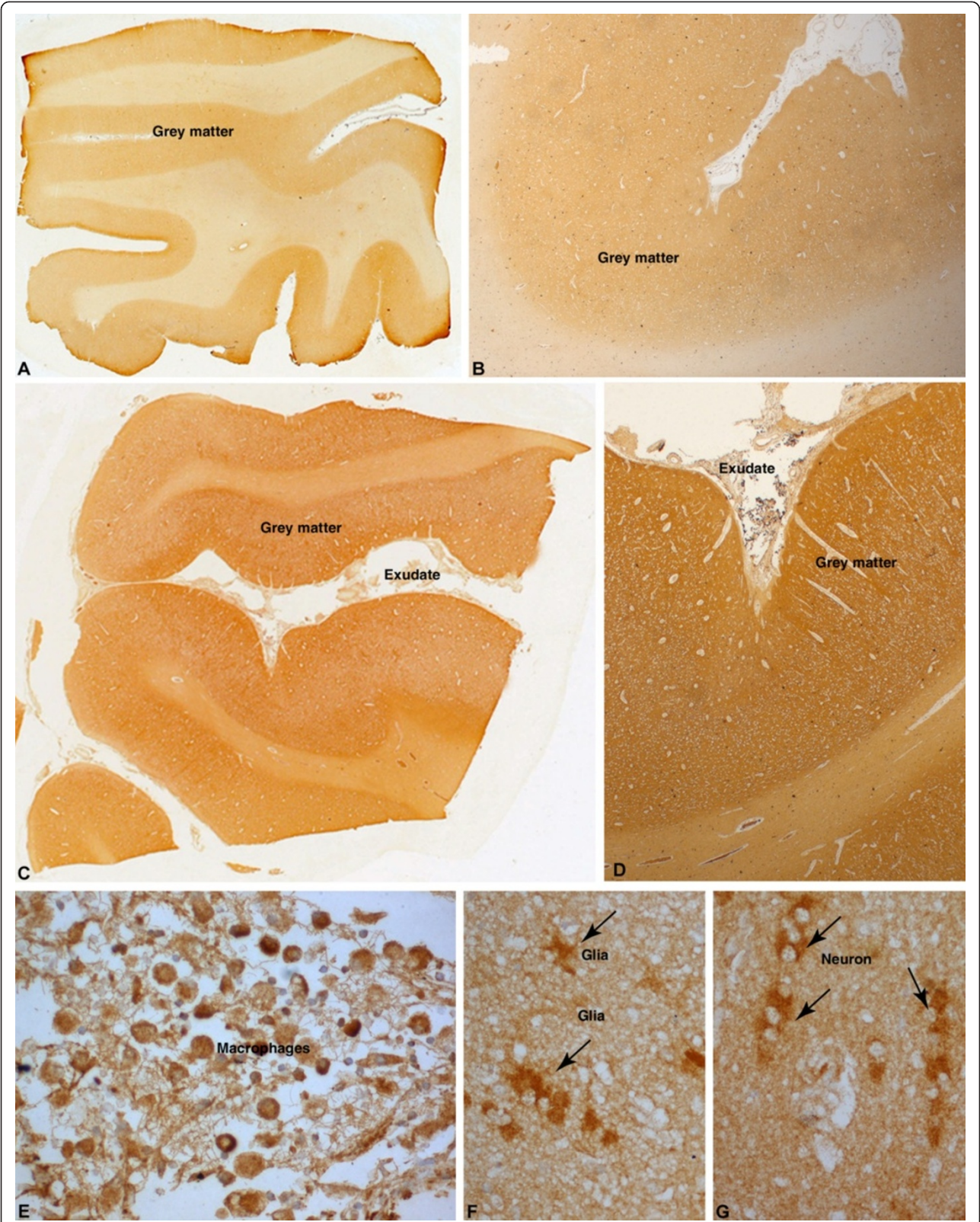


Figure 5 (See legend on next page.)

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Figure 5 Immunohistochemical labeling of NFASC protein overexpressed in TBM. Normal controls (A, B): Light labeling of the cortical ribbon (A) in the frontal cortex with synaptic pattern (B). TBM (C-G): Whole mount preparation (C) highlight the darker labeling of the cortical ribbon and the inflammatory exudate in the subarachnoid space. The macrophages (E) entrapped in the inflammatory exudate are strongly labeled. Glial cells (F, arrows) and linearly arranged neuronal cytoplasm (G, arrows) are strongly labeled in the grey matter. [A, C: x10, B, D: xObj.5, E-G: xObj.20].

in all 15 TBM cases. NFASC in control cases revealed positive labeling of neuropil in cortex (Figure 5A, B) with only occasional neurons showing cytoplasmic labeling in the lower cortical layers. No labeling of astrocytes or oligodendroglia was noted in control cases. In cases of

tuberculous meningitis, intense labeling was seen in the neuropil of grey matter in synaptic fashion reflecting upregulated expression of protein (Figure 5C, D). Several small granule neurons expressed cytoplasmic positivity in the upper cortical layers close to the surface exudates

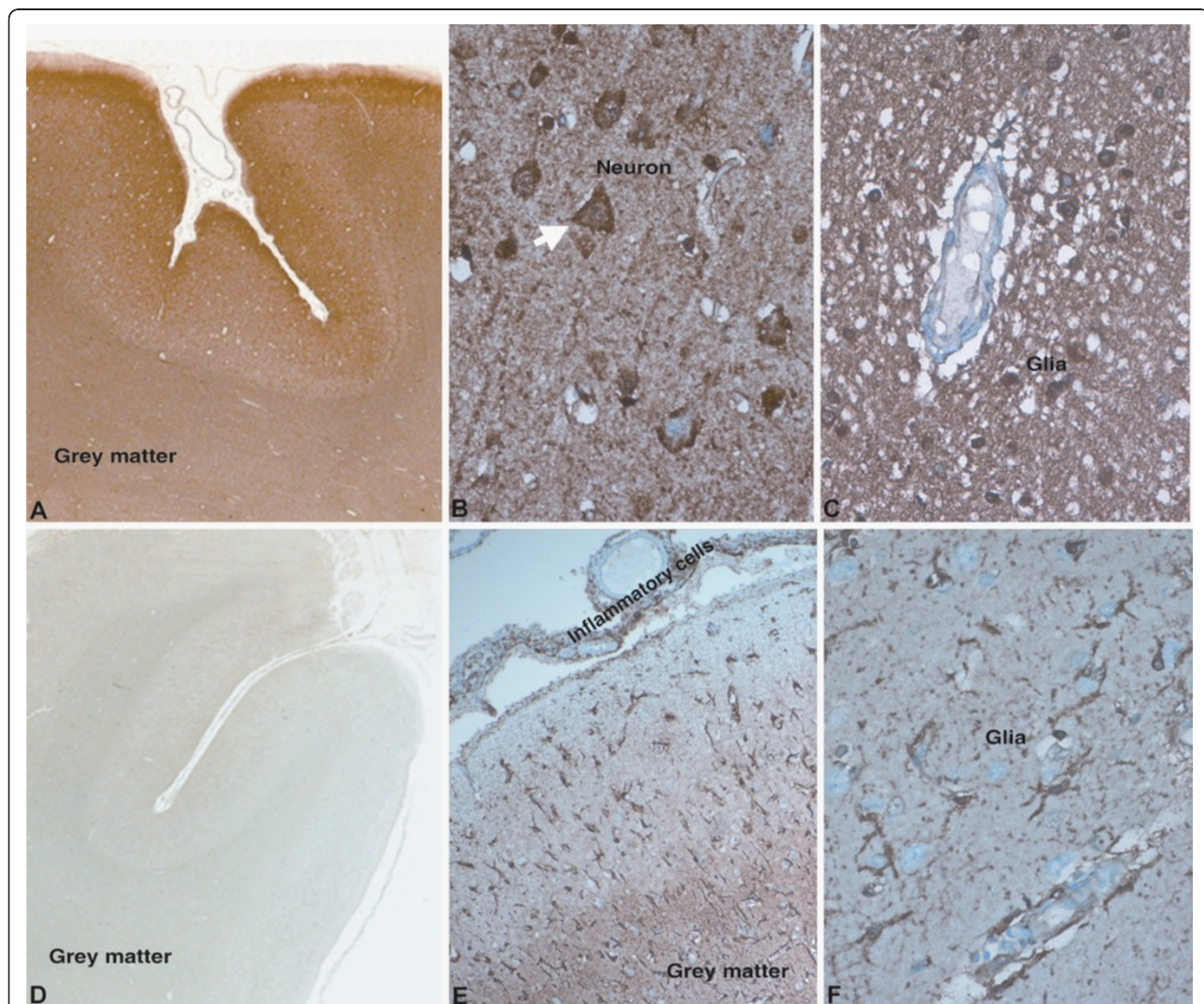


Figure 6 Immunohistochemical labeling of FTL protein downregulation in TBM. Normal controls (A-C): Intense labeling of grey and white matter (A). In the grey matter (B), the neurons (arrow) and dendritic processes are intensely labeled (B) while in the white matter (C), neuropil and oligodendroglial cells are stained intensely. TBM (D-F): In the grey and white matter of the frontal cortex, showed very light labeling in contrast to control. In the subpial zones and deep grey matter beneath the exudate and the inflammatory cells in the subarachnoid space are stained (E) while the neuropil and neurons are less intense. Microglial cells in white matter and perivascular zones show labeling (F). [A, D: whole mounts x10, B, C: xObj.20, E: xObj.x5, F: xObj.20].

(Figure 5G). In addition, several reactive hypertrophic subpial astrocytes beneath the surface exudates expressed NFASC protein in cytoplasm extending into its branching processes (Figure 5F), in addition to macrophages in the inflammatory exudates in the subarachnoid space (Figure 5E).

Ferritin light chain (FTL) is an intracellular iron storage protein and downregulation of this protein was validated by IHC in 15 TBM cases (Figure 6). Expression of this protein was observed in control cases within the grey and white matter (Figure 6A), with grey matter showing higher intensity of expression compared to white matter. In the grey matter, neurons revealed intense labelling of the cytoplasm while dendritic processes in neuropil also revealed strong labelling (Figure 6B). In the white matter, oligodendroglial cells expressed the protein in its cytoplasm in addition to neuropil showing positivity (Figure 6C). In cases of tuberculous meningitis, marked down regulation of protein expression was seen in contrast to the normal cortex (Figure 6D). Interestingly, while neuronal expression was markedly downregulated, labelling was seen in microglial cells ensheathing neurons, as also in subpial and perivascular astrocytes seen below the inflammatory exudates in the subarachnoid spaces (Figure 6E). In the white matter, neuropil and perivascular microglia revealed protein expression in the branching ramified processes of activated microglia (Figure 6F).

Public availability of proteomic data

We have submitted the peptide data to Human Proteinpedia (<http://www.humanproteinpedia.org>) [57] and raw data files to Tranche data repository (<https://proteomecommons.org/tranche/>) for easy access to the research community. The raw data is freely available in Tranche using the following hash:

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xlm5a5zp5YBqpxdJpHYWlybPCdF9Bya2AH6AZRQ4  
dppr593JwMsptPIJCyVnJdbpOA+YHk3eNNQnNHdDQ  
jelH9c7QGSAAAAAAAAAAFsYQ==
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Conclusions

The combination of mass spectrometry and quantitative proteomics is being increasingly utilized for discovering potential biomarkers. In the current study, using an iTRAQ-based quantitative proteomic approach, we were able to identify several known and novel molecules that are differentially expressed in TBM. We validated the expression levels of three novel biomarker candidates - AMPH, NFASC and FTL - by IHC on brain tissue sections from TBM cases as well as controls. These promising candidate markers warrant further evaluation in CSF from TBM patients to determine their clinical utility.

Additional files

Additional file 1: Table S1. List of TBM and control samples used in the present study.

Additional file 2: Table S2. A complete list of Proteins identified in TBM.

Additional file 3: Table S3. A complete list of peptides identified in TBM.

Abbreviation

CSF: Cerebrospinal fluid; TBM: Tuberculous meningitis; iTRAQ: Isobaric tag for relative and absolute quantitation; QTOF: Quadrupole Time-of-Flight; IHC: Immunohistochemistry; TCEP: Tris (2-carboxyethyl) phosphine; MMTS: Methyl-methanethiosulfonate; TEAB: Triethylammonium bicarbonate buffer; ACN: Acetonitrile; Poly-HRP: Polymer conjugate of horseradish peroxidase; DAB-3: 3-Diaminobenzidine; HPLC: High performance liquid chromatography; SCX: Strong cation exchange; FDR: False discovery rate.

Competing interests

The authors have declared no conflict of interest.

Authors' contributions

RC, TSKP, SKS and AP conceived the study, GSSK, YLR, PS, RC, TSKP, SKS and AP designed the experiments; GSSK, AKV, AM, SR, HCH, NAS, HP and RS, carried out the experiments; GSSK, SR, PK, RS, SRG, KW and AP analyzed the data; GSSK, SKS, and AP wrote the manuscript. All authors read and approved the final manuscript.

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