Original Article

Hepcidin and iron status among pregnant women in Bangladesh

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Although hepcidin, a recently discovered peptide hormone, is considered a major regulator of iron metabolism and anemia in chronic inflammation, its role in anemia during pregnancy has not been characterized. Our objective was to characterize the role of hepcidin in anemia during pregnancy. We examined the relationships between urinary hepcidin, iron status indicators, hemoglobin, erythropoietin, alpha-1 acid glycoprotein, and C-reactive protein in a cross-sectional study conducted among 149 pregnant rural Bangladeshi women with biospecimens obtained during home visits. Urinary hepcidin was measured using surface-enhanced laser desorption/ionization time-of-flight mass spectrometry. Urinary hepcidin, as log(intensity per mmol/L creatinine), was correlated with log ferritin ($r = 0.33, p < 0.001$), the transferrin receptor index ($r = -0.22, p = 0.007$), and log alpha-1 acid glycoprotein ($r = 0.20, p = 0.01$), but not hemoglobin ($r = 0.07, p = 0.40$), log transferrin receptor ($r = -0.07, p = 0.41$), log erythropoietin ($r = -0.01, p = 0.88$) or log C-reactive protein ($r = 0.06, p = 0.48$). The strength of the relationship between hepcidin and ferritin was maintained in multiple linear regression analyses after enhancing the sample with data from women selected for low iron stores ($n = 41$). Among pregnant women in a community-based study in rural Bangladesh, urinary hepcidin levels were related to iron status and AGP but not hemoglobin, erythropoietin, or C-reactive protein.

Key Words: anemia, hepcidin, inflammation, iron, pregnancy

INTRODUCTION

Anemia is common during pregnancy and is associated with higher perinatal and maternal morbidity and mortality in developing countries.1 Iron deficiency accounts for a large proportion of the anemia among pregnant women.2 Hepcidin, a 25 amino acid peptide, is considered a major regulator of iron metabolism and the anemia of chronic inflammation.3 Hepcidin is found in human plasma and urine,4,5 and it is synthesized primarily in the liver.4,6 It regulates iron metabolism by inhibiting duodenal iron absorption at the level of intestinal epithelium,7 and by affecting mobilization of iron from liver and spleen.8 Hepcidin binds to the iron exporter, ferroportin, inducing its internalization and degradation.9 Ferroportin is the only mammalian iron exporter identified to date and is necessary for materno-fetal iron transfer and iron efflux from duodenal enterocytes, macrophages, and hepatocytes.10

Hepcidin is expressed in an iron-replete state,11 and during iron overload.3 Urinary hepcidin levels seem to change rapidly in response to changes in iron status. For example, in normal human volunteers, urinary hepcidin levels increased up to 15-fold within 24 h after 65 mg of oral iron supplementation.11 In iron deficiency anemia, urinary hepcidin levels are low to undetectable.12 Hepcidin expression is inappropriately low in most forms of hereditary hemochromatosis.13 One particularly severe form of hemochromatosis results from mutations in the gene encoding hepcidin itself.14 The signal for the iron-replete state is still unidentified.8

Despite the central role of hepcidin in the metabolism of iron, limited data are available that link hepcidin to measures of iron status, inflammation, and anemia in pregnant women. Our goal was to examine the relationships between urinary hepcidin and ferritin, inflammation, and anemia. To address this goal, we measured urinary hepcidin and indicators of iron status and inflammation in a community-based study of women in Bangladesh at the time of pregnancy assessment, typically in the first trimester. Anemia is considered a population-wide public health problem in Bangladesh, and is particularly prevalent during pregnancy.15,16

MATERIALS AND METHODS

Study design

The study subjects consisted of a total of 190 pregnant women from northwestern Bangladesh from whom blood
and urine samples were collected in the home following pregnancy confirmation and at the time of enrollment into a larger clinical trial of micronutrient supplementation. The women represented a systematic subsample of nearly 10% (n = 149 women) of over 1500 women that contributed a blood sample. Subsequent purposive sampling of women with iron deficiency (n = 41) was done to enhance the original sample with more iron deficient subjects. This was done in order to extend the range of ferritin concentrations to lower values in the sample of women, thus allowing for the elucidation of the relationship between hepcidin and iron status indicators across a wider spectrum of iron status than that observed in the original sample. Informed consent was obtained from all subjects. The study protocol was approved by the Committee on Human Research of the Johns Hopkins Bloomberg School of Public Health and the Bangladesh Medical Research Council.

Hemoglobin (Hb) was measured from venous blood collected during the home visit using a B-Hemoglobin Analyzer (HemoCue Inc, Lake Forest, CA). Plasma and spot urine samples were aliquoted and stored in liquid nitrogen until analysis at the Johns Hopkins University and Medical Institutions for iron status indicators, C-reactive protein (CRP), alpha-1 acid glycoprotein (AGP), and urinary hepcidin.

Urinary hepcidin was measured in triplicate using surface-enhanced laser desorption/ionization time-of-flight mass spectrometry (SELDI-TOF MS) using hydrophilic normal phase chips (ProteinChip® NP20, Ciphergen Biosystems, Fremont, CA),

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\text{using Ciphergen ProteinChip Software (version 3.2.0), after calibration with hepcidin standard (Peptide Institute, Osaka, Japan) baseline subtraction and adjustment. A peak was recorded for hepcidin at the characteristic m/v of 2790 when the signal-to-noise ratio was } > 3:1. \]

Urinary creatinine was measured using a commercial analyzer (HemoCue Inc, Minoh-shi, Japan) baseline subtraction and adjustment. All data were analyzed in SAS v 9.1 (SAS Institute Inc, Cary, NC) and R (v. 2.4.1).

RESULTS

Subject characteristics for the entire sample are shown in Table 1. Nearly half the women were primiparous (48%), with 37% of parity 1-2 and 15% of parity ≥3. Women in the original sample were enrolled earlier in pregnancy than those who were selected to enhance the sample (11.1 versus 15.7 weeks gestation, \( p < 0.001 \)). Among the women selected to enhance the sample, all iron status indicators were consistent with greater degree of iron deficiency, but CRP, weight, and parity did not differ by group. Among all participants, 11% of women had CRP values > 3 mg/L, 13% of women had AGP values above 110 mg/L, \(^{21}\) and there was a moderate correlation between AGP and CRP (\( r = 0.17 \), \( p = 0.03 \)).

Frequency distributions for each iron status variable and relationships among variables are shown with scatterplots and correlation coefficients for the original sample of women in Figure 1. Urinary hepcidin was correlated with both ferritin (\( p < 0.001 \)) and TfR index (\( p = \)

| Table 1. Characteristics of pregnant Bangladeshi women (n = 190) |
|--------------|--------------|--------------|--------------|
|              | Mean         | Standard Deviation | Median       | Inter-quartile Range |
| Age (y) †    | 21.9 (5.9)   | 20.0           | (17.5-25.0)  |
| Height (cm)  | 149 (5)      | 149            | (138-164)    |
| Weight (kg)  | 42.5 (5.1)   | 41.8           | (28.9-59.9)  |
| Gestational age (wk) † | 12.0 (8.4) | 11.0           | (8.0-14.0)   |
| Haptoglobin (μg/mL) | 4.55 (5.63) | 2.35           | (0.51-7.22)  |
| Hemoglobin (g/L) | 117 (14)    | 117            | (109-126)    |
| Ferritin (μg/L) | 86 (71)     | 74             | (36-115)     |
| Transferrin receptor (μg/mL) | 4.2 (1.5)  | 3.9            | (3.2-4.8)    |
| Transferrin receptor index (TfR/log ferritin) | 0.12 (0.19) | 0.06          | (0.03-0.11)  |
| Erythropoietin (mIU/mL) † | 17.6 (17.0) | 13.5           | (10.4-19.6)  |
| C-reactive protein (mg/L) † | 1.60 (3.90) | 0.35           | (0.15-1.30)  |
| Alpha-1 acid glycoprotein (mg/dL) | 77.6 (32.8) | 72.6          | (54.4-95.0)  |

† Missing data for n = 2, 4, 5, and 6 individuals, respectively.
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There were no significant correlations between urinary hepcidin and soluble TfR, Hb, or EPO, although EPO was strongly associated with all other iron status indicators. CRP was not correlated with urinary hepcidin (r = 0.06, p = 0.48), although AGP was (r = 0.20, p = 0.01). C-reactive protein was inversely correlated with hemoglobin concentration (r = -0.21, p < 0.01) but not with other indicators of iron status. Conversely, AGP was correlated with TfR (r = 0.23, p = 0.005).

Multiple linear regression including CRP demonstrated that urinary hepcidin was positively associated with ferritin (p < 0.001) and, to a lesser extent, inversely associated with soluble TfR (p = 0.04) (Table 2). However, an overall test comparing the full model with the regression of hepcidin against ferritin alone did not reject the more parsimonious model (p = 0.18), indicating that iron stores were the overwhelming predictor of urinary hepcidin concentrations. Hepcidin was not related to EPO, Hb, CRP, or gestational age to a significant degree in the multiple linear regression model that included CRP. In the regression model that included AGP, the coefficients relating each variable to hepcidin were similar and the association between ferritin and hepcidin remained. However, the association between TfR and hepcidin was strengthened (p = 0.009), and the association between AGP and hepcidin was significant (p = 0.03). A comparison of the full model with the regression of hepcidin against ferritin alone suggested that the full model including AGP better explained urinary hepcidin concentrations (p = 0.02).

Figure 1. Pairwise scatterplots and correlations among indicators of iron metabolism among 149 Bangladeshi women. On the diagonal, the marginal distributions are shown as histograms. In the lower left triangle of the figure, we indicate the respective correlation coefficients, confidence intervals, and p-values. P-values less than 0.05 are shown in bold face. For clarity, and since correlations are scale independent, we omitted the axis labels. Urinary hepcidin, intensity/mMol creatinine, is abbreviated Hep; ferritin, μg/L, is abbreviated ferr.

Table 2. Multiple linear regression analysis† of factors associated with hepcidin (intensity per mmol/L creatinine) among Bangladeshi women

<table>
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<tr>
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<th>Model 1</th>
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<th>Model 2</th>
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<td>β</td>
<td>SE</td>
<td>p</td>
<td>β</td>
<td>SE</td>
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<tr>
<td>Ferritin (μg/L)</td>
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<td>0.13</td>
<td>&lt;0.001</td>
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<td>0.13</td>
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<td>Transferrin Receptor (μg/mL)</td>
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<td>0.04</td>
<td>-0.93</td>
<td>0.35</td>
</tr>
<tr>
<td>Erythropoietin (mIU/mL)</td>
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<td>0.22</td>
<td>0.09</td>
<td>0.35</td>
<td>0.21</td>
</tr>
<tr>
<td>Hemoglobin (g/L)</td>
<td>0.013</td>
<td>0.008</td>
<td>0.13</td>
<td>0.014</td>
<td>0.008</td>
</tr>
<tr>
<td>Gestational age (wk)</td>
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<td>0.022</td>
<td>0.64</td>
<td>0.006</td>
<td>0.022</td>
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<tr>
<td>C-reactive protein (mg/L)</td>
<td>0.052</td>
<td>0.072</td>
<td>0.48</td>
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<tr>
<td>Alpha-1 acid glycoprotein (mg/dL)</td>
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<td>---</td>
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<td>0.225</td>
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† Model 1: n = 179; F(6, 172) = 18.4, p < 0.001; R² = 0.39; Model 2: n = 181, F(6, 174) = 21.1, p < 0.001, R² = 0.42. All variables except hemoglobin and gestational age expressed as log(x)
release from splenic macrophages and the liver. Thereby reducing iron absorption in the gut and limiting iron availability for erythropoiesis, hypoxia, and inflammation. Elevated hepcidin levels were not associated with Hb concentrations among pregnant women dwelling in a rural community of Bangladesh. The four main factors that are thought to be involved in the regulation of hepcidin are iron status, metabolism and their relationships with hepcidin in a free-living population at risk of iron deficiency and anemia. Among pregnant women in Bangladesh, urinary hepcidin levels were linearly related with iron stores over a wide range of serum ferritin concentrations. To a lesser degree, tissue iron deficiency (elevated TIR) was associated with lower hepcidin concentrations. However, urinary hepcidin levels were not associated with Hb concentrations and the signal for erythropoietic activity (EPO), and the association of hepcidin with inflammatory markers was modest. It is noteworthy that women in this study were in the first trimester of pregnancy, when the demand of the body for iron is relatively low due to the cessation of menstruation. Iron requirements increase through the second and third trimesters of pregnancy to support the expansion of the red blood cell mass and tissue development of the placenta and fetus. Future work is needed to characterize changes in hepcidin in relation to iron status as pregnancy progresses and iron deficiency becomes more acute.

It is also notable that, while mild infections are common during pregnancy in this area of Bangladesh, among these community-dwelling pregnant women, CRP and AGP were not commonly elevated. Thus, the ranges of values in inflammatory markers against which to relate hepcidin were limited compared to other studies. Interleukin-6 (IL-6), which was not measured here but is involved in the acute phase response, is thought to be directly involved in the upregulation of hepcidin. However, the relationship between IL-6 and hepcidin has mostly been studied previously in subjects with more severe inflammation than what was apparent here; that is, in subjects with acute sepsis, and in healthy volunteers challenged with IL-6 or lipopolysaccharide. Although hepcidin is thought to be a major regulator of the anemia of chronic inflammation, such a role had yet to be shown definitively in a population-based study of community-dwelling adults. In this setting, AGP but not CRP showed a modest association with urinary hepcidin. The differential findings between AGP and CRP suggest that the choice of inflammatory marker is crucial for the interpretation of iron status indicators in population-based studies and may be explained by the fact that CRP rises and falls more rapidly in response to infection and inflammation than AGP.

The findings from this study are consistent with observations of Nemeth and colleagues in which serum ferritin was significantly correlated with urinary hepcidin in a mixed sample of study subjects that consisted of patients with anemia of inflammation, compensated hereditary hemochromatosis, iron overload, iron deficiency anemia, and healthy donors. Other studies of hepcidin specific to pregnancy include a rat model in which liver hepcidin expression declined throughout pregnancy as iron stores declined. The decline in liver hepcidin expression was associated with increases in duodenal iron transport proteins DMT1, dcytb, and Ireg1, implying an association of declining hepcidin with increased iron requirements and enhanced iron absorption.

Although there is considerable interest in evaluating the role of hepcidin in iron metabolism in a variety of population groups, currently the techniques for assessing bioactive hepcidin are not widely available and appear to be limited to mass spectrometric methods such as that utilized in this study. The relationship between plasma prohepcidin hormone has been explored using a commercial assay, and no association of iron or anemia status with plasma prohepcidin was demonstrated, suggesting that the commercial assay for plasma prohepcidin may not identify the active form of the hormone.

This study provides insight into the role of hepcidin in iron deficiency associated with pregnancy. Hepcidin is likely to be a key regulator of iron metabolism during pregnancy, and this study provides strong evidence that iron status in particular influences hepcidin concentrations among pregnant women of Bangladesh. Future studies are needed with larger sample sizes and longitudinal study designs to characterize changes in hepcidin.

Figure 2. Scatterplot and linear relationship between hepcidin (intensity/mmol creatinine) and ferritin (μg/L), both expressed on a log scale, among 190 Bangladeshi women. The relationship between ferritin and hepcidin was statistically significant (P < 0.001). The darker band is a confidence band for the regression line, while the lighter gray band represents pointwise prediction intervals for future observations, based on the relationship observed in this study.

DISCUSSION

This study shows that iron status is the primary determinant of urinary hepcidin concentrations among pregnant women dwelling in a rural community of Bangladesh. To our knowledge, this is the first study to report urinary hepcidin levels in pregnancy, and it provides insights into factors involved in the regulation of iron metabolism during pregnancy. The four main factors that are thought to be involved in the regulation of hepcidin are iron status, metabolism and their relationships with hepcidin in a free-living population at risk of iron deficiency and anemia. Among pregnant women in Bangladesh, urinary hepcidin levels were linearly related with iron stores over a wide range of serum ferritin concentrations. To a lesser degree, tissue iron deficiency (elevated TIR) was associated with lower hepcidin concentrations. However, urinary hepcidin levels were not associated with Hb concentrations and the signal for erythropoietic activity (EPO), and the association of hepcidin with inflammatory markers was modest.

Based on the slope of the relationship between hepcidin and ferritin (Figure 2), a one unit increase in log plasma ferritin resulted in a 1.20 unit increase in log hepcidin intensity, and 36% of the variability in log hepcidin was explained by plasma ferritin values.
over time, and a wider panel of indicators to capture the multi-factorial etiologies of iron deficiency and anemia during pregnancy and their as-yet unidentified associations with hepcidin concentrations.

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AUTHOR DISCLOSURES
None of the authors have a conflict of interest to disclose.

REFERENCES
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孟加拉懷孕婦女的鐵調素和鐵狀態

雖然近年來發現的脢肽荷爾蒙-鐵調素(hepcidin)，被認為是調控鐵代謝和慢性發炎性貧血的主要角色，但是在懷孕期貧血扮演的角色還未被證實。本研究的主要目的是描述 hepcidin 與懷孕期貧血的關聯性。我們執行一個橫斷性研究，對 149 位孟加拉鄉村的懷孕婦女做家訪，並獲得生物檢體，檢測尿中 hepcidin 和鐵狀態指標、血紅素、紅血球生成素、alpha-1 酸性糖蛋白(AGP)及 C 反應蛋白質之間的相關性。尿中 hepcidin 使用表面增強激光解析電離飛行時間質譜技術(SELDI-TOF MS)測量。尿中 hepcidin 濃度（intensity per mmol/L creatinine）轉換成對數值和鐵蛋白對數值有相關（r = 0.33, p < 0.001），和運鐵蛋白接受器指標相關（r = -0.22, p = 0.007），和 AGP 對數值亦相關（r = 0.20, p = 0.01），但是血紅素（r = 0.07, p = 0.40）和運鐵蛋白接受器對數值(r = -0.07, p = 0.41)、紅血球生成素對數值(r = -0.01, p = 0.88)或 C 反應蛋白質對數值（r = 0.06, p = 0.48）都沒有相關性。以低鐵儲存量的婦女（n = 41）來增加樣本數後，hepcidin 和鐵蛋白於複線性迴歸分析中仍維持強的相關性。在孟加拉鄉村社區研究中的懷孕婦女裡，尿中 hepcidin 濃度相關於體內鐵狀態和 AGP，但是與血紅素、紅血球生成素或 C 反應蛋白質沒有相關。

關鍵字：貧血、鐵調素、發炎、鐵、懷孕