

Acute Effect of Coffee Consumption on Urinary Proteome in Young Healthy Individuals

Abstract

Urine, which is not regulated by homeostatic mechanisms, is more likely to reflect early changes and be a source for early biomarker discovery. But urine can be affected by various factors. Coffee is widely consumed in daily life. It may have effects on urinary biomarker discovery and validation in clinical samples. An evaluation of acute coffee effect on urinary biomarker discoveries is needed. In this study, ten healthy volunteers consumed 3 bags of instant coffee in one day. Urine were collected before and after coffee intake. All urine samples were analyzed by liquid chromatography tandem mass spectrometry. After normalization of spectra, 1418 ± 148 proteins were identified, and 11 proteins were found significantly changed in all ten urine samples after coffee intake, indicating that acute coffee intake exerts impacts on urine proteome. Among all differential proteins, 1 differential protein was previously annotated as urinary candidate biomarker. Coffee is one of the most consumed beverages worldwide. Thus, when using clinical samples, the effect of coffee should be taken into consideration.

Keywords: Coffee intake; Confounding factors; Urinary proteome

Abbreviations: LC-MS/MS: Liquid Chromatography Tandem Mass Spectrometry; SBP: Systolic Blood Pressure; DBP: Diastolic Blood Pressure; HR: Heart Rate; Na: Sodium; K: Potassium; Cl: Chorion; Cr: Creatinine; SDS-PAGE: Sodium Dodecyl Sulfate Polyacrylamide Gel Electrophoresis; DTT: Dithiothreitol; IAA: Iodoacetamide

Introduction

Biomarkers are measurable changes associated with a physiological or pathophysiological process [1]. Unlike cerebrospinal fluid or blood, which is regulated by homeostatic mechanisms to remove changes, urine accumulates and reflects many changes of the whole body [2]. Thus, urine has potential to be a better biomarker source. In addition to diseases of urinary systems, the valuable information from other system diseases can be found in urine too.

As a sensitive body fluid, urine will be affected by various factors [3,4], such as diet, drugs, hormone status or other physiological conditions [5]. The differential proteins caused by confounding factors may interfere some valuable urinary information discovery. It is difficult to accurately determine whether potential biomarkers are associated with any particular pathophysiological condition or confounding factors, especially in clinical urine samples.

Coffee is one of the most consumed beverages worldwide, which is also rich in bioactive compounds [6]. Some studies suggested that coffee of both caffeinated and decaffeinated drinking plays an important role in the prevention of some diseases [7]. However, little is known regarding the influence of coffee on the urinary proteome even though coffee's beneficial effects have been studied extensively. A better understanding of its effects on the urinary proteome is necessary for urinary biomarker research.

Research Article

Volume 5 Issue 2 - 2017

Yanying Ni¹, Fanshuang Zhang¹, Manxia An¹, Wei Yin¹ and Youhe Gao^{1,2*}

¹Department of Pathophysiology, Institute of Basic Medical Sciences, Chinese Academy of Medical Sciences, China

²Department of Biochemistry and Molecular Biology, Beijing Normal University, China

*Corresponding author: Youhe Gao, Department of Biochemistry and Molecular Biology, Beijing Normal University, Gene Engineering and Biotechnology Beijing Key Laboratory, Beijing, 100875, PR. of China, Tel: 86-10-58804382; Email: gaoyouhe@bnu.edu.cn

Received: February 19, 2017 | Published: March 03, 2017

In this study, the short-term effects of coffee on the urinary proteome were investigated using label-free quantitative proteomics. Urine samples were collected before and after coffee consumption. Urinary proteins were identified using liquid chromatography tandem mass spectrometry (LC-MS/MS) (Figure 1).

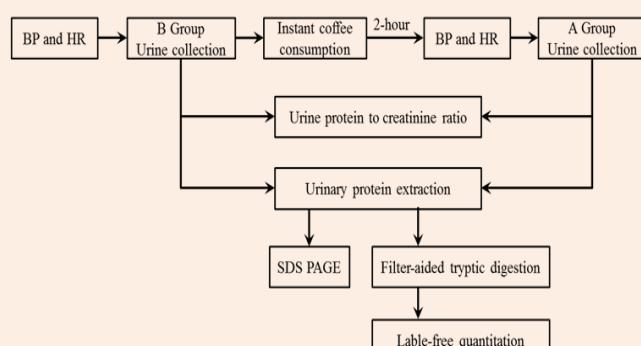


Figure 1: Workflow of effect of acute coffee consumption on urine proteome.

Urine was collected before and after coffee intake from healthy volunteers. The BP and HR were measured before urine collection. The urinary proteins were profiled by label-free LC-MS/MS.

Materials and Methods

Ethics statement

Ten healthy volunteers aged 25-30 years, including 5 men and 5 women, were recruited from Peking Union Medical College, Institute of Basic Medical Sciences, Chinese Academy of Medical Sciences, Beijing, China, during August 2016. Informed consents of all volunteers were obtained before the study. The study was

approved by the Medical Ethics Committee of Peking Union Medical College (#046-2016).

Sample collection and subject characteristics

All volunteers are not habitual coffee drinker and drank three bags of instant coffee (Nescafé Classic, Nestlé, 15 grams) in this study [8]. Urine samples were collected from ten volunteers before and two hours after coffee intake, respectively.

Before urine collecting, systolic blood pressure (SBP), diastolic blood pressure (DBP) and heart rate (HR) were measured using an automatic arm sphygmomanometer (HEM-7052, OMRON, China) 3 times, with a 1min interval between each measurement. Urinary protein, creatinine concentrations and ions in 20 samples were measured at the Peking Union Medical College Hospital.

Urinary protein preparation

Urine was centrifuged at $2000 \times g$ and $12000 \times g$ for 30 minutes at 4°C respectively to remove precipitates. Urinary proteins were extracted by ethanol and protein precipitation were dissolved in lysis buffer (8 M urea, 2M thiourea, 25mM dithiothreitol and 50mM Tris) [9].

Samples were divided into two parts. Some proteins were separated by SDS-PAGE. Some urinary proteins of each samples were digested by trypsin (Trypsin Gold, Mass Spec Grade, Promega, Fitchburg, WI, USA) in 10 kD filter units (Pall, Port Washington, NY, USA) [10]. UA (8 M urea in 0.1 M Tris-HCl, pH 8.5) was firstly added to wash the impurities, and NH4HCO3 was added to wash the remaining UA in the protein solution. After denatured by DDT and alkylated by IAA, the urinary proteins were digested by trypsin with ratio (Enzyme to protein is 1:50).

The collected peptide mixtures were desalting by Oasis HLB cartridges (Waters, Milford, MA) and dried by vacuum evaporation (Thermo Fisher Scientific, Bremen, and Germany).

Urinary proteins analysis by LC-MS/MS

The lyophilized peptides were acidified with 0.1% formic acid, and subjected to a reversed-phase micro-capillary column. The binding peptides were eluted by a gradient of buffer B (0.1% formic acid, 99.9% acetonitrile; flow rate, 0.3L/min) for 100min and were analyzed using Thermo Orbitrap Fusion Lumos (Thermo Fisher Scientific, Bremen, and Germany).

Label-free quantitation

The MS/MS data were analyzed by Progenesis software. Database was set as Uniprot_human database (*Homo sapiens* 84 888 sequences; 34,737,292 residues); the precursor mass tolerance and the fragment tolerance were set 0.05 Da; fixed modification was set carbamidomethylation of cysteines; variable modification was set oxidation of methionine and protein N-terminal acetylation. After normalization, the relative abundance was used to analyze differential expressed proteins.

Statistical analysis

Comparisons between before and after coffee intake were followed by Tukey's multiple comparison tests. P values of less than 0.05 were considered statistically significant. Statistical

analysis was performed by the Statistical Package for Social Studies version 16.0.

Results and Discussion

Transient changes on BP, HR and urine ion after consumption of coffee

The results of the present study showed that acute instant coffee consumption causes a short time decrease in heart rate but does not significantly affect the BP (Figure 2). The concentration of urinary sodium (Na), potassium (K), chorion (Cl) and creatinine (Cr) were found decrease in all urine samples after coffee intake (Figure 3), which may be caused by urine volume increased. The concentrations of urinary ions were measured in urine samples which were collected about two hours after coffee intake instead the total urine output of a specific time frame.

SDS-PAGE analysis of the urinary proteins

The urinary proteins were analyzed using SDS-PAGE (Figure 4). Compared with urinary proteins before coffee intake, the urinary patterns after coffee intake did not show any significant changes.

Urinary proteomics analysis of acute Coffee intake on the human urinary proteome

The urinary samples from ten healthy human volunteers were analyzed using Thermo Orbitrap Fusion Lumos. Each sample was analyzed twice to provide technical replicates (Table S1). After normalization of spectra, 1418 ± 148 proteins were identified (Table S2). According to the standard ($p < 0.05$, fold change ≤ 0.67 or ≥ 1.5), eleven proteins were found to be significantly altered after coffee intake (Table 1), suggesting that acute coffee consumption can affect the urinary proteome. Among these proteins, six proteins showed increased in all ten urine samples after coffee intake and five proteins showed decreased. All differential proteins were searched against the Urinary Protein Biomarker Database [11], one protein has been reported as candidate biomarkers of bladder cancer [12].

Eleven differentially expressed proteins were analyzed using the IPA tool to assess their association with important biological functions. The canonical pathway analysis demonstrated that protein changes were highly associated with Hepatic fibrosis/hepatitis stellate cell activation, Glycolysis and Inhibition of matrix metalloproteases signaling (Figure 5A). Metalloproteinase inhibitor 2 (TIMP2), involved in renal tubular epithelial cell cycle arrest in acute kidney injury, play a significant role in several pathways [13], such as Hepatic fibrosis/hepatitis stellate cell activation, Inhibition of matrix metalloproteases signaling and glioma invasiveness signaling. 14-3-3 protein gamma (YWHAG) also involved in several canonical pathways, such as cell cycle: G2/M DNA damage checkpoint regulation, ERK5 signaling, Myc mediated apoptosis signaling and HIPPO signaling.

The networks analysis showed that the cellular movement, growth and proliferation network were mostly affected (Figure 5B). YWHAG which is critical for maintaining cellular homeostasis, signal transduction and various pathophysiological processes [14,15], is a core component of this networks. In this networks,

ADP-ribosylation factor 3 and Isoform 8 of ARF GTPase-activating protein which involve in energy metabolism, are related to YWHAG.

Compared with the effects of prazosin, aspirin, or pentobarbital sodium on urine proteome [16-18], the protein changes in this study were substantially different. Changes in urine proteome may be caused by many different confounding factors, such as gender [19], diet [20] and exercise [21], which indicates that

the regulation of urinary proteins is complicated. In order to search for potential biomarkers which are truly associated with disease states or physiological variations, analyzing the effects of confounding factors on urine proteome is necessary. Based on current identification technology and quantitation strategy, the effects of acute coffee on urine proteome may not big enough. When animal models were used, the interference of confounding factors can be minimize and the association of urinary proteins with coffee may be more directly [22].

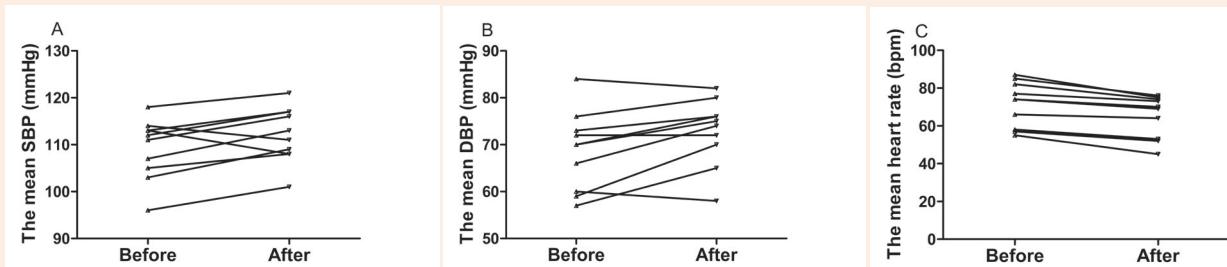


Figure 2: Effect of instant coffee on SBP, DBP and HR before and after coffee intake.

A. SBP increased in eight volunteers. B. DBP increased in seven volunteers. C. All volunteers heart rate decreased after acute coffee intake.'

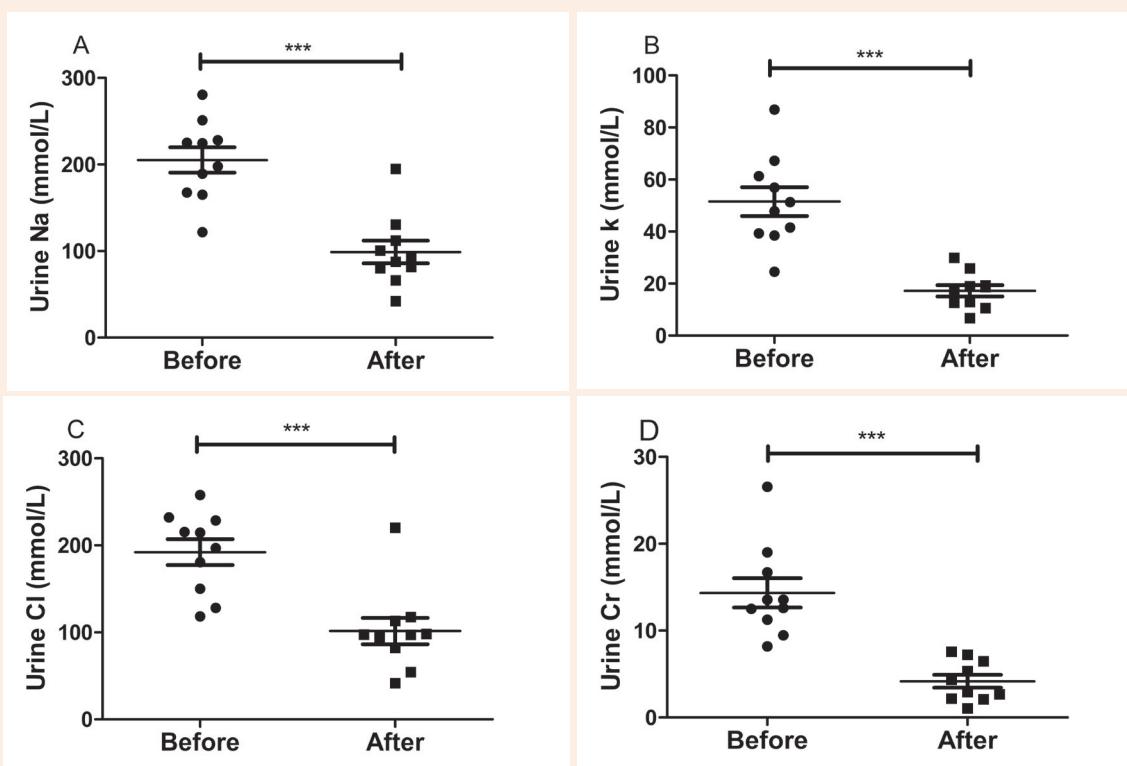
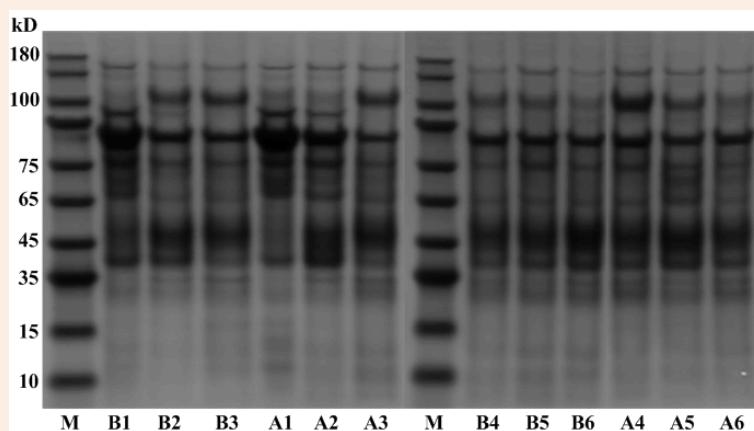
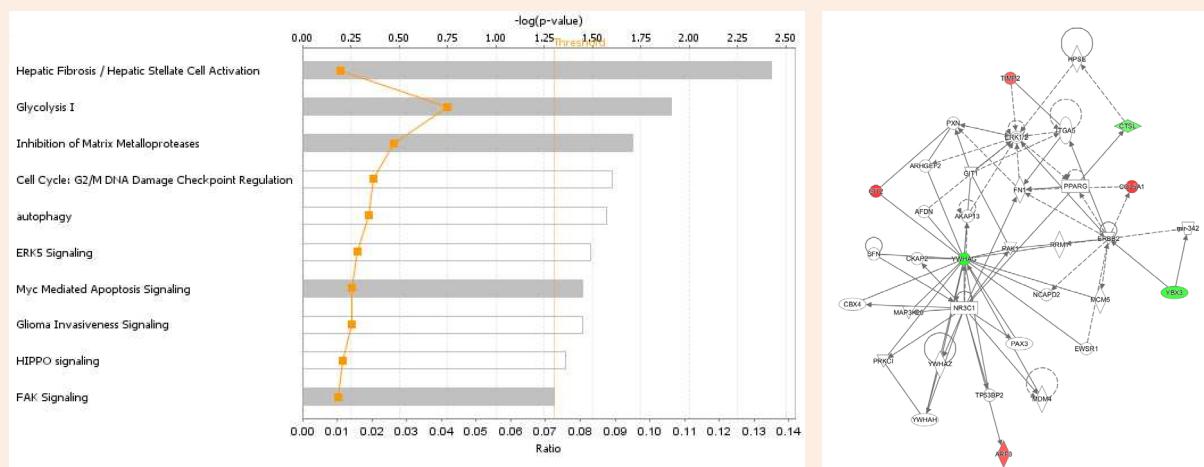


Figure 3: Effect of instant coffee on urinary ions.

A. The urinary Na; B. The urinary K; C: The urinary Cl; D. The urinary Cr.

**Figure 4:** SDS-PAGE analysis of urinary protein samples.

M: Protein marker; B1-6: Urinary protein samples before coffee intake; A1-6: Urinary protein samples after coffee intake.

**Figure 5:** Functional analysis of the protein changes induced by acute coffee intake.

A. Canonical pathways. The y axis shows the negative log of the p value. B. YWHAG network.

Table 1: Changed proteins after acute coffee intake.

Accession	Protein Name	Fold Change	P Value	Candidate Biomarkers of Diseases
Q5T5J6	Transcriptional protein SWT1	1.89	0.019	None
B4DR69	Neuronal PAS domain-containing protein 1	0.49	0.049	None
P07711	Cathepsin L1	0.66	0.014	None
P17858-2	Isoform 2 of 6-phosphofructokinase, liver type	0.48	0.046	None
P20908	Collagen alpha-1(V) chain	1.95	0.026	None
P16989	DNA-binding protein A	0.53	0.041	None
Q14161-8	Isoform 8 of ARF GTPase-activating protein GIT2	2.01	0.037	None
P61981	14-3-3 protein gamma	0.45	0.041	None
F5GWY5	Podocalyxin	1.55	0.017	None
P16035	Metalloproteinase inhibitor 2	1.54	0.042	Bladder cancer
B7ZB63	ADP-ribosylation factor 3	1.60	0.043	None

Table S1. Proteins identified from urine

Group	Biological Repeat	Technical Repeat	Identified Proteins
Before group	B1	1	929
		2	945
	B2	1	1281
		2	1321
	B3	1	1270
		2	1330
	B4	1	1268
		2	1218
	B5	1	1284
		2	1352
After group	B6	1	1267
		2	1394
	B7	1	1287
		2	1367
	B8	1	1008
		2	1054
	B9	1	1244
		2	1313
	B10	1	1241
		2	1293
After group	A1	1	864
		2	846
	A2	1	1405
		2	1445
	A3	1	1200
		2	1222
	A4	1	1119
		2	1149
	A5	1	1246
		2	1266
After group	A6	1	1197
		2	1211
	A7	1	1097
		2	1131
	A8	1	1210
		2	1815
	A9	1	1280
		2	1293
	A10	1	1029
		2	1041

Conclusion

In this study, the effects of acute coffee intake on the urine proteome were analyzed by label-free quantitative proteomics. A total of 11 proteins were found to be significantly altered, and one protein has been previously reported as candidate disease biomarker. Based on current identification technology and quantitation strategy, acute coffee intake may be able to affect the urine proteome and disturb urinary biomarker research. Thus, when using clinical samples, the effect of coffee should be taken into consideration.

Acknowledgement

This work was supported by the National Key Research and Development Program of China (2016YFC1306300); the National Basic Research Program of China (2013CB530850); Beijing Natural Science Foundation (7173264, 7172076) and funds from Beijing Normal University (11100704, 10300-310421102).

References

1. Gao Y (2013) Urine-an untapped goldmine for biomarker discovery? *Sci China Life Sci* 56(12): 1145-1146.
2. Ni Y, Gao Y (2016) Should We Search for Early Brain Disease Biomarkers in Urine. *Author Jnl* 1(1): 00003.
3. Airoldi L, Magagnotti C, Iannuzzi AR, Marelli C, Bagnati R, et al. (2009) Effects of cigarette smoking on the human urinary proteome. *Biochem Biophys Res Commun* 381(3): 397-402.
4. Afkarian M, Bhasin M, Dillon ST, Guerrero MC, Nelson RG, et al. (2010) Optimizing a proteomics platform for urine biomarker discovery. *Mol Cell Proteomics* 9(10): 2195-204.
5. Wu J, Gao Y (2015) Physiological conditions can be reflected in human urine proteome and metabolome. *Expert Rev Proteomics* 12(6): 623-636.
6. Liu JJ, Crous-Bou M, Giovannucci E, De Vivo I (2016) Coffee Consumption Is Positively Associated with Longer Leukocyte Telomere Length in the Nurses' Health Study. *J Nutr* 146(7): 1373-1378.
7. Freedman ND, Park Y, Abnet CC, Hollenbeck AR, Sinha R (2012) Association of coffee drinking with total and cause-specific mortality. *N Engl J Med* 366: 1891-904.
8. van Dam RM, Willett WC, Manson JE, Hu FB (2006) Coffee, caffeine, and risk of type 2 diabetes: a prospective cohort study in younger and middle-aged U.S. women. *Diabetes Care* 29(2): 398-403.
9. Yuan Y, Zhang F, Wu J, Shao C, Gao Y (2015) Urinary candidate biomarker discovery in a rat unilateral ureteral obstruction model. *Sci Rep* 5: 9314.
10. Wiśniewski JR, Zougman A, Nagaraj N, Mann M (2009) Universal sample preparation method for proteome analysis. *Nat Methods* 6(5): 359-362.
11. Shao C1, Li M, Li X, Wei L, Zhu L, et al. (2011) A tool for biomarker discovery in the urinary proteome: a manually curated human and animal urine protein biomarker database. *Mol Cell Proteomics* 10(11): M111.010975.
12. Eissa S, Shabayek MI, Ismail MF, El-Allawy RM, Hamdy MA (2010) Diagnostic evaluation of apoptosis inhibitory gene and tissue inhibitor matrix metalloproteinase-2 in patients with bladder cancer. *IUBMB Life* 62(5): 394-399.
13. Su Y, Gong Z, Wu Y, Tian Y, Liao X (2017) Diagnostic Value of Urine Tissue Inhibitor of Metalloproteinase-2 and Insulin-Like Growth Factor-Binding Protein 7 for Acute Kidney Injury: A Meta-Analysis. *PLoS One* 12(1): e0170214.
14. Wang Z, Nesland JM, Suo Z, Trope CG, Holm R (2011) The prognostic value of 14-3-3 isoforms in vulvar squamous cell carcinoma cases: 14-3-3beta and epsilon are independent prognostic factors for these tumors. *PLoS One* 6(9): e24843.
15. Zhang H, Zhang H, Lin Y, Li J, Pober JS, et al. (2007) RIP1-mediated AIP1 phosphorylation at a 14-3-3-binding site is critical for tumor necrosis factor-induced ASK1-JNK/p38 activation. *J Biol Chem* 282(20): 14788-14796.
16. Zhao M, Li X, Li M, Gao Y (2015) Effects of anesthetics pentobarbital sodium and chloral hydrate on urine proteome. *PeerJ* 3: e813.
17. Zhao M, Wu J, Gao Y (2016) The Specific alpha1-Adrenergic Receptor Antagonist Prazosin Influences the Urine Proteome. *PLoS One* 11(10): e0164796.
18. Li X, Gao Y (2016) Effect of Low-Dose Aspirin on Candidate Biomarkers in the Urine Proteome. *MOJ Proteomics & Bioinform* 3(3): 00087.
19. Liu X, Shao C, Wei L, Duan J, Wu S, et al. (2012) An individual urinary proteome analysis in normal human beings to define the minimal sample number to represent the normal urinary proteome. *Proteome Sci* 10(1): 70.
20. Binder H, Wirth H, Arakelyan A, Lembcke K, Tiys ES, et al. (2014) Time-course human urine proteomics in space-flight simulation experiments. *BMC Genomics* 12(15 Suppl): S2.
21. Kohler M, Franz S, Regeniter A, Ikonen A, Walpurgis K, et al. (2009) Comparison of the urinary protein patterns of athletes by 2D-gel electrophoresis and mass spectrometry-a pilot study. *Drug Test Anal* 1(8): 382-386.
22. Gao Y (2014) Roadmap to the Urine Biomarker Era. *MOJ Proteomics & Bioinform* 1: 00005.