Research Article

Cardiovascular Risk Profiles amongst Women in a Multiethnic Population in Inner City Britain: A Potential Impact of Anaemia

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The risk of diabetes is markedly reduced in men with iron deficiency anaemia (IDA). The nature of this relationship in women is not clear, nor is there information about the influence of ethnicity, given the increased susceptibility of diabetes amongst South Asians and Afro-Caribbeans. We reviewed 3563 patients with a diagnosis of anaemia from 2000 to 2007. The age-adjusted prevalence of vitamin B12 deficiency and IDA was calculated, together with cardiovascular comorbidities amongst Caucasians, South Asians, and Afro-Caribbeans. The prevalence of vitamin B12 deficiency (women only) or IDA was markedly higher in South Asians compared to Caucasians and Afro-Caribbeans. Among women with IDA, diabetes was more prevalent among South Asians (45%, 95% CI 39.0–51.0) compared to Caucasians (3.0%, 2.1–4.0); \( P < 0.001 \). Among South Asian women with vitamin B12 deficiency, the prevalence of diabetes was reduced 8.5% (5.2–12.0). South Asian women with vitamin B12 deficiency had a higher prevalence of myocardial infarction (MI) and ischemic heart disease (IHD), but this relationship was reversed in IDA. IDA is associated with a greater prevalence of diabetes in South Asian women, but it is not coordinated by a greater risk of macrovascular complications. Given the cardiovascular impact of diabetes in South Asians, this association merits further study in relation to its pathophysiological implication.

1. Introduction

Anaemia is an established risk factor for cardiovascular disease (CVD) outcomes in the general population [1], patients with chronic kidney failure, diabetes [2], acute coronary syndromes [3], and heart failure [4]. In the UK, the burden of CVD is more prominent amongst the country’s ethnic minorities [5], and this is mainly due to the preponderance of CVD risk factors, such as diabetes mellitus and hypertension, which are most evident amongst South Asian and Afro-Caribbean ethnic groups [6, 7]. In addition, the CVD risk profiles amongst ethnic minorities living in the UK are likely to have deteriorated with migration from countries of origin to the UK, specifically from changes in dietary exposures [8, 9].

A diet low in iron, folate, and vitamin B12 leading to nutrition-related anaemia could moderate CVD burden in this already high risk group [9, 10]. Vitamin B12 deficiency
results in increased oxidative stress promoting a progression of atherosclerotic vessel disease [11]. On the contrary, iron deficiency anaemia (IDA) has been reported to confer a protective effect against atherosclerotic heart disease in premenopausal women [12]. Previous epidemiological studies have not explored the prevalence of anaemia and the distribution of CVD risk factors in a multiethnic population.

The aim of the present study was to investigate the impact of anaemia on CVD risk profile among patients admitted to a multiethnic inner city teaching hospital in Birmingham, UK. We tested the null hypothesis that there were no significant differences in the effect of anaemia on CVD risk profile over a 7-year period (2000–2007). A database was compiled using data from national mortality registries and hospital records relating to the Sandwell and West Birmingham area, UK.

2. Research Design and Methods

This was a retrospective, cross-sectional analysis of hospital cases. Information from the local West Midlands Regional Health Authority computerized Hospital Activity Analysis register was used to compile a database of all patients admitted to a large multiethnic inner city hospital during the period 2000–2007 (International Classification of Disease (ICD) 10th revision, codes 430–438). The coding system has been proven to be free of errors for definitive diagnoses [13]. The data were cross-referenced with the computerized Hospital Activity Analysis register, and furthermore, hospital case notes were reviewed for detailed information on demographics, type of anaemia, and CVD risk factors (hypertension, diabetes mellitus, atrial fibrillation, myocardial infarction, ischemic heart disease, and hyperlipidaemia) to minimize any inaccuracies in coding.

2.1. Diagnostic Criteria and Definitions Used for Coding. Diagnosis was based on routine hospital practice which followed standard national guidelines [14]. A senior physician examined each patient admitted with symptoms and signs of CVD before performing any routine blood tests and further investigation. Diagnosis of anaemia was defined by hemoglobin levels <11.5 g/dL in females and <13.5 g/dL in males. IDA was defined as a Mean Corpuscular Volume (MCV) of less than 80fL along with a decreased total iron binding capacity saturation (<30%) and/or low serum iron (<11 μmol/L) and/or serum ferritin (<20 ng/mL) and/or transferrin saturation of <12%. B12 deficiency anaemia was defined as MCV > 100 fL and vitamin B12 < 211 pg/mL.

Anaemia of chronic disease was defined as anaemia in the absence of any other biochemical evidence of IDA or vitamin B12 deficiency in the presence of a chronic illness such as rheumatoid arthritis or cancer, for example. Blood pressure level >140/90 mmHg on at least two separate readings was indicative of hypertension. Diagnosis of diabetes was confirmed following patient’s clinical history and biochemical evidence of at least two measurements of fasting blood sugar readings >7.0 mmol/l. A total cholesterol level >5.2 mmol/l was indicative of hyperlipidaemia. Diagnosis of atrial fibrillation was based on relevant clinical evidence supported by a 12-lead ECG. History of myocardial infarction was confirmed by ECG evidence, further supported by elevated levels of cardiac troponin. For the diagnosis of valvular heart disease, echocardiographic evidence for diagnosis of valve lesions was considered mandatory.

2.2. Data Analysis. We tested the null hypothesis that there were no significant differences in the effect of anaemia on CVD risk profiles over a 7-year period (2000–2007). The sample size was calculated for a power of 90% and an alpha error of 0.05, where a sample size of 115 in each group would be required to detect a 10% difference in the prevalence of anaemia in subjects with CVD risk profiles.

Data were analyzed using SPSS version v14 (SPSS Inc., Chicago, IL, USA). Chi-square (χ²) test was used to compare the categorical variables (e.g., differences in the prevalence of anaemia and CVD risk factors amongst the ethnic minorities). The prevalence of CVD risk factors was age and gender adjusted using the age distribution of total hospital admissions (2000–2007). 95% CI was calculated around means/proportions.

3. Results

Between 2000 and 2007, there were 3563 hospital admissions, all diagnosed with anaemia, with complete clinical data available. Of all the patients with anaemia, 59% were women, 82% White, 13% South Asian, and 6% African Caribbean. Of these patients, 515 (14.5%) had IDA, 153 (4.3%) had vitamin B12 deficiency, 2534 (71.1%) presented with anaemia of chronic disease, and 361 (10.1%) were unspecified. Age-adjusted percentage of iron deficiency anaemia was significantly higher in South Asian women and men compared to White and African Caribbean ethnic groups ($P < 0.001$; Table 1). Vitamin B12 deficiency anaemia was more common in South Asian women compared to White women (8.2% (95% CI 4.9–11.5) and 5.5% (4.4–6.6); $P = 0.005$), respectively. Anaemia of chronic disease was significantly more common in European White groups (73.0% (71.0–75.0); $P = 0.001$; Table 1). Across the study cohort, 17 (0.5%) patients had thalassemia and 113 (3.2%) had sickle cell anaemia.

3.1. Risk Factors. Diabetes mellitus and ischemic heart disease were the most commonly encountered comorbidities at 8.2% and 10.1%. South Asian women had the highest prevalence of diabetes ($P < 0.001$). Ischemic heart disease was more prevalent amongst White and South Asian groups ($P < 0.001$) compared to Afro-Caribbeans. Whites had a higher prevalence of atrial fibrillation compared to Afro-Caribbeans and South Asians ($P < 0.001$; Table 1).

3.2. Anaemias and Cardiovascular Risk Factors. Among female patients with IDA, diabetes mellitus was significantly more prevalent among South Asian women (45% (39.0–51.0)) compared to White women (3.0% (2.1–4.0); $P < 0.001$). Among South Asian women with vitamin B12 deficiency anaemia, the prevalence of diabetes mellitus was (8.5% (5.2–12.0)). South Asian women with vitamin B12 deficiency
anaemia showed a higher prevalence of myocardial infarction (4.3% (2.0–7.0)) and ischemic heart disease (26.6% (21.3–31.9)). South Asian women with IDA showed a low prevalence of myocardial infarction and no ischemic heart disease. Amongst White women, this relationship was not seen. Amongst women with IDA, the prevalence of hypertension was (28.0% (22.4–33.2)) and (21.0% (19.0–23.0)) in South Asians and Whites, respectively. Amongst women with IDA, the prevalence of hypertension was comparable in South Asians and Whites. Across the whole cohort, patients with anaemia of chronic disease showed a higher prevalence of type 2 diabetes (𝑃 = 0.002), atrial fibrillation (𝑃 = 0.005), and ischemic heart disease (𝑃 = 0.065) irrespective of ethnicity (Table 2).

4. Discussion

The findings of this study suggest that among South Asian women with IDA, the prevalence of diabetes mellitus is exceptionally raised. This was not accompanied by an increased prevalence of ischemic heart disease—indeed, a paradoxical relationship is seen. The implication is that the prevalence of IDA in South Asian women may alleviate the risk of cardiovascular disease in a population that is susceptible to diabetes [7]. Further understanding into the pathophysiology of this relationship is required given the disturbing threat of ischemic heart disease on the Indian subcontinent [15].

Prior, community-based studies among South Asian women have demonstrated the prevalence of IDA and vitamin B12 deficiency anaemia mainly related to nutritional status [9]. The higher prevalence of IDA among women of South Asian origin living in Britain is likely to be explained by their life styles, including religious and cultural confines on certain foods [16].

Vitamin B12 deficiency anaemia is considered a major health issue in Asia especially among vegetarians. Yajnik et al. [17] have reported the risk of vitamin B12 deficiency to be greater in vegetarians compared to nonvegetarians among South Asians. Vitamin B12 is largely found in food of animal origin and is a cofactor for methionine synthase which recycles homocysteine (HCY) to methionine. Deficiency results in increased serum HCY levels, increased oxidative stress leading to increased risk of atherosclerotic vessel diseases. Levels of HCY are recognised to be increased in South Asians and are associated with an increased risk of ischaemic heart disease and associated complications [10]. The present study shows that South Asian women with vitamin B12 deficiency anaemia had a higher prevalence of myocardial infarction and ischemic heart disease which confirms that vitamin B12 deficiency confers additional cardiovascular burden in this ethnic group [9, 10].

Prospective studies have supported the hypothesis that iron plays a role in the development of type 2 diabetes and found a positive association between dietary heme-iron intake mainly from red meat and the risk of type 2 diabetes [18]. Iron is a potent metal and could convert inactive free radicals to highly active free radicals which could potentially interfere with insulin synthesis and secretion in the pancreas, leading initially to insulin resistance and later to reduced insulin secretion.
Table 2: Impact of anaemia on the distribution of CVD risk profiles in women by ethnic group.

| Risk factor               | Type of anaemia | Caucasian \((n = 1705)\) | South Asian \((n = 266)\) | Afro-Caribbean \((n = 118)\) | \(P\) value* |
|---------------------------|-----------------|---------------------------|---------------------------|---------------------------|--------------|
| Type 2 diabetes           | All anaemia     | 6.3 (5.1–7.4)             | 20.5 (16.0–25.4)          | 9.2 (4.0–14.4)            | <0.001       |
|                           | Iron deficiency| 3.0 (2.1–4.0)             | 45.0 (39.0–51.0)          | 0.0                       | <0.001       |
|                           | Vitamin B12 deficiency anaemia | 10.5 (9.1–12.0) | 8.5 (5.2–12.0) | 0.0 | 0.756 |
|                           | Anaemia of chronic disease | 7.0 (5.3–8.0) | 15.2 (11.0–20.0) | 10.1 (5.0–16.0) | 0.002 |
| Ischemic heart disease    | All anaemia     | 9.2 (8.0–11.0)             | 11.4 (8.0–15.2)          | 3.5 (0.2–7.0)            | 0.005       |
|                           | Iron deficiency| 12.0 (10.2–13.3)           | 0.0                       | 0.0                       | 0.206       |
|                           | Vitamin B12 deficiency anaemia | 12.4 (10.8–13.9) | 26.6 (21.3–31.9) | 0.0 | 0.168 |
|                           | Anaemia of chronic disease | 9.0 (7.2–10.0) | 8.4 (5.1–12.0) | 0.0 | 0.065 |
| Myocardial infarction     | All anaemia     | 1.6 (1.0–2.2)              | 2.0 (0.2–3.4)            | 0.0                       | 0.356       |
|                           | Iron deficiency| 2.3 (1.6–3.0)              | 2.4 (0.5–4.2)            | 0.0                       | 0.731       |
|                           | Vitamin B12 deficiency anaemia | 1.0 (0.5–1.4) | 4.3 (2.0–7.0) | 0.0 | 0.171 |
|                           | Anaemia of chronic disease | 1.6 (1.0–2.2) | 2.0 (0.3–4.0) | 0.0 | 0.632 |
| Hypertension              | All anaemia     | 15.3 (14.0–17.0)           | 29.2 (24.0–35.0)         | 25.0 (17.0–32.2)         | 0.918       |
|                           | Iron deficiency| 21.0 (19.0–23.0)           | 28.0 (22.4–33.2)         | 0.0                       | 0.786       |
|                           | Vitamin B12 deficiency anaemia | 24.0 (22.0–26.0) | 23.4 (18.3–28.5) | 0.0 | 0.601 |
|                           | Anaemia of chronic disease | 13.4 (12.0–15.0) | 23.0 (18.0–28.0) | 19.4 (12.3–27.0) | 0.970 |
| Atrial fibrillation       | All anaemia     | 6.6 (5.4–7.8)              | 1.1 (0–2.3)              | 3.5 (0.2–6.9)            | <0.001      |
|                           | Iron deficiency| 10.3 (9.0–12.0)            | 2.4 (0.5–4.2)            | 0.0                       | 0.024       |
|                           | Vitamin B12 deficiency anaemia | 8.8 (7.4–10.1) | 0.0 | 0.0 | 0.583 |
|                           | Anaemia of chronic disease | 6.1 (0.0–2.1) | 1.0 (0.0–2.1) | 7.1 (3.0–12.0) | 0.005 |

Data are age-adjusted percent (95% CI).

*On the basis of chi-square test across all groups.

In our study, the prevalence of diabetes mellitus was higher among South Asian women with IDA compared to those who had vitamin B12 deficiency anaemia. Early studies have demonstrated that iron metabolism influences glycosylation of hemoglobin A\(_1\)c (HbA\(_1\)c) in patients with IDA, and levels of HbA\(_1\)c levels in such patients can be misleading [19]. El-Agouza et al. [20] reported that the mean HbA\(_1\)c level reduced significantly from 6.15% ± 0.62 to 5.25% ± 0.45 after iron treatment. Thus, concluding IDA is a potential source for misdiagnosis of uncontrolled diabetes mellitus among iron-deficient patients. Haemoglobinopathies, mainly conditions such as thalassemias and sickle cell anaemia, also contribute to false HbA\(_1\)c determinations in diabetic patients [21]. In South Asian women, both IDA and thalassemia should be considered before interpreting HbA\(_1\)c levels for the diagnosis of diabetes mellitus, but further work is warranted.

South Asian women with IDA also showed a higher prevalence of hypertension along with diabetes mellitus. Reduced hemoglobin levels suggest an increase in cardiac output secondary to a higher myocardial oxygen demand which can lead to mild-to-moderate hypertension [1]. Similarly, South Asian and White women with vitamin B12 deficiency showed higher prevalence of hypertension which could be attributed to the effect of increased HCY levels [10].

In the present study, South Asian women with IDA showed lower prevalence of myocardial infarction and no cases of ischemic heart disease at all. Among premenopausal women, the reduced risk of CVD is attributed to the female sex hormones [22]. However, IDA is considered a health problem due to menstrual loss. The protective effect against atherosclerotic heart disease among iron-deficient patients could be explained by lipid metabolism where excess iron acts as a catalyst for the modification of low density lipoprotein (LDL) leading to the progression of atherosclerotic plaques [23]. This study cohort of anemic patients showed less incidence of hyperlipidemia. Our results are consistent to Özdemir et al. [12], thus indicating a lower lipid profile in iron-deficient patients’ hence, low risk of atherosclerotic diseases.

4.1. Study Limitations. This is an observational study, and the cohort consisted of anemic patients who were admitted to an inner city hospital (2000–2007). Hence, the cause or effect of anaemia cannot be proved. The diagnosis of IDA, vitamin B12 deficiency anaemia, and anaemia of chronic disease was done at a single point of time; so, duration of anaemia is not known. The prevalence of vitamin B12 deficiency anaemia among South Asian women may suggest that most of them were vegetarians, but unfortunately, in this study diet, status was not recorded. Furthermore, data on pre- or postmenopause or medications such as metformin, iron supplements, or erythropoietin which could potentially change the anaemia or cardiovascular risk profile status were not included in the analysis. In addition, the impact of \(\beta\) thalassemia and sickle cell anaemia on HbA\(_1\)c measurements is obvious, but we do...
not know the influence of trait. One possible explanation for the higher prevalence of diabetes mellitus among South Asian women with IDA could be treatment with ferrous sulfate. Finally, no data were included on inflammatory cytokine levels which could have explained the higher prevalence of anaemia of chronic disease in this study cohort among Whites.

5. Conclusions

South Asian women with IDA showed a higher prevalence of diabetes mellitus. Given the cardiovascular impact of diabetes in South Asians, this association merits further study in relation to its pathophysiological implication. Further work investigating IDA, underlying hemoglobinopathies and glycaemic control in this group, is needed. Vitamin B12 deficiency anaemia still remains a major health issue among South Asian women which needs to be tackled in order to provide a better healthcare for the management of CVD in this ethnic group.

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References

[1] M. J. Sarnak, H. Tighiouart, G. Manjunath et al., “Anemia as a risk factor for cardiovascular disease in the Atherosclerosis Risk in Communities (ARIC) study,” Journal of the American College of Cardiology, vol. 40, no. 1, pp. 27–33, 2002.

[2] M. C. Thomas, “The high prevalence of anaemia in diabetes is linked to functional erythropoietin deficiency,” Seminars in Nephrology, vol. 26, no. 4, pp. 275–282, 2006.

[3] M. S. Sabatine, D. A. Morrow, R. P. Giugliano et al., “Association of hemoglobin levels with clinical outcomes in acute coronary syndromes,” Circulation, vol. 111, no. 16, pp. 2042–2049, 2005.

[4] J. A. Ezekowitz, F. A. McAlister, and P. W. Armstrong, “Anemia is common in heart failure and is associated with poor outcomes: insights from a cohort of 12 065 patients with new-onset heart failure,” Circulation, vol. 107, no. 2, pp. 223–225, 2003.

[5] P. S. Gill, J. Kai, R. S. Bhopal, and S. Wild, “Health care needs assessment: black and minority ethnic groups,” in Health Care Needs Assessment. J. Raftery, A. Stevens, and J. Mant, Eds., The Epidemiologically Based Needs Assessment Reviews. Third Series, Radcliffe, Abingdon, UK, 2007.

[6] K. C. Ferdinando, “Managing cardiovascular risk in minority patients,” Journal of the National Medical Association, vol. 97, no. 4, pp. 459–466, 2005.

[7] J. V. Patel, H. S. Lim, A. Gunaratne et al., “Ethnic differences in myocardial infarction in patients with hypertension: effects of diabetes mellitus,” Quarterly Journal of Medicine, vol. 101, no. 3, pp. 231–236, 2008.

[8] J. K. Cruickshank and S. A. Alleyne, “Black West Indian and matched White diabetics in Britain compared with diabetics in Jamaica: body mass, blood pressure, and vascular disease,” Diabetes Care, vol. 10, no. 2, pp. 170–179, 1987.

[9] J. V. Patel, A. Vyas, J. K. Cruickshank et al., “Impact of migration on coronary heart disease risk factors: comparison of Gujaratis in Britain and their contemporaries in villages of origin in India,” Atherosclerosis, vol. 185, no. 2, pp. 297–306, 2006.

[10] M. D. Sosis, J. V. Patel, G. S. Bhatia, E. A. Hughes, R. C. Davis, and G. Y. H. Lip, “Effects of White European, African Caribbean and South Asian ethnicity on homocysteine levels in patients with systolic heart failure,” International Journal of Cardiology, vol. 129, no. 1, pp. 69–75, 2008.

[11] W. Herrmann, H. Schorr, K. Purschwitz, F. Rassoul, and V. Richter, “Total homocysteine, vitamin b12, and total antioxidant status in vegetarians,” Clinical Chemistry, vol. 47, no. 6 I, pp. 1094–1101, 2001.

[12] A. Özdemir, C. Seviç, U. Selamat, and F. Türkmen, “The relationship between iron deficiency anaemia and lipid metabolism in premenopausal women,” American Journal of the Medical Sciences, vol. 334, no. 5, pp. 331–333, 2007.

[13] A. M. George and G. B. Maddocks, “Accuracy of diagnostic content of hospital activity analysis in infectious diseases,” British Medical Journal, vol. 1, no. 6174, pp. 1332–1339, 1979.

[14] National Institute of Clinical Excellence (NICE), “National Institute of Clinical Excellence (NICE) guidelines for cardiovascular disease,” http://guidancenice.org.uk/topic/cardiovascular.

[15] K. S. Reddy and S. Yusuf, “Emerging epidemic of cardiovascular disease in developing countries,” Circulation, vol. 97, no. 6, pp. 596–601, 1998.

[16] A. H. Heald, R. Sharma, S. G. Anderson et al., “Dietary intake and the insulin-like growth factor system: effects of migration in two related populations in India and Britain with markedly different dietary intake,” Public Health Nutrition, vol. 8, no. 6, pp. 620–627, 2005.

[17] C. S. Yajnik, S. S. Deshpande, H. G. Lubree et al., “Vitamin B12 deficiency and hyperhomocysteinemia in rural and urban Indians,” Journal of Association of Physicians of India, vol. 54, pp. 775–782, 2006.

[18] R. Jiang, J. Ma, A. Ascherio, M. J. Stampfer, W. C. Willett, and F. B. Hu, “Dietary iron intake and blood donations in relation to risk of type 2 diabetes in men: a prospective cohort study,” The American Journal of Clinical Nutrition, vol. 79, no. 1, pp. 70–75, 2004.

[19] A. P. Brooks, J. Metcalfe, J. L. Day, and M. S. Edwards, “Iron deficiency and glycosylated haemoglobin A1,” The Lancet, vol. 2, no. 8186, p. 141, 1980.

[20] I. El-Agouza, A. Abu Shahla, and M. Sirdah, “The effect of iron deficiency anaemia on the levels of haemoglobin subtypes: possible consequences for clinical diagnosis,” Clinical and Laboratory Haematology, vol. 24, no. 5, pp. 285–289, 2002.

[21] W. J. Schnedl, R. Krause, G. Halwachs-Baumann, M. Trinker, W. Herrmann, R. W. Lipp, and G. J. Krejs, “Evaluation of HbA1c determination content of hospital activity analysis in infectious diseases,” American Journal of Clinical Nutrition, vol. 1094–1101, 2001.

[22] W. Herrmann, H. Schorr, K. Purschwitz, F. Rassoul, and V. Richter, “Total homocysteine, vitamin b12, and total antioxidant status in vegetarians,” Clinical Chemistry, vol. 47, no. 6 I, pp. 1094–1101, 2001.

[23] A. M. George and G. B. Maddocks, “Accuracy of diagnostic content of hospital activity analysis in infectious diseases,” British Medical Journal, vol. 1, no. 6174, pp. 1332–1339, 1979.