Review Article

Mercury Exposure and Risk of Type 2 Diabetes: A Systematic Review and Meta-Analysis

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Received 3 March 2022; Revised 2 August 2022; Accepted 5 August 2022; Published 2 September 2022

Academic Editor: Harry H. X. Wang

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Background and Aim. Previous studies have indicated that exposure to mercury (Hg) may be associated with odds of type 2 diabetes mellitus (T2DM). However, the available evidence is contradictory. This meta-analysis aimed to examine the relationship of Hg with the risk of T2DM. Methods. Scopus and PubMed databases were systematically searched from their inception to November 2021 to obtain pertinent studies. Standardized mean differences (SMDs) with corresponding 95% confidence intervals (CIs) were calculated to evaluate the difference in Hg levels between people with and without T2DM. The association of the Hg exposure with T2DM was assessed using a random-effects model by pooling the odds ratios (ORs) and 95% CIs. Results. A total of 17 studies, with 42,917 participants, aged ≥18 years, were analyzed. Overall, Hg levels were significantly higher in T2DM patients compared with non-T2DM controls (SMD = 1.07; 95%CI = 0.59 to 1.55, P ≤ 0.001), with significant heterogeneity across studies (I² = 96.1%; P = ≤ 0.001). No significant association was found between Hg exposure and risk of T2DM in the overall analysis and subgroup analysis based on the source of sample and study design. However, higher exposure to Hg was related to reduced risk of T2DM in men (OR = 0.71; 95%CI = 0.57 to 0.88), but not in women. No significant evidence for publication bias was detected. Conclusions. Although the Hg level in T2DM is significantly higher than that of nondiabetics, there was no association between Hg exposure and the overall risk of T2DM. Nevertheless, our study shows that higher exposure to Hg might reduce the risk of T2DM in men.

1. Introduction

Type 2 diabetes mellitus (T2DM) is a worldwide health concern [1]. This disease has put considerable economic burden on health systems globally that is forecasted to be increased even more in the future [2]. It is a multifactorial disease with different etiologies ranging from genetics to lifestyle [3] and is linked to further complications such as cardiovascular and renal diseases, as well as mortality [4]. The effect that exposure to heavy metals can have on T2DM...
[5, 6], obesity [7, 8], and metabolic syndrome [9] has been assessed previously. Specifically, some metals such as cadmium (Cd), Hg, and metalloid arsenic (As) are hypothesized to be related to the incidence of T2DM [5]. However, the available evidence is contradictory. The sources of these toxic metals are mainly contaminated water, polluted air, crops harvested in contaminated soil, dental care, fish consumption, and some industrial processes [10].

Hg is a heavy metal known for toxicity that exists in several forms. Inorganic Hg includes elemental or metallic mercury (Hg⁰) and mercurous (Hg²⁺) or mercuric (Hg³⁺) salts, while organic Hg includes compounds in which Hg is bonded to a structure containing carbon atoms (ethyl, methyl, phenyl, etc.) [11]. The biological behavior and toxicity of these forms vary considerably [11]. In general, Hg exposure has a broad range of toxic effects on cardiovascular, pulmonary, hematological, digestive, renal, immune, nervous, endocrine, and reproductive systems [12]. In relation to diabetes, this toxic agent can target β-cells in the pancreas and induces dysfunction and apoptosis [13]. Several mechanisms are introduced such as altering Ca²⁺ homeostasis, activation of phosphatidylinositol 3-kinase (PI3K) Akt signaling pathway, and reactive oxygen species (ROS) production [5]. Some studies have assessed the presence of this metal in diabetic patient’s scalp hair [14], urine [15], and blood [16–19] to examine any relationships between its levels with T2DM markers, but the results were heterogeneous. While some studies demonstrated positive relationships between T2DM and Hg levels in blood [20–25], urine [26], hair [27], and toenail [28], some other studies did not observe any relationship [29–36]. These discrepancies might be due to the differences in the population characteristics and sources of exposure (urine, blood, and nail). We, therefore, aimed to summarize the relationship between Hg levels in different body samples with the risk of T2DM in a comprehensive systematic review and meta-analysis.

2. Methods

In order to design and implement the present study, the guidelines of the Statement of Systematic Reviews and Preferred Reporting Meta-Analysis (PRISMA) have been considered and followed [37].

2.1. Search Strategy. The online databases PubMed and Scopus were searched extensively for related articles published before November 2021. A different combination of keywords was used in the search, which is listed as follows: (((((“Mercury”[Mesh]) OR (Mercury[Title/Abstract])) OR (methyl Mercury[Title/Abstract])) OR (Quicksilver[Title/Abstract])) OR (dimethylmercury[Title/Abstract])) OR (“colloidal Mercury”[Title/Abstract])) AND (((“DiabetesMellitus, Type 2”[Mesh]) OR (diabetes [Title/Abstract])) OR (type 2 diabetes mellitus[Title/Abstract])) OR (T2DM[Title/Abstract])) OR (“noninsulin-dependent diabetes mellitus”[Title/Abstract])). Only English studies were reviewed. All review articles and study references were checked to minimize the possibility of losing studies. To speed up the screening process, all identified studies were imported into an EndNote library, and duplicates were removed. The selection of eligible studies was independently reviewed by two researchers. First, the titles and abstracts of the studies were evaluated, and then the full text of the remaining publications was independently reviewed.

Figure 1: Flowchart of the study.
| Study                  | Country       | Year | Study design  | Sex | Total sample size | N cases with T2DM (age) | Sample source | Mean ± SD Hg in controls | Mean ± SD Hg in cases (T2DM) | Hg assessment | Type of effect size                                                                 |
|------------------------|---------------|------|---------------|-----|------------------|-------------------------|---------------|--------------------------|-----------------------------|---------------|----------------------------------------------------------------------------------|
| Anica Simić            | Norway        | 2017 | Case-control  | Both| 874              | 267 (65.4 ± 10.6)       | Blood         | 3.19 ± 1.35              | 3.6 ± 4.8                    | ICP-MS        | Mean of Hg in T2DM vs. controls OR for T2DM Adjusted for BMI, waist-to-hip ratio, family history of diabetes, smoking habits, area, education and economic status, fat fish intake |
| Yohei Hotta            | Japan         | 2019 | Case-control  | Male| 50               | 27 (66.1 ± 13.2)        | Hair          | 2.5 ± 2.94               | 2.12 ± 1.49                  | ICP-MS        | Mean of Hg in T2DM vs. controls OR for T2DM                                     |
|                        |               |      |               | Female| 46               | 15 (68.0 ± 8.5)         | Hair          | 1.37 ± 1.03              | 2.19 ± 1.83                  | —                          |                                                                  |
| Hassan Imran Afridi    | Pakistan      | 2015 | Case-control  | Male| 43               | 25 (ranged 30–50 years)| Blood         | 1.02 ± 0.07              | 1.69 ± 0.08                  | ICP-atomic emission spectrophotometer | Mean of Hg in T2DM vs. controls |
|                        |               |      |               | Female| 47               | 23 (ranged 30–50 years)| Hair          | 0.98 ± 0.03              | 1.78 ± 0.07                  |                            | —                          |
| Ailin Falkmo Hansen    | Norway        | 2016 | Case-control  | Both| 883              | 755 (65.2 ± 10.3)       | Blood         | 3.18 ± 5.26              | 3.47 ± 4                    | ICP-MS        | Mean of Hg in T2DM vs. controls OR for T2DM Age, sex, body mass index, waist-to-hip ratio, education, income, smoking, and family history of diabetes |
| Yohei Hotta            | Japan         | 2018 | Case-control  | Group 1:71 | Group 1:12 (ranged 36–86 years) | 1.94 ± 2.11 | Hair | 2.88 ± 3.52 |
|                        |               |      |               | Group 2:92 | Group 2:33 (ranged 36–86 years) | 1.94 ± 2.11 | Hair | 1.98 ± 1.55 |
| Bert B. Little         | USA           | 2020 | Case-control  | Both| 875              | 109 (55.0 ± 11.4)       | Blood         | 0.05 ± 0.27              | 0.12 ± 0.41                  | NR            | Mean of Hg in T2DM vs. controls OR for T2DM Age, gender, smoking tobacco, duration of residence, smelter worker, blood lead level, blood arsenic, cadmium level, gamma-glutamyl transpeptidase, hypertension |

Table 1: The characteristics of the included studies in meta-analysis.
| Study Name               | Country       | Year | Study design          | Sex     | Total sample size | N cases with T2DM (age) | Sample source | Mean ± SD Hg in controls | Mean ± SD Hg in cases (T2DM) | Hg assessment                         | Type of effect size |
|-------------------------|---------------|------|-----------------------|---------|-------------------|-------------------------|---------------|--------------------------|------------------------------------|----------------------|-------------------------------|
| Muhittin A. Serdar     | Turkey        | 2009 | Case-control          | Both    | 53                | 31 (59 ± 9)             | Blood         | 1.53 ± 0.69               | 1.15 ± 0.54                        | ICP-MS                            | Mean of Hg in T2DM vs. controls |
| Xin Wang               | USA           | 2020 | Prospective cohort    | Female  | 1237              | 102 (50.0 ± 3.1)        | Urine         | 1.23 ± 1.31               | 1.08 ± 1.05                        | ICP-MS                            | Mean of Hg in T2DM vs. controls |
| NEIL Io WARD           | England       | 1983 | Case-control          | Both    | 85                | 55 (59.7 ± 10.0)        | Blood         | 15 ± 5                   | 12 ± 3                             | Neutron-activation analysis (NAA) and electrothermal atomic absorption spectroscopy (EAAS) methods | Mean of Hg in T2DM vs. controls |
| Iwona Rotter           | Poland        | 2015 | Case-control          | Male    | 313               | 55 (61.3 ± 6.3)         | Blood         | 4.53 ± 2.23               | 4.45 ± 1.58                        | Inductively coupled argon plasma optical emission spectrometry (ICP-OES) | Mean of Hg in T2DM vs. controls |
| Junguo Zhang           | China         | 2021 | Case-control          | Both    | 15327             | 2132 (49.75 ± 17.88)    | Blood         | —                        | —                                  | Inductively coupled plasma dynamic reaction cell mass spectrometry | OR for T2DM                                           |
| Study                        | Country               | Year | Sex               | Total sample size | N cases with T2DM (age) | Sample source | Mean ± SD Hg in controls | Mean ± SD Hg in cases (T2DM) | Hg assessment                      | Type of effect size |
|-----------------------------|-----------------------|------|-------------------|-------------------|-------------------------|---------------|-------------------------|-------------------------------|-------------------------------|-------------------|
| KA HE                       | USA                   | 2012 | Both              | 4163              | 288 (aged 20–32 years) | Toenail        | —                       | —                            | Instrumental neutron-activation analysis | HR for T2DM |
| Tsung-Lin Tsai             | Taiwan                | 2019 | Both              | 646               | 56 (55.37 ± 12.87)     | Blood         | —                       | —                            | Cold vapor atomic absorption spectrometry | OR for T2DM | Age, sex, BMI, education, hypertension, total cholesterol, fasting glucose, cigarette smoking, alcohol consumption, saltwater fish consumption, total calorie intake, protein and fat intake, geographical strata, seasonality, C-reactive protein, and hemoglobin |
| S.-S. Moon                 | South Korea           | 2012 | Both              | 3184              | 333 (58.8 ± 10.9)      | Blood         | —                       | —                            | Gold-amalgam collection method with DMA-80 | OR for T2DM | Adjusted for age, sex, region, smoking, alcohol consumption, and regular exercise |
| DARIUSH MOZAFFARIAN        | USA                   | 2013 | Male              | 9267              | 1010 (61.2 ± 8.9)      | Toenail        | —                       | —                            | Neutron-activation analysis               | HR for T2DM | Adjusted for age, sex, race, region, month of toenail return, family history of diabetes, smoking status, BMI, hypertension, hypercholesterolemia, future cardiovascular disease case-control status (case or control), physical activity, alcohol use, and fish consumption |
| S. Cordier                 | Canada                | 2020 | Both              | 1874              | 217 (32.9 ± 4.8)       | Blood         | —                       | —                            | ICP-MS                         | OR for T2DM | Age, sex, waist circumference, smoking, omega-3 PUFAs |
| Min Kyong Moon             | South Korea           | 2021 | Both              | 3787              | NR (aged ≥19 years)    | Urine and blood| —                       | —                            | Amalgamation direct Hg analyzer     | OR for T2DM | Age, sex, cigarette smoking, alcohol drinking, exercise, and education levels were included as covariates |

NR: not reported; T2DM: type 2 diabetes; OR = odds ratio; BMI = body mass index; ICP-MS: inductively coupled plasma mass spectrometry; PUFAs: Polyunsaturated fatty acids.
2.2. Eligibility Criteria. The criteria for including eligible articles in this systematic review and meta-analysis were defined as follows: (A) cohort studies, cross-sectional, or case-control studies, (B) having a nondiabetic control group, (C) studies that reported odds ratios (ORs), hazard ratios (HRs), or relative risks (RRs) and the corresponding 95% confidence interval (CI) for the association of Hg exposure in the blood, urine, hair, and nails with T2DM, (D) studies reported the mean and standard deviation (SD) of Hg in patients with T2DM and healthy controls, (E) the full text of the article was available in English, and (F) studies were conducted on adults (aged \( \geq 18 \) years). Case reports, book chapters, conference papers, letters, editorial papers, and animal and cell culture studies were excluded.

2.3. Data Extraction. Two researchers independently reviewed the full text of the studies and extracted the data and resolved the differences through discussion with the third independent researcher. The following information was extracted: (1) study characteristics (name of the first author, design of the study, year of publication, country/ geographical location, target population, number of participants, duration of study, gender), (2) mean and SD of Hg, and (3) relevant reported risk estimates (including ORs, RRs, HRs) and the corresponding 95% CIs.

2.4. Quality Assessment of Studies. In this meta-analysis, the Newcastle–Ottawa Scale (NOS) questionnaire was used to evaluate the quality of studies in three areas (study selection, study group comparison, and exposure assessment) [38]. Studies with scores of 4 or more were considered a medium to high-quality studies.

2.5. Statistical Analysis. In this meta-analysis, a random effects model was used to test the effect of interest. Mean differences in Hg levels between the patients with T2DM and healthy controls were reported as standardized mean difference (SMD) and 95% CI. Standardized mean differences (SMD) for each original study were derived using the method of Cohen’s d [39] as the difference between means divided by the pooled standard deviation. Risk estimates reported for the relationship of Hg exposure to T2DM were pooled to estimate the overall effect size (OR and 95% CI). Heterogeneity across studies was assessed with \( I^2 \) statistics. Heterogeneity was considered significant if \( I^2 > 50\% \) (\( P < 0.1 \)). Funnel diagrams and Egger’s test were used to assess publication bias. Subgroup analysis was performed according to the type of study (prospective cohort vs. case-control), gender (both males and females), and sample source (blood, urine, and nails). STATA (version 14.0; Stata Corporation, College Station, TX) was used to perform all

| Study ID | SMD (95% CI) | Weight |
|----------|--------------|--------|
| Blood    |              |        |
| Anica Simić (2017) | 0.14 (-0.00, 0.29) | 8.81   |
| Hassan Imran Afridi (2015) | 14.40 (11.22, 17.57) | 1.83   |
| Hassan Imran Afridi (2015) | 18.51 (14.65, 22.38) | 1.32   |
| Ailin Falkmo Hansen (2016) | 0.07 (-0.12, 0.26) | 8.76   |
| Bert B. Little (2020) | 0.24 (0.04, 0.44) | 8.75   |
| Muhittin A. Serdar (2009) | -0.63 (-1.19, -0.07) | 7.93   |
| NEI Io Ward (1983) | -0.79 (-1.25, -0.32) | 8.21   |
| Iwona Rotter (2015) | -0.04 (-0.33, 0.25) | 8.60   |
| Subtotal (\( I^2 \)-squared = 96.3%, \( p = 0.000 \)) | 0.64 (0.05, 1.22) | 54.22  |
| Hair     |              |        |
| Yohei Hotta (2019) | -0.17 (-0.72, 0.39) | 7.94   |
| Yohei Hotta (2019) | 0.61 (-0.02, 1.24) | 7.71   |
| Hassan Imran Afridi (2015) | 8.81 (6.81, 10.82) | 3.50   |
| Hassan Imran Afridi (2015) | 14.97 (11.83, 18.12) | 1.85   |
| Yohei Hotta (2018) | 0.39 (-0.23, 1.02) | 7.73   |
| Yohei Hotta (2018) | 0.02 (-0.41, 0.45) | 8.30   |
| Subtotal (\( I^2 \)-squared = 96.8%, \( p = 0.000 \)) | 3.15 (1.49, 4.82) | 37.04  |
| Urine    |              |        |
| Xin Wang (2020) | -0.12 (-0.32, 0.09) | 8.75   |
| Subtotal (\( I^2 \)-squared = 96.1%, \( p = 0.000 \)) | 1.07 (0.59, 1.55) | 100.00 |

Figure 2: Forest plot for the mean levels of Hg in patients with T2DM compared with healthy controls stratified by the type of sample.
statistical tests for the current meta-analysis. *P* value < 0.05 was considered statistically significant for all statistical analyzes.

**3. Results**

3.1. Study Characteristics. The systematic search of databases yielded a total of 778 studies. After excluding duplicate publications (*n* = 168) and unrelated studies by titles/abstracts (*n* = 559), 51 publications underwent full-text screening, of which 17 studies [14–16, 18, 19, 28, 29, 32–36, 40–44], with a total sample size of 42,917 participants (5,545 cases of T2DM), published between 1984 and 2021, were eligible to be included in the current meta-analysis according to the inclusion criteria. The flowchart reporting the process of screening is presented in Figure 1. Some included studies reported different effect sizes based on their studied subgroups; for these studies, we extracted and analyzed all suitable effect sizes. Of the included studies, data on Hg levels in T2DM patients vs. healthy controls were reported in 10 studies with 15 effect sizes [14–16, 19, 32, 33, 40–42, 44], and 11 studies (3 prospective cohorts and 8 case-control) with 17 effect sizes [15, 18, 28, 29, 32, 34–36, 42–44] reported risk estimates for the association between Hg exposure and the risk of T2DM. All studies reporting risk estimates for T2DM adjusted the analysis for the potential confounders. The sample size of the analyzed studies ranged between 53 and 15,327 subjects. Regarding the sex of participants, 5 studies reported data for subgroupsof males and females separately [14, 18, 35, 36, 41], 1 study just included males [19], 1 study just included females [15], and the remaining studies were performed on a combination of both genders. The quality of studies was medium to high, with scores ranging from 4 to 9. The characteristics of the analyzed publications are presented in Table 1.

3.2. Mercury Levels in T2DM. In the pooled analysis of all eligible studies, Hg levels were significantly higher in T2DM patients compared with non-T2DM controls (random effects, SMD = 1.07; 95% CI = 0.59 to 1.55, *P* ≤ 0.001) (Figure 2), with a significant heterogeneity across studies (*I*² = 96.1%; *P* ≤ 0.001). In the stratified analysis by type of sample, compared with healthy controls, Hg levels were also significantly higher in blood (7 studies [16, 19, 32, 40–42, 44], SMD = 0.64 μg/L; 95% CI = 0.05 to 1.22, *P* = 0.03) and hair (3 studies [14, 33, 41], SMD = 3.15 μg/L; 95% CI = 1.49 to 4.82, *P* ≤ 0.001) of patients with T2DM, but not in urine samples (based on 1 study [15]) (Figure 2).
3.3. Mercury and Risk of T2DM. In the overall analysis and subgroup analysis based on the source of the sample (blood, urine, and toenail) (Figure 3) and study design (case-control vs. prospective cohort) (Figure 4), no significant association was found between Hg exposure and risk of T2DM. The pooled effect size for case-control studies was (SMD = 1.04; 95% CI = 0.83 to 1.30), and for cohort studies, it was (SMD = 0.96; 95% CI = 0.75 to 1.22), indicating no association between Hg and T2DM in both case-control and cohort studies (Figure 4). However, in the stratified analysis by the sex of participants, it was observed that higher exposure to Hg might be related to a reduced risk of T2DM in males (3 studies [18, 35, 36], OR = 0.71; 95% CI = 0.57 to 0.88), but not in females (Figure 5). Moreover, in the stratified analysis by the method used for the measurement of Hg, overall Hg exposure was not associated with T2DM (Figure 6).

3.4. Publication Bias. Egger test and funnel plot detected no significant evidence for publication bias in studies investigating the relation of Hg exposure to the risk of T2DM (Figure 7).

4. Discussion

The current study is the first systematic review and meta-analysis which aimed to investigate the association of Hg levels in different body samples with the risk of T2DM. Overall, we found consistent epidemiological evidence that Hg levels in the blood and hair samples of diabetic patients were considerably higher than in the nondiabetic control group. Nevertheless, overall, the findings of this study revealed no significant association between Hg exposure and the risk of T2DM. However, Hg exposure in the male subgroup might reduce the risk of T2DM.

The findings of the current meta-analysis demonstrated no significant association between Hg exposure and the risk of T2DM. This lack of association was even observed in the subgroups by sample source (blood, urine, and toenail) and the type of study (case-control vs. prospective cohort). A cross-sectional study conducted on 1588 men and 1596 women with age ≥30 years in the general population in South Korea indicated that blood Pb, Hg, and Cd concentrations in diabetic patients were slightly higher than the nondiabetic individuals; this difference, nonetheless, was not significant [34]. Even after controlling for age, gender, location, smoking, alcohol use, and regular exercise, the prevalence of diabetes was not affected by the blood heavy metals concentrations [34]. However, in the study by Tsai et al., in 2019 on 646 Taiwanese adults, Hg levels in red blood cells (RBC-Hg) of T2DM patients were considerably higher than the nondiabetic subjects. After controlling for the potential confounders, a significant direct association was reported between the RBC-Hg and the prevalence of T2DM [43]. A cohort study on toenail samples of 9262 American
subjects showed that toenails’ Hg concentration did not lead to a higher incidence of diabetes in women and men [18]. This finding remained constant even after separate analyzes based on classifications of Hg with higher concentrations, fish or Omega3 consumption, BMI, and age. A review study, including 34 in vivo and in vitro studies, showed a probable association between the total Hg concentration and the risk of diabetes. However, sufficient evidence for a causal and consistent relationship did not exist [45]. A case-control study showed a considerable increase in fasting blood glucose levels among individuals with high blood Hg levels (>16 mg/L) [29]. Moreover, supporting our findings, considering Hill’s causal criteria, a systematic review of 29 publications did not show sufficient evidence of any associations between Hg levels and diabetes [46]. The discrepancy among studies may result from study design, source of Hg exposure, or population characteristics such as age and sex. Moreover, because blood levels of Hg differ among ethnic groups [34], the heterogeneity in the results might be due to differences in ethnicity. The levels of Hg exposure are also dependent on the geographic region [47], and a source of the inconsistency of the available evidence may be differences in the geographic region of various studies.

Although, in the overall analysis, the current study demonstrated no association between Hg exposure and the risk of diabetes, the results of the gender-based analysis revealed that more exposure to Hg among the subgroup of men might be associated with a lower risk of T2DM. This finding resulted from the pooled analysis of only three studies [18, 35, 36]; thus, this conclusion should be interpreted with caution; however, our findings were comparable to a large case-control study (15327 subjects) by Zhang et al. which revealed a reverse association between the total Hg concentration and blood methylmercury and diabetes among adults [18]. Also, in Health Professionals Follow-Up and Nurses’ Health Study, toenails’ higher Hg levels were significantly associated with lower incidence of diabetes in both genders [35]. The mechanisms underlying the likely inverse relationship between Hg exposure and lower incidence of diabetes in men are still unknown. Previous studies reported the significance of oxidative stress in the pathogenesis of Hg toxicity [48]. As a compensatory mechanism against oxidative stress, Hg could increase the gene expression of proteins with antioxidant activity, including catalase, copper, zinc-superoxide dismutase, glutamate-cysteine ligase, thioredoxin reductase 1, manganese-superoxide dismutase, and can stimulate the antioxidant signaling pathway via direct interaction with the cysteine residues of the Keap1 and/or Akt/glycogen synthase kinase 3 beta/Fyn pathway [48], which the mentioned mechanism is protective.

![Figure 5: Forest plot for the association between Hg and risk of T2DM stratified by the sex of participants.](image-url)
against T2DM [49]. Also, some studies have found a reverse association between Hg exposure and cardiovascular disease [50–52] which might be related to the aforementioned cellular reparative function against Hg-induced oxidative stress. Although, this inverse association might be due to confoundings, such as relationships between higher Hg exposure and other factors that reduce the risk of T2DM. Nevertheless, the majority of the included studies were adjusted for fish consumption. Omega-3 fatty acids might attenuate Hg-induced adverse effects by improving the acute phase response and antioxidant status [53].

The current study is the first meta-analysis examining the association between blood, hair, toenail, and urine Hg levels and the risk of T2DM. As strength, no evidence for publication bias was identified. However, some limitations of this meta-analysis should be declared. First, significant heterogeneity was found across studies; we identified that

| Study ID | OR (95% CI) | Weight |
|----------|-------------|---------|
| Min Kyong Moon (2021) | 0.61 (0.34, 1.10) | 4.52 |
| Ailin Falkmo Hansen (2016) | 1.66 (0.79, 3.47) | 3.29 |
| Xin Wang (2020) | 0.92 (0.82, 1.03) | 11.61 |
| Junguo Zhang (2021) | 0.73 (0.55, 0.97) | 8.80 |
| Junguo Zhang (2021) | 0.82 (0.63, 1.07) | 9.14 |
| S. Cordier (2020) | 1.18 (0.42, 3.31) | 1.94 |
| S. Cordier (2020) | 1.09 (0.56, 2.15) | 3.77 |
| Subtotal (I-squared = 20.2%, p = 0.276) | 0.87 (0.76, 1.00) | 43.05 |
| NR | 1.14 (0.74, 1.74) | 6.43 |
| Subtotal (I-squared = 60.2%, p = 0.001) | 1.14 (0.74, 1.75) | 6.43 |
| NAA KA HE (2012) | 1.65 (1.07, 2.56) | 6.31 |
| DARIUSH MOZAFFARIAN (2013) | 0.86 (0.66, 1.11) | 9.22 |
| DARIUSH MOZAFFARIAN (2013) | 0.69 (0.42, 1.15) | 5.42 |
| Subtotal (I-squared = 75.8%, p = 0.016) | 0.99 (0.63, 1.58) | 20.96 |
| CVAAS Tsung-Lin Tsai (2019) | 3.34 (1.72, 6.48) | 3.84 |
| Subtotal (I-squared = 60.2%, p = 0.001) | 3.34 (1.72, 6.48) | 3.84 |
| Gold amalgamation direct Hg analyzer S.-S. Moon (2012) | 1.08 (0.76, 1.53) | 7.65 |
| Min Kyong Moon (2021) | 0.64 (0.37, 1.13) | 4.81 |
| Min Kyong Moon (2021) | 1.91 (1.02, 3.55) | 4.18 |
| Min Kyong Moon (2021) | 0.74 (0.40, 1.39) | 4.18 |
| Min Kyong Moon (2021) | 1.07 (0.62, 1.86) | 4.90 |
| Subtotal (I-squared = 48.1%, p = 0.103) | 1.01 (0.73, 1.40) | 25.71 |
| Overall (I-squared = 60.2%, p = 0.001) | 1.00 (0.85, 1.17) | 100.00 |

NOTE: Weights are from random effects analysis.
this heterogeneity did not originate from the sample type or study design, but the gender of participants. Second, the majority of the included studies were case-control in design, which might be affected by the unexamined confounding factors and suffer from a higher probability of bias than cohort studies. Other weaknesses of the present study include the difference in Hg concentration of various body samples, the difference in the methods of measuring Hg levels, and the biomarkers’ measurement precision. Hg levels in hair, nail, and urine are usually reflective of longer aggregation, and its levels in the blood sample are usually indicative of short-term exposure, which this could be a source of the observed heterogeneity [54].

Overall, this meta-analysis indicated that although the Hg level in diabetic individuals is significantly higher than the nondiabetics, there existed no association between Hg exposure and the risk of T2DM. However, exposure to Hg in men might reduce the risk of T2DM; however, additional studies are required to confirm this finding.

Data Availability
All data and codes are available upon request to the corresponding author.

Conflicts of Interest
The authors declare that there are no conflicts of interest.

Authors’ Contributions
BGN, TR, PJ, FM, MZ, AD, and NB participated in the study design and in writing the manuscript. MHD, FA, and MM participated in the data collection. SA and BGN analyzed the data. TR and PR interpreted the data and critically reviewed the paper under the supervision of SA. All authors read and approved the final manuscript.

References
[1] S. Chatterjee, K. Khunti, and M. J. Davies, “Type 2 diabetes,” The Lancet, vol. 389, no. 10085, pp. 2239–2251, 2017.
[2] C. Bommer, V. Sagalova, E. Heesemann et al., “Global economic burden of diabetes in adults: projections from 2015–2030,” Diabetes Care, vol. 41, no. 5, pp. 963–970, 2018.
[3] E. R. Pearson, “Type 2 diabetes: a multifaceted disease,” Diabetologia, vol. 62, no. 7, pp. 1107–1112, 2019.
[4] Y. Zheng, S. H. Ley, and F. B. Hu, “Global aetiology and epidemiology of type 2 diabetes mellitus and its complications,” Nature Reviews Endocrinology, vol. 14, no. 2, pp. 88–98, 2018.
[5] T. L. M. Hectors, C. Vanparys, K. van der Ven et al., “Environmental pollutants and type 2 diabetes: a review of mechanisms that can disrupt beta cell function,” Diabetologia, vol. 54, no. 6, pp. 1273–1290, 2011.
[6] Y. W. Chen, C. Y. Yang, C. F. Huang, D. Z. Hung, Y. M. Leung, and S. H. Liu, “Heavy metals, islet function and diabetes development,” Islets, vol. 1, no. 3, pp. 169–176, 2009.
[7] H. Hernández-Mendoza, M. J. Rios-Lugo, H. E. Álvarez-Loredo et al., “Serum lead levels and its association with overweight and obesity,” Journal of Trace Elements in Medicine & Biology, vol. 72, Article ID 126984, 2022.
[8] H. Hernández-Mendoza, H. E. Álvarez-Loredo, E. T. Romero-Guzmán et al., “Relationship between serum levels of arsenic, cadmium, and mercury and body mass index and fasting plasma glucose in a Mexican adult population,” Biological Trace Element Research, vol. 200, pp. 1–8, 2022.
[9] A. Planchart, A. Green, C. Hoyo, and C. J. Mattingly, “Heavy metal exposure and metabolic syndrome: evidence from human and model system studies,” Current environmental health reports, vol. 5, no. 1, pp. 110–124, 2018.
[10] P. B. Tchounwou, C. G. Yedjou, A. K. Patolla, and D. J. Sutton, “Heavy metal toxicity and the environment,” EXPERIMENTA Supplementum, vol. 101, pp. 133–164, 2012.
[11] R. A. Bernhoff, “Mercury toxicity and treatment: a review of the literature,” Journal of Environmental and Public Health, vol. 2012, Article ID 460508, 10 pages, 2012.
[12] K. M. Rice, E. M. Walker, M. Wu, C. Gillette, and E. R. Blough, “Environmental mercury and its toxic effects,” J Prev Med Public Health, vol. 47, no. 2, pp. 74–83, 2014.
[13] Y. W. Chen, C. F. Huang, K. S. Tsai et al., “Methylmercury induces pancreatic beta-cell apoptosis and dysfunction,” Chemical Research in Toxicology, vol. 19, no. 8, pp. 1080–1085, 2006.
[14] Y. Hotta, R. Fujino, O. Kimura, Y. Fujii, K. Haraguchi, and T. Endo, “Assessment of diabetics by the quantification of essential elements and stable isotope ratios of carbon and nitrogen in scalp hair,” Obesity Medicine, vol. 15, Article ID 100106, 2019.
[15] X. Wang, C. A. Karvonen-Gutierrez, W. H. Herman, B. Mukherjee, S. D. Harlow, and S. K. Park, “Urinary metals and incident diabetes in midlife women: study of Women’s Health across the Nation (SWAN),” BMJ open diabetes research & care, vol. 8, no. 1, Article ID e001233, 2020.
[16] M. I. Ward and B. Pim, “Trace element concentrations in blood plasma from diabetic patients and normal individuals,” Biological Trace Element Research, vol. 6, no. 6, pp. 469–487, 1984.
[17] M. Ahlqwist, C. Bengtsson, L. Lapidus, I. A. Gerdgahd, and A. Schütz, “Serum mercury concentration in relation to survival, symptoms, and diseases: results from the prospective population study of women in Gothenburg, Sweden,” Acta Odontologica Scandinavica, vol. 57, no. 3, pp. 168–174, 1999.
[18] J. Zhang, J. Wang, J. Hu, J. Zhao, J. Li, and X. Cai, “Associations of total blood mercury and blood methylmercury concentrations with diabetes in adults: an exposure-response analysis of 2005–2018 NHANES,” Journal of Trace Elements in Medicine & Biology, vol. 68, Article ID 126845, 2021.
[19] I. Rotter, D. Kosik-Bogacka, B. Dolegowska, K. Safranow, A. Lubirowska, and M. Laszczyńska, “Relationship between the concentrations of heavy metals and bioelements in aging men with metabolic syndrome,” International Journal of Environmental Research and Public Health, vol. 12, no. 4, pp. 3944–3961, 2015.
[20] J.-W. Chang, H.-L. Chen, H.-J. Su, P.-C. Liao, H.-R. Guo, and A. Lubkowska, and M. Laszczyńska, “Relationship between the concentrations of total blood mercury and blood methylmercury and body mass index and fasting plasma glucose in a Mexican adult population,” Biological Trace Element Research, vol. 200, pp. 1–8, 2022.
[21] R. Durak, Y. Gülen, M. Kurudirek, M. Kaçal, and I. Capoglu, “Heavy metal exposure and metabolic syndrome: evidence from human and model system studies,” Current environmental health reports, vol. 5, no. 1, pp. 110–124, 2018.
[22] P. B. Tchounwou, C. G. Yedjou, A. K. Patolla, and D. J. Sutton, “Heavy metal toxicity and the environment,” EXPERIMENTA Supplementum, vol. 101, pp. 133–164, 2012.
[22] A. S. Ettinger, P. Bovet, J. Plange-Rhule et al., “Distribution of metals exposure and associations with cardiometabolic risk factors in the “Modeling the Epidemiologic Transition Study,””, Environmental Health, vol. 13, no. 1, p. 90, 2014.

[23] C. Jeppesen, B. Valera, N. O. Nielsen, P. Bjerringaard, and M. E. Jørgensen, “Association between whole blood mercury and glucose intolerance among adult Inuit in Greenland,” Environmental Research, vol. 143, pp. 192–197, 2015.

[24] P. Dufault, Z. Berg, R. Crider et al., “Blood inorganic mercury is directly associated with glucose levels in the human population and may be linked to processed food intake,” Integrative molecular medicine, vol. 2, no. 3, 2015.

[25] S. Pal, J. M. Blais, M. A. Robidoux et al., “The association of type 2 diabetes and insulin resistance/secretion with persistent organic pollutants in two First Nations communities in northern Ontario,” Diabetes and Metabolism, vol. 39, no. 6, pp. 497–504, 2013.

[26] Z. Ghaedrahmat, B. Cheraghian, N. Jaafarzadeh, A. Takdastan, H. B. Shahbazi, and M. Ahmadi, “Relationship between urinary heavy metals with metabolic syndrome and its components in population from Hoveyze b cohort study: a case-control study in Iran,” Journal of Trace Elements in Medicine & Biology, vol. 66, Article ID 126757, 2021.

[27] M. G. Skalnaya and V. A. Demidov, “Hair trace element contents in women with obesity and type 2 diabetes,” Journal of Trace Elements in Medicine & Biology, vol. 21, pp. 59–61, 2007.

[28] S. Cordier, E. Anassour-Laouan-Sidi, M. Lemire, N. Costet, M. Lucas, and P. Ayotte, “Association between exposure to persistent organic pollutants and mercury, and glucose metabolism in two Canadian Indigenous populations,” Environmental Research, vol. 184, Article ID 109345, 2020.

[29] G. Forte, B. Bocca, A. Peruzzu et al., “Blood metals concentration in type 1 and type 2 diabetics,” Biological Trace Element Research, vol. 156, no. 1-3, pp. 79–90, 2013.

[30] M. Futatsuka, T. Kitano, and J. Wakamiya, “An epidemiological study on diabetes mellitus in the population living in a methyl mercury polluted area,” Journal of Epidemiology, vol. 6, no. 4, pp. 204–208, 1996.

[31] A. F. Hansen, A. Simić, B. O. Asvold et al., “Trace elements in early phase type 2 diabetes mellitus—a population-based study. The HUNT study in Norway,” Journal of Trace Elements in Medicine & Biology, vol. 40, pp. 46–53, 2017.

[32] Y. Hotta, R. Fujino, O. Kimura, and T. Endo, “Essential and non-essential elements in scalp hair of diabetics: correlations with glycated hemoglobin (HbA1c),” Biological and Pharmaceutical Bulletin, vol. 41, no. 7, pp. 1034–1039, 2018.

[33] S.-S. Moon, “Association of lead, mercury and cadmium with diabetes in the Korean population: the Korea national health and nutrition examination survey (KNHANES) 2009-2010,” Diabetic Medicine 2009–2010, vol. 30, no. 4, pp. e143–e148, 2013.

[34] D. Mozaffarian, P. Shi, J. S. Morris et al., “Methylmercury exposure and incident diabetes in U.S. men and women in two prospective cohorts,” Diabetes Care, vol. 36, no. 11, pp. 3578–3584, 2013.

[35] M. K. Moon, I. Lee, A. Lee et al., “Lead, mercury, and cadmium exposures are associated with obesity but not with diabetes mellitus: Korean National Environmental Health Survey (KoNEHS) 2015–2017,” Environmental Research, vol. 204, Article ID 111888, 2022.

[36] D. Moher, A. Liberati, J. Tetzlaff, D. G. Altman, and P. Group, “Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement,” PLoS Medicine, vol. 6, no. 7, Article ID e100097, 2009.

[37] J. Peterson, V. Welch, M. Losos, and P. Tugwell, The Newcastle-Ottawa Scale (NOS) for Assessing the Quality of Non-randomised Studies in Meta-Analyses, pp. 1–12, Ottawa Hospital Research Institute, Ottawa, Canada, 2011.

[38] J. Cohen, Statistical Power Analysis for the Behavioral Sciences, pp. 20–26, Lawrence Erlbaum Associates, Hillsdale, NJ, USA, 1988.

[39] M. A. Serdar, F. Bakir, A. Haşimi et al., “Trace and toxic element patterns in nonsmoker patients with noninsulin-dependent diabetes mellitus, impaired glucose tolerance, and fasting glucose,” International Journal of Diabetes in Developing Countries, vol. 29, no. 1, p. 35, 2009.

[40] H. I. Afridi, F. N. Talpur, T. G. Kazi, and D. Brabazon, “Effect of trace and toxic elements of different brands of cigarettes on the essential status of Irish referent and diabetic mellitus consumers,” Biological Trace Element Research, vol. 167, no. 2, pp. 209–224, 2015.

[41] A. Simić, A. F. Hansen, B. O. Asvold et al., “Trace element status in patients with type 2 diabetes in Norway: the HUNT3 Survey,” Journal of Trace Elements in Medicine & Biology, vol. 41, pp. 91–98, 2017.

[42] T.-L. Tsai, C.-C. Kuo, W.-H. Pan, T.-N. Wu, P. Lin, and S.-L. Wang, “Type 2 diabetes occurrence and mercury exposure—from the national nutrition and health survey in Taiwan,” Environmental International, vol. 126, pp. 260–267, 2019.

[43] B. B. Little, R. Reilly, B. Walsh, and G. T. Vu, “Cadmium is associated with type 2 diabetes in a superfund site lead smelter community in Dallas, Texas,” International Journal of Environmental Research and Public Health, vol. 17, no. 12, p. 4558, 2020.

[44] C. Roy, P.-Y. Tremblay, and P. Ayotte, “Is mercury exposure causing diabetes, metabolic syndrome and insulin resistance? A systematic review of the literature,” Environmental Research, vol. 156, pp. 747–760, 2017.

[45] C.-C. Kuo, K. Moon, K. A. Thayer, and A. Navas-Acien, “Environmental chemicals and type 2 diabetes: an updated systematic review of the epideimiologic evidence,” Current Diabetes Reports, vol. 13, no. 6, pp. 831–849, 2013.

[46] A. Lie, N. Gundersen, and K. J. Korsgaard, “Mercury in urine—sex, age and geographic differences in a reference population,” Scandinavian Journal of Work, Environment & Health, vol. 8, no. 2, pp. 129–133, 1982.

[47] M. Fujimura and F. Usuki, “Methylmercury-mediated oxidative stress and activation of the cellular protective system,” Antioxidants, vol. 9, no. 10, p. 1004, 2020.

[48] J. A. Scott and G. L. King, “Oxidative stress and antioxidant treatment in diabetes,” Annals of the New York Academy of Sciences, vol. 1031, no. 1, pp. 204–213, 2004.

[49] D. Mozaffarian, P. Shi, J. S. Morris et al., “Mercury exposure and risk of cardiovascular disease in two US cohorts,” New England Journal of Medicine, vol. 364, no. 12, pp. 1116–1125, 2011.

[50] E. Guallar, M. I. Sanz-Gallardo, P. V. Veer et al., “Mercury, fish oils, and the risk of myocardial infarction,” New England Journal of Medicine, vol. 347, no. 22, pp. 1747–1754, 2002.

[51] J. K. Virtanen, S. Voutilainen, T. H. Rissanen et al., “Mercury, fish oils, and risk of acute coronary events and cardiovascular
disease, coronary heart disease, and all-cause mortality in men in eastern Finland,” *Arteriosclerosis, Thrombosis, and Vascular Biology*, vol. 25, no. 1, pp. 228–233, 2005.

[53] M. Karapehlivan, M. Ogun, I. Kaya, H. Ozen, H. A. Deveci, and M. Karaman, “Protective effect of omega-3 fatty acid against mercury chloride intoxication in mice,” *Journal of Trace Elements in Medicine & Biology*, vol. 28, no. 1, pp. 94–99, 2014.

[54] W. I. Mortada, M. A. Sobh, M. M. El-Defrawy, and S. E. Farahat, “Reference intervals of cadmium, lead, and mercury in blood, urine, hair, and nails among residents in Mansoura city, Nile delta, Egypt,” *Environmental Research*, vol. 90, no. 2, pp. 104–110, 2002.