Quantitative profiling of endocannabinoids and related N-acyl-ethanolamines in human CSF using nano LC-MS/MS

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Abstract Endocannabinoids, a class of lipid messengers, have emerged as crucial regulators of synaptic communication in the CNS. Dysregulation of these compounds has been implicated in many brain disorders. Although some studies have identified and quantified a limited number of target compounds, a method that provides comprehensive quantitative information on endocannabinoids and related N-acyl-ethanolamines (NAEs) in cerebrospinal fluid (CSF) is currently lacking, as measurements are challenging due to low concentrations under normal physiological conditions. Here we developed and validated a high-throughput nano LC-ESI-MS/MS platform for the simultaneous quantification of endocannabinoids (anandamide and 2-arachidonoylglycerol), ten related NAEs, and eight additional putatively annotated NAEs in human CSF. Requiring only 200 μl of CSF, our method has limits of detection from 0.28 to 61.2 pM with precisions of relative SD <15% for most compounds. We applied our method to CSF from 45 healthy humans and demonstrated potential age and gender effects on concentrations of endocannabinoids and NAEs. Notably, our results show that docosahexaenoyl-ethanolamide concentrations increase with age in males. Our method may offer new opportunities to gain insight into regulatory functions of endocannabinoids in the context of (ab)normal brain function.—Kantae, V., S. Ogino, M. Noga, A. C. Harms, R. M. van Dongen, G. L. J. Onderwater, A. M. J. M. van den Maagdenberg, G. M. Terwindt, M. van der Stelt, M. D. Ferrari, and T. Hankemeier. Quantitative profiling of endocannabinoids and related N-acyl-ethanolamines in human CSF using nano LC-MS/MS. J. Lipid Res. 2017. 58: 615–624.

Supplementary key words brain lipids • quantitation • tandem mass spectrometry • cerebrospinal fluid • liquid chromatography • lipids • miniaturization

The endocannabinoid system has emerged as a crucial regulator of synaptic communication in the CNS (1–3). Dysregulation of this system is implicated in various neurological and psychiatric disorders, such as neuroinflammation, stroke, brain trauma, anxiety, and depression (4–8). The endocannabinoid system consists of endogenous lipid messengers (endocannabinoids) that activate two distinct cannabinoid (CB) (CB1 and CB2) receptors and the enzymes responsible for the synthesis and degradation of the endocannabinoids. Anandamide (AEA; N-arachidonoyl-

Abbreviations: ACN, acetonitrile; AEA, anandamide; 2AG, 2-arachidonoylglycerol; 2-AGE, 2-arachidonoylglycerol ether; CB, cannabinoid; CSF, cerebrospinal fluid; DEA, N-docosatetraenoyl-ethanolamine; DGLA, N-docosahexaenoylethanolamine; DHEA, N-docosahexaenoylethanolamine; FAAH, fatty acid amide hydrolase; ISTD, internal standard; LEA, N-linoleoyl-ethanolamine; LOD, limit of detection; LOQ, limit of quantification; MAGL, monoacylglycerol lipase; MeOH, methanol; MBM, multiple reaction monitoring; NADA, N-arachidonoyl-dopamine; NAE, N-acyl-ethanolamine; OEA, N-oleoyl-ethanolamine; O-AEA, (O-arachidonoyl ethanolamine; PEA, N-palmitoyl-ethanolamine; QC, quality control; RSD, relative standard deviation; SE, N-stearoyl-ethanolamine.

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The online version of this article (available at http://www.jlr.org) contains a supplement.
ethanolamine) and 2-arachidonoylglycerol (2-AG) are the two best studied endocannabinoids (9). They are synthesized on demand from membrane lipids in a stimulus-dependent manner and are degraded by fatty acid amide hydrolase (FAAH) and monoacylglycerol lipase (MAGL), respectively (10, 11).

In addition to AEA and 2-AG, there are endogenous lipid compounds capable of activating the CB1 and CB2 receptors, i.e., virodhamine [O-arachidonoyl ethanolamine (O-AEA)], N-arachidonyl dopamine (NADA), and noladin ether [2-arachidonoylglycerol ether (2-AEG)], but their physiological relevance is largely unknown. Related bioactive lipids, such as N-vinylethanolamine (LEA), N-palmitoylethanolamine (PEA), N-oleylethanolamine (OEA), and N-stearoyl ethanolamine (SEA) are produced through the same biosynthetic pathway as AEA. These N-acyl ethanolamines (NAEs) have negligible or only weak affinity for the CB1 and CB2 receptors, but are able to indirectly modulate CB receptor activity by interfering with endocannabinoid metabolism, the so-called “entourage effect” (12). Of note, some of the NAEs may exert biological activities through other proteins, such as transient receptor potential cation channel subfamily V member 1 (TRPV1), G-coupled protein receptors (GPR55, GPR18, and GPR119), or nuclear receptors (peroxisome proliferator-activated receptors α and γ) (13, 14). It is worth mentioning that the endocannabinoids, arachidonoylglycerols, and fatty acid ethanolamides are not only metabolized by FAAH or MAGL, but are oxidized by several other enzymes, such as cyclo-oxygenase-2 (COX-2), cytochrome P450, and lipoygenase (LOX) enzymes. These enzymes convert endocannabinoids into a new array of oxygenated intermediates, such as prostaglandin ethanoloamides (called as prostamides) and prostaglandin glycerol esters, or the lipids can be lipoxigenated to many hydroxylated end-products. However, these intermediates are pharmacologically distinct from their parent endocannabinoids and possess different biological functions (15, 16).

Quantification of endocannabinoid levels in the CNS is important to understand their role in human brain disease. Because the cerebrospinal fluid (CSF) is exchanged with brain interstitium, it is regarded as the most relevant accessible body fluid that reflects free endocannabinoid and NAE levels in the CNS (17). However, limited information is currently available on endocannabinoid and NAE levels in the CSF from healthy humans or patients. Previous studies have quantified the endocannabinoids, 2-AG and AEA (18–21), and two NAEs (OEA and PEA) (22–24), but not the related lipid messengers.

Under normal physiological conditions, endocannabinoid concentrations are very low (in the nanomolar to femtomolar range) in CSF compared with their levels in blood and tissues. This puts extra demands on the sensitivity of the quantification method. Advanced powerful techniques, such as ultra-HPLC, coupled to MS have been employed to quantify endocannabinoids in blood (25–27) and other tissue extracts (28), but these techniques are less suitable for endocannabinoid quantification in CSF due to its limited availability. We envisioned that nano LC would offer the opportunity to increase the sensitivity of the analytical system; a microfluidic chip-based approach would provide the required sensitivity by allowing injection of a larger sample volume by coupling an on-line enrichment column to a small internal diameter analytical column, resulting in a higher concentration in the eluting peak. In addition, the reduced flow rate of nano LC compared with conventional LC can improve sensitivity through an increase in the ionization efficiency (29). Using microfluidic chip-based technology, incorporating an enrichment column, a six-port valve, a separation column, and a nano electrospray tip allows the coupling of miniaturized separation with a nano ESI source (30). This design results in lower dead volumes and fewer connections compared with classical nano LC equipment, thereby leading to increased sensitivity and robustness by less peak broadening and dilution effects during transfer of the sample to the LC column and later to the LC-MS interface (31).

By making use of these advantages, we developed and validated a nano LC-ESI-MS/MS method for the simultaneous quantification of endocannabinoid and related NAEs in clinical (human) CSF. By combining a simple liquid-liquid extraction method for sample preparation, a nano LC separation, and multiple reaction monitoring (MRM) for detection, we could show that an extended range of endocannabinoids and related molecules can be measured. To show the applicability of this novel method, CSF samples of 45 healthy volunteers were measured to study variation in endocannabinoid levels in relation to gender and age.

MATERIAL AND METHODS

Materials

UPLC-grade acetonitrile (ACN) and methanol (MeOH) were purchased from Actu-All Chemicals (Os, The Netherlands). Formic acid was purchased from Acros Organics (Morris Plains, NJ). Toluene was obtained from Biosolve (Valkenswaard, The Netherlands). EDTA disodium salt was obtained from Sigma-Aldrich Chemie B.V. (Zwijndrecht, The Netherlands). Purified Millipore water was obtained from a Milli-Q PF Plus system (Merck Millipore). NADA, 1-AG, 2-AG, N-docosatetraenylethanolamid (DEA), N-docosahexaenylethanolamide (DHEA), 2-AGE, N-dihomo-γ-linolenylethanolamide (DGLEA), AEA, SEA, OEA, LEA, PEA, O-AEA, and deuterated standards, NADA-d8, 2-AG-d8, DHEA-d4, AEA-d8, SEA-d3, OEA-d4, LEA-d4, and PEA-d4 were purchased from Cayman Chemical (Ann Arbor, MI).

Preparation of standards and internal standards

Pure standards (>98% purity) in ethanol or ACN at different stock concentrations were provided from the manufacturer. The standard stock solutions were diluted to a concentration of 1 mM using ACN. A final mix of working solution at 100 nM for compounds in group A (NADA, AEA, DEA, DHEA, DGLEA, 2-AGE, LEA, and O-AEA) and at 1,000 nM for compounds in group B (2-AG, 1-AG, SEA, OEA, and PEA) were prepared. The deuterated internal standard (ISTD) solutions were diluted to 100 μM except for 2-AG-d8 at 250 μM. A final working solution was prepared with 100 nM AEA-d8, NADA-d8, DHEA-d4 and 1,000 nM 2-AG-d8, SEA-d3, PEA-d4, LEA-d3, and OEA-d4.
Calibration curve preparation

On the day of analysis, the standard working solution was further diluted to seven additional calibration levels as shown in Table 1. Each calibration point was mixed with ISTD-mix at a 1:1 ratio. The concentrations of ISTDs were chosen to be near the endogenous concentrations of the analytes in CSF.

Sampling protocol for clinical CSF samples used for method development and validation

CSF was sampled via lumbar puncture between L3/L4, L4/L5, or L5/S1. CSF was collected in 12 ml polystyrene tubes (numbers 160172 and 160172; Greiner Bio-One B.V., Alphen aan den Rijn, The Netherlands) and processed within 30 min. A small portion was used for cell count, and the remaining CSF was centrifuged at 21°C for 5 min (500 g). A portion of the supernatant was used for clinical measurements (i.e., glucose, total protein), and the remaining CSF was stored in 2.0 ml polyethylene cryotubes (Micro tube, number 368632; Nunc, Nürnberg, Germany) and stored at –20°C. All samples remained at –20°C until they were pooled in a 50 ml polypolypropylene tube (number 210261; Greiner Bio-One B.V.). The pooled sample was aliquoted into 2 ml Eppendorf tubes (number 211-2129; VWR, Amsterdam, The Netherlands) and stored at –80°C. These clinical CSF samples were used for the purpose of method validation and are termed “pooled CSF samples”. On the day of analysis, 1,000 µl of ethanol was added to each aliquot (0.5 ml CSF) to model conditions of study samples, mixed thoroughly, and placed on ice.

Sampling of healthy subjects

CSF study samples were collected for an ongoing biochemical study, which was approved by the local medical ethics committee (LUMC). All subjects provided written informed consent before sampling. Prior to CSF sampling, 6 ml of cold ethanol (ethanol absolute, number 8098; J.T. Baker, Phillipsburg, NJ) was added to a 15 ml polypolypropylene sampling tube (number 188271; Greiner Bio-One B.V.) and placed on ice. Ethanol was added for denaturation of enzymes present in CSF upon sampling. Subsequently, 3 ml of CSF was sampled directly into the tube and inverted several times to mix CSF and ethanol thoroughly. Immediately after sampling, the CSF/ethanol mixture was divided in 1.5 ml aliquots into 7 x 1.8 ml cryotubes (number 368632; Nunc, Harenberg, The Netherlands) and placed on dry ice within 30 min from sampling. All aliquots were stored at –80°C within 60 min from sampling. No extra freeze-thaw cycles were allowed before sample preparation.

Sample extraction

Sample extraction was performed on ice. For CSF samples, 600 µl of the CSF/ethanol mixture (200 µl/400 µl) was transferred into 2 ml Eppendorf tubes and 100 µl of 100 mM citrate buffer (pH 5.0) with 25 mM EDTA and 10 µl of deuterated ISTD mixture were added. After adding 550 µl of ice-cold tolouene, the tubes were thoroughly mixed for 3 min using a bullet blender (Next Advance, Inc., Averill Park, NY), followed by a centrifugation step (4°C, 5,000 g, 5 min). Then 475 µl of the upper layer (tolouene) were aspirated and transferred into 500 µl Eppendorf tubes. Samples were dried under a gentle stream of nitrogen and placed on ice followed by reconstitution with 10 µl of water/MEOH (50/50). The reconstituted samples were centrifuged (4°C, 10,000 g, 5 min) before transferring into vials for LC-MS.

Chip-based nano LC-MS/MS analysis

Endocannabinoids and related NAEs were measured using a targeted method on an Agilent 6460 triple quadrupole MS system in conjunction with an Agilent 1100/1200 series nano-LC system consisting of a G1375 capillary pump, a G2226 nano pump, and a G1376 uWPS autosampler with thermostat (Agilent Technologies, Waldbronn, Germany). Separation was carried out with an Agilent Polarimetric Chip Cube MS interface (Agilent Technologies) operated at ambient temperatures. This chip has a 360 nL enrichment column with a 150 mm (length) x 75 µm (width) separation column and a particle size of 3 µm. The mobile phase was 10 mM formic acid/water (A) and ACN (B). The analytical gradient was performed in two steps: first, 8 µl of sample were loaded on the enrichment column using the capillary pump delivering a mobile phase in isocratic mode composed of 40% B at a flow rate of 2 µl/min. After the sample was trapped on the enrichment column, it was switched in line with the analytical column. The nano pump gradient started at 45% B and linearly increased up to 80% B in 12 min at a flow rate of 400 nl/min. The column was then held at 95% B for 2 min before returning to 45% B. The column was re-equilibrated for 5 min prior to the next injection, giving a total analysis time of 20 min. During the analytical gradient, the enrichment column was flushed with 16 µl of 100% B for 8 min. At the same time, the injection needle was washed with ACN/water (1/1, v/v) using an injection program. The temperature in the autosampler was maintained at 8°C. ESI was operated in positive ion mode with a capillary spray voltage set at 1,800 V and nitrogen nebulizer gas flow rate of 41/min and a temperature of 365°C. For endocannabinoids and related NAEs, individually optimized MRM parameters were determined for each transition for target compounds and ISTDs. The detailed list of MRM parameters and transitions is shown in supplemental Table S1.

To maximize the potential of the platform, MRM transitions for other NAEs were also included. Targets were selected based on: 1) combinations of major fatty acids and ethanolamine (general structure of NAEs) and 2) measurements of CSF by precursor ion scan mode for the characteristic product ion of NAEs, which is the protonated ethanolamine ion at m/z 62 (data not shown). These putatively annotated NAEs are listed as EA (from ethanolamine) followed by a molecular mass (m/z) value. Details of MRM transitions included in the method are listed in supplemental Table S1.

Data preprocessing

Peak area integration was performed with Mass Hunter quantitative analysis (version B.05.01; Agilent, Santa Clara, CA) and manually inspected. Peak areas of analytes were corrected by the

| Compound Group | C0 | C1 | C2 (Low) | C3 | C4 (Medium) | C5 | C6 (High) | C7 |
|----------------|----|----|----------|----|-------------|----|-----------|----|
| A              | 0  | 0.01 | 0.03 | 0.1 | 0.3 | 1 | 3 | 10 |
| B              | 0  | 0.1  | 0.3  | 1   | 3  | 10 | 30 | 100 |

*Seven levels of standard solutions (C1 to C7) are spiked to build the calibration line. These standards will be diluted 21 times in CSF matrix for their final concentrations.*
showed high concentrations for LEA, OEA, PEA, and SEA and were detected as statistical outliers. These concentrations were possibly caused by (an) exogenous source(s) (33). After removal of the outliers, the final QC set was used to determine optimal ISTDs for compounds without an isotopically labeled equivalent as an ISTD, as previously described (34). Values from QC samples were used for monitoring data quality and correcting for shifts in MS sensitivity over time (34). After correction with QC samples, RSDs of study QC samples were calculated for all compounds. Next, corrected data of biological samples (identified compounds only) were analyzed with principle component analysis to detect outliers. In the current data set, one outlier was detected and was removed from further statistical analysis. Finally, RSDs of duplicates were calculated and afterwards one of two samples was randomly selected from the remaining duplicate pairs to analyze variability. One duplicate pair was identified as an outlier due to contamination and was excluded.

Statistical analysis
Prior to statistical analysis, study QC-corrected data were log-transformed for data normalization. To study whether endocannabinoid and related NAE concentrations depend on gender or age, multiple linear regression analysis was applied. For each compound, its concentration was set as the dependent variable with gender and age as predictors. To study whether potential age effects are gender-specific, linear regression was repeated after splitting data by gender. Statistical analysis was performed with R software (version 3.0.3, function lm). All reported P values are without multiple testing corrections. P < 0.05 was considered significant.

RESULTS AND DISCUSSION

Method development
Although some analytical methods for profiling endocannabinoids and related NAEs in human plasma have been reported (25, 27), methods for CSF are limited to AEA, 2-AG, PEA, and OEA (22, 23). To achieve a high-throughput and sensitive method for detection and quantification of these compounds, two important parameters were optimized. First, LC conditions, including gradient and solvent composition, were optimized to obtain an adequate separation of molecules, especially isomers that are indistinguishable by fragmentation pattern, while at the same time maintaining a short chromatographic run time (Fig. 1). For example, DGLEA and EA350 are structural isomers, but are chromatographically separated (Fig. 2). Next, optimal transitions were determined for each compound to maximize sensitivity, and for each MS/MS transition it was confirmed with standards so that there was no cross-talk between compounds. This method uses dynamic MRM, which calculates optimal MRM parameters for multiple analytes using user inputs of retention time windows (delta RT) and cycle time and these parameters remain constant throughout all runs. The individual MRM dwell time is adjusted to maintain constant cycle time, which is selected to provide adequate sampling speed for quantitation of narrow LC peaks. When compared with traditional time segmented methods, dynamic MRM methods include fewer ion transitions per unit time during a typical MS
scans, which results in longer dwell times for individual ion transitions. The MRM was operated in positive mode as the analytes were more efficiently ionized in that mode. For selective detection and quantification of co-eluting metabolites, specific precursor and fragment ions were used for their separation. The fragmentation pattern for NAE derivatives included the characteristic loss of m/z 62.1 resulting from the fragmentation of the amide bond (supplemental Fig. S3A). Our chip-based nano LC-MS/MS method allowed quantification of 12 endocannabinoids and related NAEs, along with eight putatively identified NAEs in CSF (supplemental Table S1).

The isomerization from 2-AG to 1-AG due to acyl migration complicates the measurements of the two regio-isomers. As shown in Table 2, the sum of 2-AG and 1-AG showed a superior RSD (5–11%) during batch measurement compared with 2-AG only (3–38%), which indicated that acyl migration might have occurred during sample preparation and measurement. Another contribution to that error could be problems with peak integration, as the two isomers are not separated at baseline. Based on this result, the sum of 1-AG and 2-AG was used for further data analysis in this work. In liquid-liquid extraction, tolune was previously reported to suppress acyl migration during the evaporation step to a large extent (35). However, moderate levels of acyl migration were observed in the current study and this is likely due to a difference in matrices and sample handling procedures compared with previous studies. Moreover, this is the first study to use a toluene extraction method for CSF; whereas, the previous studies were demonstrated in plasma (25, 35). Furthermore, it has been reported that elevated temperatures, presence of serum albumin in the sample, and high pH values accelerate acyl migration (35, 36). Interestingly, a recent study reports that, in plasma, 1-AG is not an endogenous compound, but rather the result of chemical isomerization during sample storage and, therefore, summing the concentrations of both the isomers will provide meaningful data for biological interpretation (27).

It should be noted that nonnegligible amounts of SEA and, to a lesser extent, other NAEs (PEA, SEA, and OEA) were observed in the ethanol that was used during sample collection. This indicates that ethanol used in the sample collection protocol is a source of contaminants. We also observed contamination of NAEs, especially SEA, OEA, and PEA, at low levels when using MeOH instead of ACN (data not shown). However, only low levels of SEA compared with the endogenous concentration in CSF were seen in the blank (MeOH/water) showing the soundness of the analytical method (supplemental Fig. S1). The consequence of the contaminants present in ethanol, from solvents and other potential sources for PEA, SEA, and OEA, was a decline of analytical characteristics (linearity, LOD, LOQ, precision, recovery, and matrix effect). Previous studies identified basal concentrations of these compounds in organic solvents, plastics, and glassware (33). A recommendation to minimize contamination is to avoid addition of ethanol during CSF sampling or to check for contamination prior to use.

The stability of endocannabinoids and NAEs and their ex vivo generation, i.e., during and after sampling, needs greater attention to avoid variability and inaccuracy of the measurements. To date, the stability of endocannabinoids in CSF was not reported, but in blood, NAE concentrations dramatically increase when withdrawn blood samples are improperly stored. The 2-AG and 1-AG levels were shown to be stable for 2 h, but a gradual reduction of 2-AG was detected after 4 h (25). Here, the pooled CSF used for validation was not stored immediately at cold conditions after withdrawal, which might cause increased
NAE and 2-AG/1-AG levels. This has no negative consequences for this study because these pooled samples were only used to assess the validation parameters by spiking the ISTDs, and no biological conclusions were drawn from this pooled sample. The study samples, on the other hand, were processed and stored within 30 min after sample collection to improve the stability and accuracy of the measurements.

**Method validation**

Validation was performed using pooled CSF samples. Linearity and sensitivity (LOD and LOQ) are summarized in Table 3. Precision (intra- and inter-day) values are summarized in Table 4. Recovery and matrix effect are summarized in Fig. 3.

**Fig. 2.** Endogenous peaks of endocannabinoids and related NAEs in CSF. The x-axis is retention time and the y-axis is response signal.

**TABLE 2.** Difference in percent RSD expressed in percentage between 2-AG alone and combined 2-AG and 1-AG

| Calibration Levels | 2-AG (%) | 1-AG + 2-AG (%) |
|--------------------|----------|-----------------|
| C0                 | 27       | 5               |
| C1                 | 58       | 8               |
| C2                 | 20       | 3               |
| C3                 | 15       | 11              |
| C4                 | 19       | 6               |
| C5                 | 16       | 8               |
| C6                 | 8        | 5               |
| C7                 | 3        | 4               |

**Linearity and sensitivity.** Pooled CSF samples were spiked with seven levels of calibration standards (Table 1) to determine linearity, LOD, and LOQ for each compound (nine targets). The goodness-of-fit, $R^2$, ranges from 0.995 to 0.999 for all compounds [with the exception of SEA ($R^2$ 0.976)], demonstrating that the method is linear in the selected calibration range (supplemental Fig. S4). The LOQs are between 0.9 and 61.2 pM for most of the compounds, except for those with a contamination problem (PEA and SEA), which shows that the method is suitable for measuring endogenous levels in CSF (Table 5).

**Precision.** The intra- and inter-batch variability was assessed using the standards spiked at three different concentrations [low (C2), medium (C4), and high (C6)] in pooled CSF samples (Table 4). Quadruplicate sample preparations were performed in each batch, and a total of three batches were processed. The intra-batch RSDs ranged from 1% to 23%. Possible batch-to-batch effects were calculated using the three calibration levels (C2, C4, and C6) measured on three different days. Inter-batch effects ranged from 3% to 21%.

**Recovery and matrix effect.** Recovery and matrix effect were evaluated using deuterated compounds otherwise used as ISTDs (Fig. 3). Recovery was calculated as the ratio of the response of the ISTD of samples spiked before
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CSF endocannabinoids and related NAEs in healthy subjects

To demonstrate the applicability of the method, we measured 45 human CSF samples from 22 healthy females (mean age 30.9 ± 12.8 years SD) and 23 healthy males (mean age 39.3 ± 16.5 years SD). Nine compounds were detected and quantified (Fig. 2).

Endogenous concentrations and ranges of endocannabinoids measured in healthy subjects are listed in Table 5. In addition, the RSDs of QCs and duplicates used during the analysis of healthy subjects are reported in Table 5. The concentrations of AEA and 2-AG are in line with the previous studies (37). In addition to the compounds shown in Table 3, several putatively annotated NAEs were observed in human CSF (supplemental Fig. S2). Of note, an interesting observation in the current CSF samples is the ratio of DHEA/AEA being around 10, which is contrary to plasma, where the ratio is almost equal to 1 (27). This could be because the availability of DHA is greater in CSF than arachidonic acid, precursors for DHEA and AEA, which are then processed downstream at equal rates. If this is not the case, this could indicate that the synthesizing enzymes have a preference for DHA to generate DHEA or, alternatively, it could be that AEA is a better substrate for FAAH than DHEA. However, further investigation is required to elucidate the exact reason, but this was not the main aim of this study.

**TABLE 3.** Validation parameters: calibration range, retention time, LODs, and LOQs

| Compound | C | C=C | m/z | Calibration Ranges (pM) | Retention Time (min) | R² | LOD (pM) | LOQ (pM) |
|----------|---|-----|-----|-------------------------|----------------------|----|---------|---------|
| PEA      | 16| 0   | 299.5 | 4.7–4761               | 13.29                |    | 0.996   | 35.4    | 107.4   |
| LEA      | 18| 2   | 323.5 | 0.47–476               | 11.99                |    | 0.996   | 3.2     | 9.6     |
| OEA      | 18| 1   | 325.5 | 4.7–4761               | 13.83                |    | 0.997   | 14.4    | 43.7    |
| SEA      | 18| 0   | 327.6 | 4.7–4761               | 16.07                |    | 0.976   | 146.9   | 445     |
| AEA      | 20| 4   | 347.5 | 0.47–476               | 12.90                |    | 0.997   | 0.28    | 0.86    |
| DGLEA    | 20| 3   | 349.6 | 0.47–476               | 12.89                |    | 0.996   | 0.53    | 1.6     |
| DHEA     | 22| 6   | 371.6 | 0.47–476               | 11.84                |    | 0.995   | 0.9     | 2.8     |
| DEA      | 22| 4   | 375.6 | 0.47–476               | 13.88                |    | 0.995   | 0.31    | 0.94    |
| 1-AG + 2-AG | 20| 4   | 378.6 | 4.7–4761               | 13.44 (1-AG)         |    | 0.997   | 20.2    | 61.2    |

The LOD and LOQ values of SEA are high due to the contaminant peak present in ethanol added during sample collection.

*"C" and "C=C" are the number of carbon atoms and the number of double bonds in the fatty acid chain of the molecule, respectively. R², correlation coefficient.

The LOD and LOQ values of SEA are high due to the contaminant peak present in ethanol added during sample collection.

**TABLE 4.** Analysis of pooled CSF samples showing precision (intra- and inter-batch effect)

| Compound | Low (C2) | Middle (C4) | High (C6) | Low (C2) | Middle (C4) | High (C6) |
|----------|----------|-------------|-----------|----------|-------------|-----------|
| PEA      | 9        | 8           | 3         | 14       | 13          | 4         |
| LEA      | 23       | 12          | 11        | 21       | 18          | 7         |
| OEA      | 9        | 8           | 5         | 13       | 12          | 5         |
| SEA      | 14       | 7           | 9         | 15       | 17          | 7         |
| AEA      | 6        | 5           | 6         | 6        | 6           | 5         |
| DGLEA    | 14       | 5           | 10        | 13       | 5           | 8         |
| DHEA     | 5        | 1           | 3         | 4        | 5           | 3         |
| DEA      | 9        | 3           | 5         | 10       | 7           | 4         |
| 1-AG + 2-AG | 11      | 5           | 2         | 14       | 9           | 11        |

The precision values are expressed in RSD. Low, middle, and high are three different concentrations (for concentrations see Table 1) used to show intra- and inter-batch precision.

**TABLE 3.** Validation parameters: calibration range, retention time, LODs, and LOQs

**TABLE 4.** Analysis of pooled CSF samples showing precision (intra- and inter-batch effect)
The identity of one of the putatively annotated NAEs, EA350 peak, was assigned as 5(Z),8(Z),11(Z)-eicosatrienoic acid ethanolamide using a commercially available authentic standard and the identity was further confirmed by comparing m/z, retention time, and fragmentation pattern of the authentic standard. The MS/MS fragmentation is shown in supplemental Fig. S3B. In a similar manner, the identities of all the remaining putatively annotated NAEs should be confirmed by purchasing authentic standards; however, these standards are still not commercially available. Although further work needs to be done for annotating all metabolites and understanding the biological implications, this highly sensitive method clearly shows applicability in measuring a broad range of endocannabinoids and related NAEs.

Although we could not confirm the identity for most of the putatively identified compounds and observed some matrix effect due to ion suppression, we believe that our method with this broader coverage of endocannabinoids and their congeners will help us to understand the biological insights of the endocannabinoid system as a “whole” in future studies, and to possibly understand the biological role of these putatively identified compounds. Moreover, this method depicted acceptable reproducibility and sensitivity of these molecules in the application to clinical samples, such as the migraine samples.

**Gender and age differences in healthy controls**

It is important for designing large case-control studies to know whether endocannabinoid and related NAE concentrations show a gender or age dependence. In our study, these factors were analyzed using data from healthy subjects using multiple linear regression analysis. Significant effects of age were present for DHEA. DHEA concentrations were higher in older males (Fig. 4). In blood, higher concentrations of DHA, the fatty acid of DHEA, were reported in the elderly (≥65 years, gender not reported) (38). Intake of DHA can alter DHEA concentrations in blood and brain (39). Additionally, age-related changes in DHA metabolism could play important roles as well (40). There were no significant gender differences in endocannabinoids or related NAEs when corrected for the age imbalance (supplemental Table S2). This study proves the importance of proper age and sex matching of cases and controls to obtain significant results in studies directed at discovery of endocannabinoid-related biomarkers and/or to obtain novel insights in health and disease.

**CONCLUSIONS**

A nano LC-ESI-MS/MS method for the simultaneous quantification of classical endocannabinoids and the related NAEs was developed and validated in human (clinical) CSF. We measured AEA, 2-AG, ten annotated NAEs, and eight putatively annotated NAEs, whereas previously only four endocannabinoids had been measured in CSF (22, 23). Requiring only 200 μl of CSF per sample, this sensitive method has LODs reaching picomolar-femtomolar levels, while maintaining good linearity and precision. Application of this method in CSF from healthy humans resulted in novel insights on gender and age effects, i.e., concentrations of DHEA increased with age in males, illustrating the possibilities of this method to study the regulatory functions of endocannabinoids in the context of brain (dys)function.

The authors acknowledge Agilent Technologies for their advice regarding the Chip Cube system.

**TABLE 5. Concentrations of endocannabinoids and related NAEs quantified in 45 CSF samples of healthy subjects with the RSD of QCs and duplicates**

| Compound Name | Minimum (pM) | Maximum (pM) | Median (pM) | SD | QC_RSD (%) (n = 18) | Duplicate_RSD (%) (n = 4 Pairs) |
|---------------|-------------|-------------|-------------|----|-------------------|-------------------------------|
| PEA           | 41.4        | 270.4       | 83.7        | 48.7 | 10                | 48                            |
| LEA           | 3.6         | 28.8        | 9.6         | 6.2  | 10                | 33                            |
| OEA           | 22.0        | 116.6       | 44.4        | 19.0 | 15                | 20                            |
| SEA           | 67.9        | 310.3       | 108.4       | 54.5 | 13                | 57                            |
| AEA           | 0.5         | 2.7         | 1.6         | 0.5  | 10                | 14                            |
| DGLEA         | 0.4         | 2.2         | 1.0         | 0.4  | 5                 | 6                             |
| DHEA          | 3.61        | 24.89       | 8.48        | 3.92 | 4                 | 2                             |
| DEA           | 1.2         | 5.6         | 2.9         | 0.9  | 5                 | 6                             |
| 1-AG + 2-AG   | 36.5        | 235.3       | 82.6        | 44.7 | 11                | 6                             |

The RSD expressed in percentage of QCs and duplicates used during the analysis of healthy subjects.
Fig. 4. Relationships between DHEA concentrations and age. Linear regression lines are plotted separately for males (solid line) and females (dotted line). Solid circles represent males and solid triangles represent females. The y-axis shows relative concentrations (in picomoles) before log transformation.

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