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Comparison of LC-MS/MS and chemiluminescent immunoassays for immunosuppressive drugs reveals organ dependent variation in blood cyclosporine A concentrations

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Abstract:

Introduction: Life-long monitoring of immunosuppressive drugs (ISDs) in blood is essential after organ transplantation. However, the ISD concentrations vary depending on the assay employed. ISDs are strongly bound to cytoplasmic proteins in erythrocytes in circulation. Therefore, the relatively rapid sedimentation of blood cells in whole blood samples may affect the results when using liquid handling robots.

Methods: We used 1115 blood samples from outpatients and ward patients with kidney (n=373), liver (n=101), heart (n=29) and bone marrow (n=155) transplant. Whole blood samples were pretreated by protein precipitation. Alternatively, the samples were hemolyzed by freezing prior to precipitation. ISDs were analyzed by a 2-plexing liquid chromatography tandem mass spectrometry (LC-MS/MS) assay and commercial chemiluminescent microparticle immunoassays (CMIA).

Results: The difference between the two sample preparation practices was negligible (<2%). Overall, the measured ISD concentrations in patient samples were lower by LC-MS/MS than by CMIA. The difference was the largest (20.2%) and the smallest (9.1%) in samples from liver and from heart transplant patients, respectively.

Conclusions: CMIA overestimates blood ISD concentrations as compared to LC-MS/MS. The extent of the difference was found to be organ transplant dependent. The ISDs can be quantitated either from intact or hemolyzed blood samples.

Abbreviations

LC-MS/MS, liquid chromatography tandem mass spectrometry
CMIA, chemiluminescent immunoassay

ISD, Immunosuppressive drugs
CsA, cyclosporine A
TAC, tacrolimus
SIR, sirolimus
EVE, everolimus
TDM, therapeutic drug monitoring
HUSLAB, Helsinki University Hospital laboratory
CRM, certified reference material
ACN, acetonitrile
MeOH, methanol
FA, formic acid
QA, quality assurance
rpm, revolutions per minute
RCF, relative centrifugal force
LOD, limit of detection
LOQ, limit of quantitation
RE%, relative error
CV%, coefficient of variation
CE, collision energy
CXP, collision cell exit potential

1 Introduction

Human organ transplantation has been a crucial help saving lives for over 60 years [1].

Immunosuppressive drugs (ISD) are needed after transplant operation to prevent organ rejection.

Due to dose- and concentration-dependent toxicity, the therapeutic range of ISDs is narrow and monitoring of their concentrations in blood is thus a life-long procedure after organ transplantation.

Over the years, analytical methods for determination of ISDs have undergone major developments from chromatographic assays to immunoassays [2,3]. Today, liquid chromatography-tandem mass spectrometry (LC-MS/MS) is considered the golden standard, while the disadvantages of immunoassays are well recognized. For example, non-specific matrix effects and cross-reactions between immunoassay antibodies and drug metabolites can lead to overestimation of the drug concentration. This in turn can lead to critical mistreatment of the patients.

ISDs cyclosporine A (CsA), tacrolimus (TAC), sirolimus (SIR) and everolimus (EVE) are strongly bound to cytoplasmic proteins in erythrocytes. The proportion of bound ISDs has been reported to be 41 – 58% for CsA, 85% for TAC, 95% for SIR and >75% for EVE [4]. Therefore, whole blood instead of plasma specimens are routinely used for therapeutic drug monitoring (TDM) of ISDs. Protein precipitation by robotic sample handling prior to LC-MS/MS is a widely employed approach [5-7]. However, the variable and relatively rapid sedimentation of blood cells is a common challenge [6] and may result in false results. With traditional liquid handling robots, one way to mix patient samples is repeated aspirating and dispensing, which is time-consuming and sometimes inadequate. Alternatively, the samples can be mixed with a separate mixer prior to loading them into the robot and pipet the sample before sedimentation. In this approach, the extent of

sedimentation is difficult to define. To overcome these challenges we optimized an alternative sample preparation technique by hemolyzing the samples prior to pipetting.

More than one hundred ISD monitoring samples are analyzed daily at Helsinki University Hospital laboratory HUSLAB. In addition to the relatively high amount of samples, the clinical need to obtain test results within few hours at the wards sets challenges for the monitoring laboratory. In this study, we describe the development and validation of a high throughput LC-MS/MS assay for CsA, TAC, SIR and EVE with pretreatment by protein precipitation of intact or hemolyzed blood samples, and the comparison of the LC-MS/MS assay with Abbott Architect CMIA assay for CsA, TAC and SIR employing a representative clinical sample comprising different transplant recipients.

2 Experimental

2.1 Reagents and materials

TAC, SIR, EVE and the respective stable isotope-labeled internal standards ($^{13}\text{C}_2, ^2\text{H}_4$ – TAC, $^{13}\text{C}_2, ^2\text{H}_4$ – EVE and $^{13}\text{C}_2, ^2\text{H}_3$ -SIR) were purchased from Alsachim (Illkirch Graffenstaden, France). CsA and its internal standard ($^{13}\text{C}_2, \text{d}_4$ - CsA) were purchased from TRC (Toronto Research Chemicals, Toronto, Canada). Certified reference materials (CRM) were purchased from Cerilliant (1.0 mg/mL of CsA in ACN, 1.0 mg/mL of TAC in ACN and 1.0 mg/mL of rapamycin *i.e.* SIR in ACN, Sigma-Aldrich). Calibrators (Multilevel Calibrator set with six concentrations and XL variant with seven concentrations) and quality assurance (QA) samples (two concentrations) were purchased from Chromsystems Instruments and Chemicals GmbH (Gräfelfing, Germany). LC-MS-grade methanol (MeOH), 2-propanol, zinc sulfate, ammonium formate and formic acid (FA) were purchased from Honeywell Fluka and Sigma-Aldrich. All reagents were of the highest analytical grade.

2.2 Samples and patients

Blood samples were drawn into EDTA-tubes (BD Vacutainer, Franklin Lakes, New Jersey, USA). We used samples from healthy volunteers (n=4) and from organ transplant patients (n=1115). Informed consent was obtained from all volunteers. Patient samples were analyzed as part of their diagnostic workup at Helsinki University Hospital, Helsinki, Finland. In this study, the samples were stripped of identifiers. Samples were from patients treated with CsA (n=561), tacrolimus (n=389) and SIR (n=125). Patients treated with CsA had undergone kidney (n=163), liver (n=63), bone marrow (n=155) and heart (n=12) transplantation. Likewise, patients treated with tacrolimus had kidney (n=210), liver (n=38) and heart (n=17) transplants. The type of transplant of SIR treated outpatients remained unknown. In addition, 40 samples from pediatric patients treated with CsA due to bone marrow transplantation were included.

2.3 Instrumentation

Samples were prepared with a Freedom Evo 150 liquid handling robot with Evoware2 Standard software (Tecan, Männedorf, Switzerland) equipped with a plate shaker and centrifuged with a Rotixa 50 RS centrifuge (Hettich Zentrifugen, Geldermalsen, Netherlands). Transcend II LX-2 multiplexing system consisting of two channels and two columns working simultaneously, Ultimate 3000 RSLC pumps, CTC-PAL autosampler and Aria OS software (Aria OS1.7) were used as the front end in the analytical system (Thermo Fisher Scientific, Waltham, USA). API4000 or API4000 QTRAP triple quadrupole mass spectrometers (AB Sciex, Toronto, Canada) with a Turbo-V electrospray ion source and Analyst software (Analyst 1.6.2) were used for detection. For chromatographic separation, we used a cyano column (XSelect HSS CN 2.5 μ m, 2.1x50 mm, Waters) eluted with a linear gradient of 2 mmol/l ammonium formate and 0.1% formic acid in water (A) and in methanol (B). The flow rate was 600 μ l/min, runtime 3.2 mins and column temperature 45 °C. For the MS/MS detection the next transitions were followed: m/z 1220.0 $\rightarrow$ 1203.0 (CsA), m/z 1227.0 $\rightarrow$ 1210.0 ($^{13}\text{C}_2$, d₄- CsA), m/z 821.5 $\rightarrow$ 768.5 (tacrolimus), m/z 826.6 $\rightarrow$ 773.7 (^{13}C , $^2\text{H}_4$ – tacrolimus), m/z 931.7 $\rightarrow$ 864.7 (SIR), m/z 935.6 $\rightarrow$ 864.7 (^{13}C , $^2\text{H}_3$ -SIR), m/z 975.7 $\rightarrow$ 908.7 (EvE) and m/z 981.6 $\rightarrow$ 914.8 ($^{13}\text{C}_2$, $^2\text{H}_4$ – EvE). Ionization was carried out in positive ion mode with the ion source spray voltage set at 4250 V and the temperature at 325 °C. The injection volume was 5 μ l and a bypass valve was used to divert the first 60 sec of the flow to waste and the following 36 sec to mass detection. Four wash cycles of the syringe and the injector were performed between sample injections.

2.4 Sample preparation

Unless otherwise stated, all samples were carefully mixed horizontally rotating at 100 rpm for at least 15 mins (Heidolph Unimax 2010, Schwabach, Germany). Samples were treated essentially as described previously [7]. Briefly, blood samples, calibrators and controls were pipetted (85 μ l) into the wells of a 96-deep well plate (Thermo Fisher Scientific, Waltham, USA) along with 85 μ l of deionized water. The plate was shaken for 30 sec (1300 rpm). Then, 300 μ l of precipitation solution (MeOH:0.1 mol ZnSO₄; 80:20) containing the internal standards (50 μ g/l $^{13}\text{C}_2$, d₄- CsA, 20 μ g/l ^{13}C , $^2\text{H}_2$ - TAC, 7,5 μ g/l ^{13}C , $^2\text{H}_3$ -SIR and 15 μ g/l $^{13}\text{C}_2$, $^2\text{H}_4$ – EVE) was added. The plate was covered with a sealing tape (Thermo Fisher Scientific, Waltham, USA) and shaken for five mins (1300 rpm). The plate was then centrifuged for ten mins at +4 °C at 2890 RCF and the supernatant was used for analysis. Alternatively, samples were prepared by freezing the whole blood samples at -80 °C ethanol bath for five mins, letting the samples thaw for 10 min in front of a fan or a water bath and reach room temperature, vortex-mixing them shortly, followed by pipetting of samples and reagents as above.

To test the effect of blood cell sedimentation on measured ISD concentration, blood samples (n=15) were first prepared and analyzed as routinely by mixing horizontally for ≥ 15 min as described above. Then, the same samples were left in an upright position for ten mins to allow eventual blood cell sedimentation to occur before the analysis was repeated.

2.5 Immunoassay of blood CsA, TAC and SIR

Samples were analyzed by CMIA employing Architect Reagent Kit for CsA, TAC and SIR with an Abbott Architect i2000SR analyzer (Abbott Diagnostics Division, Santa Clara, USA) routinely used at HUSLAB Helsinki University Hospital. The assays were performed according to the instructions of the manufacturer with manual sample precipitation prior to the automated immunoassay, except that CsA was analyzed in duplicate and reported as the mean. The intra- and inter-assay variation of the assays were 4.8% and $\leq 13.3\%$ for CsA, 4.1% and $\leq 6.3\%$ for TAC and 2.8% and $\leq 4.4\%$ for SIR, respectively.

2.6 Validation of the LC-MS/MS assay

HUSLAB is an accredited laboratory according to the SFS-EN ISO/IEC 17025:2005 and SFS-EN ISO 15189:2013 standards. The assay was validated following HUSLAB standard procedures. Limit of detection (LOD) was defined as the concentration corresponding to the mean intensity of 14 blank samples (sodium chloride 9 mg/ml solution, Braun) plus three SD's. Linearity and limit of quantification (LOQ) were determined by diluting calibrators with sodium chloride solution to obtain a nine-point calibration curve (CsA 3.2 – 1840 $\mu\text{g/l}$, TAC 0.3 – 42.3 $\mu\text{g/l}$, EVE 0.3 – 44.5 $\mu\text{g/l}$ and SIR 0.3 – 49.0 $\mu\text{g/l}$) to be measured six times on three different days. The calibration curves were derived using $1/x^2$ weighted linear least-squares regression. Linearity and limit of quantification (LOQ) was defined as the range and the lowest concentration, respectively, that could be measured with an inaccuracy (relative error, RE) and imprecision (coefficient of variation, CV) $< 15\%$. Intra-assay variation was measured by repeating two levels of QA samples 16 times in one batch and inter-assay variation was assessed by collecting five results on five different days (n=25). Process efficiency was determined according to Marchi *et al* [8] from calculated concentrations in blood and mobile phase spiked with two concentrations of the drugs (CsA 100 $\mu\text{g/l}$ and 500 $\mu\text{g/l}$, TAC, EVE and SIR 6 $\mu\text{g/l}$ and 20 $\mu\text{g/l}$). Accuracy was determined similarly using certified reference materials for spiking. Certified reference material for EVE was not available at the time of this study and accuracy was thus not examined. Signal suppression/enhancement i.e. matrix effect was determined by comparing the peak areas obtained from blank blood supernatant spiked with the drugs (n=4) to those from mobile phase spiked with the same amount of the drugs (CsA 100 $\mu\text{g/l}$ and 500 $\mu\text{g/l}$, TAC, EVE and SIR 6 $\mu\text{g/l}$ and 20 $\mu\text{g/l}$, n=4).

Carry-over was investigated by analyzing 50% methanol after the highest calibrator routinely used (CsA 971 $\mu\text{g/l}$, TAC 41.6 $\mu\text{g/l}$, SIR 49.6 $\mu\text{g/l}$ and EVE 44.1 $\mu\text{g/l}$, n=5). Linearity of sample dilution

was evaluated by diluting high ISD concentration samples with sodium chloride solution (9 mg/ml) 1:2, 1:4, 1:8 and 1:16 (CsA) or 1:2, 1:4, and 1:8 (TAC) or 1:2 and 1:4 (SIR) and multiplying the measured concentration by the respective dilution factor. Stability of the ISDs in blood samples (n=16) at room temperature and at +4 °C was studied during eight consecutive days. Samples with <15% change from the initial concentration were considered stable.

2.7 Statistical methods

Methods were compared and statistically evaluated by Passing-Bablok correlation, Bland-Altman regression, paired t-test and Wilcoxon test using Analyse-it software for Microsoft Excel 4.80.9 (Analyse-it software Ltd., <http://www.analyse-it.com>). A p-value <0.05 was considered statistically significant.

3 Results

3.1 Analytical and preanalytical validation of the LC-MS/MS method

The intra- and inter-assay CVs of the LC-MS/MS assay were $\leq 3.6\%$ and $\leq 3.5\%$ for CsA, $\leq 4.5\%$ and $\leq 6.2\%$ for TAC, $\leq 5.3\%$ and $\leq 5.0\%$ for SIR, and $\leq 5.7\%$ and $\leq 5.8\%$ for EVE, respectively. Carry-over of each ISD was $\leq 0.3\%$. Patient samples with concentrations higher than the highest calibrator were linearly diluted up to 1:16 (CsA), 1:8 (TAC), and 1:4 (SIR). All ISDs in blood samples were stable for eight days at room temperature and at +4 °C. Other validation results are presented in Table 1. No interfering peaks were discovered to disturb selectivity (Figure 1).

3.2 Comparison of sample preparation procedures

Routinely, patient blood samples are mixed for ≥ 15 mins prior to analysis. An alternative sample preparation procedure was to freeze the samples in a -80°C ethanol bath for 5 min prior to protein precipitation. The results of frozen (y) and fresh (x) patient samples were compared. The CsA concentration results in patient samples (n=95) correlated according to the formula $y = -0.2 + 1.0x$ with a t mean difference -0.2% (-0.8 µg/L, p=0.960). Similarly, the TAC results in patient samples (n=51) correlated according to $y = -0.2 + 1.0x$ with a mean difference of 1.8% (0.2 µg/L, p=0.175). In addition, CsA (n=15) and TAC (n=15) blood samples were analyzed immediately after mixing and after a 10 min sedimentation time. The measured blood CsA concentrations after mixing and sedimentation were 192.1 µg/L (range 42 - 396 µg/L) and 157.1 µg/L (range 32.2 - 376 µg/L), respectively. Likewise, blood TAC concentrations were 8.5 µg/L (range 5.3 – 11.2 µg/L) and 9.2 µg/L (range 5.8 – 12.1 µg/L), respectively (Figure 2). In contrast to CsA (mean difference -21.0% or -35.0 µg/L, p=0.171), the mean difference in TAC concentration was statistically significant (8.3% or 0.7 µg/L, p<0.05). However, the mean difference in CsA concentration was statistically significant (3.1% or -5.0 µg/L, p=0.009) when the three outliers were excluded (Figure 2).

3.3 Comparison of LC-MS/MS and CMIA

Our newly developed LC-MS/MS method and the immunoassays were compared using 1115 patient samples (Figure 3 and Tables 2 and 3). The mean CsA and TAC results obtained by the CMIA were 14.8% and 22.9%, respectively, higher than those by the LC-MS/MS assay. SIR (mean 6.2 µg/L by CMIA and 8.4 µg/L by LC-MS/MS) in patient samples (n=125) correlated according to the formula $CMIA = -0.2 + 1.4 LC-MS/MS$ and the mean difference was 35.8% (2.2 µg/L). The difference between CsA, TAC and SIR concentrations by LC-MS/MS and CMIA within all patient groups was statistically significant ($p < 0.05$).

4 Discussion

Our LC-MS/MS assay is fast and suitable for simultaneous quantification of CsA, TAC, SIR and EVE in blood samples. In our laboratory, we run two Transcend II LX-2 multiplexing systems daily for ISD samples. After start-up procedures and calibration (45 min), a result is completed every 1.6 min from both instruments. Samples from the hospital campus area are reported within five hours from sample draw. With the workflow of four to five batches and 100 to 200 samples daily we achieved a comparable turnaround time with another recently published studies [4,7].

According to Mika *et al.*, the most common analytical columns used in current ISD assays are C18 and phenyl-hexyl columns [9]. Cyano C18 columns are designed especially for hydrophobic bases. We found the cyano C18 column chemistry to be more suitable for analysis of hydrophobic ISDs enabling rapid elution, as well as better resolution and peak shape compared to traditional C18 columns.

Today's laboratory analysis of ISD's can roughly be divided into two groups: chromatographic separation with mass-to-charge ratio based detection (LC-MS/MS) and biological recognition of antibody-antigen immunoassays. Both are widely used. LC-MS/MS is considered the gold standard assay with high sensitivity and accuracy, but despite improved user friendliness, it is still quite challenging for the operator. Immunoassay systems may be user friendly, but they lack specificity and sometimes repeatability, and are highly dependent on the reagent supplier. We found that the CMIA overestimates ISD concentrations by 10.0—35.8%. This finding is well in line with previously published studies [10,11].

In this study the smallest mean difference between the LC-MS/MS and CMIA assays was for CsA (14.8%, n=561). Our results agree with those of Vogeser *et al.* using samples from 697 CsA treated patients and of Hetu *et al.* observing a positive average bias of 15.6% (n=36) [10,11]. To the best of our knowledge, LC-MS/MS and CMIA CsA assay comparisons with different transplant cohorts have not been reported before. We found that the largest difference between LC-MS/MS and CMIA concentrations for CsA was within the liver transplant group (25.9%, n=63) and the smallest in our limited heart transplant group (10.0%, n=12). Enzymes like cytochromes P450

located in the endoplasmic reticulum of hepatocytes and enterocytes metabolize hydrophobic CsA to various water-soluble metabolites (AM1, AM9 and AM4N) that are either metabolized further or excreted to bile or urine [12]. According to Wang *et al.*, the concentrations of CsA metabolites can be particularly high in liver transplant patients [13]. The CMIA assay is considered fairly specific but still with $\leq 3.3\%$ cross-reactivity with AM1, AM1c, AM4N, AM9 and AM19 [14]. The liver cohort samples in this study were mainly from patients with a fairly recent transplantation, when CsA metabolite concentrations can be high [13] and elimination of them may be impaired [15]. This in combination with the known CMIA cross-reactivity is likely to explain the observed discrepancy between CMIA and LC-MS/MS results within our liver transplant group. Hanwi *et al.* found high metabolite levels in bone marrow transplant patients. Thus, the 17.0 -19.1% positive CsA bias in our kidney, bone marrow and pediatric cohorts by CMIA as compared to LC-MS/MS assay may also be due to cross-reaction of CsA metabolites in the CMIA. CsA metabolites, physiological changes affecting pharmacokinetics and time after transplantation are likely to explain these differences and should be considered when interpreting results obtained with the CMIA especially with liver transplantation patients.

All TAC treated patient samples (n=389) gave on average 22.9% higher concentrations by the CMIA than by the LC-MS/MS assay. The smallest difference of 1.4 $\mu\text{g/L}$ (20.4%, n=210) was within the kidney cohort and the largest within the heart cohort (2.2 $\mu\text{g/L}$, 28.0%, n=17). Bazin *et al.* reported similar mean biases of 1.06 ng/ml, 1.55 ng/ml and 1.11 ng/ml with heart, liver and kidney transplant cohorts, respectively [17]. Others have reported positive biases of 14 to 20% as well [11,18-20]. Wallemac *et al.* have studied the metabolite cross-reactivity of TAC in *in vitro* samples [20]. Four main metabolites M-I (13-O-demethylTAC), M-II (31-O-demethyltacrolimus), M-III (15-O-demethyltacrolimus), and M-IV(12-hydroxytacrolimus) were measured by CMIA and LC-MS/MS and significant cross-reactivity was proven with M-II (94%) and M-III (45%).

A major mean bias of 35.8% between the LC-MS/MS and CMIA was found in our samples from patients treated with SIR (n=125). Others have observed similar positive biases ranging from 14% to 41.4% [11,21,22]. Schmid *et al.* found significant cross-reactivity of two SIR metabolites in the CMIA assay, mainly that caused by F4 (11-hydroxy-sirolimus, 37%) and by F5 (41-O-demethyl-sirolimus, 20%) [21].

Automation of sample pretreatment by liquid handling robots helps to achieve good repeatability and improved turn-around time by LC-MS/MS assays. However, whole blood sedimentation may be an issue. Our liquid handling robot is programmed to aspirate the sample from ten mm depth. Most of the CsA and all TAC concentrations were higher when samples were left standing for 10 mins before pipetting. This is likely to be explained by sedimentation, *i.e.* concentrated layer of blood cells at the depth where aspiration takes place. Sedimentation rate varies as can be seen

from three samples with clearly decreased CsA concentration. Measured blood TAC concentration increased significantly after sample sedimentation while that of CsA increased in twelve and decreased in three samples. This shows that the extent of sedimentation can result in falsely elevated or decreased measured ISD concentration. These findings may be partly explained by a previous report according to which TAC is 85% bound to erythrocytes and CsA is only 41 – 58% bound [4,23]. Taken together, the effect of blood cell sedimentation cannot be ignored. We developed an alternative sample pretreatment procedure, where blood cells are hemolyzed in -80 °C prior to pipetting. ISD concentrations in frozen and fresh samples showed excellent correlation. Thus, sample hemolysis may be the method of choice when analyzing large sample batches.

5 Conclusion

We compared our newly developed LC-MS/MS assay with a CMIA assays for CsA, TAC and SIR in blood samples. In agreement with previous studies, we found that CMIA overestimates the ISD concentrations. Particularly for CsA, the bias varied between different transplant groups. Moreover, to overcome false ISD results caused by blood cell sedimentation, sample hemolysis prior to pipetting may be a useful approach.

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Table 1. Analytical validation results of the LC-MS/MS assays. n.d., not determined.

	CsA	TAC	EVE	SIR
Retention time, min	1.4	1.2	1.3	1.3
Linear range, µg/l	3.2 – 1840	1.1 – 42.3	0.3 – 44.5	0.3 -49.0
Accuracy, %	91.6	80.3	n.d.	96.4
Process efficiency, % (mean)	94.8	93.9	94.0	89.3
Matrix effect, %	-10.8	-20.7	-11.6	-18.5
LOD, µg/l	1.6	0.3	0.02	0.2
LOQ, µg/l	3.1	1.1	0.3	0.3

Table 2. Comparison of LC-MS/MS (x) and CMIA (y) results (µg/l) in CsA-treated organ transplant patient samples and mean concentrations. Mean difference is from Bland-Altman difference plot.

	Passing-Bablok	Mean difference	Mean concentration LC-MS/MS / CMIA (µg/l)
All (n=561)	$y = -11 + 1.3x$	14.8 %, CI 13.8-15.9% (24.9 µg/l , CI 22.6-27.3 µg/l)	149.4 / 174.3
Kidney (n=163)	$y = -17.1 + 1.3x$	19.5%, CI 17.1-21.8% (29.5 µg/l, CI 25.2-33.7 µg/l)	140.1 / 169.5
Liver (n=63)	$y = -5.5 + 1.3x$	25.9%, CI 21.6-30.3% (47.9 µg/l, CI 39.6-56.2 µg/l)	182.3 / 231.1
Bone marrow (n=155)	$y = -15.5 + 1.3x$	17.0 %, CI 15.0-18.9% (33.2 µg/l, CI 28.1-38.2 µg/l)	181.0 / 214.1
Pediatric (n=40)	$y = -29.3 + 1.4x$	18.5% CI 14.2-22.7% (28.1 µg/l, CI 21.0-35.2 µg/l)	137.5 / 165.6
Heart (n=12)	$y = -6.8 + 1.1x$	10.0%, CI 3.8-16.2% (10.4 µg/l, CI 4.5-16.4 µg/l)	106.7 / 117.2

Table 3. Comparison of LC-MS/MS (x) and CMIA (y) results (µg/l) in TAC-treated organ transplant patient samples and mean concentrations. Mean difference is from Bland-Altman difference plot.

	Passing-Bablok	Mean difference	Mean concentration LC-MS/MS / CMIA (µg/l)
All (n=389)	$y = -0.2 + 1.2x$	22.9%, CI 21.8-24.1% (1.7 µg/l, CI 1.6-1.8 µg/l)	7.7 / 9.4
Kidney (n=210)	$y = -0.2 + 1.2x$	20.0%, CI 18.7-21.2% (1.4 µg/l, CI 1.4-1.6 µg/l)	7.2 / 8.6
Liver (n=38)	$y = 0.3 + 1.2x$	22.7%, CI 20.1-25.3% (1.9 µg/l, CI 1.6-2.2 µg/l)	8.6 / 10.5
Heart (n=17)	$y = 1.6 + 1.1x$	28.0%, CI 21.7-34.3% (2.2 µg/l, CI 1.8-2.6 µg/l)	8.3 / 10.5

Figure 1. Ion chromatograms of (A) CsA calibrator (476 µg/l), (B) CsA patient sample (415 µg/l), (C) TAC calibrator (26.9 µg/l), (D) TAC patient sample (11.1 µg/l), (E) SIR calibrator (13.5 µg/l), (F) SIR patient sample (6.7 µg/l), (G) EVE calibrator (12.2 µg/l) and (H) EVE patient sample (7.3 µg/l). The peak width varies between 6 - 9 secs.

Figure 2. Comparison of CsA (A) and TAC (B) concentrations before and after 10 min sedimentation of the samples (n=15). Three outliers of CsA results are shown with dotted line.

Figure 3. Passing-Bablok (A, B) and Bland-Altman (C, D) analysis of CsA (A, C; n=561) and TAC results (B, D; n=389) in patient samples by LC-MS/MS and CMIA. In Bland-Altman regression analysis uniform line: mean, dotted line: 95% limit of agreement.

Highlights:

- LC-MS/MS and chemiluminescent immunoassay results of four immunosuppressive drugs in patient samples were compared.
- Cyclosporine A concentration in blood from liver transplantation patients was found assay dependent.
- Sedimentation of blood cells effects the measured concentration.
- Hemolyzed whole blood is an alternative sample form.

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