The Impact of Delayed Storage on the Measured Proteome and Metabolome of Human Cerebrospinal Fluid

Therese Rosenling,1 Marcel P. Stoop,2 Agnieszka Smolinska,3 Bas Muilwijk,4 Leon Coulier,4 Shanna Shi,5 Adrie Dane,5 Christin Christin,1 Frank Suits,6 Peter L. Horvatovich,1 Sybren S. Wijmenga,3 Lutgarde M.C. Buydens,3 Rob Vreeken,7 Thomas Hankemeier,5,7 Alain J. van Gool,8 Theo M. Luider,2 and Rainer Bischoff1*

BACKGROUND: Because cerebrospinal fluid (CSF) is in close contact with diseased areas in neurological disorders, it is an important source of material in the search for molecular biomarkers. However, sample handling for CSF collected from patients in a clinical setting might not always be adequate for use in proteomics and metabolomics studies.

METHODS: We left CSF for 0, 30, and 120 min at room temperature immediately after sample collection and centrifugation/removal of cells. At 2 laboratories CSF proteomes were subjected to tryptic digestion and analyzed by use of nano-liquid chromatography (LC) Orbitrap mass spectrometry (MS) and chipLC quadrupole TOF-MS. Metabolome analysis was performed at 3 laboratories by NMR, GC-MS, and LC-MS. Targeted analyses of cystatin C and albumin were performed by LC–tandem MS in the selected reaction monitoring mode.

RESULTS: We did not find significant changes in the measured proteome and metabolome of CSF stored at room temperature after centrifugation, except for 2 peptides and 1 metabolite, 2,3,4-trihydroxybutanoic (threonic) acid, of 5780 identified peptides and 93 identified metabolites. A sensitive protein stability marker, cystatin C, was not affected.

CONCLUSIONS: The measured proteome and metabolome of centrifuged human CSF is stable at room temperature for up to 2 hours. We cannot exclude, however, that changes undetectable with our current methodology, such as denaturation or proteolysis, might occur because of sample handling conditions.

The stability we observed gives laboratory personnel at the collection site sufficient time to aliquot samples before freezing and storage at −80 °C.
© 2011 American Association for Clinical Chemistry

Conditions between the time of collection of a biological sample and its arrival at the analytical research laboratory might not always be adequate for subsequent proteomics and metabolomics analyses, especially in cases in which the sample collection was not originally performed with these large-scale analyses in mind. To detect reliable molecular biomarkers, it is imperative to handle biological fluids according to standardized procedures and to evaluate the effects of preanalytical parameters on the final result to avoid artifacts (1, 2). Earlier studies on urine, plasma, and cerebrospinal fluid (CSF)9 have shown that sample handling can affect the stability of proteins as well as metabolites (3–11). Standardized sample handling is also important when trying to compare results between different laboratories (12–14). In the search for molecular biomarkers related to disorders of the central nervous system, CSF is the most promising biofluid because of its close contact to the affected tissue (13, 15–20).

In this study, we analyzed a set of human CSF samples to assess protein and metabolite stability at room temperature after a low-speed centrifugation step to remove cells. To cover a wide range of proteins and metabolites, we combined the results from several analytical platforms, including LC-MS, GC-MS, and nuclear magnetic resonance (NMR).

9 Nonstandard abbreviations: CSF, cerebrospinal fluid; NMR, nuclear magnetic resonance; QTOF, quadrupole TOF; MS/MS, tandem mass spectrometry; NSC, nearest shrunken centroid; PCA, principal component analysis; SRM, selected reaction monitoring; TSP-d4, 3-(trimethylsilyl)propionic acid-d4.
Materials and Methods

SAMPLE SET
We obtained 6 human CSF samples from the Department of Neurology at the Erasmus University Medical Center (Rotterdam, the Netherlands). The CSF samples were collected as part of routine clinical examination of patients with various symptoms (Table 1). All samples were collected via lumbar puncture between the third and fourth lumbar vertebrae with a Spinocan needle (0.90 mm × 88 mm). The medical ethics committee of the Erasmus University Medical Center approved the study protocol, and all patients gave their informed consent. Samples were centrifuged (10 min at 956 g) within 5 min after collection to remove cells. Aliquots were directly snap-frozen in liquid nitrogen or left at room temperature for 30 and 120 min before snap-freezing and storage at −80 °C. Routine CSF diagnostics, including total protein and albumin concentration measurements as well as intrathecal cell count, were performed. To eliminate the possibility that samples were contaminated with blood, hemoglobin and apolipoprotein B100 were measured and confirmed to be absent in all specimens. Sample H1 was analyzed only with respect to proteomics.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Age, years</th>
<th>Sex</th>
<th>Diagnosis</th>
<th>Protein concentration, mg/L</th>
<th>Clinic albumin concentration, mg/L</th>
<th>SRM albumin concentration, mg/L</th>
<th>Cells/µL after centrifugation, n</th>
</tr>
</thead>
<tbody>
<tr>
<td>H1b</td>
<td>49</td>
<td>M</td>
<td>Migraine</td>
<td>415</td>
<td>193</td>
<td>192.3</td>
<td>0</td>
</tr>
<tr>
<td>H2</td>
<td>56</td>
<td>M</td>
<td>Idiopathic intracranial hypertension</td>
<td>472</td>
<td>247</td>
<td>237.5</td>
<td>0</td>
</tr>
<tr>
<td>H3</td>
<td>69</td>
<td>F</td>
<td>Headache</td>
<td>395</td>
<td>236</td>
<td>221.0</td>
<td>0</td>
</tr>
<tr>
<td>H4</td>
<td>48</td>
<td>M</td>
<td>Idiopathic intracranial hypertension</td>
<td>436</td>
<td>241</td>
<td>225.6</td>
<td>0</td>
</tr>
<tr>
<td>H5</td>
<td>29</td>
<td>F</td>
<td>Clinical isolated syndrome (neuromyelitis optica)</td>
<td>387</td>
<td>226</td>
<td>213.9</td>
<td>0</td>
</tr>
<tr>
<td>H6</td>
<td>38</td>
<td>F</td>
<td>Epilepsy</td>
<td>381</td>
<td>184</td>
<td>194.2</td>
<td>0</td>
</tr>
</tbody>
</table>

Average over 3 time points (see online Supplementary Table S4).

Table 1. Description of CSF samples used for stability studies.

We processed the data using a pipeline developed in C++ as described (21, 22). MzData-XML data were converted to ASCII format over a mass range of 200–1600 m/z (outside this range no multiply charged ions were detected), a retention time range of 3–80 min (peptide elution range), and an intensity threshold of 300 counts. We applied a double cross-validated nearest shrunkened centroid (NSC) algorithm to the complete peak matrix; for a certain shrinkage value, the NSC comparison gives a cross-validation error between 0 and 1, where 1 implies that both classes are assigned to the wrong class, 0.5 is a random class assignation, and 0 means that both classes are correctly assigned (23). NSC-selected features were compared by

(21). Enrichment and separation was done by using an LC chip (G4240-63001 SPQ110, Agilent Technologies) (separation column, 150 mm × 75 μm Zorbax 300SB-C18, 5 μm; trap column, 160-nL Zorbax 300SB-C18, 5 μm). The LC separations were carried out as described by use of the following gradient: 80-min linear gradient from 3% to 40% B (0.1% formic acid in acetonitrile) in A (0.1% formic in ultrapure water), followed by 10-min linear gradient from 40% to 50% B and 10-min linear gradient from 50% to 3% B (21). We performed MS analysis under the following conditions: mass range, 200–2000 m/z in profile mode; acquisition rate, 1 spectrum/s; fragmenter voltage, 175 V; skimmer voltage, 65 V; OCT 1 RF Vpp, 750 V. The spray voltage was approximately 1800 V, and the drying gas (N2) was 6 L/min at a temperature of 325 °C. Each spectrum was mass corrected by use of internal standards (methyl stearate, m/z 299.294457, and HP-1221, m/z 1221.990637) evaporating from a wetted wick inside the spray chamber. Reproducibility was monitored on selected cytochrome C peaks in the QC samples; mass difference between the theoretical and the measured values was within 4 ppm. The selected peaks showed a peak area CV of <20% and a retention time CV of <2%.

ChipLC QTOF-MS PROTEOMICS ANALYSIS
In quintuplicate, we randomly injected 0.5 μL trypsin-digested CSF (H1–H6; T0, T30, T120), and we injected 0.5 μL digested QC samples (pooled CSF spiked with cytochrome C; Fluka, final concentration, 375 fmol/μL) and blanks between every tenth sample for LC-MS analysis on an Agilent chipLC QTOF-MS system as reported...
univariate statistical analysis (Student t-test with Bonferroni correction for multiple comparisons) and ANOVA (Microsoft Excel 2007 and SPSS 16.0). The features were considered significantly different with a P value <0.05 (T0 vs T120 and T0 vs T30) in at least 5 of 6 samples (T0 vs T120 and T0 vs T30). Each discriminant feature was analyzed by targeted MS/MS for identification. We applied principal component analysis (PCA) (24) to the complete peak matrix (10 000 peaks) as well as to the NSC-selected features (MatLab, R2009a). For visualization, we created box-and-whisker plots in Origin 7.0.

nanoLC ORBITRAP-MS/MS SHOTGUN PROTEOMICS ANALYSIS
Trypsin-digested CSF samples (H1–H6: T0, T30, T120) were injected in random order and analyzed by MS/MS (shotgun approach) on an Ultimate 3000 nanoLC system (Dionex) online coupled to a hybrid linear ion trap/Orbitrap mass spectrometer (LTQ Orbitrap XL; Thermo Fisher Scientific) as described (21).

Data files were analyzed and preprocessed using the Progenesis LC-MS software package (Nonlinear Dynamics). Retention times were aligned, and the intensities of the ions were normalized based on the total ion current. To assess interpatient variability, all identified peaks were analyzed by PCA. All identified peaks were also analyzed by the NSC algorithm for classification (23). Peptides were analyzed for differential abundance between the groups by ANOVA. P values <0.01 were considered significant.

We searched all MS/MS spectra against the UniProt/SwissProt database (version 57.6, taxonomy Homo sapiens, 20 070 sequences) using Mascot (version 2.2.06). Search parameters were as follows: parent ion tolerance, 2 ppm; amino acid modifications, carbamidomethylation of cysteine (fixed) and oxidation of methionine (variable).

nanoLC-MS/MS ANALYSES IN THE SELECTED REACTION MONITORING MODE
Trypsin-digested CSF samples (H1–H6: T0, T30, T120) were spiked with known concentrations of stable isotope-labeled peptide standards corresponding to sequences 427–434 (FQNALLVR) of human serum albumin and 52–62 (ALDFAVGGEYNK) of human cystatin C for quantification by selected reaction monitoring (SRM) (see Supplementary Tables S1 and S2, which accompany the online version of this article at http://www.clinchem.org/content/vol57/issue12).

We performed chromatographic separation of spiked CSF digests on an Ultimate 3000 nanoLC system (Dionex). Spiked CSF digest (1 µL) was loaded onto a C18 trap column (PepMap C18, 300-µm i.d. × 5 mm, 5-µm particle size and 100-Å pore size; Dionex) and washed for 5 min at a flow rate of 20 µL/min 0.1% trifluoroacetic acid in H2O. Next, the trap column was switched in line with the analytical column (PepMap C18, 75-µm i.d. × 150 mm, 3-µm particle size and 100-Å pore size; Dionex). Peptides were eluted at a flow rate of 300 nL/min with the following gradient: 0%–45% solvent B in 30 min, where solvent A is H2O/acetonitrile 98%/2% (vol/vol), 0.1% formic acid and solvent B is H2O/acetonitrile 20%/80% (vol/vol), 0.1% formic acid. The separation of the peptides was monitored with a UV detector (absorption at 214 nm).

Multiple reaction monitoring detection was performed by means of a triple quadrupole tandem mass spectrometer (4000 QTRAP; AB Sciex) in the positive ion mode. As technical control for the measurements, a single spiked CSF digest was measured after every sixth run. A technical control for the enzymatic digestion (1 sample digested at 3 separate times) was also included in this quantitative analysis. We performed data analysis using the multiple reaction monitoring data analysis program Skyline (version 0.7) (25), with the ratio of the analyte peptide to the known concentration of the spiked isotope-labeled internal peptide standard to calculate the concentrations of the original peptides. For the cystatin C peptide, we used the mean of both fragment ion ratios for the determination of the protein concentration, and we used a paired, 2-sided t-test to test for differences in peptide concentrations between the time points.

GC-MS METABOLICOMICS ANALYSIS
CSF samples (H2–H6: T0, T30, T120) were treated with an oximation reagent followed by silylation before GC-MS analysis (21, 26). Each sample was injected twice in random order and analyzed on an Agilent 6890 gas chromatograph coupled to an Agilent 5973 quadrupole mass spectrometer as described (21).

Peaks were characterized by retention time and m/z ratio and identified by comparison with a spectral database (TNO) (21). We analyzed all detected metabolites by PCA. A 2-tailed Student t-test was applied to all known metabolites (T0 vs T30 and T0 vs T120). Metabolites with a P value <0.05 were considered to be significantly affected by storage time.

NMR METABOLICOMICS ANALYSIS
Samples (H2–H6: T0, T30, T120) were randomized before sample preparation and analysis. We diluted 50 µL CSF in 200 µL D2O (heavy water) (99.99% D). We added 25 µL of 8.8 mmol/L TSP-d4 [3-(trimethylsilyl)propionic acid-d4 sodium salt, 99% D] stock solution in D2O to 250 µL CSF to a final concentration of 0.8 mmol/L TSP as internal standard and as chemical shift reference (80.00). The TSP-d4 stock solution was prepared from dry TSP-d4. The pH was adjusted (7.0–7.1) by adding phosphate buffer (9.7 µL of 1 mol/L stock solution) to a final concentration of 35
Finally, the sample (284.7 μL) was transferred to a Shigemi microcell NMR tube for measurements. Each sample was analyzed once.

One-dimensional $^1$H NMR spectra were acquired on an 800-MHz Inova (Varian) system equipped with a 5-mm triple-resonance, Z-gradient HCN cold probe. Suppression of water was achieved by using Watergate (delay, 85 μs) (28). For each spectrum, 256 scans of 18,000 data points were accumulated with a spectral width of 9000 Hz. The acquisition time for each scan was 2 s. An 8-s relaxation delay was used between scans.

Before spectral analysis, all acquired free induction decays were zero-filled to 32,000 data points, multiplied with a 0.3-Hz line-broadening function, Fourier transformed, and manually phased. Calibration of the chemical shift scale was done on the external reference standard TSP-d$_4$ by use of ACD/SpecManager software (Advanced Chemistry Development). Spectra were transformed to MatLab, version 7.6 (R2008b) (Mathworks), for further analysis.

We preprocessed NMR spectral data by baseline correction using the asymmetric least squares method (29) and aligned the data with the correlation-optimized warping method (30). Each spectrum was divided (along the chemical shift axis) into equally sized bins (0.04 ppm), and each data point was averaged over each bin. The areas of the bins were summed to provide an integral so that the intensities of the peaks in such defined spectral regions could be extracted. Each NMR spectrum was reduced to 210 variables, calculated by integrating regions of equal width (0.04 ppm) corresponding to the regions of 0.7–9.

To remove effects of variation in water resonance suppression, spectral regions between 4.4 and 5.4 were removed. All spectra thus reduced were normalized to unit area.

The data were further processed by supervised VAST (variable stability) scaling, to determine group-specific scaling factors (31). To visualize possible systematic variation, grouping, trends, and outliers, we applied PCA to the entire data set. To remove biological (patient-to-patient) variation, data were further mean centered per patient and VAST scaled.

**Fig. 1.** Multivariate statistical analysis (PCA) of the 10,000 most intense peaks selected from chipLC QTOF-MS proteomic data (quintuplicate sample analysis).

No separation based on time between sampling and freezing [T0 (▲)/T30 (*)/T120 (○)] is visible, whereas data from individual samples cluster together, indicating that the interindividual differences are larger than those related to time. (A), All samples. (B), Samples H2 (●) and H5 (■). (C), Samples H1 (+) and H6 (▲). (D), Samples H3 (×) and H4 (†).
reaction mixture in duplicate on an Acquity UPLC™ system (Waters Chromatography B.V.) coupled to a Quattro Premier Xe tandem quadrupole mass spectrometer (Waters) operated under the MassLynx data acquisition software (version 4.1; Waters). Quantification and preanalysis of the data were done using LC-QuanLynx (Waters) and Microsoft Excel 2003, respectively. The complete set of amino acids in all samples was examined by PCA. We analyzed the data using a 2-tailed Student t-test (T0 vs T30 and T0 vs T120); amino acids with P values <0.05 were considered discriminatory.

Results

Proteomics Analysis
The Orbitrap-MS/MS shotgun analysis resulted in a list of 55 421 peaks, from which 5780 peptides were identified. All identified peptides from the Orbitrap-MS/MS data and the 10 000 most intense QTOF-MS peaks (complete peak matrix) were used for unsupervised multivariate statistical analysis (PCA). No trend with respect to delay before storage was visible (Figs. 1A and 2A). Run-to-run variability on the LC-MS/MS platforms for proteomics was within 20% with respect to peak area and within 0.5% with respect to retention time. PCA results showed that biological variation is more prominent than the effect of time at room temperature between sampling and freezing, because data points clustered according to the individual patients rather than according to time points (Fig. 1, B–D and Fig. 2, B–D).

ANOVA comparison by time variance on the Orbitrap-MS/MS data resulted in 56 peaks (6 identified peptides) with a P value <0.01, which is well below 554, the number of peaks that would receive this P value due to chance alone when identical samples were comparing (null hypothesis) (1% of all 55 421 detected peaks). NSC analysis pointed also to only random differences between the time groups, with a cross-validation error of 0.5. This led to the conclusion that for the observed proteins there was no significant discrimination between the samples stored at \(-80 ^\circ\)C immediately after centrifugation.

Fig. 2. Multivariate statistical analysis (PCA) of 5780 identified peaks from nanoLC Orbitrap-MS/MS proteomic data (single sample analyses).

No separation based on time between sampling and freezing [T0 (▼)/T30 (*)/T120 (○)] is visible, whereas data from individual samples cluster together, indicating that the interindividual differences are larger than those related to time. (A), All samples. (B), Samples H2 (*) and H5 (■). (C), Samples H1 (+) and H6 (▲). (D), Samples H3 (×) and H4 (♦).
and samples left at room temperature for 30 or 120 min before being frozen and stored.

NSC analysis of the QTOF-MS data revealed that differences between T0 and T120 were random, with a double cross-validation error of 0.5. Comparison of T0 vs T30 by NSC reached a minimal average cross-validation error of 0.34 and resulted in only 11 selected peaks. PCA on the NSC-selected peaks (T0 vs T30) from the QTOF data showed no clear discrimination but a weak tendency of clustering according to time groups (Fig. 3A).

The concentrations of 2 proteins in CSF, albumin and cystatin C, were measured by targeted MS analysis in the SRM mode. These proteins are exemplary of the vast majority of CSF proteins, which we found remained unchanged after 120 min at room temperature. Albumin was chosen because it represents the largest part of CSF total protein, a parameter that is often used in CSF-based diagnosis of disease, and cystatin C was chosen as a protein that is sensitive to storage conditions (1, 2). Concentrations of cystatin C and albumin were calculated on the basis of the measured ratios of the corresponding spiked isotope-labeled internal peptide standards to their biological counterparts, confirming that variation between the different time points was not statistically significant (see online Supplementary Tables S3 and S4). The measured concentrations of both proteins were both found to agree with reported CSF concentrations (32, 33). The albumin concentrations measured by SRM were also in agreement with albumin concentrations measured by standard clinical chemistry techniques (Table 1). In addition, we assessed the trypsin cleavage efficiency by monitoring the release of a tag from the lysine end of the cystatin C peptide during the regular digestion procedure. After the overnight digestion procedure, the entire peak in the LC-MS data corresponding to the peptide, including the tag, had completely disappeared,

Fig. 3. (A), Multivariate statistical analysis by PCA based on 11 NSC-selected peaks derived from chipLC QTOF-MS proteomics data [T0 (▲) vs sT30 (●)]. (B, C), Univariate statistical analysis of 2 peaks that decreased significantly with respect to delay time between CSF sampling and freezing at room temperature. Data are represented as box-and-whisker plots with significant P values marked (P < 5 × 10⁻⁵) (T0 vs T30 and T0 vs T120). (B), Peak detected at m/z 656.335 that decreased significantly after 30 and 120 min at room temperature. (C), Peak at m/z 736.383 that decreased significantly after 30 and 120 min at room temperature. Statistical analysis was based on 2-tailed Student t-tests with Bonferroni correction of the combined data from 5 repetitive analyses of 6 human CSF samples (H1–H6).
The observed relative SDs for the cystatin C measurements were <10%, and those for albumin were <4%. Technical variability with sample pretreatment was <4%, and without sample pretreatment, <2% (Table 2).

**Table 2. Relative SDs (RSDs) of the SRM measurements of cystatin C and albumin.**

<table>
<thead>
<tr>
<th>Sample</th>
<th>Samples, n</th>
<th>RSD of cystatin C, %</th>
<th>RSD of albumin, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>H1</td>
<td>3</td>
<td>5.20</td>
<td>2.19</td>
</tr>
<tr>
<td>H2</td>
<td>3</td>
<td>6.62</td>
<td>1.78</td>
</tr>
<tr>
<td>H3</td>
<td>3</td>
<td>7.72</td>
<td>3.13</td>
</tr>
<tr>
<td>H4</td>
<td>3</td>
<td>7.55</td>
<td>3.05</td>
</tr>
<tr>
<td>H5</td>
<td>3</td>
<td>9.15</td>
<td>2.59</td>
</tr>
<tr>
<td>H6</td>
<td>3</td>
<td>2.72</td>
<td>1.51</td>
</tr>
<tr>
<td>Technical variation, sample pretreatment</td>
<td>3</td>
<td>3.74</td>
<td>2.34</td>
</tr>
<tr>
<td>Technical variation, MS measurement</td>
<td>4</td>
<td>0.82</td>
<td>1.74</td>
</tr>
</tbody>
</table>

* Delayed storage at −80 °C caused slightly higher RSDs of protein concentrations in individual patients than did technical controls of measuring a single sample multiple times. The values also depend on the measured protein. Mean RSD for cystatin C in the 6 patient samples within ±SD = 6.25 ± 4.99; mean RSD for albumin in the 6 patient samples within ±SD = 2.40 ± 0.66.

**METABOLICOMICS ANALYSIS**

GC-MS analysis detected 86 peaks of which 63 could be (tentatively) identified on the basis of spectral libraries. Repeatability of the GC-MS metabolite and LC-MS amino acid analyses was within 20% with respect to peak area (see online Supplementary Tables S7 and S8). This analysis was complemented by targeted LC-MS of 19 natural amino acids. NMR analysis identified and quantified 51 metabolites with a repeatability of 2%–8% (see Supplementary Table S6). PCA of the data from the different analytical platforms showed that clustering occurs primarily according to the individual patients rather than to the time points when all data are considered (Fig. 4, A–C). Mean centering the NMR data per patient and VAST scaling showed, that there is no variation in the metabolome according to delay time (see online Supplementary Fig. S1). Statistical analysis revealed that the concentration of 2,3,4-trihydroxybutanoic acid (threonic acid) detected by GC-MS (Fig. 4D) increased in all samples with increased time left at room temperature. In sample H2, the increase of this metabolite was extremely high after 120 min. Nonparametric ANOVA (Kruskal–Wallis) showed that the discrimination between T0 and T120 was clearly significant (P < 0.005). The Bonferroni-corrected Student t-test, as well as 1-way ANOVA, resulted in highly significant time–group separation when the sample providing the outlier value was left out (P < 0.00005 for both t-test and ANOVA).

**Discussion**

We present a study of the stability of the measured proteome and metabolome when human CSF samples are left at room temperature for up to 2 h between lumbar puncture and storage at −80°C, to mimic delayed storage in clinical routine practice.

Unsupervised multivariate statistical analysis (PCA) showed that patient-to-patient variation was the most prominent, overriding variation that was due to delayed storage time. After variable selection based on preclassification of the samples according to delay time, we found that only 2 peptides and 2 metabolites changed significantly over time among approximately 6000 detected peptides (2755 unique peptides; see online Supplementary Table S5) and 88 detected metabolites. Our results demonstrate that human CSF prepared according to the described procedure is suitable for proteomics and metabolomics analysis even when left at room temperature for 2 h, provided that all cells have been removed by centrifugation. Quantification of albumin and cystatin C by targeted MS in the SRM mode using stable isotope–labeled internal standard peptides showed that there is no statistically significant difference over 2 h of delay time, confirming our proteomics results.

Another study on the stability of the proteome in CSF at room temperature pointed in the same direction, with the detection of only 2 polypeptides that changed concentrations after storage (32). These samples were contaminated with blood, however, because both polypeptides were derived from hemoglobin. Another study showed that blood contamination decreases the stability of the CSF proteome (11), corroborating our earlier results (21). One explanation for the decreased level of the 2 unidentified peaks in the proteome analysis is the possible adsorption to the vial surface, e.g., via hydrophobic or van der Waals interactions (21, 33, 34). Metabolomics revealed increased concentrations of threonic acid after storage at room temperature. This increase may be caused by oxidative degradation of ascorbic acid (35–37), because here the ascorbic acid concentrations were slightly decreased with increased time at room temperature. This decrease was too small to be a significant factor, however. Interestingly, threonic acid decreased in CSF containing white blood cells (21), a result that might be due to
further metabolism of the acid by enzymes released from white blood cells. The concentration of threonic acid was too low for NMR detection.

The stability of metabolites and proteins measured with the most common analytical profiling methods is an important factor to take into consideration when handling biofluids and designing biomarker studies. A previous study from our team shows that the biological variation of some proteins and peptides has large variability (sometimes exceeding 100%), which requires high discriminating power for compounds considered as biomarker candidates (38).

In conclusion, we assessed the stability of CSF using 5 different analytical platforms, showing that overall there were very minor changes in peptides (2 of approximately 6000 in trypsin-digested CSF) and metabolites (1 metabolite of 93). Earlier studies showed that blood or white blood cell contamination reduces CSF stability considerably, highlighting the importance of the initial centrifugation step. Because we did not add antioxidants, we cannot draw conclusions about other oxygen-sensitive metabolites such as the catecholamines. The observed increase in threonic acid over time, however, indicates that oxygen-sensitive metabolites require specific protective measures during sample preparation and storage.

Author Contributions: All authors confirmed they have contributed to the intellectual content of this paper and have met the following 3 requirements: (a) significant contributions to the conception and design, acquisition of data, or analysis and interpretation of data; (b) drafting or revising the article for intellectual content; and (c) final approval of the published article.

Authors’ Disclosures or Potential Conflicts of Interest: Upon manuscript submission, all authors completed the Disclosures of Potential Conflict of Interest form. Potential conflicts of interest:

Employment or Leadership: None declared.
Consultant or Advisory Role: None declared.
Stock Ownership: None declared.
Honoraria: None declared.
Research Funding: The study was performed within the framework of the Top Institute Pharma project number D4-102. T. Rosenling, the project BioRange 2.2.3 from the Netherlands Proteomics and the Netherlands Bioinformatics Center; C. Christin, the project

Fig. 4. Statistical analysis of metabolomics data derived from human CSF (H2–H6).

(A, B, and C), Multivariate statistical analysis by PCA based on all detected metabolites. (A), GC-MS (90 metabolites, duplicate sample analysis). (B), NMR (51 metabolites, single analysis). (C), LC-MS targeting the 19 natural amino acids (sextuplicate analysis). (D), Univariate statistical analysis (Kruskal–Wallis nonparametric ANOVA) of combined data from duplicate analyses of 5 human CSF samples (H2–H6) by GC-MS. The analysis was visualized by box-and-whisker plots with significant P value (T0 vs T120) marked (P < 5 × 10\(^{-3}\)). Threonic acid was significantly increased (Kruskal–Wallis) after storage at room temperature for 120 min (P < 5 × 10\(^{-3}\)).
Stability of CSF Proteome and Metabolome

BioRange 2.2.3 from the Netherlands Proteomics and the Netherlands Bioinformatics Center; P. Horvatovich, the project BioRange 2.2.3 from the Netherlands Proteomics and the Netherlands Bioinformatics Center; R. Bischoff, the project BioRange 2.2.3 from the Netherlands Proteomics and the Netherlands Bioinformatics Center.

Expert Testimony: None declared.

Role of Sponsor: The funding organizations played no role in the design of study, choice of enrolled patients, review and interpretation of data, or preparation or approval of manuscript.

Acknowledgments: The authors thank Prof. Dr. Rogier Hintzen from the Department of Neurology, Erasmus University Medical Center, Rotterdam, the Netherlands, for providing the CSF samples.

References