Prevalence and Determinants of Anemia in Older People With Diabetes Attending an Outpatient Clinic: A Cross-Sectional Audit

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The prevalence of anemia increases with increasing age, affecting ~ 10% of the general population ≥ 65 years of age. However, the prevalence of anemia in patients with diabetes could be double this figure. Anemia in patients with diabetes is likely to be related to renal insufficiency. However, it has been shown that diabetes increases the risk for developing anemia by two- to threefold compared to those without diabetes and similar renal function, suggesting that diabetes is associated with other underlying causes of anemia in addition to renal impairment.

In the elderly, anemia is an independent predictor of adverse outcomes such as poor quality of life, falls, decreased physical function, impaired cognition, increased infections, and mortality. It has also been shown that anemia is associated with impaired activities of daily living, increased disability, and hospitalization. Therefore, recognizing and treating the underlying cause of the anemia is an important aspect of patient care.

Underlying causes of anemia in older people with diabetes are likely to be multiple, including decreased renal function, inflammation, bone marrow suppression, and nutritional deficiencies. Although the prevalence of anemia increases with age, there is a dearth of literature investigating the burden of anemia in old (≥ 75 years of age) and very old (≥ 85 years of age) individuals with diabetes. The aim of this study was to investigate the prevalence and determinants of anemia among older people with diabetes (≥ 75 years of age) attending a long-term follow-up in a single outpatient clinic.

Design and Methods

The setting for this cross-sectional audit was an outpatient diabetes clinic for older people (≥ 75 years of age) in a District General Hospital in the United Kingdom. The clinic is run by a geriatrician with a special interest in diabetes. Older patients attending the clinic have access to care provided by a diabetes specialist nurse, physiotherapist, occupational therapist, dietitian, and chiropodist to provide a holistic approach to patients’ needs as well as onsite vascular surgery and psychogeriatric services. The clinic runs weekly and accepts referrals from the community, other hospital-based outpatient clinics, and inpatient departments as follow-up after hospital discharge.

The study population included all patients in long-term follow-up (> 2 years) who attended the outpatient clinic during a consecutive 3-month period.

Data collection

Baseline characteristics such as age, sex, ethnic origin, comorbidities, number of medications, and social circumstances were collected during patient interviews, and laboratory results for each patient were downloaded from the central database of the pathology laboratory using patient details. To avoid acute fluctuations in the laboratory results, averages during the past 24 months were calculated. Laboratory values collected included hemoglobin (Hb) level (g/l), estimated glomerular filtration rate (eGFR; ml/min/1.73 m² based on the Modified Diet in Renal Disease study equation), A1C (%), and hematonic studies such as iron profile, vitamin B₁₂, and folic acid levels.

Anemia detection

The prevalence of anemia was determined if the average Hb level during the past 24 months was below
the normal range used by the hospital laboratory (< 120 g/l for women and < 140 g/l for men).

**Determinants of anemia**

We examined five potential variables as determinants of anemia: older age (≥ 85 years of age), longer duration of diabetes (> 15 years), uncontrolled diabetes (A1C > 7.5%), multiple comorbidities (> 4), and presence of chronic kidney disease (CKD), defined as an eGFR < 60 ml/min/1.73 m².

**Statistical analysis**

Continuous variables are presented as means and standard deviations (SDs), and categorical variables are presented as percentages. A χ² test was used to compare categorical variables, Fisher’s exact test was used when the expected cell value was < 5, a t test was used for continuous variables, and a two-sided P of < 0.05 was considered significant. We performed an initial univariate and then a multivariate logistic regression analysis using the presence of anemia as the outcome or the dependent factor and the above-mentioned five variables individually and then in combination as the predictors. Odds ratios (ORs) and 95% confidence intervals (CIs) were used to estimate the association between outcome and variables. Statistical analysis was performed using the statistical software package Stata, version 10 (StataCorp LP, College Station, Tex.)

**Study Results**

**Baseline characteristics**

A total of 142 patients attended the outpatient clinic during the 3-month period. Twenty-seven patients had been attending the clinic for < 2 years and were excluded from the study. The remaining 115 patients had had long-term follow-up in the clinic (> 2 years) and formed the study population. Baseline characteristics are summarized in Table 1. All patients were > 75 years of age (mean age 83.6 [SD 5.2] years), and 51 (44%) were ≥ 85 years of age. Most patients (88%) lived in their own home and had a long duration of diabetes (mean duration 16.0 [SD 6.4] years). CKD was prevalent in 66.1% of patients. Three patients had a history of treated breast cancer, two had prostate cancer, and two had bowel cancer. No patients in the cohort had active or terminal cancer. Only six patients reported drinking a little alcohol socially.

| Table 1. Baseline Characteristics of All Patients (n = 115) |
|-----------------------------------------------------------|
| **Age (years)** | Mean (SD) | 83.6 (5.2) | |
| **Sex [n (%)]** | Male | 57 (49.6) | |
| **Ethnic origin [n (%)]** | White | 108 (94) | |
| **Number of comorbidities [n (%)]** | Hypertension | 72 (62.6) | |
| **Number of medications** | Mean (SD), range | 8.4 (2.5), 3–14 | |
| **Social factors [n (%)]** | Living alone | 53 (46) | |
| **Duration of diabetes (years)** | Mean (SD), range | 16 (6.4), 6–46 | |
| **Insulin therapy (%)** | 57 (49.6) | |
| **A1C (%)** | Mean (SD), range | 7.8 (1.2), 5.2–10.9 | |
| **eGFR (ml/min/1.73 m²)** | Mean (SD), range | 52.1 (20.4), 10.8–90 | |
| **< 60 [n (%)]** | 76 (66.1) | |
Prevalence of anemia

Anemia was prevalent in 68 patients (59%), of whom 35 were men and 33 were women. There was no history of hereditary anemias such as thalassemia or sickle cell disease. The majority of anemia (80%) was normocytic; anemia was microcytic in 17% and macrocytic in 3% of patients. There was no significant difference between patients with or without anemia regarding sex, number of comorbidities, number of medications, insulin therapy, hematocrit therapy, social factors, or functional status (Table 2). The use of metformin, pioglitazone, angiotensin-converting enzyme (ACE) inhibitors, angiotensin receptor blockers (ARBs), or antiplatelets was similar in both groups.

Determinants of anemia

Patients with anemia were significantly older (84.6 vs. 82.1 years, \( P = 0.01 \)), had a longer duration of diabetes (17.7 vs. 13.5 years, \( P = 0.03 \)), and had lower eGFRs (47.8 vs. 58.1 ml/min/1.73 m², \( P = 0.01 \)) than patients without anemia. In the univariate logistic regression analysis, three of the five potential determinants significantly predicted anemia: older age \( \geq 85 \) years (OR 4.8, 95% CI 2.1–7.8, \( P = 0.001 \)), longer duration of diabetes > 15 years (OR 2.9, 95% CI 1.3–to 6.2, \( P = 0.01 \)), and CKD (OR 3.4, 95% CI 1.5–7.6, \( P = 0.002 \)), whereas number of comorbidities and diabetes control were not significant. However, in the multivariate logistic regression analysis, older age (OR 4.6, 95% CI 1.9–8.1, \( P = 0.001 \)) and CKD (OR 2.9, 95% CI 1.2–6.9, \( P = 0.01 \)) remained significant, whereas the effect of CKD was reduced to borderline significance (OR 2.4, 95% CI 0.96–5.7, \( P = 0.06 \)) (Table 3).

Table 2. Comparison of Patients With and Without Anemia

|                             | Anemia       | No Anemia    | Difference (95% CI), \( P \)           |
|-----------------------------|--------------|--------------|----------------------------------------|
| \( n \) (%)                 | 68 (59.1)    | 47 (40.1)    |                                        |
| Hb [g/l; mean (SD)]         | 11.3 (0.9)   | 13.4 (0.8)   | 2.1 (1.8 to 2.6), < 0.001              |
| Age [years; mean (SD)]      | 84.6 (4.5)   | 82.1 (5.7)   | 2.5 (0.6 to 4.4), 0.01                 |
| Sex [male; \( n \) (%)]     | 35 (51.5)    | 22 (47)      | 4.5 (–13.0 to 22.0), 0.6               |
| Comorbidities [mean number (SD)] |              |              |                                        |
| Hypertension [\( n \) %]    | 3.9 (1.1)    | 4.0 (1.6)    | 0.1 (–0.4 to 0.6), 0.8                 |
| Vascular disease [\( n \) %] | 24 (62)      | 30 (63)      | 1.0 (–15.0 to 20.0), 0.8               |
| Dementia [\( n \) %]        | 50 (74)      | 32 (68)      | 6.0 (–11.0 to 22.0), 0.5               |
| Falls [\( n \) %]           | 14 (21)      | 9 (19)       | 2.0 (–0.2 to 6.0), 0.6                 |
| Depression [\( n \) %]      | 15 (22)      | 12 (25)      | 3.0 (112.0 to 13.0), 0.7               |
| Hematocrit deficiency [\( n \) %]* | 17 (25)  | 8 (17)       | 0.1 (–0.1 to 0.3), 0.3                 |
| Hypothyroidism [\( n \) %]  | 10 (9)       | 7 (6)        | 3.0 (–4.0 to 11.0), 0.5                |
| Medications [mean number (SD)] |              |              |                                        |
| Insulin therapy [\( n \) %] | 33 (49)      | 24 (51)      | 2.0 (–11 to 16), 0.8                   |
| Hematocrit supplement [\( n \) %] | 19 (28)  | 7 (15)       | 0.2 (–0.1 to 0.4), 0.1                 |
| Social factors and function [\( n \) %] |              |              |                                        |
| Lives alone                 | 31 (46)      | 22 (47)      | 1.0 (–12 to 17), 0.9                   |
| Carers help                 | 27 (38)      | 20 (43)      | 5.0 (–12 to 19), 0.6                   |
| Mobile independent          | 22 (32)      | 13 (28)      | 4 (–12 to 21), 0.6                     |
| ADL independent             | 31 (46)      | 23 (49)      | 3 (–13 to 21), 0.7                     |
| Duration of diabetes [years; mean (SD)] | 17.7 (7.9) | 13.5 (6.1)  | 4.2 (0.3 to 7.8), 0.03                 |
| A1C [%; mean (SD)]          | 7.6 (1.2)    | 8 (1.1)      | 0.4 (–0.3 to 6.8), 0.1                 |
| eGFR [ml/min/1.73 m²; mean (SD)] | 47.8 (7.9) | 58.1 (12.4) | 10.3 (2.7 to 14.6), 0.01               |

*Hematocrit deficiency encompasses iron, vitamin \( B_12 \), and folic acid.

ADL, activities of daily living.
Table 3. Univariate and Multivariate Analysis of Possible Predictors of Anemia

| Predictor                        | Unadjusted OR (95% CI), P   | Adjusted OR (95% CI), P   |
|---------------------------------|-----------------------------|---------------------------|
| Older age ≥ 85 years            | 4.8 (2.1–7.8), 0.001        | 4.6 (1.9–8.1), 0.001      |
| Longer duration of diabetes > 15 years | 2.9 (1.3–6.2), 0.01         | 2.9 (1.2–6.9), 0.01       |
| Low eGFR < 60 ml/min/1.73 m²    | 3.4 (1.5–7.6), 0.002        | 2.4 (0.96–5.7), 0.06      |
| Comorbidities > 4               | 1.4 (0.6–3.0), 0.4          | 1.04 (0.4–2.6), 0.9       |
| Uncontrolled diabetes (A1C > 7.5%) | 0.6 (0.4–1.1), 0.1          | 0.6 (0.2–1.3), 0.2        |

Discussion

This study demonstrates a higher prevalence of anemia (59%) in older people (> 75 years of age) with diabetes attending an outpatient clinic. Previous studies of a younger population (average age 65 years) with similar renal function to our population (CKD in ~70% of subjects) have reported a lower prevalence (~20%) of anemia in patients with diabetes, suggesting that our higher prevalence is likely to be the result of the older age of our population. Anemia prevalence increases with age, reaching up to 26% of the general older population (≥ 85 years of age), but is significantly higher in people with diabetes, reaching up to 78.6% of diabetic centenarians, confirming the contributing effect of diabetes to the prevalence of anemia.

The cause of diabetes’ contribution to anemia development is still unclear. In the Cardiovascular Health Study, although diabetes predicted hemoglobin decline by −0.9 g/l (95% CI −0.18 to −0.02), it did not predict the development of anemia (OR 1.04, 95% CI 0.78–1.38), whereas baseline increasing age and CKD predicted both hemoglobin decline and anemia over a 3-year period (−1.5 g/l, 95% CI −2.4 to −0.6, and −0.9 g/l, 95% CI −1.6 to −0.3, for hemoglobin decline and 1.96, 95% CI 1.45–2.64, and 1.39, 95% CI 1.13–1.71, respectively, for anemia development). The discrepancy in the effect of diabetes causing hemoglobin decline but not leading to anemia development could be because of the short (3-year) duration of the study.

In our study, long duration of diabetes (> 15 years), CKD, and older age (≥ 85 years) were the main predictors of anemia. However, in the multiple logistic regression analysis, the effect of CKD on anemia prediction almost disappeared after adjustment for diabetes duration and older age. This suggests that CKD acts as a mediator rather than a direct cause of anemia in older people with diabetes. In other words, anemia in older people with diabetes is mainly driven by age and duration of diabetes rather than by renal function. Our population was old enough and had a long enough duration of diabetes to show this effect, which has not been seen in previous studies of younger populations. This may explain why the prevalence of anemia was still higher in people with diabetes than in those without diabetes who had similar renal function. Diabetes, independent of renal function, has recently been shown to increase the risk of anemia (OR 2.15, 95% CI 1.07–4.31). The results of our study question the accepted concept that anemia in diabetes is mainly caused by a decline in renal function and suggests that, in older people with diabetes, anemia is mainly related to older age and longer duration of diabetes, which may have different or additional mechanisms for anemia.

Although drug therapy such as metformin, pioglitazone, ACE inhibitors, ARBs, and antiplatelets may be associated with anemia, we did not find this association in our study. Although we did not measure nutritional status, it is unlikely that the anemia in this study was caused by chronic blood loss or nutritional deficiencies because the anemia was normochromic in the majority of patients (80%), there were no patients in our cohort suffering from chronic bowel disease, and there was no association with hematinic deficiency or hematinic supplement use.

The anemia shown in this study is likely to be related to aging and to be of unexplained etiology. This finding may explain the failure of interventions with erythropoiesis-stimulating agents to normalize hemoglobin levels in older patients with diabetes and CKD that also showed increased incidence of vascular complications and even mortality.

Our study also suggests that anemia in this age-group of older patients with diabetes could be a marker of the severity or long duration of diabetes, especially when investigations for anemia reveal no obvious cause. Whether novel interventions to increase hemoglobin will have a role in ameliorating the
adverse health outcomes of diabetes requires future studies.

**Conclusion**
Anemia is highly prevalent in older people with diabetes. Older age and duration of diabetes were identified as significant predictors of anemia, whereas CKD was found to act as a mediator rather than a direct cause. In some very old patients, anemia appears to be related to aging, and its etiology remains unexplained. Therefore, no treatment has been well studied. It remains to be determined whether interventions to improve hemoglobin concentration will be associated with improved function or survival.

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