Serum uric acid: Association with anthropometric, lipid and complete blood count parameters in Iranian healthy population

CURRENT STATUS: UNDER REVIEW

BMC Cardiovascular Disorders ▶ BMC Series

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DOI: 10.21203/rs.3.rs-20016/v1

SUBJECT AREAS
   Cardiothoracic Surgery   Cardiac & Cardiovascular Systems

KEYWORDS
   Complete blood count, lipid profile, anthropometric index, serum uric acid
Abstract
Objectives A relationship between elevated serum uric acid (SUA) and hypertension, metabolic syndrome and cardiovascular disease has been established. In this study, the relationship of SUA levels and anthropometric measures, serum lipid profile and neutrophil to lymphocyte ratio (NLR) and platelet to lymphocyte ratio (PLR) was examined.

Methods Anthropometric parameters including body-mass index (BMI), waist circumference (WC), waist to height ratio (WHtR), waist to hip ratio (W/H), waist to pelvic ratio (W/P), neck circumference (NC), body fat mass (BFM) were obtained, and serum lipid profile containing, low-density lipoprotein cholesterol (LDL-C), high-density lipoprotein cholesterol (HDL-C), total cholesterol, triglycerides (TG), also, NLP, PLR, basal metabolic rate (BMR), visceral fat level (VFL) and percent body fat (PBF) were measured among 2921 healthy subjects of young and middle-aged Iranian people.

Statistical analysis was performed using SPSS 21 software. To assess the normality of data, the Kolmogorov–Smirnov test was used. Logarithmic transformation was performed for some variables with non-normal distribution. The association between 2 quantitative variables was measured using bivariate correlation (Pearson or Spearman). Pearson correlations and multiple regression analysis were performed to assess the correlation between variables. Simple and multiple regression analyses were performed to predict some variables. P-value < 0.05 was considered significant.

Results There were 1113 males (mean age, 41.49 ± 8.62 years) and 1808 females (mean age, 42.36 ± 9.07 years) in this study. The male group had a mean SUA level of 4.81 ± 1.2 mg/dl and the female group had a mean of 4.76 ± 1.1 mg/dl. The results of data analysis showed all studied factors were correlated with SUA level except VFL, BFM, and PLR. The highest correlation was related to skinfold fat thickness, BMR and HDL.

Conclusion According to the finding of this study, SUA level measurement might be advisable in healthy population to identify those at increased risk of health problems who might benefit from further evaluation.

Introduction
SUA is produced by metabolism of purine nucleotides, it is one of the major hydrophilic antioxidant in
human blood plasma and inhibits free radicals, so, there were some theories that increased SUA levels would be good based on antioxidant activities [1, 2, 3, 4, 5].

On the other hand, SUA increase correlates and predicts development of conditions associated with oxidative stress. [6] Various studies have shown the relationship between elevated SUA levels and an increased risk of hypertension, metabolic syndrome, obesity and renal disease [7-14].

Moreover, recent evidence has suggested that UA is an inflammatory factor, which can induce endothelial dysfunction and elevated SUA can be considered as a cardiovascular risk factor, therefore, SUA has been lionized as a cardiovascular risk factor in recent prospective and cohort studies [15, 16].

The contribution of dyslipidemia and disturbed anthropometric indices to cardiovascular risk are well established [17-19]. BMI and central adiposity have been identified as potential risk factors of major metabolic problems. In several studies on metabolic syndrome have reported that SUA is related to BMI, WC, and dyslipidemia [16, 20, 21].

NLR and PLR are novel inflammatory markers and associated with worse results in various disease states [22].

The purpose of this study was to evaluate the overall of SUA level in young and middle age healthy population and to explore the association between SUA and anthropometric indices, serum lipid profile, NLR and PLR.

**Method**

**Participants**

This cross sectional study is a pilot study of a health care employee’s prospective cohort study among staff at Shahid Beheshti University of Medical Sciences in Tehran, Iran, which was initiated in October 2017 [23].

In this study, there are 3256 employees who participate from 2017 to 2019. The documents are collected using self-report data, interview, clinical and laboratory findings.

The examinations include to complete a health and medical history questionnaire, taking a blood sample and measurement of anthropometric parameters. Eligibility criteria included age18-65 years,
voluntarily participation, and related medical history. Patients with a history of disease including diabetes mellitus, hypertension, heart disease, hepatic, kidney, gastrointestinal, hematological and respiratory disease, cancer and stroke were excluded from this study. Ultimately, in total 2921 subjects were enrolled in this study. The participants were divided to 3 groups based on their BMI (Kg/m2). Normal group 18.5=<BMI<24.9, over weight:25=<BMI<29.9, and obese group: BMI≥30. Participants were informed about the study by written informed consent. The confidentiality of participants’ information was respected.

**Data collection**

Complete cell count, fasting sugar, lipids (including total cholesterol, HDL, LDL-C, triglyceride), aspartate aminotransferase (AST), alanine aminotransferase (ALT), blood urine nitrogen (BUN), serum uric acid, creatinine (Cr), and routine examination of the urine were measured after fasting for at least 12 hours. The blood samples were examined within 4 hours of collection.

Anthropometric parameters including height, weight, BMI, arm circumference (AC), wrist circumference, hip circumference (HC), pelvic circumference (PC), WC, NC, W/H and W/P were measured using the InBody 770 analyzer.

The triceps skinfold site was used for the assessment of skinfold fat thickness using caliper device. Nursing staffs educated for the survey measured anthropometry and supervised by a trained doctor. Height was measured with a fixed standard stadia meter following study protocols without shoes and weight was assessed in light clothing. BMI was computed as weight (kg) divided by height in meters squared (m²).

**Statistical analysis**

Data analyzing was done using SPSS 21 software (Version 18.0; SPSS Inc. Chicago, IL, USA). For descriptive statistics of qualitative variables, number and percent, and for describing quantitative variables, mean and standard deviation (SD) were used. Mann-Whitney test was used for comparing continuous variable with non-normal distribution. Chi-square test was done for evaluating association between categorical variables. For assessing correlation between non-normal distributed
continuous variable, Spearman’s test was used. Simple and multiple regression analyses were performed to predict some variables.

Results
A total of 2921 employees were participated in study. Among all participants, 1113 (38.1%) were male and 1808 (61.9%) were female. The mean age of female and male was 42.36 ± 9.07 and 41.49 ± 8.62 years, respectively. There was a significant difference for age between male and female (P-value= 0.02) and the median age for both gender was 42 years.

Most participants had bachelor degree (36%) also education level between male and female was different (P- value <0.001). The number of subjects with collegiate education was 1990 (72.5%) and subjects without collegiate was 754 (27.5%). The BMI for all participants was 27.03 ±4.33 Kg/m2 and there was not any difference for both gender (P- value= 0.63). The participants were evaluated for laboratory and anthropometric components and also were compared for both gender.

Details of comparison are presented in table 1. Most of parameters were the same for male and female and only were different in some items. In the male, cholesterol (182.28), LDL (102.45) and platelet (264.68) counts were higher than female (P-value <0.001). In the female, HCT (42.65) and RBC (5.07) were higher than male (P-value <0.001). In the case of anthropometric parameters, height of male (160.43) was higher than female (157.51) (P- value= 0.01). For male, HC (P- value= 0.01) and skinfold fat thickness was higher than female (P- value < 0.001) (Table 1).

Discussion
In this cohort-based study, we observed that all studied factors except VFL, BFM and PLR were correlated with SUA level, although NLR and total cholesterol were correlated only in female subjects and the highest correlation was related to BMR, skinfold fat thickness and HDL. Strengths of the current study was its focus on healthy population without any known medical diseases, that was based on the patient’s view of their condition, physical exams and laboratory tests.

Hyperuricemia is associated with cardiovascular disease, but it is commonly assumed a marker of organ dysfunction rather than a pathogenic role for progression and prognosis. SUA is produced by endogenous and exogenous metabolism of urine in human [1]. Although UA is a hydro-philic
antioxidant agent, the correlation between SUA levels and various diseases is considered [2-4]. Due to this controversy, UA was no longer regarded as a real cardiovascular risk factor [24]. In the recent years with improved knowledge about the function of UA in cardiorenal disease, discussion about the role of UA has increased.

Recent evidence has proposed that UA is not only an inflammatory marker but also is a direct cause of cardiovascular disease development through atherogenic effect. However, these results come from limited information and still there is the lack of sufficient knowledge of UA function. Several studies have been examined the association of SUA and the risk for hypertension, type 2 diabetes and metabolic syndrome in different population but studies in healthy adults are little [5,25,26]. In recent years, the prevalence of hyperuricemia has been increased in the world population, especially in the developing countries [27-29].

Obesity is known as strong risk factor and independent predictor for the development of hyperuricemia. [30, 31] Obesity is a well-established risk factor for cardiometabolic disorders, and its prevalence in adults appears to have increased in recent times [32,33]. Anthropometry is simple, inexpensive and effective method for screening of obesity and metabolic disorders [34]. The most useful anthropometric index of obesity that associates strongly with cardiometabolic disorders remains controversial. BMI is promulgated by the World Health Organization as the most useful epidemiological measure of overall obesity, but it is not suitable to account for body fat distribution [35]. In recent years, various new anthropometric indices have described to evaluate obesity and fat distribution in humans. BMI is a useful marker of body fat based on measurement of weight and height, while WC, WHtR and W/H reflect abdominal or central visceral fat [36-39].

In our study, based on BMI categories, 45.2% of the patients were overweight (BMI: 25-29.9) and 22.8% were obese (BMI≥30). (Table 1) WHtR ≥0.5 is a simple marker of central (visceral) obesity and in our study mean of WHtR was 0.56± 0.06.

Several studies have demonstrated association between SUA levels and body weight and positive correlation between SUA level and BMI in healthy subjects, which was also noted in this study. [40-42] Based on some studies, correlation between WC and SUA is higher than BMI. [43, 44] In our study, as
presented in table 2, in addition to BMI, WC and WHtR have positive correlation with SUA level, also, the higher BMI patients had higher mean SUA levels.

In previous studies, NC was found as an indicator of upper body obesity that can be used to determine overweight and obese individuals and is correlated positively with the markers of the metabolic syndrome, insulin resistance and high risk of type 2 diabetes. [45,46] Three cross-sectional studies evaluated the association between NC and SUA level [47-49]. Of note, these all studies were conducted among Chinese adults. Jiang et al reported that NC had positive association with UA level in hyperuricemia and non-hyperuricemia population. [47] The other two studies found NC was significantly associated with likelihood of having hyperuricemia when comparing the highest NC quartile group with the lowest NC quartile group. [48,49] In our study, NC had positive correlation with SUA level, that was compatible with previous studies results. These findings suggest that upper body fat, as estimated by NC, may have a unique association with SUA level even in normal BMI groups. 

Bosy-Westphal et al showed that the measurement of BFM as a direct measure of adiposity in comparison with BMI and WC had no further advantage to predict of obesity-related metabolic risk [50]. The VFL is a simple method that correlates with the fat mass. [51] In our study, BFM and VFL were not correlated with SUA level in both genders. These findings may suggest that central adiposity markers have closer associations with SUA level than markers of general adiposity.

Basal metabolic rate is dependent to physical exercise and appropriate dietary behavior [52]. The results of this study showed that BMR had a high negative correlation with SUA level in both genders. Similar to this one was found for skinfold fat thickness.

In present investigation, hyperuricemia was explained if participants have SUA levels more than 7.0 mg/dL in men or more than 6.0 mg/dL in women [53, 54]. Based on this definition, all men participants were nonhyperuricemic and the prevalence of hyperuricemia in women was 18.1% that was nearly matched with the universal prevalence rate reported to be between 2.6% and 36% [55].

We further divided subjects into male and female groups. The male group had a mean SUA level of 4.81 mg/dl (SD: 1.28 mg/dl), and the female group had a mean of 4.76 mg/dl (SD: 1.19 mg/dl). It is observed that serum UA levels were similar in males and females. Although this result was not
accordance with previous studies [56, 57]. it was anticipated that SUA level in man be more than females, because estrogen promotes excretion of UA [58,59].

Atherogenic dyslipidemia, include high LDL-C and TG levels with low HDL-C levels and anthropometric parameters, such as obesity are modifiable risks in both genders. The contribution of atherogenic dyslipidemia and disturbed anthropometric indices to cardiovascular risk are well established based on many epidemiological studies [17-19].

Hyperuricemia is suggested to play a role in adipose tissue as a mediator of proinflammatory endocrine unbalance which may have an important role in inflammatory process and dyslipidemia that leads to atherogenesis [60]. A few investigations described the association between SUA level and lipid profiles [61-63].

Based on our findings, SUA levels were positively correlated with serum TG and LDL-C level, also there was an inverse correlation between SUA and HDL-C level in both genders. Correlation between total cholesterol and SUA level was found only in female. These findings are in line with previous studies [61,63, 64]. Lin et al reported among components of the metabolic syndrome, abnormal TG had the maximum effect on SUA level [65]. However, the mechanism of the association of TG and SUA level has not been clear whether genetic factors or other factors such as the patient lifestyle are influencing. [66-68] In our study, based on linear regression model, the severity of correlation of TG and LDL was similar.

Moreover, uric acid is known as an inflammatory marker [15]. Whole blood count is an usual clinical test that components of it including neutrophils and lymphocytes are immune system elements which have important roles in inflammatory process. Ratios between these immune system factors (NLR and PLR) have been recorded as inflammatory markers that have been used more often in recent years [69,70].

Many studies have reported the correlation between NLR and disease activity or the relationship with other inflammatory markers in patients with primary Sjögren’s syndrome, rheumatoid arthritis, ankylosing spondylitis, and familial Mediterranean fever [71-74]. These studies have been performed mainly in inflammatory diseases, but, the association between SUA level and these novel
inflammatory markers is not well investigated in healthy population. The present paper is one of the first studies that performed to evaluate NLR and PLR values in healthy population, and investigating the correlation between those values and UA level. In this study, NLR and PLR values were similar between men and women. NLR value was correlated with SUA level (P <0.001) but no correlation was found between PLR and UA level. So, in healthy population, high UA level may act as an inflammatory marker.

Regarding prevention of disease is prior to treatment and prevention is impossible without a knowledge of the relative causes of disease, this study focused on young and middle-aged healthy people without evidence of any disease, also, in addition to lipid and traditional anthropometric indices, additional novel anthropometric measures and new inflammatory markers were considered. This study had some limitations. First, the participants were drawn from a particular employee group. This may not be a true representation of general population, therefore, there are limits for application of our results to other populations. Further studies are required to replicate our results in other groups. Second, the cross-sectional study design cannot demonstrate cause-effect relationships between studied factors and SUA.

Conclusion
Our findings showed SUA level to be correlated with anthropometric indices, lipid parameters and NLR in healthy population. whereas, the data of the role of SUA level and its use in healthy subjects is limited, the results of this study suggest that SUA can be used as a marker to trigger the screening for underlying factors of disease occurrence in healthy adults.

Declarations

**Ethical Approval and Consent to participate**
Data collection in this study was approved by the Ethics Commission of Shahid Beheshti University of Medical Science. The patients included in this study provided signed informed consent.

**Consent for publication**
No images, videos or other personal data of the participants are included in this manuscript.

**Availability of data and materials**
All available data are included in the manuscript. Raw data are not for public use because personal data are included.

**Competing interests**

The authors declare that they have no competing interests.

**Funding**

Not applicable.

**Author’s contributions**

TA: analyzed data, drafted, did background research, and revised the manuscript, HS: data collection and revised the manuscript, PR: data collection and drafted manuscript, HN: analyzed data and drafted manuscript.

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Tables

Table 1. Demographic, laboratory and anthropometric parameters of participants by sex

| Variable             | Total        | Male         | Female       |
|----------------------|--------------|--------------|--------------|
|                      | Mean (S.D)   | Mean (SD)    | Mean (SD)    |
| Demographic parameters |              |              |              |
| Age (years)          | 41.82 (8.80) | 41.49 (8.62) | 42.36 (9.07) |
| Education level, N (%)|              |              |              |
| Diploma and lower       | 754 (27.5) | 476 (45.4) | 278 (16.4) |
|------------------------|------------|------------|------------|
| Associate              | 299 (10.9) | 121 (11.5) | 178 (10.5) |
| Bachelor               | 995 (36.3) | 245 (23.4) | 750 (44.2) |
| Master and Doctorate   | 696 (25.4) | 207 (19.7) | 489 (28.8) |

**Laboratory parameters**

| Parameter                  | Normal range | Abnormal range | Extremely abnormal range |
|----------------------------|--------------|----------------|--------------------------|
| Uric acid (mg/dl)          | 4.80 (1.21)  | 4.81 (1.28)   | 4.76 (1.19)              |
| TG (mg/dl)                 | 128.07 (87.11) | 126.61 (83.54) | 132.56 (97.13)          |
| Total cholesterol (mg/dl)  | 180.55 (38.74) | 184.28 (38.45) | 169.27 (37.46)          |
| HDL-c (mg/dl)              | 47.31 (8.99)  | 47.19 (8.96)  | 47.69 (9.03)            |
| LDL-c (mg/dl)              | 100.43 (25.82) | 102.45 (27.11) | 94.23 (20.22)           |
| Neutrophil, ×10⁹/L         | 57.10 (7.72)  | 57.03 (7.89)  | 57.22 (7.41)            |
| Lymphocyte, ×10⁹/L         | 35.30 (7.41)  | 35.46 (7.60)  | 35.04 (7.08)            |
| NLR                       | 1.74 (0.66)   | 0.68 (0.01)   | 0.63 (0.02)             |
| Platelet, ×10⁹/L           | 261.59 (61.62) | 264.68 (62.15) | 256.38 (60.40)          |
| PLR                       | 7.88 (2.69)   | 7.97 (2.72)   | 7.74 (2.64)             |
| HCT (%)                   | 42.12 (4.40)  | 41.81 (4.31)  | 42.65 (4.51)            |
| RBC (10⁶/mm³)             | 5.03 (0.56)   | 5.00 (0.55)   | 5.07 (0.56)             |
| Cr (mg/dl)                | 0.98 (0.18)   | 0.98 (0.17)   | 0.96 (0.17)             |
| Anthropometric parameters |
|---------------------------|
| **Weight (kg)**           | 71.29 (18.69) | 71.47 (17.30) | 71.01 (20.74) |
| **Height (cm)**           | 159.33 (28.85) | 160.43 (25.36) | 157.51 (33.70) |
| **BMI (kg/m2)**           | 27.03 (4.33) | 27.06 (4.38) | 26.97 (4.21) |

| BMI categorize, N (%)     |
|---------------------------|
| <18.5                     | 19 (0.70) | 14 (0.80) | 5 (0.50) |
| 18.5-24.9                 | 881 (31.4) | 565 (32.0) | 320 (30.3) |
| 25-29.9                   | 1269 (45.2) | 790 (45.1) | 479 (45.3) |
| >=30                      | 640 (22.8) | 387 (22.1) | 253 (23.9) |

| **WC (cm)**               | 89.85 (18.93) | 90.28 (17.16) | 89.15 (21.49) |
| **WHR**                   | 0.56 (0.06) | 0.06 (0.001) | 0.07 (0.002) |
| **HC (cm)**               | 99.64 (18.73) | 100.36 (16.63) | 98.48 (21.67) |
| **W/H**                   | 1.45 (0.09) | 0.92 (0.06) | 2.63 (1.71) |
| **NC (cm)**               | 35.03 (7.14) | 35.20 (6.41) | 34.75 (8.18) |
| **AC (cm)**               | 29.24 (6.06) | 29.42 (5.74) | 28.95 (6.91) |
| **Wrist circumference (cm)** | 37.70 (16.22) | 16.29 (2.79) | 16.10 (3.61) |
| **PBF**                   | 33.72 (7.86) | 33.83 (7.92) | 33.47 (7.74) |
| **BMR (kcal)**            | 1407.19 (227.109) | 1406.28 (220.20) | 1409.21 (241.921) |
| **skinfold fat thickness(mm)** | 29.17 (10.24) | 29.77 (9.99) | 28.18 (10.57) |
The mean of uric acid for all subjects was $4.80 \pm 1.21$ and this mean between under and upper median age groups, male and female groups and also among education categories did not show any significant differences, while the mean among BMI levels showed some differences. Consequently, uric acid was increased by BMI increasing (P-value < 0.001).

In table 2, the relation between uric acid level and laboratory as well as anthropometric parameters, were evaluated. Most of parameters showed significant relation (all P-value < 0.05) except PLR, Cr, VFA and BFM.

| Variable            | Correlation | P-value |
|---------------------|-------------|---------|
| Laboratory parameters |             |         |
| TG (mg/dl)          | 0.43        | <0.001  |
| TC (mg/dl)          | 0.16        | <0.001  |
| HDL-C (mg/dl)       | -0.26       | <0.001  |
| LDL-C (mg/dl)       | 0.14        | <0.001  |
| Neutrophil, $\times 10^9$/L | -0.14   | <0.001  |
| Lymphocyte, $\times 10^9$/L | 0.12   | <0.001  |
| Parameter                      | Beta  | p value  |
|-------------------------------|-------|----------|
| NLR                           | -0.13 | <0.001   |
| Platelet, $\times 10^9$/L     | 0.06  | 0.03     |
| PLR                           | -0.04 | 0.14     |
| HCT                           | 0.45  | <0.001   |
| RBC ($10^6$/mm$^3$)           | -0.10 | <0.001   |
| Cr (mg/dl)                    | -0.005| 0.82     |
| Anthropometric parameters     |       |          |
| Weight (kg)                   | 0.37  | <0.001   |
| Height (cm)                   | 0.34  | <0.001   |
| BMI (kg/m$^2$)                | 0.24  | <0.001   |
| WC (cm)                       | 0.39  | <0.001   |
| WHtR                          | 0.18  | <0.001   |
| HC (cm)                       | 0.22  | <0.001   |
| W/H                           | 0.27  | <0.001   |
| NC (cm)                       | 0.45  | <0.001   |
| AC (cm)                       | 0.36  | <0.001   |
| Wrist Circumference (cm)      | 0.38  | <0.001   |
| PBF                           | -0.18 | <0.001   |
| BMR (kcal)                    | 0.43  | <0.001   |
Continuously, relation between uric acid and laboratory and anthropometric parameters according to sublevels of demographic and BMI variables are shown in details in table 3 and 4.

Table 3. Relation between uric acid and laboratory parameters by sex, age, education and BMI levels
| Laboratory parameters | Sex | Age | Education | BMI |
|-----------------------|-----|-----|-----------|-----|
| Triglyceride (mg/dl)  | Female | Male | Under median | Upper median | Non-collegiate | Collegiate | <18.5 |
| Cholesterol (mg/dl)   | 0.43* | 0.42* | 0.41* | 0.45* | 0.42* | 0.43* | 0.34** |
| HDL (mg/dl)           | 0.16* | 0.13** | 0.17* | 0.15* | 0.11*** | 0.17* | 0.51** |
| LDL (mg/dl)           | -0.27* | -0.24* | -0.24* | -0.29* | -0.29* | -0.25* | -0.59* |
| Neutrophil×10^9/L     | -0.17* | -0.06*** | -0.11* | -0.17* | -0.08*** | -0.13* | 0.12** |
| Lymphocyte×10^9/L     | 0.15* | 0.04*** | 0.09** | 0.15* | 0.09*** | 0.10* | -0.25* |
| NLR                   | -0.16* | -0.04*** | -0.10** | -0.16* | -0.08*** | -0.11* | 0.19** |
| Platelet×10^9/L       | 0.06** | 0.034*** | 0.01*** | 0.09** | 0.04*** | -0.02*** | -0.19** |
| PLR                   | -0.05*** | -0.004*** | -0.03*** | -0.06*** | -0.01*** | -0.05*** | 0.08** |
| HCT                   | 0.45* | 0.43* | 0.45* | 0.44* | 0.43* | 0.45* | 0.39* |
| RBC (10^6/mm3)        | -0.11* | -0.08*** | -0.12* | -0.08** | -0.13** | -0.08** | 0.22** |
| Cr (mg/dl)            | -0.01*** | 0.03*** | -0.008*** | 0.00*** | 0.05*** | -0.02*** | - |

Table 4. Relation between uric acid and anthropometric parameters by sex, age, education and BMI levels
| Variable                      | Weight(kg) | Height(cm) | BMI(kg/m2) | WC(cm) | WHtR | HC(cm) | W/H | NC(cm) | AC(cm) | Wrist Circumference(cm) | Protein | PBF | BMR(kcal) | Skinfold fat thickness (mm) | VFL | BFM |
|------------------------------|------------|------------|------------|--------|------|--------|------|--------|--------|--------------------------|---------|-----|----------|--------------------------------|-----|-----|
|                              | 0.35*      | 0.31*      | 0.24*      | 0.29*  | 0.16*| 0.19*  | 0.25*| 0.41*  | 0.33*  | 0.35*                     | 0.41*   | -0.16*| 0.41*    | 0.14*                                         | 0.03***| 0.04***|
|                              | 0.47*      | 0.37*      | 0.32*      | 0.41*  | 0.24*| 0.30*  | 0.34*| 0.57*  | 0.47*  | 0.48*                     | 0.52*   | -0.23*| 0.49*    | 0.18*                                         | -0.04***| -0.05**|
|                              | 0.37*      | 0.34*      | 0.23*      | 0.33*  | 0.19*| 0.22*  | 0.26*| 0.44*  | 0.36*  | 0.38*                     | 0.43*   | -0.17*| 0.42*    | 0.16*                                         | -0.02***| -0.02***|
|                              | 0.38*      | 0.33*      | 0.24*      | 0.30*  | 0.16*| 0.22*  | 0.28*| 0.45*  | 0.37*  | 0.38*                     | 0.45*   | -0.18*| 0.44*    | 0.14*                                         | -0.01***| -0.01***|
|                              | 0.39*      | 0.34*      | 0.24*      | 0.31*  | 0.16*| 0.22*  | 0.27*| 0.47*  | 0.38*  | 0.36*                     | 0.47*   | -0.19*| 0.45*    | 0.16*                                         | -0.04***| -0.03***|
|                              | 0.36*      | 0.31*      | 0.24*      | 0.30*  | 0.16*| 0.20*  | 0.47*| 0.43*  | 0.35*  | 0.36*                     | 0.59*   | -0.17*| 0.41*    | 0.17*                                         | -0.01***| -0.02|

*S P-value < 0.001

**Significant P-value: 0.001 - <0.05

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The factors related with uric acid using regression model, are shown as table 5. With the aid of backward method, through 9 steps, the best model with adjusted R square of 0.4, revealed that Triglyceride, LDL, HDL, Neutrophil, Lymphocyte, platelet, HCT, BMR and skinfold fat thickness, were factors related with uric acid (all p-values < 0.05).

Table 5. Results of linear regression model

| Variable               | Beta  | 95 % CI          | P- value |
|------------------------|-------|------------------|----------|
| TG (mg/dl)             | 0.004 | (0.003, 0.005)   | <0.001   |
| HDL-C (mg/dl)          | -0.011| (-0.018, -0.003 )| 0.008    |
| LDL-C (mg/dl)          | 0.004 | (0.001, 0.006)   | 0.006    |
| Neutrophil, ×10^9/L    | -0.028| (-0.53, -0.003)  | 0.030    |
| Lymphocyte, ×10^9/L    | 0.029 | (-0.055, -0.003) | 0.032    |
| Platelet, ×10^9/L      | 0.001 | (0.000, 0.002)   | 0.018    |
| HCT (%)                | 0.077 | (0.059, 0.095)   | <0.001   |
| BMR                    | -0.008| (-0.014, -0.001) | 0.017    |
| skinfold fat thickness (mm) | -0.008| (-0.016, -0.001) | 0.024    |