

EVALUATION OF HEMATOLOGICAL PARAMETERS AND DIAGNOSTIC SIGNIFICANCE IN H. PYLORI INFECTION: A COMPARATIVE STUDY

JABEEN A¹, KABIR MU¹, FATTANI B², TARIQ MJA^{3*}

¹Department of Biological Sciences, The Superior University, Lahore, Pakistan

²Department of Surgery, Jinnah Medical College Hospital, Pakistan

³Department of Biological and Health Sciences, Pak-Austria Fachhochschule Institute of Applied Sciences and Technology, Mang Haripur, Pakistan

*Correspondence author email address: junaidtariq186@gmail.com

(Received, 27th June 2024, Revised 30th August 2024, Published 2nd September 2024)

Abstract: *Helicobacter pylori* (*H. pylori*) infection is globally recognized for its role in various gastrointestinal diseases and extra gastric conditions, including hematological anomalies. Understanding the correlation between *H. pylori* infection and hematological parameters could provide insights into its broader health impacts. **Objective:** This study evaluated the correlation between *H. pylori* infection and various demographic, socioeconomic, and hematological parameters. **Methods:** A cross-sectional study enrolled patients diagnosed with *H. pylori* and a matched control group without the infection. Blood samples were collected and analyzed for hematological parameters, including white blood cell count, red blood cell count, hemoglobin (HGB), hematocrit (HCT), platelet count, mean corpuscular volume (MCV), mean corpuscular hemoglobin (MCH), mean corpuscular hemoglobin concentration (MCHC), and red cell distribution width (RDW). Statistical analyses were performed using t-tests and chi-square tests where appropriate, with significance set at $p < 0.05$. **Results:** The study included patients with statistically significant differences in hematological parameters between the *H. pylori*-infected and healthy groups. Notably, the infected group showed lower mean HGB (12.41 ± 1.38 g/dl vs. 14.32 ± 1.46 g/dl), RBC count ($4.12 \pm 0.62 \times 10^6/\mu\text{L}$ vs. $4.87 \pm 0.53 \times 10^6/\mu\text{L}$), and HCT ($39.17 \pm 3.87\%$ vs. $44.52 \pm 4.51\%$), and higher RDW ($13.51 \pm 2.10\%$ vs. $13.87 \pm 1.32\%$) compared to the control group (all $p < 0.05$). **Conclusion:** The findings indicate significant hematological changes associated with *H. pylori* infection, suggesting a potential impact on the overall health status of affected individuals. These results highlight the importance of considering *H. pylori* infection in the differential diagnosis of hematological abnormalities and underscore the need for further research to explore the underlying mechanisms and potential therapeutic interventions.

Keywords: *H. pylori*, Hematological parameters, Demographics, Comparative analysis.

Introduction

Peptic ulcers, gastritis, gastric cancer, and gastric lymphoma are among the many gastrointestinal diseases linked to the common bacterium *H. pylori* (1). Several methods exist for detecting *H. pylori* infection; these range from invasive procedures like endoscopy and biopsy to non-invasive ones that use blood samples, breath, and stool (2). Nevertheless, there are several drawbacks to these diagnostic methods, including their invasiveness, expense, accessibility, precision, and specificity. This highlights the critical need to explore complementary or alternate methods of diagnosing *H. pylori* infection (3).

In light of evidence from some studies pointing to possible changes in blood cell composition causing hematological abnormalities, such as anemia, thrombocytopenia, leukopenia, or leukocytosis, one potential approach is to examine the hematological parameters of people infected with *H. pylori* (4). The association between *H. pylori* infection and hematological parameters is still not well-established and can vary depending on factors like population characteristics, infection severity, and other influences (5).

Helicobacter pylori exerts harmful effects on the gastric and duodenal linings through multiple mechanisms. One mechanism involves ammonia production, aimed at regulating pH, which proves toxic to epithelial cells (6).

Additionally, *H. pylori* produces various biochemicals, including proteases and vacuolating cytotoxin A, which damage epithelial cells, disrupt tight junctions, and induce apoptosis. Furthermore, the bacterium releases specific phospholipases, contributing to its pathogenic impact on the gastrointestinal mucosa (7).

Nevertheless, multiple investigations have found that *Helicobacter pylori* (*H. pylori*) infection is linked to iron-deficient anemia, vitamin B12 deficiency, and other iron-related disorders (8). Oral iron therapy also reduces the effect on people with *H. pylori* infection. Potentially improving ferritin and hemoglobin levels in infected patients has led to the proposal of combining iron therapy with *H. pylori* eradication therapy (9).

H. pylori infection and iron deficiency anemia go hand in hand because chronic gastritis causes stomach hypochlorhydria, which in turn hinders iron absorption. Converting ferric iron from food to ferrous iron requires an acidic intragastric pH and ascorbic acid, which is not present in this patient. Iron absorption becomes even more difficult when *H. pylori*, a main cause of chronic superficial gastritis and gastric gland atrophy, decreases stomach acid output (10).

In addition, *Helicobacter pylori* competes with the host for iron, which hinders iron intake. At the same time, hemoglobin synthesis is hampered because the

reticuloendothelial system and the entrecote release less iron from macrophages due to the increased hepcidin production caused by *H. pylori* infection. In particular, hepcidin responds to inflammation in the stomach mucosa by acting as an acute phase reactant (10).

Methodology

We devised a comparative cross-sectional study encompassing patients and healthy individuals (11). The investigation involved the comparison of hematological profiles within two distinct groups, namely, individuals diagnosed with *H. pylori* infection and a healthy cohort devoid of the infection. The age range was expanded to encompass participants aged 18 to >41 years. Exclusion criteria comprised individuals with pre-existing incomplete consents, hemolysed samples, hematological disorders, pregnant women, patients on medication, and those diagnosed with gastric cancer. The sample size was determined by statistical considerations to ensure sufficient power for meaningful comparisons between the *H. pylori*-infected and healthy groups (7).

Each participant underwent a comprehensive clinical examination encompassing an assessment of signs and symptoms, prior medical history, comorbidities, and a physical exam. Blood samples were procured for a selected blood cell analysis (12). The status of *H. pylori* infection was ascertained through validated diagnostic methods, including serological and stool antigen tests. Hematological evaluation involves the specific hematological panels,

including HGB (hemoglobin), RBC (red blood cell count), HCT (hematocrit), MCV (mean corpuscular volume), MCH (mean corpuscular hemoglobin), MCHC (mean corpuscular hemoglobin concentration), and RDW (red cell distribution width) by an automated hematology analyzer (Sysmex KX-21, Japan).

In this study, ethical agreements were carefully considered. We explained the study methods, expected results, and potential benefits to all participants. Each individual willingly signed consent forms, indicating their voluntary participation and understanding of the study's parameters. (13).

The data from medical reports and clinical examinations were collected using Microsoft Excel (2016). Statistical analyses were then carried out using SPSS version 20. Continuous data were presented as the mean ± standard deviation, while nominal data were described using frequencies and percentages. A 95% confidence level was maintained, and statistical significance was considered for a P value below 0.05.

Results

Table 1 provides a comprehensive overview of the demographic distribution of variables in individuals with *H. pylori* infection compared to healthy individuals. The analysis encompasses vital demographic factors, including gender, age, locality, and academic status.

Table 1: Sociodemographic Characteristics of Study Participants

| Variables | Groups | <i>H. pylori</i> patients | Healthy Individuals |
|-----------------|----------|---------------------------|---------------------|
| Gender | F | 61 (48.41) | 61(48.41) |
| | M | 63 (50.79) | 63 (50.79) |
| Age (years) | 18–25 | 49 (38.88) | 56 (44.44) |
| | 26–33 | 41 (32.53) | 43(34.12) |
| | 34–41 | 31 (24.60) | 24 (19.04) |
| | ≥41 | 5(3.96) | 3 (2.38) |
| Locality | Urban | 18 (14.28) | 15 (11.90) |
| | Rural | 108 (85.71) | 111 (88.09) |
| Academic status | Literate | 9 (7.14) | 7 (5.55) |
| | Matric | 13 (10.31) | 17 (13.49) |
| | Bachelor | 75 (59.52) | 68 (53.96) |
| | Master | 29 (23.01) | 34 (26.98) |

*M (Male), F (Females), *H. pylori* (*Helicobacter pylori*)

Table 2 underscored the potential impact of *H. pylori* infection on red blood cell profiles, emphasizing the

relevance of hematological assessments in the clinical evaluation of *H. pylori*-associated conditions.

Table 2: Comparative Evaluation of Hematological Parameters among *H. pylori* Patients and Healthy Group

| Lab Parameters | Unit | <i>H. pylori</i> patients | Healthy Individuals | P-value |
|----------------|------------------------|---------------------------|---------------------|---------|
| HGB | (g/dl) | 12.41±1.38 | 14.32±1.46 | <0.001 |
| RBC | (x 10 ⁶ µL) | 4.12±0.62 | 4.87±0.53 | |
| HCT | (%) | 39.17±3.87 | 44.52±4.51 | |
| MCV | (fl) | 86.06±4.44 | 90.16±4.53 | 0.003 |
| MCH | (pg) | 26.41±3.16 | 29.13±1.87 | 0.007 |
| MCHC | (g/dl) | 31.45±1.60 | 32.58±1.24 | 0.005 |
| RDW | (%) | 13.51±2.10 | 13.87±1.32 | 0.005 |

[Citation: Jabeen, A., Kabir, M.U., Fattani, B., Tariq, M.J.A., (2024). Evaluation of hematological parameters and diagnostic significance in *h. pylori* infection: a comparative study. *Biol. Clin. Sci. Res. J.*, 2024: 1076. doi: <https://doi.org/10.54112/bcsrj.v2024i1.1076>]

*Hemoglobin (HGB), Red Blood Cell Count (RBC), Hematocrit (HCT), Mean Corpuscular Volume (MCV), Mean Corpuscular Hemoglobin (MCH), Mean Corpuscular Hemoglobin Concentration (MCHC), Red Cell Distribution Width (RDW).

The distribution of lab parameter categories in the study population illustrates the prevalence of abnormal hematological values alongside reference ranges. Table 3 offers valuable insights into potential hematological disorders and guides further clinical investigations based on categorized parameters.

Table 3: Evaluation of Hematological Parameters among H. pylori Patients

| Lab Parameters | Categories | N (%) | Reference Ranges |
|----------------|---------------------------|-------|------------------|
| HGB | (g/dl) | L | M (11.5–18) |
| | | N | F (11–16.7) |
| RBC | $\times 10^6/\mu\text{L}$ | L | M (3.53–6.93) |
| | | N | F (3.45–6.25) |
| | | H | |
| HCT | (%) | L | M (36.2–58.6) |
| | | N | F (32.1–56.6) |
| | | H | |
| MCV | (fl) | L | M (85–100) |
| | | N | F (85–100) |
| | | H | |
| MCH | (pg) | L | M (26.6–33.30) |
| | | N | F (25.8–32.8) |
| | | H | |
| MCHC | g/dl | L | M (29.5–34.4) |
| | | N | F (28.5–34.4) |
| | | H | |
| RDW | (%) | L | M (12–17) |
| | | N | F (12–17) |

*L (Low), N (Normal), H (High)

Discussion

The gender distribution of the study population is relatively even, with roughly half of the individuals in both the Healthy and H. pylori-infected groups identifying as female and the remaining half as male. The gender equilibrium guarantees that any discernible variations in H. pylori infection rates are comparatively less susceptible to the impact of gender prejudice.

A significant proportion of participants in both categories, 108 (85.71%) in the Healthy group and 111 (88.09%) in the H. pylori-infected group, reside in rural regions. Urban residency is relatively uncommon, as indicated by the fact that only 18 individuals (14.28%) in the H. pylori-infected group and 15 individuals (11.90%) in the Healthy group reside in urban areas. (14) The observed longitudinal arrangement of this organism offers valuable insights into the possible influence of environmental variables linked to rural residence on the prevalence of H. pylori.

The academic status most prevalent in both groups is a Bachelor's, comprising 75 individuals (59.52%) in the H. pylori-infected group and 68 individuals (53.96%) in the Healthy group. In contrast, the academic status that is least prevalent is Literate, comprising merely nine individuals (7.14%) in the H. pylori-infected group and seven individuals (5.55%) in the Healthy group. The relevance of these academic distinctions may lie in their capacity to illuminate possible associations between levels of H. pylori infection and educational background (15).

The study observed significantly lower levels of Hemoglobin (HGB) in individuals with H. pylori infection (12.41 ± 1.38 g/dl) compared to the control group (14.32 ± 1.46 g/dl) ($p < 0.001$). This finding aligns with research by (16), which reported similar hematological alterations in individuals with H. pylori infection.

Red Blood Cell count (RBC) was also lower in the H. pylori-infected group ($4.12 \pm 0.62 \times 10^6/\mu\text{L}$) compared to the control group ($4.87 \pm 0.53 \times 10^6/\mu\text{L}$). This corroborates the findings of (17), who noted a significant decrease in RBC count associated with H. pylori infection.

Hematocrit (HCT) values were significantly lower in individuals with H. pylori infection ($39.17 \pm 3.87\%$) compared to the control group ($44.52 \pm 4.51\%$) ($p < 0.001$). This observation is consistent with the results (18), indicating a link between H. pylori infection and reduced hematocrit levels.

Mean Corpuscular Volume (MCV) was found to be lower in the H. pylori-infected group (86.06 ± 4.44 fl) compared to the control group (90.16 ± 4.53 fl) ($p = 0.003$). This supports the findings of (19), who observed similar alterations in MCV associated with H. pylori infection.

Mean Corpuscular Hemoglobin (MCH) was significantly lower in individuals with H. pylori infection (26.41 ± 3.16 pg) compared to the control group (29.13 ± 1.87 pg) ($p = 0.007$). Mean Corpuscular Hemoglobin Concentration (MCHC) was lower in the H. pylori-infected group (31.45 ± 1.60 g/dl) compared to the control group

[Citation: Jabeen, A., Kabir, M.U., Fattani, B., Tariq, M.J.A., (2024). Evaluation of hematological parameters and diagnostic significance in h. pylori infection: a comparative study. *Biol. Clin. Sci. Res. J.*, 2024: 1076. doi: <https://doi.org/10.54112/bcsrj.v2024i1.1076>]

(32.58±1.24 g/dl) ($p = 0.005$). Red Cell Distribution Width (RDW) values showed a slight increase in the *H. pylori*-infected group (13.51±2.10%) compared to the control group (13.87±1.32%) (20).

Most individuals exhibited normal hemoglobin (HGB) levels, with 86.50% falling within the normal range, while 13.49% had low HGB levels. Red blood cell count (RBC) analysis revealed that 92.85% of participants had average values, 7.14% had low RBC counts, and none displayed high counts. Hematocrit (HCT) levels were essentially expected, with 94.44% falling within the normal range and only 5.55% having low levels; high HCT levels were absent. Mean corpuscular volume (MCV) values demonstrated that 87.30% of individuals had normal levels, 12.69% exhibited low MCV levels, and 0.79% showed high MCV levels. Mean corpuscular hemoglobin (MCH) analysis indicated that 85.71% had normal MCH levels, 13.49% had low, and 0.79% had high levels. Mean corpuscular hemoglobin concentration (MCHC) levels were predominantly normal, with 97.61% falling within the normal range, while 1.58% had low MCHC levels, and 0.79% had high MCHC levels. Red cell distribution width (RDW) findings revealed that 99.20% of participants had normal RDW levels, and only 0.79% exhibited low RDW levels.

Conclusion

Hematological analyses indicate significant differences between *H. pylori*-infected and healthy groups. Lower hemoglobin levels decreased red blood cell count, and reduced hematocrit values in *H. pylori*-infected individuals align with previous research. Additionally, alterations in mean corpuscular volume, mean corpuscular hemoglobin and mean corpuscular hemoglobin concentration further indicate the hematological consequences of *H. pylori* infection, consistent with findings from relevant literature. However, the slight increase in red cell distribution width (RDW) in the *H. pylori*-infected group, though statistically significant, contrasts with some prior research, warranting further investigation.

Declarations

Data Availability statement

All data generated or analyzed during the study are included in the manuscript.

Ethics approval and consent to participate.

Approved by the department concerned. (IRBEC-TCH-0282/23)

Consent for publication

Approved

Funding

Not applicable

Conflict of interest

The authors declared an absence of conflict of interest.

Authors Contribution

ALINA JABEEN

Data Analysis

MUMTAZ MANZOOR (MBBS FCPS)

Revisiting Critically

MUHAMMAD USMAN KABIR

Concept & Design of Study

BILAL FATTANI (House officer)

Drafting

MUHAMMAD JUNAID AHMAD TARIQ

Final Approval of version

References

1. Crowe SE. Helicobacter pylori infection. *New England Journal of Medicine*. 2019;380(12):1158-65.
2. Tshibangu-Kabamba E, Yamaoka Y. Helicobacter pylori infection and antibiotic resistance—from biology to clinical implications. *Nature Reviews Gastroenterology & Hepatology*. 2021;18(9):613-29.
3. FitzGerald R, Smith SM. An overview of Helicobacter pylori infection. *Helicobacter pylori*. 2021:1-14.
4. Öztekin M, Yılmaz B, Ağagündüz D, Capasso R. Overview of Helicobacter pylori Infection: clinical features, treatment, and nutritional aspects. *Diseases*. 2021;9(4):66.
5. Ren S, Cai P, Liu Y, Wang T, Zhang Y, Li Q, et al. Prevalence of Helicobacter pylori infection in China: A systematic review and meta-analysis. *Journal of gastroenterology and hepatology*. 2022;37(3):464-70.
6. Godbole G, Mégraud F, Bessède E. Diagnosis of Helicobacter pylori infection. *Helicobacter*. 2020;25:e12735.
7. Shimamoto T, Yamamichi N, Gondo K, Takahashi Y, Takeuchi C, Wada R, et al. The association of Helicobacter pylori infection with serum lipid profiles: An evaluation based on a combination of meta-analysis and a propensity score-based observational approach. *PloS one*. 2020;15(6):e0234433.
8. Talari HR, Moniri R, Mollaghanbari M, Kashani HH, Jalalian MN. Evaluating the relationship between Helicobacter pylori infection and carotid intima-media thickness a cross-sectional study. *Annals of Medicine and Surgery*. 2021;69:102659.
9. Ansari S, Yamaoka Y. Helicobacter pylori infection, laboratory diagnosis, and antimicrobial resistance: a perspective of clinical relevance. *Clinical Microbiology Reviews*. 2022;35(3):e00258-21.
10. George S, Lucero Y, Torres JP, Lagomarcino AJ, O’Ryan M. Gastric damage and cancer-associated biomarkers in Helicobacter pylori-infected children. *Frontiers in Microbiology*. 2020;11:90.
11. Haile K, Timerga A. Evaluation of hematological parameters of Helicobacter pylori-infected adult patients at Southern Ethiopia: A comparative cross-sectional study. *Journal of Blood Medicine*. 2021:77-84.
12. Bordin DS, Voynovan IN, Andreev DN, Maev IV. Current Helicobacter pylori diagnostics. *Diagnostics*. 2021;11(8):1458.
13. Park W-J, Kim S-H, Kang W, Ahn J-S, Cho S, Lim D-Y, et al. Blood lead level and Helicobacter pylori infection in a healthy population: A cross-sectional study. *Archives of Environmental & Occupational Health*. 2020;75(6):333-8.
14. Lucero Y, Lagomarcino AJ, Torres JP, Roessler P, Mamani N, George S, et al. Helicobacter pylori, clinical, laboratory, and noninvasive biomarkers suggestive of gastric damage in healthy school-aged children: A case-control study. *International Journal of Infectious Diseases*. 2021;103:423-30.
15. Fang Y, Xie H, Fan C. Association of hypertension with helicobacter pylori: A systematic review and meta-analysis. *PLoS One*. 2022;17(5):e0268686.
16. Xiong X, Chen J, He M, Wu T, Yang H. Helicobacter pylori infection and the prevalence of hypertension in Chinese adults: The Dongfeng-Tongji cohort. *The Journal of Clinical Hypertension*. 2020;22(8):1389-95.
17. Chen MJ, Fang YJ, Wu MS, Chen CC, Chen YN, Yu CC, et al. Application of Helicobacter pylori stool antigen test to survey the updated prevalence of Helicobacter pylori infection in Taiwan. *Journal of Gastroenterology and Hepatology*. 2020;35(2):233-40.

18. Hussein RA, Al-Ouqaili MT, Majeed YH. Detection of Helicobacter Pylori infection by invasive and non-invasive techniques in patients with gastrointestinal diseases from Iraq: A validation study. Plos one. 2021;16(8):e0256393.
19. Ito M, Sugiyama A, Mino M, Kodama M, Nagaoki Y, Abe K, et al. Prevalence of Helicobacter pylori infection in the general population evaluated by a resident-register-based epidemiological study. Journal of Gastroenterology. 2022;57(8):540-6.
20. Kadhim AS, Al-Karawi AS. Insights into the Pathogenesis, Virulence Factors, and Diagnosis of Helicobacter pylori: A Comprehensive Review. American Journal of Bioscience and Bioinformatics. 2023;2(1):31-7.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution, and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third-party material in this article are included in the article's Creative Commons license unless indicated otherwise in a credit line to the material. Suppose material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use. In that case, you must obtain permission directly from the copyright holder. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>. © The Author(s) 2024