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Citation: Biomedical chromatography (2015), 29(9): 1309-1316

Issue Date: 2015-09

URL: http://hdl.handle.net/2433/202094

This is the peer reviewed version of the following article: Kawanishi, M., Yano, I., Yoshimura, K., Yamamoto, T., Hashi, S., Masuda, S., Kondo, T., Takaori-Kondo, A., and Matsubara, K. (2015) Sensitive and validated LC-MS/MS methods to evaluate mycophenolic acid pharmacokinetics and pharmacodynamics in hematopoietic stem cell transplant patients. Biomed. Chromatogr., 29: 1309–1316, which has been published in final form at http://dx.doi.org/10.1002/bmc.3423. This article may be used for non-commercial purposes in accordance with Wiley Terms and Conditions for Self-Archiving. This is not the published version. Please cite only the published version.

Type: Journal Article

Textversion: author
Sensitive and Validated LC-MS/MS Methods to Evaluate Mycophenolic Acid Pharmacokinetics and Pharmacodynamics in Hematopoietic Stem Cell Transplant Patients

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Short title: LC-MS/MS assessment of MPA pharmacokinetics and pharmacodynamics

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This work was supported in part by JSPS KAKENHI Grant Numbers 24928023, 25460210, and 26927010.

Conflict of Interest: none
ABSTRACT:

Monitoring of pharmacodynamics in addition to pharmacokinetics is one of strategies to individualize mycophenolate mofetil (MMF) therapy. The purpose of this study was to develop sensitive liquid chromatography-tandem mass spectrometry (LC-MS/MS) methods for evaluation of the pharmacokinetics and pharmacodynamics of mycophenolic acid (MPA). Concentrations of mycophenolic acid glucuronide (MPAG), mycophenolic acid acyl-glucuronide (AcMPAG), as well as unbound MPA and MPAG, were determined, and inosine-5′-monophosphate dehydrogenase (IMPDH) activity was calculated by measuring concentrations of produced xanthosine-5′-monophosphate (XMP) and intracellular adenosine-5′-monophosphate (AMP) after incubation of peripheral blood mononuclear cell (PBMC) lysates. A metal-free Mastro™ column and 2 gradient patterns were used to improve the quantification limit of XMP to 0.1 µM. In the clinical MPA concentration range, the linearity of the calibration curve, inter- and intra-day precision, and accuracy satisfied the relevant FDA guidelines. The MPA concentrations in hematopoietic stem cell transplant (HSCT) patients determined by the enzyme assay and the present LC-MS/MS method showed a good correlation ($r^2 = 0.95$, $p < 0.001$). In this study, we report sensitive and validated LC-MS/MS methods to evaluate the pharmacokinetics and pharmacodynamics of...
MPA, which are sufficiently sensitive to assess small quantities of PBMC lysates collected shortly after HSCT.

**Key words:** mycophenolic acid, IMPDH, LC-MS/MS, therapeutic drug monitoring
INTRODUCTION

Mycophenolate mofetil (MMF) is an immunosuppressive agent used worldwide for the prevention of rejection after solid-organ transplantations (Kaufman et al., 2004), and it is also used in hematopoietic stem cell transplant (HSCT) recipients for graft-versus-host disease (GVHD) prophylaxis (Minagawa et al., 2012). The therapeutic effect of mycophenolic acid (MPA), the active and hydrolyzed form of MMF, is based on the potent, selective, and reversible inhibition of inosine-5′-monophosphate dehydrogenase (IMPDH). IMPDH is the rate-limiting enzyme involved in de novo synthesis of guanosine nucleotides; IMPDH catalyzes the oxidation of inosine-5′-monophosphate (IMP) to xanthosine-5′-monophosphate (XMP) by a nicotinamide adenine dinucleotide (NAD⁺)-dependent reaction (Allison et al., 2000). Because this XMP synthesis pathway is essential for the mitogenic function of lymphocytes, the inhibitory effect of MPA on IMPDH stops T cell proliferation and thereby leads to immunosuppression (Allison et al., 2000).

MPA is glucuronidated in the liver mainly to an inactive metabolite, mycophenolic acid glucuronide (MPAG), and partially to a pharmacologically active metabolite, mycophenolic acid acyl-glucuronide (AcMPAG) (Shipkova et al., 2003). Fig. 1 shows the structure of each compound. MPAG undergoes enterohepatic recirculation (Bullingham et al., 1998). MPA is
highly bound to albumin and the unbound fraction of MPA is approximately 1–3% of the total concentration (Nowak et al., 1995). MMF has large inter-individual variability in pharmacokinetics and requires therapeutic drug monitoring (TDM) (Staatz et al., 2007). While the recommended target range for the MPA area under the curve from 0 to 12 hours (AUC$_{0–12}$) in renal transplant recipients is 30–60 mg · h/L (Shaw et al., 2001), the optimal MMF dose for GVHD prophylaxis in HSCT patients has not been established (Minagawa et al., 2012). Moreover, it was reported that pre-transplant IMPDH activity is associated with clinical outcome in renal transplant patients, which suggests that IMPDH activity should be investigated as a suitable marker in the field of transplantation (Glander et al., 2004). Recently, it was reported that the area under the effect curve (AUEC) of IMPDH activity on day 21 after HSCT was associated with non-relapse mortality and overall mortality, but was not associated with GVHD (Li et al., 2014). Further studies are needed on the pharmacokinetics and pharmacodynamics of MPA in HSCT patients.

To evaluate IMPDH activity in peripheral blood mononuclear cells (PBMCs), various methods such as liquid chromatography-tandem mass spectrometry (LC-MS/MS) have been described (Glander et al., 2009; Maiguma et al., 2010; Laverdière et al., 2012). However, there remain several limitations in established IMPDH assay methods: calibration standards
are prepared by dilution without PBMC matrix (Glander et al., 2009; Laverdière et al., 2012), and the quantification limit of XMP is 0.25 µM, which is insufficient to determine the concentrations of XMP and AMP in samples obtained from HSCT patients.

Therefore, we developed sensitive LC-MS/MS methods to measure plasma concentrations of MPA, its 2 metabolites, and unbound MPA and MPAG, as well as IMPDH activity in PBMC lysates. We used PBMC matrix as a calibration standard for our LC-MS/MS analyses according to the Bioanalytical Method Validation (BMV) guideline (FDA, 2013), and aimed to improve the sensitivity of XMP by measurements using a specific column and 2 gradient patterns. We evaluated and validated the presently reported LC-MS/MS methods in samples from 3 healthy volunteers and 6 HSCT patients.

**EXPERIMENTAL**

*Chemicals and reagents*

MPA was purchased from Wako Pure Chemical Industries Ltd. (Osaka, Japan). MPAG was purchased from Analytical Services International Ltd. (London, UK). AcMPAG and XMP disodium salt were purchased from Santa Cruz Biotechnology Inc. (Santa Cruz, CA, USA). Mycophenolic acid-d3 (MPA-D₃) was purchased from Toronto Research Chemicals Inc.
(North York, Canada). Adenosine-5′-monophosphate (AMP) sodium salt from yeast, IMP disodium salt from yeast, and NAD\(^+\) were purchased from Nacalai Tesque Inc. (Kyoto, Japan). 8-Bromo-adenosine-5′-monophosphate (Br-AMP) sodium salt was purchased from Jena Bioscience (Jena, Germany). All the chemicals used were of the highest grade available.

**Stock solutions, working solutions, calibration standards, and quality control samples**

For analysis of MPA, its metabolites, and unbound MPA and MPAG, stock solutions were prepared by dissolving MPA (1 mg/mL), MPAG (10 mg/mL), AcMPAG (1 mg/mL) and MPA-D\(_3\) (1 mg/mL) in methanol. All solutions were stored at -80 °C. Working solutions were prepared by diluting stock solutions with methanol to the appropriate concentrations (200 \(\mu\)g/mL MPA, 1600 \(\mu\)g/mL MPAG, and 40 \(\mu\)g/mL AcMPAG) for analysis of MPA and its metabolites, and further diluting solutions to 0.5 \(\mu\)g/mL MPA and 50 \(\mu\)g/mL MPAG for analysis of unbound MPA and MPAG. All diluted working solutions were stored at -20 °C.

Calibration standards were prepared from the working solutions to yield target concentrations of MPA (0.1 to 20 \(\mu\)g/mL), MPAG (0.8 to 160 \(\mu\)g/mL), and AcMPAG (0.02 to 4 \(\mu\)g/mL) by diluting solutions with acidified plasma for analysis of MPA and its metabolites. For analysis of unbound MPA and MPAG, MPA and MPAG calibration solutions were prepared at
concentrations ranging from 5 to 200 ng/mL and 500 to 20000 ng/mL, respectively, using purified water. The working solution of internal standard MPA-D₃ was prepared from stock solution at the time of the assay by dilution with methanol. Quality control (QC) samples of MPA, MPAG, and AcMPAG were prepared at low, medium, and high concentrations in acidified plasma of healthy subjects and kept at -20 °C.

For analysis of IMPDH activity, XMP (1 mM) and Br-AMP (1 mM) stock solutions were prepared in water, and AMP (1 mM) stock solution was prepared in 0.5% ammonium hydroxide. All solutions were stored at -80 °C. Working solutions of 100 μM XMP and 500 μM AMP were stored at -20 °C. Calibration standards were prepared for XMP (0.1 to 10 μM) and AMP (0.5 to 50 μM) by dilution with a mixture of the supernatant from the PBMC lysate, incubation buffer, and stop solution. The working solution for internal standard Br-AMP was prepared from stock solution at the time of assay by dilution with 95% water and 5% methanol.

Clinical sample collection

Blood samples were collected in 3 mL heparinized collection tubes from 6 HSCT patients before and 1, 2, 4, and 8 hours after MMF administration on day 7 and 21 after
transplantation. Blood samples were centrifuged at 15000 g for 10 minutes, and 20 µL of 10% acetic acid was added to 1 mL of the separated plasma sample for stabilization of AcMPAG. Unbound MPA was separated by ultrafiltration of 500 µL plasma using Amicon Ultra 30 K centrifugal filter devices (Merck Millipore Ltd., Carrigtwohill, Ireland) at 14000 g for 10 min. MPA concentrations of all clinical samples were also determined using an enzyme assay (Roche total MPA assay on Cobas 6000; Roche Diagnostics, Tokyo, Japan), and the results of this enzyme assay were compared to those obtained using the LC-MS/MS method described herein.

PBMCs were isolated from the remaining blood sample by density gradient centrifugation using Ficoll-Paque PLUS (GE Healthcare Bio-Sciences AB, Uppsala, Sweden), and frozen at -20 °C until analysis. Area under the plasma concentration time curve from 0 to 8 hours (AUC\textsubscript{0,8}) of MPA and its metabolites was determined using the linear trapezoidal method. This clinical study was performed in accordance with the Declaration of Helsinki and its amendments and was approved by the Kyoto University Graduate School and Kyoto University Hospital Ethics Committee. Written informed consent was obtained from each patient.
**Assay condition**

IMPDH activity was evaluated in PBMC lysates based on previous reports (Glander *et al.*, 2009; Maiguma *et al.*, 2010; Laverdière *et al.*, 2012) with slight modifications. First, 300 µL of water was added to a sample of thawed PBMC lysate, after which the mixture was agitated by vortexing to break down cell membranes and centrifuged at 1000 g for 2 min. The reaction was initiated by adding 50 µL of the supernatant to 52 µL of incubation mixture consisting of 1.6 mM phosphate buffer (consisting of sodium dihydrogen phosphate and sodium monohydrogen phosphate, pH 7.4), 3.9 mM potassium chloride, 1.2 mM of IMP, and 0.6 mM of NAD\(^+\) (final concentrations). The incubation assay was performed at 37 °C for 150 min. The reaction was stopped by the addition of 5 µL of 2.5 M perchloric acid followed by 10 µL of 3 M potassium monohydrogen phosphate. The mixture was centrifuged for 5 min and the supernatant was used for determination. AMP concentrations were determined using another sample without the addition of IMP and NAD\(^+\). IMPDH activity was calculated based on XMP formation and normalized to intracellular AMP using the following equation, in which produced XMP (\(XMP_{produced}\)) and AMP (\(AMP_{measured}\)) concentrations are expressed in micromoles per liter and incubation time (\(t_i\)) is expressed in seconds:

\[
\text{IMPDH activity (µmol \cdot s}^{-1} \cdot \text{mol}^{-1} \text{AMP}) = \frac{XMP_{produced} \cdot 10^6}{(t_i \cdot AMP_{measured})}.
\]
**Internal standards for LC-MS/MS measurements**

MPA-D₃ and Br-AMP were chosen as internal standards for LC-MS/MS assays (Delavenne et al., 2011; Laverdière et al., 2012). Fifty µL of acidified plasma sample was mixed with 100 µL of 0.2 µg/mL MPA-D₃ internal standard for analysis of MPA and its metabolites, 50 µL of ultra-filtrated plasma sample was mixed with 10 µL of 0.2 µg/mL MPA-D₃ internal standard for analysis of unbound MPA. On one hand, 60 µL of incubation mixture was mixed with 10 µL of 100 µM Br-AMP internal standard for analysis of IMPDH activity. Each sample was filtered through a Cosmonice Filter W (Nacalai Tesque Inc., Kyoto, Japan) with a pore size of 0.45 µm for LC-MS/MS determination.

**LC-MS/MS conditions**

The LC-MS/MS system (LCMS-8040; Shimadzu, Kyoto, Japan) was run in multiple reactions monitoring mode with positive ion mode for determinations. The dwell time was 0.1 sec for all compounds. The columns were kept at 40 °C. The autosampler was kept at 4 °C, and 1 µL sample solution was injected for each determination. LabSolutions LCMS software (Shimadzu) was used to control the instruments and to process the data.
For analysis of MPA, its metabolites and unbound MPA and MPAG, chromatographic separation was achieved with a Gemini C18 column (150 mm × 2.00 mm, particle size 5 µm; Phenomenex, Torrance, CA, USA). The column was eluted with combinations of water (A) and methanol (B) containing 0.1% formic acid and 2 mM ammonium acetate, respectively. The flow rate was 0.2 mL/min. The elution gradient was: 0–5 min, linear from 30 to 90% B; 5–9 min, 90% B; 9–12 min, 100% B; 12–15 min, 30% B. The retention times were as follows: MPA, 6.92 min; MPAG, 5.86 min; AcMPAG, 6.78 min; MPA-D3, 6.92 min. The molecules were detected according to the following mass transitions: 321.1 > 207.1 (MPA), 514.1 > 207.1 (MPAG), 514.2 > 207.1 (AcMPAG), and 324.2 > 210.0 (MPA-D3).

For analysis of IMPDH activity, a metal-free Mastro™ column (150 mm × 2.1 mm, particle size 3 µm; Shimadzu GLC, Tokyo, Japan) was used, and the solvents were water (A) and methanol (B) containing 1 mM ammonium acetate, respectively. The flow rate was 0.2 mL/min, and we used 2 elution gradient patterns. The gradient for XMP analysis was: 0–4.5 min, linear from 10 to 50% B; 4.5–5.0 min, 100% B; 5.0–5.5 min, linear from 100 to 10% B; 5.5–10.0 min, 10% B. The gradient for AMP analysis was: 0–4.5 min, linear from 0 to 50% B; 4.5–5.0 min, 100% B; 5.0–5.5 min, linear from 100 to 0% B; 5.5–10.0 min, 0% B. Retention times in the XMP assay were 1.64 min for XMP and 3.60 min for Br-AMP, and
retention times in the AMP assay were 4.11 min for AMP and 4.83 min for Br-AMP. The molecules were detected according to the following mass transitions: 365.1 > 97.2 (XMP), 348.1 > 136.2 (AMP), and 428.0 > 216.0 (Br-AMP).

**Method validation**

The intra- and inter-day precision levels were assessed based on the coefficient of variation (CV, %), whereas the accuracy (%) was determined by the following equation:

\[
\text{Accuracy} \, (\%) = \left( \frac{\text{QC}_{\text{measured}}}{\text{QC}_{\text{reference}}} \right) \times 100
\]

where \( \text{QC}_{\text{measured}} \) is the measured QC concentration and \( \text{QC}_{\text{reference}} \) is the reference QC concentration. Assessments were validated by analyzing 5 replicates of QC samples. The stability of MPA and its metabolites in acidified plasma samples from patients was examined at 4 °C continuously for the first 24 hours after blood sampling. XMP and AMP stability in PBMC lysate samples from healthy subjects was examined at -20 °C continuously for 10 weeks after PBMC extraction. Seven- or 8-point calibration curves were prepared by adding standard products to blank matrix. The coefficient of determination (\( r^2 \)) was used to evaluate the linearity of the calibration curve. Matrix effects were determined by comparing the peak areas (corrected by internal standards) of compounds added to acidified plasma or PBMC.
matrix with those added to water.

The internal levels of XMP or AMP in the reagents of IMP and NAD$^+$ or PBMC samples were determined by analyzing incubation samples with or without them. The produced XMP concentrations at 0, 60, 120, and 150 min after incubation were determined using PBMCs from 3 healthy subjects. The concentrations of AMP and protein in PBMC lysate samples of various concentrations from 3 healthy subjects were determined. Protein concentrations were measured by the Bradford method.

**Inhibition assay of MPA and AcMPAG**

MPA-mediated inhibition of IMPDH activity was assessed in PBMC lysate samples from 3 healthy subjects. MPA and AcMPAG were added to each PBMC lysate sample to a final concentration ranging from 0 to 300 ng/mL and from 0 to 3000 ng/mL, respectively, and incubated as previously described. MPA- or AcMPAG-mediated inhibition of IMPDH activity was assayed by measuring residual IMPDH activity after the addition of MPA or AcMPAG. Pharmacodynamic parameters were calculated by the following sigmoid inhibitory maximum effect model:

$$E = E_0 \cdot \left[1 - I_{\text{max}} \cdot \frac{C_p^\gamma}{(IC_{50}^\gamma + C_p^\gamma)}\right]$$
where $E_0$ is baseline IMPDH activity, $I_{\text{max}}$ is maximal IMPDH inhibition, $C_p$ is the MPA plasma concentration, $IC_{50}$ is the MPA concentration that causes 50% of maximal IMPDH inhibition, and $\gamma$ is the Hill coefficient that governs the slope of the MPA concentration versus IMPDH activity. The $E_0$ values were taken as the original IMPDH activity of each subject without the addition of MPA and AcMPAG.

**RESULTS**

*Method validation*

We initially confirmed the selectivity of the chromatographic method for the targeted analytes and the internal standard. Fig. 2 shows the chromatograms of every tested analytes in acidified plasma matrix (A–D) and in PBMC matrix (E and F). As shown in Table 1, the calibration curve was linear in the examined range. Matrix effects of each compound and each concentration were stable (Table 1). Table 2 shows the inter- and intra-day precision (CV, %) and accuracy (%). All CVs were less than 15% and accuracy ranged from 85% to 115%, with the exception of the lower limit of quantification for XMP (in this case, CVs were less than 20% and accuracy ranged from 80% to 120%). MPA, MPAG, and AcMPAG were stable at 4 °C under acidic conditions until 24 hours after blood sampling (101.9 ± 4.6%
of initial values, mean ± SD, n = 3). IMPDH activity in PBMC lysate samples stored at -20 °C was stable until 10 weeks after extraction (105.8 ± 14.2% of initial values, mean ± SD, n = 3).

We further measured potential contamination by XMP and AMP in our commercial sources of IMP and NAD+ as previously suggested (Laverdière et al., 2012). As shown in Fig. 3, the IMP and NAD+ used for the incubation medium contained 0.1 µM XMP and 6.1 µM AMP, respectively. The combination of 1.2 mM IMP and 0.6 mM NAD+ in the incubation medium led to linear formation of XMP in 3 healthy subjects under the assay conditions described above (Fig. 4A). The AMP and protein concentrations in PBMC lysates from 3 healthy subjects showed a good correlation, but a certain amount of protein was contained in each sample at 0 µM AMP (Fig. 4B). The addition of MPA (0–300 ng/mL) or AcMPAG (0–3000 ng/mL) to the PBMC lysates showed a similar inhibition curve in 3 healthy subjects (Fig. 5). IC50, Imax, and γ were 1.65 ± 0.92 ng/mL, 0.99 ± 0.03, and 1.08 ± 0.52, respectively, for MPA, and 9.48 ± 1.52 ng/mL, 0.98 ± 0.01, and 1.00 ± 0.23, respectively, for AcMPAG (mean ± SD, n = 3 for each result).

Measurement of clinical samples
Time course data was obtained from 6 HSCT patients on day 7 and 21. The MPA concentrations determined by the enzyme assay and LC-MS/MS showed a good correlation ($r^2 = 0.95$, $p < 0.0001$, Fig. 6A). The AUC$_{0-8}$ ratio of MPAG/MPA varied from 10 to 40 in each individual, but this ratio was similar across measurements in each patient, and thus showed low intra-individual variability (Fig. 6B). The AUC$_{0-8}$ ratio of AcMPAG/MPA showed high inter- and intra-individual variability (Fig. 6C). Fig. 7 shows a typical time profile of MPA concentration and IMPDH activity on day 7 after oral administration of 30 mg/kg/day, three times per day of MMF.

**DISCUSSION**

Based on a previous report (Delavenne *et al.*, 2011), we developed a method to analyze plasma concentrations of MPA, its metabolites MPAG and AcMPAG, and unbound MPA and MPAG. Analytical methods for IMPDH activity have been published previously (Glander *et al.*, 2009; Maiguma *et al.*, 2010; Laverdière *et al.*, 2012), but there are limitations on these established methods: they did not use PBMC matrix for calibration standards, and the quantification limit of XMP was 0.25 µM, which was not sufficient to determine the amount of XMP shortly after HSCT. The LC-MS/MS method described here used PBMC matrix for
calibration standards and allowed the sensitive measurement of IMPDH activity. Our LC-MS/MS method can determine XMP concentrations as low as 0.1 µM by using a metal-free column that suppresses the adsorption of phosphate compounds onto the column surface. Furthermore, the analytical LC-MS/MS methods reported herein were fully validated with regard to correct peak shapes, linearity within the range of clinical concentrations, and reproducibility and accuracy of inter- and intra-day measurement (Fig. 2; Tables 1, 2). We demonstrated that XMP and AMP could be quantified even in the small quantities of PBMC lysates collected from patients on day 7 and 21 after HSCT.

As stated in a previous report (Laverdière et al., 2012), XMP and AMP have been detected as contaminants in commercially available IMP and NAD⁺ (Fig. 3). Therefore, we determined the amount of XMP in each sample by subtracting the XMP contamination (0.1 µM) from the measured values. AMP concentrations were measured without IMP and NAD⁺.

It has been reported that the IC₅₀ of MPA for IMPDH activity in human PBMC lysates in vitro is 5–6 nM (1.6–1.9 ng/mL) (Maiguma et al., 2010) or 2–3 ng/mL (Glander et al., 2001), but no reports have compared the IC₅₀ of MPA to that of AcMPAG. We added 0–300 ng/mL of MPA or 0–3000 ng/mL of AcMPAG to PBMC lysates from 3 healthy subjects and incubated them for 150 min; the IC₅₀ of MPA (1.65 ± 0.92 ng/mL) coincided with previously
reported values, and the IC$_{50}$ of AcMPAG (9.48 ± 1.52 ng/mL) was about 6 times greater than that of MPA (Fig. 5). IMPDH activity was completely inhibited by 100 ng/mL MPA and 1000 ng/mL AcMPAG. Although small inter-individual variability in IMPDH activity was observed in the 3 healthy subjects, large inter-individual variability in IMPDH activity in HSCT patients is expected, as shown in a previous report (Li et al., 2014).

IMPDH activity decreased as the MPA plasma concentration increased, and there was no time delay between the peak MPA concentration and the lowest level of IMPDH activity (Fig. 7). Previous studies show that monitoring pharmacodynamics in addition to pharmacokinetics might be useful for individualized therapy with MPA (Glander et al., 2004). Recently, pharmacokinetic and pharmacodynamic analyses were performed in 56 nonmyeloablative HSCT patients (Li et al., 2014), but IMPDH activity was determined only on day 21, and the activity on day 7 could not be determined because of myelosuppression resulting from even the nonmyeloablative-conditioning regimen. In addition, determination of IMPDH activity not only at a stable time point but also at an early time point after HSCT is important, because IMPDH activity changes as the chimerism progresses. Utilizing the sensitive analytical method reported herein, it is possible to measure IMPDH activity at time points as early as day 7 after HSCT, which is not possible with previously reported analysis
methods. Therefore, our method should allow for precise evaluations of pharmacokinetics and pharmacodynamics that will benefit HSCT patients.

Because each sample contains a different number of PBMCs, IMPDH activity must be corrected by cell number. Although enzyme activity can be normalized using cell count, protein concentration, and AMP concentration, use of cell count for normalization causes high intra- and inter-observer variability. We simultaneously determined the amount of AMP and protein in samples in order to determine the appropriate correction for the number of PBMCs. Although AMP concentrations and protein concentrations showed a good linear correlation (Fig. 4B), each sample included a certain amount of excessive protein, which may have been due to extracellular protein that could not be washed out by the process of PBMC isolation that was performed on the final samples. It has been reported that the use of multiple washing steps after cell isolation may remove MPA from cells, leading to an underestimation of \textit{in vivo} MPA-induced IMPDH inhibition (Glander \textit{et al.}, 2012). However, a minimal washing procedure resulted in residual protein and erythrocyte contamination that may have influenced the estimation of IMPDH activity when corrected by protein concentration (Glander \textit{et al.}, 2012). Normalization of the results from each sample based on its respective intracellular AMP concentration minimizes the impact of contaminants on the determination
of IMPDH activity. Therefore, in our study we used a minimal washing procedure and selected intracellular AMP as the factor by which to correct the number of PBMCs.

The good correlation between MPA plasma concentrations measured using the LC-MS/MS method reported here and the enzyme assay normally used for routine TDM in our hospital indicated that our LC-MS/MS method was comparable to the routine automated method. However, the MPA concentrations measured using the enzymatic method were overestimated in comparison with those measured using LC-MS/MS, possibly because of the cross-reaction of AcMPAG in the enzymatic method. Figs. 6B and C show the inter-individual variability in the AUC ratios of MPAG/MPA and AcMPAG/MPA. Although the reason for the intra-individual variability in the AcMPAG/MPA ratio is unclear, it might be caused by differences in the activity of the UDP-glucuronosyl transferases responsible for metabolism of MPAG and AcMPAG. In future studies, we will investigate the influence of enterohepatic circulation of MPAG and the effect of AcMPAG on the pharmacodynamics of MPA using the newly developed LC-MS/MS methods reported here.

CONCLUSION

We developed sensitive and validated LC-MS/MS methods for the analysis of MPA
pharmacokinetics and pharmacodynamics in HSCT patients. Because we needed to improve the analytical sensitivity of measurements of IMPDH activity for the analysis of samples in HSCT patients, we used a specific column and 2 gradient patterns, and the quantification limit of XMP was improved to 0.1 µM, which allowed us to determine XMP in small samples of PBMCs collected on day 7 after HSCT. The newly developed methods reported here provide good precision and accuracy according to the BMV guideline of the FDA, and are convenient for large-scale clinical studies. Precise MPA pharmacokinetic/pharmacodynamic monitoring will allow clinicians to further optimize MPA-induced immunosuppression for various diseases and transplantation procedures.
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Figure legends

Fig. 1. Structures of MPA (A), MPAG (B), and AcMPAG (C).

Fig. 2. Chromatograms obtained from acidified plasma matrix containing 6.7 ng MPA (A), 53.3 ng MPAG (B), 1.3 ng AcMPAG (C) and 1.3 ng MPA-D3 (D). Chromatograms obtained from PBMC matrix containing 6.9 pmol XMP and 14 pmol Br-AMP (E) and 34.3 pmol AMP and 14 pmol Br-AMP (F).

Fig. 3. Internal levels of XMP and AMP in a buffer solution only (A), containing IMP and NAD⁺ (B), containing PBMCs (C), and containing IMP, NAD⁺, and PBMCs (D). PBMCs were obtained from healthy subjects. Each bar shows the mean ± SD (n = 3).

Fig. 4. Time profiles of XMP production in 3 healthy subjects (A), and the correlation between AMP and protein concentrations in PBMC lysates from 3 healthy subjects (B). Each symbol represents the mean ± SD (n = 3) for each subject. XMP production increased linearly with the time of incubation (A: r² = 0.89–0.97; p < 0.001). The AMP concentration was strongly correlated with the protein concentration (r² = 0.97; p < 0.001).

Fig. 5. *In vitro* inhibition of IMPDH activity by MPA (A) and AcMPAG (B). After PBMC lysates from 3 healthy subjects were incubated for 150 min with several concentrations of MPA or AcMPAG, IMPDH activity was determined. Each figure shows a typical curve from
each subject.

**Fig. 6.** Correlation of MPA concentrations measured using the enzyme assay with those determined using the LC-MS/MS method (A). AUC$_{0.8}$ ratio of MPAG/MPA (B) and AcMPAG/MPA (C) in 6 patients. Time course data obtained from 6 HSCT patients treated with MMF at the first and third week after transplantation were used in the analyses.

**Fig. 7.** A typical time course of MPA concentration and IMPDH activity after oral administration of 30 mg/kg/day, three times per day of MMF on day 7 after HSCT.
<table>
<thead>
<tr>
<th>Calibration range</th>
<th>$r^2$</th>
<th>Matrix effect ($n = 3$)</th>
<th>Mean ± SD (%)</th>
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<tr>
<td></td>
<td></td>
<td>Low</td>
<td>Middle</td>
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<td>MPA 0.1–20 µg/mL</td>
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<td>104 ± 5</td>
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<td>75.0 ± 11.0</td>
<td>94.1 ± 6.7</td>
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<td>free MPA 5–200 ng/mL</td>
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<td>98.7 ± 4.1</td>
<td>99.7 ± 3.2</td>
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<tr>
<td>free MPAG 0.5–20 µg/mL</td>
<td>0.999</td>
<td>97.0 ± 4.2</td>
<td>97.8 ± 3.0</td>
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<td>XMP 0.1–10 µM</td>
<td>0.997</td>
<td>70.3 ± 9.0</td>
<td>76.8 ± 6.4</td>
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<td>AMP 0.5–50 µM</td>
<td>0.999</td>
<td>90.1 ± 8.9</td>
<td>92.1 ± 2.2</td>
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<td>Intra-day (n = 5)</td>
<td></td>
<td>Inter-day (n = 5)</td>
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<tr>
<td></td>
<td>CV (%)</td>
<td>Accuracy (%)</td>
<td>CV (%)</td>
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<td>Middle</td>
<td>High</td>
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<td>MPAG</td>
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<td>5.37</td>
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<td>AcMPAG</td>
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<td>free MPAG</td>
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<td>0.71</td>
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<tr>
<td>XMP</td>
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<td>1.67</td>
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<tr>
<td>AMP</td>
<td>4.47</td>
<td>1.57</td>
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Fig. 2

A. MPA

B. MPAG

C. AcMPAG

D. MPA-D₃

E. Br-AMP

F. AMP
**Fig. 3**

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<th>IMP and NAD⁺ (-)</th>
<th>IMP and NAD⁺ (+)</th>
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<td><strong>PBMCs (-)</strong></td>
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<td>6.05</td>
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<td><strong>PBMCs (+)</strong></td>
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<tr>
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<td>2.54</td>
<td>8.96</td>
</tr>
</tbody>
</table>

**A**

PBMCs (-)

**B**

PBMCs (+)
Fig. 4

A

Incubation time (min)

Produced XMP (µM)

B

AMP (µM)

Protein (mg/dL)
Fig. 5

A

MPA concentration (ng/mL) vs. IMPDH activity (µmol/s/mol AMP), showing a downward trend with increasing concentration.

B

AcMPAG concentration (ng/mL) vs. IMPDH activity (µmol/s/mol AMP), also showing a downward trend with increasing concentration.
Fig. 6

A

$Y = 1.18X + 0.72$

$r^2 = 0.95$

MPA concentration by enzyme assay (μg/mL)

MPA concentration by LC-MS/MS (μg/mL)

B

AUC ratio of MPAG / MPA

Pt. 1  Pt. 2  Pt. 3  Pt. 4  Pt. 5  Pt. 6

0.0  0.1  0.2  0.3  0.4  0.5

AUC ratio of AcMPAG / MPA

Pt. 1  Pt. 2  Pt. 3  Pt. 4  Pt. 5  Pt. 6
Fig. 7

![Graph showing MPA concentration and IMPDH activity over time.](image-url)