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Prevalence of Metabolic Syndrome in Iran: A systematic review and metaanalysis

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| ARTICLE INFO | ABSTRACT |
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| Article history: Received: 08 March 2017 Revised: 04 July 2017 Accepted: 22 July 2017 | Background: Metabolic syndrome is a set of metabolic disorders, including abdominal obesity, hypertension, increased fasting blood sugar, increased serum triglyceride level, and decreased high-density lipoprotein (HDL) cholesterol level. This condition elevates the risk of cardiovascular diseases and diabetes. Regarding this, the present study aimed to evaluate the prevalence of metabolic syndrome in Iran. Methods: This systematic review was conducted on the articles published within 2000- |
| Key words: Iran Meta-analysis Metabolic syndrome Prevalence Systematic review | 2015. The search was performed using the international databases, including Google Scholar, Science Direct, PubMed, and Scopus, and Persian databases of SID, IranMedex, and MagIran. The keywords employed during the searching process entailed: "Prevalence", "Frequency", and "Metabolic syndrome". The heterogeneity between the studies was assessed using I2 index. Data analysis was performed through meta-analysis technique (random-effects model) in Stata version 12. Results: In the 32 studies conducted in Iran with the sample size of 74,440 cases, the total prevalence of metabolic syndrome was reported to be 32% (95% CI: 28-35). Based on the National Cholesterol Education Program/Adult Treatment Panel III, International Diabetes Federation, and Joint Interim Statement criteria, the prevalence rates of metabolic syndrome were reported to be 30% (95% CI: 25-34), 34% (95% CI: 29-40), and 39% (95% CI: 33-45) in 23, 13, and 5 studies, respectively. Conclusion: As this study indicated, metabolic syndrome has a high prevalence in Iran. However, no significant changes have been observed in the prevalence of metabolic syndrome can be prevented by focusing on the reduction of risk factors for this disorder. |

1. Introduction

Metabolic syndrome was first used in 1920 to refer to three conditions of hypertension, high blood sugar, and gout. In 1988, Reaven introduced insulin resistance as the main feature of this disorder, calling it syndrome X.¹ After about one year, Kaplan added the most important component of this disorder to this list, i.e., abdominal obesity and abdominal subcutaneous adipose tissue. He recognized this series (i.e., hypertriglyceridemia, glucose intolerance, abdominal obesity, and hypertension) as the "Deadly Quartet".²

In 1998, the World Health Organization provided a definition for metabolic syndrome. Subsequently, the National Cholesterol Education Program/American Adult Therapy Panel Ш III) and (NCEP/ATP International Diabetes Federation (IDF) presented the diagnostic criteria of this disorder.³⁻⁶ In all of the mentioned definitions, the presence of three abnormal factors out of abdominal obesity, hypertension, high triglyceride, low high-density lipoprotein (HDL), and high blood sugar is indicative of metabolic syndrome.

Specific cut-points have been determined for all components of metabolic syndrome. However, the

waist circumference, which is used for the measurement of abdominal obesity, requires more evaluation and depends on the national and regional cutoffs and definitions.⁷ Several studies have indicated that metabolic syndrome is associated with diabetes and mortality due to cardiovascular diseases.⁸

The prevalence of coronary diseases and diabetes is respectively 2-3 and 3-5 times higher in the individuals with metabolic syndrome than in the people without such syndrome.^{9, 10} Several studies investigated the prevalence of metabolic syndrome have reported various results based on different diagnostic criteria. According to the NCEP criteria, the prevalence rates of metabolic syndrome are 32.1%, 18.3%, 14.9%, and 24.5% in Iran, India, Japan, and China, respectively.¹¹⁻¹⁴

In a systematic review and meta-analysis conducted by Maleki et al. in Iran, the prevalence rates of metabolic syndrome were reported to be 36% and 27% within 2003-2011 according to the IDF and ATPII criteria, respectively.¹⁵ In another meta-analysis, which reviewed the published studies on the prevalence of metabolic syndrome in Iran within 2000-2013, a high prevalence was reported.¹⁶ However, in the mentioned study, the prevalence of metabolic syndrome was not assessed based on the NCEP/ATPIII criteria.

The growing prevalence of hypertension, hyperlipidemia, obesity, and metabolic disorder in Iran¹⁷⁻¹⁹ over the past few years requires proper planning and policy-making for the intervention and modification of factors affecting metabolic syndrome in this country. Regarding this, it is essential to determine the prevalence of metabolic syndrome in Iran. With this background in mind, this study was conducted to determine the prevalence of metabolic syndrome in Iran based on the NCEP/ATPIII, IDF, and JIS criteria.

2. Methods

2.1. Design

This systematic review and meta-analysis was conducted based on the IDF, JIS, and NCEP/ATPII criteria disaggregated by gender in 2016 in Iran.

2.2. Data Sources

This systematic review was conducted on the articles published within 2000-2015. The search was performed using the international databases, including Google Scholar, Science Direct, PubMed, and Scopus, as well as Persian databases of SID, IranMedex, Medlib, and MagIran. The keywords employed during the searching process entailed: "Prevalence", "Frequency", "Metabolic syndrome",

and "Iran". The Persian translations of these keywords were applied along with their possible combinations for searching the Persian databases.

2.3. Study Selection

At first, a list was prepared by the researchers from the titles and abstracts of all the available articles in the mentioned databases. At this stage, all of the cross-sectional studies, which entailed the "Prevalence" keywords of and "Metabolic syndrome", were added to the primary list. After the elimination of the articles with repeated titles, the abstracts of the listed articles were evaluated to find the suitable papers. All of the cross-sectional studies, which investigated the prevalence of metabolic syndrome in Iran based on the NCEP/ATPII, IDF, and JIS criteria and published in Persian and English languages without a time limitation were evaluated.

The exclusion criteria included: 1) nonrepresentativeness of the study population, 2) use of non-random sampling technique, 3) investigation of the high-risk groups (e.g., pregnant women as well as patients with cirrhosis, hepatitis, and certain diseases), and 4) use of non-standard measurement tools.

2.4. Instrument

The quality of the related articles was determined using a checklist developed by Mousazadeh.²⁰ This checklist records such data as study objective, type of study, sample size, sampling method, data collection tool, evaluation of variables, study groups, and data analysis. Each article has a score range of 0-12 (once score is given to each part), and a score of ≥ 8 is indicative of acceptable methodological quality.

2.5. Data Extraction

In order to avoid bias, the process of searching for articles was performed by two separate researchers, and the research director's opinion was applied in case of disagreement on an article. Considering the inclusion and exclusion criteria, the intended data of the selected articles were recorded in the data collection form that was prepared in the EXCEL software. The recorded data included the first author, publication year, study setting, gender of the study population, sample size, diagnostic criteria, and number of patients with metabolic syndrome.

2.6. Data analysis

The reported prevalence of metabolic syndrome in each study was estimated using point estimation

with a confidence interval of 95% desegregated by gender and different provinces. The heterogeneity of the studies was evaluated through the Cochran's Q test and I² index. In this study, the I² of < 25%, 25-75%, and > 75% were considered as low, medium, and high heterogeneity, respectively.

Due to the observation of heterogeneity between the studies that was indicated by obtaining a statistically significant I^2 index, the DerSimonian and Laird's random effects model was used to combine studies and obtain the point estimate at the significance level of 0.1. In addition, the forest plot applied for the visual assessment of was heterogeneity between the selected studies. In addition, the Egger's test and funnel plot were employed to evaluate the publication bias and effect of studies with small sample size. Moreover, the sensitivity analysis was used to assess the role of each study in the final result.

The relationship of metabolic syndrome prevalence with publication year and sample size of the studies was evaluated using meta-regression analysis. On the other hand, the subgroup analysis was run to estimate the prevalence based on the diagnostic criteria of metabolic syndrome and gender differentiation. The prevalence rate and confidence interval based on diagnostic criteria were presented in cumulative flow diagrams. Data analysis was performed in the Stata software, version 12.

2.7. Ethical considerations

This article was derived from a research project approved by the Kurdistan University of Medical Sciences, Sanandaj, Iran.

3. Results

Out of the 335 retrieved articles, 130 studies were found relevant to the topic of interest by the title evaluation. In the next stage, the abstract of these 130 papers were evaluated. Finally, 32 articles were entered into the study, which were published within 2006-2015 (Diagram 1).

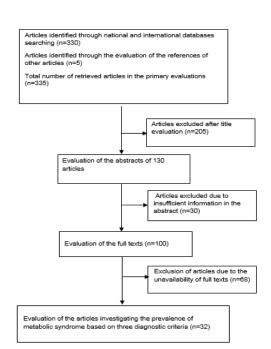


Diagram 1. Flow diagram of the stages of article inclusion to this systematic review

The selected articles were published within 2006-2015. The total sample size in the 32 evaluated articles was 74,440 individuals (mean population in each article: 2,327 cases). In all of the studies, the eligible individuals were selected through random sampling technique, and all of the selected studies were cross-sectional. The characteristics of the selected articles are presented in Table 1. Out of the 32 reviewed studies, the

prevalence rates of metabolic syndrome were estimated based on the IDF, JIS, and NCEP/ATPIII criteria in 13, 5, and 23 articles, respectively. Furthermore, the total prevalence of metabolic syndrome was estimated as 32% (95% CI: 28-35).

Since the heterogeneity index of the studies (I2=99.1%) was significant, the random effects model was applied. The forest plot of the present meta-analysis is presented in Figure 1. The highest

and lowest prevalence rates of metabolic syndrome based on the IDF criteria were 65% and 4% in Kerman (95% CI: 62-68) and Zanjan, Iran (95% CI: 2-6), respectively.

According to the JIS criteria, the highest (44%) and lowest (31%) prevalence rates of metabolic syndrome were related to Tehran. In addition, the highest and lowest prevalence rates of this disease according to the NCEP/ATPII criteria were reported to be 45% (CI 95%: 41-49) and 9% (CI 95%: 8-10) in Hamedan and Ahvaz, respectively (Figure 1).

The prevalence rates of metabolic syndrome were 36% and 27% in women and men, respectively, which was indicative of the higher prevalence of this disease in the females as stated in all definitions. The related forest plot based on gender differentiation according to diagnostic criteria is presented in figures 2 and 3. The results of the sensitivity analysis revealed that the one-by-one elimination of the selected articles from the analysis process based on the three given diagnostic criteria led to no significant changes in the shared estimation of the prevalence in none of the criteria.

The publication bias of the reviewed articles was assessed using the Egger's regression test, indicating that according to the IDF (P=0.496) and NCEP/ATPIII (P=0.496) criteria, the publication

bias was not statistically significant. However, based on the JIS criteria, a significant publication bias was obtained (P=0.030). The Egger diagram of publication bias related to the evaluated diagnostic criteria is presented in Figure 4.

With regard to the meta-regression results, no significant relationship was observed between the total prevalence of metabolic syndrome based on the IDF (P=0.260), JIS (P=0.932), and NCEP/ATPIII (P=0.540) criteria and the year of publication. The constant slope of meta-regression lines in all diagnostic criteria of Figure 5 is also indicative of this lack of association.

According to Figure 6, it can be inferred that with regard to the constant meta-regression slope, the prevalence of metabolic syndrome based on the IDF (P=0.504) and NCEP/ATPIII (P=0.617) criteria had no significant relationship with sample size. Nevertheless, there was a significant association between the prevalence of metabolic syndrome and increased sample size based on the JIS criteria. In other words, the prevalence of this syndrome decreased with increased sample size (P=0.012). In these diagrams, the circles signify the weight of the studies, i.e., the larger circles indicates bigger sample size.

| | | | | | | | analyeie | | | |
|--------|---------------|------|----------------------|--------------------|------------------------|---|-------------|------------|--------------|-----------------|
| _ | Ţ | | | Age c | Gender | Diagno metabo | Sa | Pr | | idence erval |
| Number | First author | Year | City | Age of participant | Gender of participants | Diagnostic criteria for metabolic syndrome | Sample size | Prevalence | Bottom limit | Upper limit |
| 1 | Jalali 21 | 2009 | Akbarabad (Kavar, | >19 | Both | IDF | 1402 | 31 | 28 | 33 |
| I | Jaiali 21 | 2009 | Fars) | 219 | Doun | NCEP | 1402 | 29 | 27 | 31 |
| 2 | Hadayegh 22 | 2009 | Tehran | >65 | Both | IDF | 720 | 42 | 38 | 46 |
| 2 | Kaukhaai 22 | 2012 | Zahadan | . 10 | Deth | IDF | 1900 | 25 | 23 | 27 |
| 3 | Keykhaei 23 | 2012 | Zahedan | >19 | Both NCEI | NCEP | 1802 | 21 | 19 | 23 |
| | | | | | 5.4 | IDF | (| 32 | 31 | 33 |
| 4 | Hadayegh 24 | 2007 | Tehran | >20 | Both | NCEP | 10368 | 33 | 32 | 34 |
| _ | | | | | | IDF | | 65 | 62 | 68 |
| 5 | Forouzanfar25 | 2015 | Kerman | Unknown | Both | NCEP | 950 | 73 | 71 | 76 |
| | | | _ | | | IDF | | 36 | 34 | 37 |
| 6 | Ghorbani 26 | 2012 | Semnan | 70-30 | Both | NCEP | 3799 | 28 | 27 | 30 |
| | | | | | | IDF | | 12 | 10 | 13 |
| | | | Zahedan | >16 | Both | NCEP | 2243 | 12 | 11 | 13 |
| 7 | Ostovaneh 27 | 2014 | | | | IDF | | 27 | 26 | 28 |
| | | | Amol | >16 | Both | NCEP | 5826 | 28 | 27 | 29 |

Table 1. Characteristics of articles entered into the meta-analysis

| | | | | Dalvand S | et al. | | | | | |
|----|-------------------------|-------|------------|-----------|---------|------|-----------|----------|----------|----------|
| 8 | Ebrahimi Mamghani 28 | 2011 | Tabriz | Unknown | Males | IDF | 76 | 57 | 45 | 68 |
| | | | | | | | 73 434 | 60 53 | 49 49 | 72 58 |
| 9 | Maharlouei 29 | 2013 | Shiraz | >40 | Females | IDF | 490 | 32 | 28 | 36 |
| 10 | Mohebi 30 | 2012 | Zanjan | 67-20 | Males | IDF | 12138 | 32 | 32 | 33 |
| 11 | Ebrahimi 31 | 2009 | Shahr Reza | 49-15 | Females | IDF | 1501 | 17 | 15 | 19 |
| 12 | Kazemi 32 | 2008 | Zanjan | 21-17 | Both | IDF | 507 | 4 | 2 | 6 |
| | Esmailzadeha | | | | | IDF | | 34 | 31 | 37 |
| 13 | 33 | 2013 | Qazvin | 78-20 | Both | JIS | 1107 | 39 | 36 | 42 |
| | Zeduech 04 | 004.0 | Tahara | 10 | Dath | NCEP | 005 | 31 | 28 | 33 |
| 14 | Zarkesh 34 | 2012 | Tehran | >19 | Both | JIS | 365 | 44 | 39 | 49 |
| 15 | Hosseinpanah 35 | 2012 | Tehran | 84-21 | Both | JIS | 347 | 38 | 33 | 43 |
| 16 | Faam 36 | 2013 | Tehran | 70-20 | Both | JIS | 4665 | 31 | 30 | 33 |
| 17 | Amiri 37 | 2014 | Tehran | >20 | Females | JIS | 603 | 44 | 40 | 48 |
| 18 | Keikhah 38 | 2013 | Isfahan | 60-30 | Both | NCEP | 3228 | 36 | 34 | 37 |
| 19 | Rashidi 39 | 2014 | Ahvaz | 19-10 | Both | NCEP | 2246 | 9 | 8 | 10 |
| 20 | Marjani 40 | 2012 | Gorgan | >20 | Female | NCEP | 160 | 21 | 14 | 27 |
| 21 | Esmail Nasab 41 | 2012 | Kurdistan | 64-25 | Both | NCEP | 1194 | 29 | 27 | 32 |
| 22 | Mehrabian 42 | 2011 | Isfahan | 90-53 | Females | NCEP | 539 | 25 | 21 | 29 |
| 23 | Fakhrzadeh 11 | 2006 | Tehran | 64-25 | Both | NCEP | 1480 | 28 | 25 | 30 |
| 24 | Marjani 43 | 2012 | Gorgan | >45 | Females | NCEP | 100 | 31 | 22 | 40 |
| 25 | Gharipour 44 | 2011 | Isfahan | >19 | Both | NCEP | 12514 | 23 | 22 | 24 |
| 26 | Jamshidi 45 | 2014 | Hamedan | 83-40 | Both | NCEP | 550 | 45 | 41 | 49 |
| 27 | Hajian-Tilki 46 | 2014 | Babol | 70-20 | Both | NCEP | 1000 | 42 | 39 | 45 |
| 28 | Tabatabaei 47 | 2015 | Shiraz | >20 | Both | NCEP | 377 | 27 | 22 | 31 |
| 29 | Saberi 48 | 2009 | Kashan | >30 | Males | NCEP | 429 | 36 | 31 | 40 |
| 30 | Delavar 49 | 2009 | Babol | 50-30 | Females | NCEP | 916 | 31 | 28 | 34 |
| 31 | Mahjoub 50 | 2012 | Babol | >20 | Both | NCEP | 933 | 24 | 21 | 26 |
| 32 | Moeini 51 | 2012 | Tehran | 40-15 | Both | NCEP | 282 | 23 | 18 | 28 |

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IDF: International Diabetes Federation, NCEP: National Cholesterol Education Program, JIS: Joint Interim Statement

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|---------|---|----|-----|--|
|---------|---|----|-----|--|

| Study D | ES (95% CI) | % Weight |
|--|--|-------------|
| DF | | |
| Hadaegh F,2007 (Tehran) | 0.32 (0.31, 0.33) | 2.35 |
| Hadaegh F,2009 (Tehran) | 0.42 (0.38, 0.46) | 2.28 |
| Kaykhaei MA,2012 (Zahedan) | 0.25 (0.23, 0.27) | 2.33 |
| arahimi H,2009 (Shahreza) | | 2.33 |
| | • 0.17 (0.15, 0.19) | |
| brahimi-Mamghani M,2011 (Tabriz) | 0.57 (0.45, 0.68) | 1.82 |
| brahimi-Mamghani M,2011 (Tabriz) | 0.60 (0.49, 0.72) | 1.81 |
| oroozanfar Z,2015 (Kerman) | • 0.65 (0.62, 0.68) | 2.30 |
| horbani R,2012 (Semnan) | 0.36 (0.34, 0.37) | 2.34 |
| alali R,2009 (Akbar Abad) | 0.31 (0.28, 0.33) | 2.32 |
| azemi SA,2008 (Zanjan) | 0.04 (0.02, 0.06) | 2.33 |
| Nohebi I,2012 (Zanjan) | • 0.32 (0.32, 0.33) | 2.35 |
| Ostovaneh MR,2014 (Zahedan) | 0.12 (0.10, 0.13) | 2.34 |
| Ostovaneh MR.2014 (Amol) | 0.27 (0.26, 0.28) | 2.34 |
| smailzadehha N.2013 (Qazvin) | 0.34 (0.31, 0.37) | 2.31 |
| Aharlouei N, 2013 (Shiraz) | 0.53 (0.49, 0.58) | 2.24 |
| Aaharlouei N, 2013 (Shiraz) | 0.32 (0.28, 0.36) | 2.24 |
| Subtotal (I-squared = 99.4% , p = 0.000) | 0.32 (0.29, 0.30) | 36.05 |
| Jubiotal (roqualeu = 39.4%, p = 0.000) | 0.34 (0.29, 0.40) | 30.05 |
| IIS | | |
| Esmailzadehha N,2013 (Qazvin) | l 🛨 0.39 (0.36, 0.42) | 2.31 |
| Hosseinpanah F,2012 (Tehran) | 0.38 (0.33, 0.43) | 2.22 |
| Amiri P,2014 (Tehran) | 0.44 (0.40, 0.48) | 2.27 |
| Zarkesh M.2012 (Tehran) | 0.44 (0.39, 0.49) | 2.22 |
| Faam B,2013 (Tehran) | 0.31 (0.30, 0.33) | 2.34 |
| Subtotal (I-squared = 93.9% , p = 0.000) | 0.39 (0.33, 0.45) | 11.35 |
| | | |
| | | |
| Mahjoub S,2012 (Babol) | 0.24 (0.21, 0.26) | 2.31 |
| Fabatabaie AH,2015 (Shiraz) | 0.27 (0.22, 0.31) | 2.25 |
| Delavar MA,2009 (Babol) | 0.31 (0.28, 0.34) | 2.30 |
| Esmailnasab N,2012 (Kurdistan) | 0.29 (0.27, 0.32) | 2.31 |
| akhrzadeh H,2006 (Tehran) | • 0.28 (0.25, 0.30) | 2.32 |
| Gholipour M,2011 (Isfahan) | • 0.23 (0.22, 0.24) | 2.35 |
| lajian Tilaki K,2014 (Babol) | 0.42 (0.39, 0.45) | 2.30 |
| lamshidi L,2014 (Hamedan) | 0.45 (0.41, 0.49) | 2.26 |
| Keykhah M,2013 (Isfahan) | 0.36 (0.34, 0.37) | 2.34 |
| <i>A</i> arjani A,2012 (Gorgan) | 0.21 (0.14, 0.27) | 2.15 |
| Marjani A,2012 (Gorgan) | 0.31 (0.22, 0.40) | 1.97 |
| | | 2.34 |
| Rashidi H,2014 (Ahvaz) Saberi HR,2009 (Kashan) | | 2.34 |
| | 0.36 (0.31, 0.40) | |
| Moini A,2012 (Tehran) | • 0.23 (0.18, 0.28) | 2.23 |
| Mehrabian F,2011 (Isfahan) | •••••••••••••••••••••••••••••••••••••• | 2.28 |
| smailzadehha N,2013 (Qazvin) | 0.31 (0.28, 0.33) | 2.31 |
| Ostovaneh MR,2014 (Zahedan) | 0.12 (0.11, 0.13) | 2.34 |
| Ostovaneh MR,2014 (Amol) | 0.28 (0.27, 0.29) | 2.34 |
| ladaegh F,2007 (Tehran) | 0.33 (0.32, 0.34) | 2.35 |
| Ghorbani R,2012 (Semnan) | 0.28 (0.27, 0.29) | 2.34 |
| lalali R,2009 (Akbar Abad) | 0.29 (0.27, 0.31) | 2.32 |
| Foroozanfar Z,2015 (Kerman) | • 0.73 (0.70, 0.76) | 2.31 |
| Kaikhaei MA,2012 (Zahedan) | • 0.21 (0.19, 0.23) | 2.33 |
| Subtotal (I-squared = 99.3% , p = 0.000) | 0.30 (0.25, 0.34) | 52.60 |
| Dverall (I-squared = 99.3%, p = 0.000) | 0.32 (0.29, 0.36) | 100.00 |
| NOTE: Weights are from random effects analysis | | |
| VOTE, weigns are non-random enects analysis | I | |
| 758 0 | .758 | |
| | | |

Figure 1. Forest plot of metabolic syndrome prevalence in Iran based on three diagnostic criteria

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| Study ID | | ES (95% CI) | % Weigh |
|--|-------------|---------------------|------------|
| | | ES (95% CI) | weigi |
| IDF | | | |
| Ebrahimi-Mameghani M,2011 (Tabriz) | | 0.60 (0.49, 0.72) | 3.42 |
| Mohebi I,2012 (Zanjan) | ۲ | 0.32 (0.32, 0.33) | 4.56 |
| Hadaegh F,2009 (Tehran) | | 0.26 (0.22, 0.30) | 4.36 |
| Ebrahimi-Mameghani M,2011 (Tabriz) | | 0.57 (0.45, 0.68) | 3.43 |
| Ostovaneh MR,2014 (Zahedan) | | 0.11 (0.09, 0.13) | 4.53 |
| Foroozanfar Z,2015 (Kerman) | | 0.47 (0.42, 0.52) | 4.29 |
| Hadaegh F,2007 (Tehran) | | 0.21 (0.20, 0.22) | 4.55 |
| Kaykhaei MA,2012 (Zahedan) | - | 0.20 (0.17, 0.23) | 4.47 |
| Ostovaneh MR,2014 (Amol) | | 0.26 (0.24, 0.27) | 4.54 |
| Subtotal (I-squared = 98.9%, p = 0.000) | | > 0.32 (0.25, 0.38) | 38.15 |
| | | | |
| | | | |
| Keykha M,2013 (Isfahan) | | 0.31 (0.28, 0.34) | |
| Shorbani R,2012 (Semnan) | | 0.17 (0.15, 0.19) | |
| Saberi HR,2009 (Kashan) | | - 0.36 (0.31, 0.40) | |
| Mahjoub S,2012 (Babol) | | 0.16 (0.11, 0.21) | |
| Ostovaneh MR,2014 (Zahedan) | | 0.07 (0.05, 0.08) | |
| Foroozanfar Z,2015 (Kerman) | | - 0.65 (0.60, 0.70) | |
| Jamshidi L,2014 (Hamedan) | | 0.26 (0.21, 0.31) | |
| Esmailnasab N,2012 (Kurdistan) | * | 0.17 (0.14, 0.20) | |
| Tabatabaie AH,2015 (Shiraz) | | 0.17 (0.11, 0.22) | |
| Kaykhaei MA,2012 (Zahedan) | * | 0.15 (0.13, 0.18) | |
| Rashidi H,2014 (Ahvaz) | <u>₩</u> | 0.11 (0.09, 0.13) | |
| Hajian Tilaki K,2014 (Babol) | 1 | 0.37 (0.32, 0.41) | 4.34 |
| Ostovaneh MR,2014 (Amol) | • | 0.21 (0.19, 0.22) | 4.54 |
| Hadaegh F,2007 (Tehran) | | 0.24 (0.23, 0.25) | 4.55 |
| Subtotal (I-squared = 98.7%, p = 0.000) | | 0.24 (0.19, 0.29) | 61.85 |
| Overall (I-squared = 98.9%, p = 0.000) | • | 0.27 (0.23, 0.31) | 100.0 |
| NOTE: Weights are from random effects analysis | | | |
| 715 | 0 | l .715 | |

Proportion (95% Confidence Interval)

Figure 2. Forest plot of metabolic syndrome prevalence in males based on three diagnostic criteria

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|---------|---|----|-----|--|
|---------|---|----|-----|--|

| Study | | % |
|--|---|--------|
| ID | ES (95% CI) | Weigh |
| IDF | | |
| Maharlouei N,2013 (Shiraz) | 0.32 (0.28, 0.36) | 3.56 |
| Ostovaneh MR,2014 (Amol) | 0.28 (0.26, 0.30) | 3.62 |
| Maharlouei N,2013 (Shiraz) | 0.53 (0.49, 0.58) | 3.55 |
| Hadaegh F,2009 (Tehran) | 0.63 (0.58, 0.69) | 3.52 |
| Ostovaneh MR,2014 (Zahedan) | 0.13 (0.11, 0.14) | 3.61 |
| Foroozanfar Z,2015 (Kerman) | 0.78 (0.74, 0.81) | 3.58 |
| Kaykhaei MA,2012 (Zahedan) | 0.28 (0.25, 0.31) | 3.60 |
| Hadaegh F,2007 (Tehran) | 0.41 (0.40, 0.42) | 3.62 |
| barahimi H,2009 (Shahreza) | 0.17 (0.15, 0.19) | 3.62 |
| Subtotal (I-squared = 99.5%, p = 0.000) | 0.39 (0.28, 0.50) | 32.29 |
| JIS | 1 | |
| Amiri P,2014 (Tehran) | 0.44 (0.40, 0.48) | 3.57 |
| Subtotal (I-squared = .%, p = .) | • 0.44 (0.40, 0.48) | 3.57 |
| NCEP/ATPIII | | |
| Mehrabian F, 2011 (Isfahan) | 0.25 (0.21, 0.29) | 3.58 |
| Rashidi H,2014 (Ahvaz) | 0.07 (0.06, 0.09) | 3.62 |
| Ghorbani R,2012 (Semnan) | 0.38 (0.36, 0.40) | 3.61 |
| Marjani A,2012 (Gorgan) | 0.31 (0.22, 0.40) | 3.34 |
| Moini A,2012 (Tehran) | 0.23 (0.18, 0.28) | 3.54 |
| Esmailnasab N,2012 (Kurdistan) | 0.41 (0.37, 0.45) | 3.57 |
| Ostovaneh MR.2014 (Amol) | • 0.35 (0.33, 0.37) | 3.62 |
| Keykha M,2013 (Isfahan) | ■ ■ | 3.61 |
| Hadaegh F,2007 (Tehran) | 0.41 (0.39, 0.42) | 3.62 |
| Marjani A,2012 (Gorgan) | 0.21 (0.14, 0.27) | 3.48 |
| Delavar MA,2009 (Babol) | €.21 (0.14, 0.27) €.1 (0.28, 0.34) | 3.60 |
| Hajian Tilaki K,2014 (Babol) | 0.31 (0.28, 0.34) | 3.56 |
| Mahjoub S,2012 (Babol) | 0.22 (0.19, 0.26) | 3.59 |
| Tabatabaie AH,2015 (Shiraz) | 0.37 (0.30, 0.44) | 3.46 |
| | | |
| Foroozanfar Z,2015 (Kerman) | - 0.79 (0.76, 0.83) | 3.59 |
| Jamshidi L,2014 (Hamedan) | 0.19 (0.14, 0.24) | 3.54 |
| Kaykhaei MA,2012 (Zahedan) | 0.25 (0.22, 0.27) | 3.60 |
| Ostovaneh MR,2014 (Zahedan) | 0.17 (0.15, 0.19) | 3.61 |
| Subtotal (I-squared = 99.3%, p = 0.000) | 0.32 (0.25, 0.39) | 64.14 |
| Overall (I-squared = 99.4%, p = 0.000) | 0.35 (0.29, 0.41) | 100.00 |
| NOTE: Weights are from random effects analysis | | |
| 828 0 | .828 | |
| Proportion (95% Confidence | | |

Figure 3. Forest plot of metabolic syndrome prevalence in females based on three diagnostic criteria

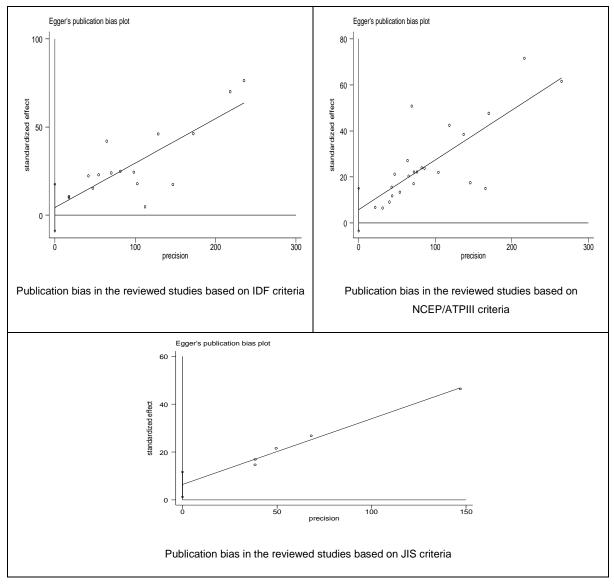


Figure 4. Funnel plot for the evaluation of publication bias in the reviewed studies based on three diagnostic criteria

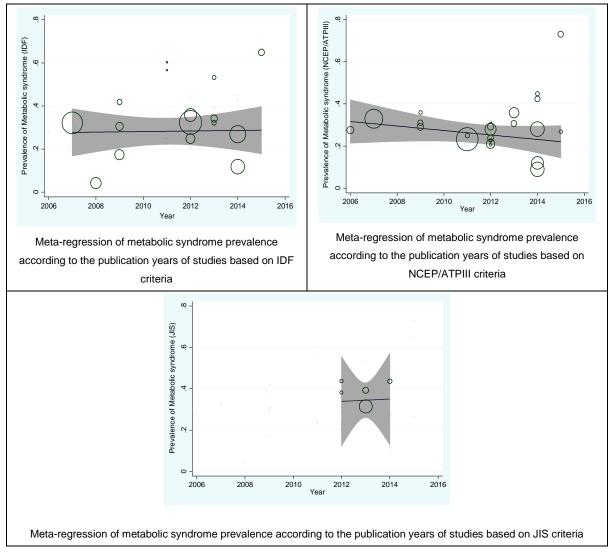


Figure 5. Meta-regression of metabolic syndrome prevalence according to the publication years of studies based on three diagnostic criteria

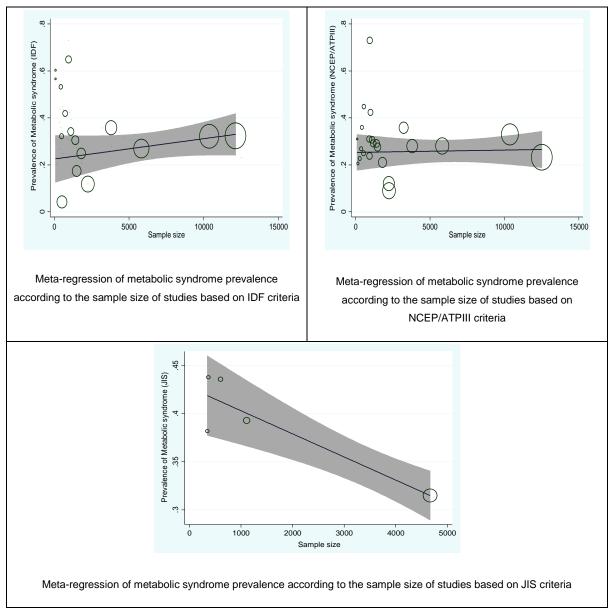


Figure 6. Meta-regression of metabolic syndrome prevalence according to the sample size of studies based on diagnostic criteria

4. Discussion

In the 32 evaluated studies with the total sample size of 74,440 cases, the total prevalence of metabolic syndrome was reported to be 32% in Iran, which was higher compared to those of other countries. In addition to such factors as inactivity, changes in individuals' lifestyle, and increased urbanization, other factors, such as larger waist size and lower HDL cholesterol were the cause of increased prevalence of metabolic syndrome in the Iranian population, compared to that in the Western countries.²¹

In a meta-analysis conducted by Maleki et al. (2014), the total prevalence of metabolic syndrome

was reported to be 36% based on the IDF criteria.¹⁵ In another systematic review and meta-analysis carried out by Amir Kalali et al. (2015) to determine the prevalence of metabolic syndrome based on the IDF criteria in Iran, this rate was reported as 34.6%.¹⁶ In the present study, the prevalence of this syndrome was reported to be 33% according to the IDF criteria. The majority of the published systematic reviews on the prevalence of metabolic syndrome (13 studies) have used the IDF criteria. With regard to the increased accumulation of sample size, our results could provide a more accurate estimation of the prevalence of this disorder, according to the diagnostic criteria in the country, compared to the studies conducted by Amir Kalali (seven studies) and Maleki (five studies). The total prevalence of metabolic syndrome according to the JIS criteria was lower in the present study, compared to the research carried out by Ami Kalali et al. (39% versus 41.5%, respectively). Similar to the mentioned study, the prevalence of metabolic syndrome according to the JIS in the present study was higher, compared to the other criteria.^{15, 16} However, it should be noted that limited numbers of studies have reported the prevalence of this syndrome according to the JIS criteria. In the present study, only five articles used the JIS criteria. Therefore, further evaluations are required for more accurate detection of metabolic syndrome prevalence based on this criterion. According to the results of the present study, the prevalence of metabolic syndrome based on the NCEP/ATPIII criteria was lower, compared to the other criteria (30%). In this regard, no meta-analysis has been performed to assess the prevalence of metabolic syndrome according to the NCEP/ATPIII criteria. Therefore, it is not possible to compare the shared estimate of metabolic syndrome prevalence according to this diagnostic criteria with other metaanalysis in the Iranian population. According to the results of a study conducted on 26,000 Hindi adults, the prevalence rates of metabolic syndrome were reported to be 25.8% and 18.3% according to the IDF and NCEP criteria, respectively.²¹ In another study carried out in China, the prevalence rate of this syndrome was 25.8% and 15.7% according to the definitions of IDF and NCEP, respectively.²² The higher prevalence rate of this syndrome based on the IDF criteria, compared to the NCEP criteria, might be due to the presence of lower cut-off point for the waist size in the definition of the IDF.

In line with other studies,²³⁻²⁸ the current review revealed that the prevalence rate of metabolic syndrome based on all diagnostic criteria was higher in females, compared to that in males. Similarly, in a meta-analysis performed by Maleki et al. and Amir Kalali et al, this syndrome was more prevalent in the females, compared to that in males based on all the three criteria.^{15, 16} However, in some countries, the metabolic syndrome was reported to have an equal prevalence rate in both genders.²⁹⁻³¹ Tabatabaei et al. conducted a study to determine the relationship between gender and prevalence of metabolic syndrome. They reported the prevalence of this disorder as 15.9% and 29.1% in male and female participants, respectively, which was indicative of a statistically significant difference in this regard.³² The higher prevalence of metabolic syndrome among the women is attributed to abdominal obesity, which is mainly due to low physical activity, higher birth rate, presence of estrogen receptors, and menopause.^{17, 33} For instance, abdominal obesity

was reported to be more prevalent in women, compared to that in men in the studies conducted in South Korea (27% versus 0.2%) and Turkey (54.8% versus 17.2%).^{28, 34}

In the studies conducted on the European societies, the mean prevalence of this syndrome was reported to be about 24% according to various diagnostic criteria, age groups, and geographical situation.³⁵ In the Latin American countries, about one-fourths of the people are diagnosed with this syndrome.^{36, 37} This difference in reports might be due to various etiologies, including insulin resistance, obesity (especially abdominal obesity), lipid disorders, glucose intolerance, hypertension, pre-inflammatory condition, genetics, intrauterine growth retardation, fast urbanization, nutrition transition, inactivity, social-mental stresses as well as economic, social, and cultural factors.^{17, 38} One of the major drawbacks of this study was the lack of access to full texts and insufficient information of some articles.

5. Conclusion

According to the results of the reviewed studies, metabolic syndrome has a high prevalence in Iran according to the IDF, JIS, and NCEP/ATPIII criteria. In addition, this prevalence is higher in the females based on the three given criteria, compared to that in the males. The high prevalence of metabolic syndrome in Iran can be ascribed to modern lifestyle, low level of activity, increased use of processed food with little nutritional value, increased urbanization, and more tendency toward western lifestyle.

On the other hand, a healthy life style can prevent the high prevalence of metabolic syndrome by focusing on reducing the risk factors for this disorder. Therefore, to obtain a global health level, we should perform more studies in this regard, optimally inform the policy-making organizations, and allocate more resources for the development of health in Iran.

Conflicts of interest

The authors declare no conflicts of interest.

Authors' contributions

Sahar Dalvand: Study concept and design and Statistical analysis and interpretation of data and Manuscript drafting and Technical and material support. Enayatollah Bakhshi: Study concept and design and Critical revision of the manuscript. Masoud Taheri-Asl: Data collection. Mozhdeh Zarei: Data collection and Manuscript drafting. Reza Ghanei Gheshlagh: Manuscript drafting and Technical and material support.

Acknowledgments

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