Vitamins and *Helicobacter pylori*: An Updated Comprehensive Meta-Analysis and Systematic Review

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Background: Over recent decades, epidemiological studies have shown relationships between vitamins and *Helicobacter pylori* (*H. pylori*) infection and eradication, but the results are controversial.

Methods: A comprehensive meta-analysis and systematic review were conducted to clarify the relationships between common types of vitamins and *H. pylori*. We applied meta-regression, subgroup analysis and sensitivity analysis to obtain available evidence. Articles published from January 1991 to June 2021 in PubMed, EMBASE, and the Cochrane Library were searched.

Results: In total, we identified 48 studies. The results indicate that *H. pylori*-positive patients had lower serum vitamin B₁₂ (standardized mean difference (SMD) = −0.30; 95% confidence interval (CI): −0.53 – −0.08), folate (SMD = −0.69; 95% CI: −1.34 – −0.04), vitamin C (SMD = −0.37; 95% CI: −0.57 – −0.18) and vitamin D (SMD = −0.34; 95% CI: −0.49 – −0.18) levels than *H. pylori*-negative patients. Patients in which *H. pylori* had been successfully eradicated had higher serum vitamin D levels (SMD = 1.37; 95% CI: 0.37–2.38) than in patients in which eradication had been unsuccessful. The serum vitamin B₁₂ levels of *H. pylori*-positive patients improved after successful *H. pylori* eradication therapy (SMD = 1.85; 95% CI: 0.81–2.90), and antioxidant vitamin supplementation to an *H. pylori* eradication regimen improved the eradication rate (risk ratio = 1.22; 95% CI: 1.02–1.44 for per-protocol analysis; risk ratio = 1.25; 95% CI: 1.06–1.47 for intention-to-treat analysis).

Conclusions: *H. pylori* infections decrease the serum levels of several types of vitamins, eradication of *H. pylori* could rescue its adverse effects, and antioxidant vitamin supplementation may improve the *H. pylori* eradication rate.

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Keywords: vitamins, helicobacter pylori, meta-analysis, systematic review, relationship
INTRODUCTION

Helicobacter pylori (H. pylori) is a gastric gram-negative, spiral-shaped microaerophilic pathogen (1). About half of the global population is infected with H. pylori, and the infection rate in developing countries is higher than in developed countries (2, 3). H. pylori is the main risk for chronic gastritis, gastric ulcer, gastric cancer and mucosa-associated lymphoid tissue–associated lymphoma (4–6). It can damage the gastric mucosa and affect the absorption of trace elements, especially vitamins (7). Vitamin deficiency can upset the internal balance of the human body and cause a variety of diseases outside of the digestive system (8–10).

Vitamins are members of a huge family. At present, there are dozens of known vitamins that may be divided into fat-soluble and water-soluble categories. The relationships between H. pylori infection and various vitamins have attracted attention worldwide. Some studies found that H. pylori infections reduce serum vitamin levels (11, 12); several studies have revealed that after H. pylori eradication, the serum vitamin levels increase (13, 14). Some randomized controlled trials (RCTs) found that vitamin supplementation combined with standard anti-H. pylori therapy increase the H. pylori eradication rate (15, 16). However, the results have been inconsistent.

Meta-analyses on the relationships between vitamins and H. pylori have been published (17–19). These studies involved one vitamin or a certain aspect of the relationships between vitamins and H. pylori, or the number of included studies was limited. Yang et al. (17) found vitamin D could improve the success rate of H. pylori eradication. However, they only identified three relevant studies to support this conclusion. Afsar et al. (18) reported a relationship between H. pylori infection and micronutrient (vitamin B12 and folate) levels in pregnant women. Nevertheless, the effects of H. pylori on the population excluding pregnant women were not evaluated. Li et al. (19) assessed the effects of antioxidant vitamins supplementation on the rate of H. pylori eradication. However, only three supporting studies were referenced in that work. In recent years, many excellent articles have been published. To update the results and obtain more credible conclusions, we conducted this systematic review and meta-analysis to evaluate the relationships more comprehensively, thereby providing a theoretical basis for clinical practice and public health policy-making.

METHODS

Data Sources and Search Strategy

This meta-analysis was registered on PROSPERP (No. CRD42021268127) (20) and compliant with the main PRISMA statement (21). A comprehensive and systematic search was carried out for relevant studies describing relationships between vitamins and H. pylori in biologic and medical databases (Medline, Web of Science, Embase, Chinese Biomedical Database and Cambridge Scientific Abstracts databases). We developed a search strategy using following keywords: “vitamins,” “vitamin A,” “vitamin B,” “vitamin C,” “vitamin D,” “vitamin E,” “β-Carotene,” “retinol,” “cobalamin” “folate,” “folic acid,” “tocopherol,” “antioxidants,” “micronutrient,” and “Helicobacter pylori” (as shown in Supplementary Table 1). Duplicate works were collapsed into a single entry. Additionally, we scanned the reference lists of all the relevant published studies and reviews. The two blinded reviewers (Xianlei Cai and Xueying Li) selected the studies and specified the exclusion criteria.

Inclusion and Exclusion Criteria

The inclusion criteria were as follows: (1) Observational or experimental research; (2) Comparisons of serum vitamin levels between H. pylori - positive and H. pylori - negative patients; (3) Comparisons of serum vitamin levels between successful and failed H. pylori eradication patients; (4) Comparisons of serum vitamin levels before and after successful H. pylori eradication therapy; (5) Comparisons of H. pylori eradication rate between antioxidant vitamin supplementation groups and controlled groups for H. pylori – positive patients; and (6) Original studies in English or Chinese indexed up to June 2021.

The exclusion criteria were as follows: (1) Original studies did not involve the relationships between vitamins and H. pylori; (2) Studies did not provide sufficient data for a meta-analysis; (4) reviews, comments, letters, and animal studies; and (5) low-quality studies [Newcastle-Ottawa scale and Cochrane collaboration’s tool were used to assess the quality and bias risk for cross-sectional, cohort and random-controlled studies as described in a previous study (22)].

Data Extraction

All the data were extracted by three researchers independently (Xianlei Cai, Xueying Li and Yangli Jin) using standardized form. The characteristics of the identified relevant works were records as follows: name of first author, year of publication, country, study design (cross-sectional, case series, cohort and RCTs), age, number of subjects, gender, type of vitamins (vitamins A, B, C, D and E) and presentation of effect magnitude [mean ± standard deviation (SD), mean ± standard error of mean (SEM), median (interquartile range), median (range), odds risk (OR), risk ratio (RR), or hazard ratio (HR) with 95% confidence interval (CI)].

Statistical Analyses

Mean serum vitamin levels and the SDs were used in pooling analyses. If the original studies provided other estimates [mean ± SEM, median (interquartile range) or median (range)], we converted the data using a common method described by Hozo et al. (23). Because of the inconsistent units used in different studies, the pooled results were expressed in terms of standardized mean difference (SMD) with 95% CI. If the RCTs and cohort studies only provided 2 × 2 table data, we calculated the responding RRs. Additionally, RRs and 95% CIs were used to show the differences in the H. pylori eradication rates between vitamin supplementation and control groups.

A meta-regression was performed to examine the sources of heterogeneity from disparate types of vitamin supplementation (vitamin C or vitamin C plus vitamin E), and we identified influence factors having positive coefficients (p ≤ 0.05). Q-test and I² were used to assess heterogeneity. If the results
showed notable heterogeneity ($p \leq 0.05$ and $I^2 > 50\%$), then pooled estimates were calculated using random-effects models (DerSimonian and Laird method) (24). Otherwise, fixed-effects models were used (Mantel-Haenszel method) (25). Subgroup analyses were performed to evaluate relationships between diverse types of vitamins and H. pylori. Forest and funnel plots were drawn and publication bias was tested by a weighted Egger and Begg’s tests (26, 27). Sensitivity analyses were performed by omitting one estimate at a time to assess the relative influence of each work on pooled results. If the included estimates were less than four, then we did not carry out the meta-analysis and conducted systematic review instead. All the analyses were performed using STATA version 12.0 (StataCorp LP).

RESULTS

Study Characteristics
The flow-through of the study selection process is described using a modified PRISMA diagram (Figure 1). In total, forty-eight high quality studies (39 observational studies and 9 RCTs) with 73 independent estimates of relationships between vitamins and H. pylori in four fields were included in this meta-analysis and systematic review. In total, 31 studies compared serum vitamin levels between H. pylori - positive and - negative patients; 5 studies compared serum vitamin levels between successful and failed H. pylori eradication patients; 6 studies compared patient serum vitamin levels before and after successful H. pylori eradication therapy; and 10 studies focused on the effects of vitamin supplementation on the H. pylori eradication rate. Of these, three studies (28–30) considered more than one effect of H. pylori on vitamins. There were 173,013 participants from Turkey (12 studies), the United Kingdom (6 studies), China (4 studies), Italy (4 studies), Brazil (2 studies), Iran (2 studies), Israel (2 studies), Japan (2 studies), the USA (2 studies), and one each from Argentina, Australia, Egypt, Germany, Greece, India, Lebanon, Netherlands, Pakistan, Palestine, Poland, Saudi Arabia and Switzerland. Overall, Tables 1–4 present the summaries of all the studies included in the meta-analysis. The original estimates reported in the articles are summarized in Supplementary Tables 2–5.
**TABLE 1 | Basic characteristics of studies comparing serum vitamin levels between *H. pylori* + groups and *H. pylori* - negative groups.**

| Study | Year | Area | Design | Age of HP + groups | No. of HP + groups (male/female) | Age of HP - groups | No. of HP - groups (male/female) | Quality |
|-------|------|------|--------|---------------------|----------------------------------|-------------------|----------------------------------|---------|
| **Vitamin A** | | | | | | | | |
| Phull et al. (31) | 1998 | UK | Cross-sectional | Mean 46.0 y | 25 (18/7) | Mean 54 y | 18 (7/11) | 7* |
| Zhang et al. (32) | 2000 | UK | Cross-sectional | 19–89 y | 41 (N/A) | 19–89 y | 27 (N/A) | 7* |
| Toyonaga et al. (33) | 2000 | Japan | Cross-sectional | Mean 47.0 y | 37 (13/24) | Mean 44.7 y | 40 (14/28) | 8* |
| **Vitamin B12** | | | | | | | | |
| Tamura et al. (34) | 2002 | Japan | Cross-sectional | Mean 64 y | 57 (44/13) | Mean 63 y | 36 (25/11) | 8* |
| Cenerelli et al. (35) | 2002 | Italy | Cross-sectional | Mean 54.7 y | 31 (19/12) | Mean 50.9 y | 42 (23/19) | 8* |
| Shuvat-Sudai et al. (36) | 2003 | Israel | Cross-sectional | Mean 52.8 y | 96 (N/A) | Mean 49.2 y | 37 (N/A) | 7* |
| Trinarchi et al. (37) | 2004 | Argentina | Cross-sectional | Mean 56.8 y | 8 (3/5) | Mean 62.4 y | 21 (9/12) | 7* |
| Ojien et al. (38) | 2004 | Netherlands | Cross-sectional | N/A | 29 (N/A) | N/A | 60 (N/A) | 7* |
| Sarari et al. (39) | 2008 | Palestine | Cross-sectional | Mean 43.4 y | 43 (24/19) | Mean 49.2 y | 29 (14/15) | 7* |
| Stettin et al. (40) | 2008 | Germany | Cross-sectional | Mean 50.8 y | 69 (27/42) | Mean 47.3 y | 21 (8/13) | 8* |
| Kakehasi et al. (41) | 2002 | Brazil | Cross-sectional | Mean 63.7 y | 34 (0/37) | Mean 62.5 y | 27 (0/27) | 8* |
| Gerig et al. (42) | 2013 | Switzerland | Cross-sectional | Mean 42.3 y | 85 (21/64) | Mean 40.9 y | 319 (85/234) | 8* |
| Oijen et al. (43) | 2004 | Argentina | Cross-sectional | Mean 46.4 y | 31 (12/19) | Mean 45.2 y | 19 (8/11) | 8* |
| **Folate** | | | | | | | | |
| Tamura et al. (34) | 2002 | Japan | Cross-sectional | Mean 64 y | 57 (44/13) | Mean 63 y | 36 (25/11) | 8* |
| Cenerelli et al. (35) | 2002 | Italy | Cross-sectional | Mean 54.7 y | 31 (19/12) | Mean 50.9 y | 42 (23/19) | 8* |
| Shuvat-Sudai et al. (36) | 2003 | Israel | Cross-sectional | Mean 52.8 y | 96 (N/A) | Mean 49.2 y | 37 (N/A) | 7* |
| Stettin et al. (39) | 2008 | Germany | Cross-sectional | Mean 50.8 y | 69 (27/42) | Mean 47.3 y | 21 (8/13) | 8* |
| Gerig et al. (41) | 2013 | Switzerland | Cross-sectional | Mean 42.3 y | 85 (21/64) | Mean 40.9 y | 319 (85/234) | 8* |
| Ulasoglu et al. (42) | 2019 | Turkey | Cross-sectional | Mean 44.8 y | 213 (N/A) | Mean 44.8 y | 76 (N/A) | 7* |
| Sarry et al. (43) | 2019 | Turkey | Cross-sectional | Mean 46.4 y | 31 (12/19) | Mean 45.2 y | 19 (8/11) | 8* |
| **Vitamin C** | | | | | | | | |
| Banerjee et al. (28) | 1994 | UK | Cross-sectional | N/A | 19 (N/A) | N/A | 10 (N/A) | 7* |
| Rokkas et al. (47) | 1999 | Greece | Cross-sectional | Mean 42.0 y | 30 (17/13) | Mean 42.5 y | 10 (6/4) | 7* |
| Jarosz et al. (48) | 2000 | Poland | Cross-sectional | Mean 45.5 y/39.0 y | 21 (11/10) | 32 (18/14) | 17 (10/7) | 16 (9/7) | 8* |
| Toyonaga et al. (33) | 2000 | Japan | Cross-sectional | Mean 47.0 y | 37 (13/24) | Mean 44.7 y | 40 (14/28) | 8* |
| Woodward et al. (49) | 2001 | UK | Cross-sectional | 25–74 y | 766 (N/A) | 25–74 y | 403 (N/A) | 8* |
| Rokkas et al. (47) | 1999 | Greece | Cross-sectional | Mean 42.0 y | 30 (17/13) | Mean 42.5 y | 10 (6/4) | 7* |
| **Vitamin D** | | | | | | | | |
| Antico et al. (53) | 2012 | Italy | Cross-sectional | 20–80 y | 21 (N/A) | 20–80 y | 163 (N/A) | 7* |
| Gerig et al. (41) | 2013 | Switzerland | Cross-sectional | Mean 42.3 y | 85 (21/64) | Mean 40.9 y | 319 (85/234) | 8* |
| Han et al. (54) | 2019 | China | Cross-sectional | Mean 47.1 y | 496 (236/260) | Mean 48.1 y | 257 (127/300) | 8* |
| Surmeli et al. (43) | 2019 | Turkey | Cross-sectional | Mean 74.7 y | 43 (11/32) | Mean 78.2 y | 211 (91/120) | 9* |
| Assad et al. (55) | 2019 | Lebanon | Cross-sectional | Mean 39.3 y | 225 (88/137) | Mean 41.9 y | 235 (88/137) | 8* |
| Gao et al. (56) | 2020 | China | Cross-sectional | Mean 12.1 y | 2113 (1202/911) | Mean 12.4 y | 4783 (2865/2098) | 9* |
| Shafrir et al. (44) | 2021 | Israel | Cross-sectional | Mean 41.0 y | 7564 (3875/3764) | Mean 42.2 y | 74843 (37421/37422) | 9* |

(Continued)
TABLE 1 | Continued

| Study                  | Year | Area   | Design       | Age of HP + groups | No. of HP + groups (male/female) | Age of HP - groups | No. of HP - groups (male/female) | Quality |
|------------------------|------|--------|--------------|--------------------|----------------------------------|--------------------|----------------------------------|---------|
| Vitamin E              |      |        |              |                    |                                  |                    |                                  |         |
| Phull et al. (31)      | 1998 | UK     | Cross-sectional | Mean 46 y          | 25 (18/7)                       | Mean 54 y          | 18 (7/11)                        | 7*      |
| Zhang et al. (32)      | 2000 | UK     | Cross-sectional | 19–89 y           | 41 (N/A)                        | 19–89 y            | 27 (N/A)                         | 7*      |
| Toyonaga et al. (33)   | 2000 | Japan  | Cross-sectional | Mean 47.0 y        | 37 (13/24)                      | Mean 44.7 y        | 40 (14/26)                       | 8*      |

UK, United Kingdom; N/A, not available; y, years; m, months; “The “star system” of the Newcastle–Ottawa scale.

TABLE 2 | Basic characteristics of studies comparing serum vitamin levels between the successful H. pylori eradication groups and the failed groups.

| Study                  | Year | Area   | Design       | Age of successful groups | No. of successful groups (male/female) | Age of failed groups | No. of failed groups (male/female) | Quality |
|------------------------|------|--------|--------------|--------------------------|----------------------------------------|----------------------|------------------------------------|---------|
| Vitamin D              |      |        |              |                          |                                  |                      |                                    |         |
| Yildirim et al. (57)   | 2017 | Turkey | Cross-sectional | N/A                      | 170 (N/A)                            | N/A                  | 50 (N/A)                           | 7*      |
| Shahawy et al. (58)   | 2018 | Egypt  | Cross-sectional | 18–80 y                 | 105 (N/A)                           | 18–80 y              | 45 (N/A)                           | 7*      |
| Maggi et al. (2)       | 2021 | Pakistan | Cross-sectional | 18–60 y                 | 88 (42/46)                           | 18–60 y              | 36 (18/18)                        | 8*      |
| Shatla et al. (59)    | 2021 | Saudi Arabia | Cross-sectional | N/A                      | 109 (N/A)                           | N/A                  | 42 (N/A)                           | 7*      |
| Shafrir et al. (56)   | 2021 | Israel | Cross-sectional | N/A                      | 45821 (N/A)                         | N/A                  | 29722 (N/A)                       | 9*      |

N/A, not available; y, years; “The “star system” of the Newcastle–Ottawa scale.

TABLE 3 | Basic characteristics of studies comparing serum vitamin levels before and after H. pylori eradication therapy.

| Study                  | Year | Area   | Design       | Age             | No. before eradication (male/female) | No. after eradication (male/female) | Test time after eradication | Quality |
|------------------------|------|--------|--------------|-----------------|--------------------------------------|-------------------------------------|-------------------------------|---------|
| Vitamin B<sub>12</sub> |      |        |              |                 |                                      |                                     |                               |         |
| Kaptan et al. (13)     | 2000 | Turkey | Case series  | Mean 59.5 y     | 31 (19/12)                           | 31 (19/12)                         | 3 or 6 m                     | 8*      |
| Serin et al. (60)      | 2002 | Turkey | Case series  | Mean 43 y       | 65 (N/A)                             | 65 (N/A)                           | 2–3 m                        | 7*      |
| Ozer et al. (61)       | 2005 | Turkey | Case series  | Mean 41 y       | 41 (N/A)                             | 41 (N/A)                           | 1 m                           | 7*      |
| Marino et al. (14)     | 2007 | Brazil | Case series  | Mean 72.8 y     | 59 (N/A)                             | 59 (N/A)                           | 6 m / 12 m                   | 7*      |
| Folate                 |      |        |              |                 |                                      |                                     |                               |         |
| Kaptan et al. (13)     | 2000 | Turkey | Case series  | Mean 59.5 y     | 31 (19/12)                           | 31 (19/12)                         | 3 or 6 m                     | 8*      |
| Ozer et al. (61)       | 2005 | Turkey | Case series  | Mean 41 y       | 41 (N/A)                             | 41 (N/A)                           | 1 m                           | 7*      |
| Vitamin C              |      |        |              |                 |                                      |                                     |                               |         |
| Banerjee et al. (28)   | 1994 | UK     | Case series  | N/A             | 11 (N/A)                             | 11 (N/A)                           | 1 m / 6 m                    | 7*      |
| Annibale et al. (29)   | 2003 | Italy  | Case series  | Median 47 y     | 5 (N/A)                              | 5 (N/A)                            | 6 m                           | 7*      |

N/A, not available; y, years; m, months; “The “star system” of the Newcastle–Ottawa scale.

Vitamin Levels Discrepancies Between H. pylori - Positive and - Negative Patients

The number of studies on the relationships between H. pylori and vitamin B<sub>12</sub>, folate, vitamin C and vitamin D was sufficient for a meta-analysis (Table 1, Supplementary Table 2).

For vitamin B<sub>12</sub>, 12 estimates were included in the pooled analysis. The results indicated that H. pylori – positive patients had lower serum vitamin B<sub>12</sub> levels than H. pylori – negative patients (SMD = −0.30; 95% CI: −0.53 – −0.08; Figure 2), with heterogeneity ($P < 0.001; I^2 = 71.4\%$) and publication bias (Begg’s test $z_{c} = 2.26, P = 0.024$; Egger’s test $t = 0.05$; Figure 3A). The sensitivity analysis showed that the results were stable and reliable (Supplementary Figure 1A).

For folate, eight estimates were incorporated into the meta-analysis. Similarly, the results showed that H. pylori – positive patients had lower serum folate levels than H. pylori-negative patients (SMD = −0.69; 95% CI: −1.34 – −0.04; Figure 2), with obvious heterogeneity ($P < 0.001; I^2 = 95.8\%$). Publication
bias was not found (Begg’s test $z_c = 0.87, P = 0.386$; Egger’s test $t = 0.254$; Figure 3A). The sensitivity analysis showed that the results were influenced by some positive data (Supplementary Figure 1B).

For vitamin C, 14 estimates were incorporated into the pooled analysis. The results revealed that $H. \text{pylori}$ – positive patients had lower serum vitamin C levels than $H. \text{pylori}$-negative patients (SMD $= -0.37; 95\% \text{ CI: } -0.57 - -0.18$; Figure 2), with obvious heterogeneity ($P < 0.001; I^2 = 87.9\%$). There was no publication bias (Begg’s test $z_c = 0.44, P = 0.661$; Egger’s test $t = 0.130$; Figure 3A). The sensitivity analysis showed that the result was stable (Supplementary Figure 1C).

For vitamin D, seven estimates were included in the meta-analysis. The result also found that $H. \text{pylori}$ – positive patients had lower serum vitamin D levels than $H. \text{pylori}$-negative patients (SMD $= -0.34; 95\% \text{ CI: } -0.49 - -0.18$; Figure 2), with heterogeneity ($P < 0.001; I^2 = 95.0\%$). Publication bias was not found (Begg’s test $z_c = 0.60, P = 0.548$; Egger’s test $t = 0.412$; Figure 3A). The sensitivity analysis revealed that the results were robust (Supplementary Figure 1D).

For vitamins A and E, only three studies were identified; therefore, we did not conduct a pooled analysis. Among these three studies, Phull et al. (31) indicated that there was no relationship between $H. \text{pylori}$ infection and serum vitamin A and E levels, and their conclusion were similar to those of Zhang et al. (32) and Toyonaga et al. (33).

### Vitamin Levels Discrepancies Between Successful and Failed $H. \text{pylori}$ Eradication Patients

Overall, five studies focusing on vitamin D level discrepancies between successful and failed $H. \text{pylori}$ eradication patients were included in the meta-analysis (Table 2, Supplementary Table 3). The results indicated that the patients with successful $H. \text{pylori}$ eradication had higher serum vitamin D levels than the failed patients (SMD $= 1.37; 95\% \text{ CI: } 0.37 - 2.38$; Figure 4), with heterogeneity ($P < 0.001; I^2 = 98.4\%$). Because all five studies reported positive estimates, the funnel plot was asymmetric (Figure 3B). We did not assess publication bias using weighted Egger test and Begg’s tests owing to the insufficient numbers of estimates. The sensitivity analysis revealed that the results were robust (Supplementary Figure 2).

There were no studies exploring other vitamin levels (vitamin A, B, C and E) discrepancies between successful and failed $H. \text{pylori}$ eradication patients. More relevant research is recommended to clarify the relationships.

### Vitamin Level Discrepancies Before and After Successful $H. \text{pylori}$ Eradication Therapy

The numbers of studies for vitamins B$_{12}$ and C were sufficient for a meta-analysis (Table 3, Supplementary Table 4). For vitamin B$_{12}$, five estimates were incorporated into the pooled analysis. The result indicated that after successful $H. \text{pylori}$ eradication, serum vitamin B$_{12}$ increased (SMD $= 1.85; 95\% \text{ CI: } 0.81 - 2.90$; Figure 5), with heterogeneity ($P < 0.001; I^2 = 96.0\%$). The funnel plot was symmetric (Figure 3C). The sensitivity analysis showed that the results were stable and reliable (Supplementary Figure 3). For vitamin C, four estimates were included in the meta-analysis. The results showed that $H. \text{pylori}$ eradication did not increase the serum vitamin C level (SMD $= -0.32; 95\% \text{ CI: } -1.56 - -0.91$; Figure 5), with heterogeneity ($P = 0.002; I^2 = 79.7\%$). The funnel plot was symmetric (Figure 3C). Because the four estimates were extracted from two studies, we
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FIGURE 2 | Forest plot of the meta-analysis on comparison of serum vitamin levels between *H. pylori* - positive and *H. pylori* - negative patients (n1: number of *H. pylori* – positive patients; n2: number of *H. pylori* – negative patients); *H. pylori* – positive patients had lower serum vitamin B12 levels than *H. pylori* – negative patients (SMD = −0.30; 95% CI: −0.53 – −0.08; *P* < 0.001; I² = 71.4%); *H. pylori* – positive patients had lower serum folate levels than *H. pylori* - negative patients (SMD = −0.69; 95% CI: −1.34 – −0.04; *P* < 0.001; I² = 95.8%); *H. pylori* – positive patients had lower serum vitamin C levels than *H. pylori* - negative patients (SMD = −0.37; 95% CI: −0.57 – −0.18; *P* < 0.001; I² = 87.9%); *H. pylori* – positive patients had lower serum vitamin D levels than *H. pylori* - negative patients (SMD = −0.34; 95% CI: −0.49 – −0.18; *P* < 0.001; I² = 95.0%).
did not conduct a sensitivity analysis and the results should be interpreted cautiously.

For folate, we only identified two relevant studies; therefore, a pooled analysis was not performed. Both Kaptan et al. (13) et al. and Ozer et al. (61) concluded that successful H. pylori eradication was not associated with an increased serum folate level.

**The Effects of Antioxidant Vitamin Supplementation on H. pylori Eradication**

A total of 10 trials on the effects of antioxidant vitamin supplementation on H. pylori eradication were identified (Table 4 and Supplementary Table 4). Among them, nine studies were RCTs, and one study was cohort research. The risks of bias were summarized in Supplementary Table 6. To avoid heterogeneity derived from different study designs, we did not include the cohort research (69) in the meta-analysis. We calculated the pooled results of the per-protocol analysis and intention-to-treat analysis from RCTs.

For the per-protocol analysis, eight estimates were included. A meta-regression was performed to assess potential sources of heterogeneity from eradication therapy (triple or quadruple therapy) and types of vitamin supplementation (vitamin C or vitamin C plus vitamin E). We found that eradication therapy and types of vitamin supplementation were not influencing factors ($P = 0.441$ for eradication therapy; $P = 0.707$ for types of vitamin supplementation). Therefore, all eight estimates were incorporated into the meta-analysis. The results indicated that combining antioxidant vitamin supplementation with standard therapy could increase the H. pylori eradication rate ($RR = 1.22; 95\% CI: 1.02–1.44$; Figure 6), with heterogeneity ($P < 0.001$; $I^2 = 81.0\%$). There was no publication bias (Begg's test $z_c =$...
FIGURE 4 | Forest plot of the meta-analysis on the comparison of serum vitamin levels between *H. pylori* successful and failed eradication patients (n1: number of successful eradication patients; n2: number of failed eradication patients); The patients with successful *H. pylori* eradication had higher serum vitamin D levels than the failed patients (SMD = 1.37; 95% CI: 0.37 – 2.38; *P* < 0.001; I² = 98.4%).

0.12, *P* = 0.902; Egger's test t = 0.662), and the funnel plot was symmetric (Figure 3D). The sensitivity analysis revealed that the results were robust (Supplementary Figure 4).

For the intention-to-treat analysis, 10 estimates were included. A meta-regression was conducted, and the result revealed that eradication therapy and types of vitamin supplementation were not influencing factors (*P* = 0.832 for eradication therapy; *P* = 0.510 for types of vitamin supplementation). The results of the pooled analysis showed that antioxidant vitamins supplementation increased the *H. pylori* eradication rate (RR = 1.25; 95% CI: 1.06–1.47; Figure 6), with heterogeneity (*P* < 0.001; I² = 75.1%). Publication bias was not found (Begg's test za = 0.18, *P* = 0.858; Egger's test t = 0.973; Figure 3D). The sensitivity analysis showed that the results were stable.

Only the Kockar et al. study (67) explored the effects of vitamin A supplementation on *H. pylori* eradication. They determined that vitamin A was ineffective in *H. pylori* eradication. More relevant RCTs are recommended to clear the relationship.

**DISCUSSION**

The debate over the *H. pylori*-vitamin association has been persistent. The controversy mainly focuses on the following four aspects: (1) vitamin level discrepancies between *H. pylori*-positive and -negative patients; (2) vitamin level discrepancies between successful and failed *H. pylori* eradication patients; (3) vitamin level discrepancies before and after successful *H. pylori* eradication therapy; and (4) the effects of vitamin supplementation on *H. pylori* eradication. Phull et al. (31), Zhang et al. (32) and Toyonaga et al. (33) found that *H. pylori* infection was not associated with serum vitamin A and E levels. The vitamin B12 results from Trimarchi et al. (37), Sarari et al. (11) and Ulasoglu et al. (42) indicated that *H. pylori* had an adverse effect on serum vitamin B12 levels, whereas other observational studies (35, 36, 40, 44) presented a null association between *H. pylori* and vitamin B12. The findings of Tamura et al. (34), Shuval-Sudai et al. (36) and Ulasoglu et al. (42) showed that *H. pylori*-negative patients had higher serum folate levels than *H. pylori*-positive patients; however, some studies (35, 39, 41, 43, 44) did not present similar results. A similar controversy exists regarding the associations between serum vitamin C or D levels and *H. pylori*. The effects of *H. pylori* eradication on serum vitamin levels have aroused extensive interest. Several studies (2, 57–59) found that patients that underwent successful *H. pylori* eradication had higher serum vitamin D levels than patients with in which eradication failed. Several studies (13, 14, 60, 61) described vitamin B12 level discrepancies before and after successful *H. pylori* eradication therapy; nevertheless, there were also inconsistent results. Vitamin supplementation was hypothesized to aid in *H. pylori* eradication. Some high-quality RCTs were performed to assess the assisting role of antioxidant vitamin...
supplementation on *H. pylori* eradication. Sezikli et al. (65) found that supplementation with vitamin C plus vitamin E increases the *H. pylori* eradication rate; whereas Chuang et al. (68) and Zojaji et al. (16) indicated that simple vitamin C supplementation increases the *H. pylori* eradication rate. In contrast, studies from other researchers (66, 67) showed that vitamin supplementation has no effects on the *H. pylori* eradication rate. Several meta-analyses had also been published. Yang et al. (17) found that 25-hydroxyvitamin D levels in *H. pylori* - negative patients were higher than in *H. pylori* - positive patients, and patients with vitamin D deficiency had lower eradication rates of *H. pylori*. Lahner et al. (70) reported a comprehensive meta-analysis in 2012 focused on the relationships between micronutrients and *H. pylori*. This study found that *H. pylori* was associated with ascorbic acid levels in gastric juice, which were increased by the eradication treatment. At present, many relevant articles have been published, and we have the opportunity to update the results and obtain reliable conclusions.

The results of the meta-analysis suggested negative effects of *H. pylori* on serum vitamin B<sub>12</sub>, folate, vitamin C and vitamin D levels. *Helicobacter pylori*-induced atrophic gastritis impairs stomach acidification and secretion functions and causes the malabsorption of nutrients. Vitamin B<sub>12</sub> plays indispensable roles in promoting the development and maturation of red blood cells and in maintaining normal hematopoietic functions (71). A deficiency in an intrinsic factor caused by *H. pylori* infection would aggravate vitamin B<sub>12</sub> absorption-related disorders. Folate participates in the metabolism of genetic materials and proteins, and it affects mammal reproduction (72). A folate deficiency may cause neural tube malformation, megaloblastic anemia, depression and malignant tumor formation (71, 73). A vitamin B<sub>12</sub> deficiency may cause a folate metabolic disorder and
aggravate the folate deficiency. Vitamin C and E, as vitamins with antioxidant effects, play roles in eradicating oxygen radicals and in maintaining the body’s steady state (74, 75). Vitamin C may also promote the formation of tetrahydrofolic acid (75, 76). A shortage of vitamin C aggravates a folate deficiency. Therefore, the effects of *H. pylori* infections on human vitamin levels are not independent, and there are interactions among the various vitamins.

Our study found that patients that had undergone successful *H. pylori* eradication had higher serum vitamin D levels than patients in which *H. pylori* eradication failed. We speculated that vitamin D is a protective factor for *H. pylori*. In terms of mechanism, a combination of vitamin D and the vitamin D receptor may activate immune responses and participate in the anti-*H. pylori* process (77, 78). Yang et al. (17) also performed a related meta-analysis, but they only included three papers; consequently, the limited number of studies was not suitable for a pooled analysis. Our work included five relevant studies, including the study of Shafrir et al. (56) that covered data from more than 70,000 patients, which improved the reliability of the results. Apart from those on vitamin D, we did not find any studies that explored the relationships between other vitamins and the success of *H. pylori* eradication. Further research is needed to investigate the differences in other vitamin levels between patients with successful *H. pylori* eradication compared with those that failed.

This work revealed that serum vitamin B12 levels in patients after *H. pylori* eradication were significantly higher than those before *H. pylori* eradication. We reaffirmed the adverse effects of *H. pylori* on vitamin B12. For patients with a vitamin
B12 deficiency, aggressive H. pylori eradication therapy and additional vitamin B12 supplementation are necessary. For vitamin C, this meta-analysis did not produce a statistically significant result. However, the four estimates included in this study were from only two reports. Thus, there were too few relevant studies included, and the results should be interpreted cautiously.

The pooled results of the RCTs showed that both the per-protocol analysis and intention-to-treat analysis suggested that antioxidant vitamin supplementation could improve the eradication rates of H. pylori of standard regimens. In the selection of standard eradication regimens, Kockar et al. (67), Chuang et al. (68) and Zojaei et al. (16) chose vitamin C for additional supplementation, whereas other studies used vitamin C combined with vitamin E for supplementation. Although both vitamin C and vitamin E have antioxidant effects, we were still concerned that inconsistent supplementation selection would introduce bias. Moreover, except the studies of Sezikli et al. (15) and Demirci et al. (66), which adopted the quadruple regimen, other studies adopted the triple regimen. The choices of eradication therapy (triple or quadruple therapy) also could introduce bias. We used a meta-regression to eliminate the possible bias caused by different kinds of vitamin supplementation and different eradication schemes from the statistical field, and then, we conducted pooled analyses to improve the reliability of the results. The meta-analysis of Li et al. (19) found that supplementation with antioxidant vitamins did not benefit the eradication rate. However, this analysis included only three studies. Although Ochoa et al. (79) also conducted a meta-analysis on a similar topic in 2018, there were errors and shortcomings in the data extraction and chart construction. Moreover, the above two articles only performed intention-to-treat analyses of the data. We believed that the per-protocol analysis should not be ignored in the RCTs. The meta-analysis of Yang-Ou et al. (80) focused on the effects of antioxidants on H. pylori eradication. However, this study combined antioxidant vitamins with other antioxidant (curcumin and cranberry). Our study incorporated intention-to-treat and per-protocol analyses and updated the results. Interestingly, the cohort study by Kaboli et al. (69) found that vitamin C supplementation reduces the dosage of clarithromycin. Li et al. (81) revealed that the H. pylori treatment and vitamin supplementation reduced the incidence of gastric cancer.

A complete and comprehensive search is necessary to find all the published data relevant to the meta-analysis. In addition to the usual English databases (Medline, Web of Science and Embase), we also searched for Chinese studies using the Chinese Biomedical Database and for conference papers using the Cambridge Scientific Abstracts databases to identify suitable studies. Although the search results did not ultimately expand the number of relevant articles, the processes remained essential. Unfortunately, it was hard for us to search manuscript with other languages, which might have resulted in the exclusion of some suitable articles. At last, we identified 48 relevant articles in this meta-analysis. We ran a better selection of studies and the number of included articles was larger than previous meta-analysis.

The heterogeneity of the selected studies cannot be ignored. The different methods of vitamin detection may result in variations in the results. The different areas, different numbers of patients enrolled in each study, some positive data and large amounts of included estimates may have increased the heterogeneity. Because the number of included studies was limited, it was hard for us to conduct subgroup analyses to reduce the influence of heterogeneity. Nevertheless, we performed meta-regression and sensitivity analyses to reduce the effects of heterogeneity on the credibility of the conclusions. The results of the meta-regression helped us eliminate the possibility that heterogeneity was a result of different eradication methods or vitamin supplementations. The sensitivity analyses helped us eliminate the influence of some positive data on the conclusions. In addition to the meta-regression and sensitivity analyses, we used random-effects models to establish relationships among the variables with high heterogeneity. Therefore, this work is appropriate to provide evidence, but the conclusions should be interpreted cautiously.

The study provides a comprehensive analysis of the interrelations between H. pylori and the most common vitamins from four aspects. Nevertheless, this work has some shortcomings. First, Turkish scholars have done a great deal of work on the effects of vitamin supplementation on H. pylori eradication. However, the contributions of researchers from other countries are limited. Different races and dietary habits may influence the results. It is difficult to conduct subgroup analyses to address this issue. Second, quadruple anti-H. pylori therapy is recommended currently. Some studies included in the work still used the triple approach. Moreover, the dose of vitamin supplementation is also controversial. Thus, the results should be interpreted cautiously. More RCTs with large samples focusing on the effects of vitamin supplementation on H. pylori eradication are needed to confirm our results and explore the appropriate dose.

CONCLUSIONS

In summary, this meta-analysis demonstrates that H. pylori infections can reduce the serum levels of several vitamins. The eradication of H. pylori rescues its adverse effects. Antioxidant vitamin supplementation may increase the rate of H. pylori eradication. Aggressive H. pylori eradication therapy is necessary, and the advantages of multivitamin supplementation for H. pylori-positive patients outweigh the disadvantages.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding authors.

AUTHOR CONTRIBUTIONS

XC, WY, and XiL conceived and designed the study. XC and XuL acquired and analyzed the data. XC, YJ, MZ, YX, YW, and CL
interpreted the data and drafted the manuscript. XL and WY reviewed and corrected the manuscript. All authors approved the final version to be published.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fnut.2021.781333/full#supplementary-material

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