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blood, urine, and manure by high-performance liquid chromatography/in ductively coupled plasma mass spectrometry. A method for speciation of arsenic species in blood, urine, and manure was developed using highperformance liquid chromatography (HPLC) coupled with inductively

coupled plasma mass spectrometry (ICP-MS). The separation step included an anion exchange column for the separation of inorganic arsenic species such as arsenite and arsenate and an anion exchange column followed by a cation exchange column for the separation of

organic arsenic species, such as methylarsonic acid, dimethylarsinic acid, and monomethylarsonic acid. The detection limits for inorganic arsenic species and organic arsenic species were 10 ng/L and 10 ng/L, respectively. The recovery percentages for inorganic arsenic species

in blood and urine were greater than 95%, and those for organic arsenic species in blood and urine were less than 90%. Accuracy of these analyses for blood and urine were confirmed by comparing them with the analyses using ultra-high performance liquid chromatography coupled

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with ICP-MS. For organic arsenic species in manure, the recovery percentage was less than 80%. The recoveries of arsenic species in blood and urine were greater than 95%. However, the recoveries for organic arsenic species in manure and total arsenic were less than 90%. The

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analysis was developed as a screening test, so the number of samples was limited to eight samples. The results demonstrated that the proposed method is applicable for speciation of inorganic and organic arsenic species in blood and urine. In vivo antiangiogenic potential of

azithromycin: a role for azithromycin-induced upregulation of adenosine and P2Y1 receptors. The antiinflammatory, antiangiogenic, and proapoptotic actions of the macrolide antibiotic, azithromycin (AZM) are well-established, yet its role in the anti-

angiogenic action of interferon-alpha (IFN- α) in in vitro models is unknown. This study tested the hypothesis that AZM upregulates the expression of genes important to antiangiogenic and proapoptotic actions of IFN- α . We compared the antiangiogenic 6d1f23a050

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