The absence of hemoglobin Hb H band in the proven hemoglobin H disease patient due to the iron deficiency anemia

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Abstract

Introduction: Iron deficiency anemia has not only been rarely seen among hemoglobin H disease patients but can mask the diagnosis of hemoglobin H disease also.

Case Presentation: A 50-year-old Thai woman had marked and progressive anemia with just splenomegaly on the physical examination. Her blood tests showed: Hb 3.8 g%, MCV 50.4 fl, MCH 15.4 pg, ferritin 5.0 ng/ml, serum iron 33 ug/dl, TIBC 123 ug/dl, the Hb analysis using the high-performance liquid chromatography method found only A₂A, Hb A₂ 2.1 %. Iron deficiency anemia was diagnosed and continuously treated with the oral iron tablets. After two months of treatment, her blood showed: Hb 7.7 g%, MCV 69.3 fl, MCH 20.5 pg, reticulocyte 6.0 %, ferritin 6.2 ng/ml, Hb analysis showed: A₂AH Bart, Hb A₂ 1.7 %. The diagnosis of Hb H disease was added. Later Hb H was confirmed by the positive PCR for alpha thalassemia-1, Southeast Asian deletion and alpha thalassemia-2, 3.7 kb deletion, genes. After five months of the iron treatment, Hb 8.8 g%, MCV 59.3 fl, MCH 16.7 pg, ferritin 110.7 ng/ml, Hb analysis: A₂ABartH, Hb A₂ 1.4 %. The band of Hb H was found after Hb concentration was raised even though the ferritin level was still low.

Conclusion: The absence or presence of Hb H band of the Hb H disease patient on Hb electrophoresis seemingly depended on the hemoglobin concentration more than the sufficiency of the iron storage.

Introduction

Hemoglobin H (Hb H) disease is a genetic disease resulted from the co-inheritance of alpha-thalassemia-1 and alpha-thalassemia-2 heterozygosity. Its main clinical manifestation is mild to moderate hemolytic microcytic anemia, Hb 10.0 ± 1.2 g%, MCV 67 ± 7 fl, MCH 19 ± 2 pg [1] or thalassemia intermedia. As other chronic hemolytic diseases [2], the iron storage, represented by the serum ferritin, is usually normal or increased in this disease [3,4], as compared with the normal control [5]. Moreover 73.9 % of Hb H disease patients have an iron overload [6] although most cases do not need regular blood transfusion [7].

On the contrary, iron deficiency anemia has been rarely reported among Hb H disease patients. Besides the rarity, iron deficiency anemia can mask the diagnosis of Hb H disease by causing the disappearance of Hb H band on the Hb analysis [8]. Now we report a new case of the iron deficiency anemia in the patient with Hb H disease that was confirmed by the genetic approval.

Case presentation

A 50-year-old Thai menopausal woman presented gradually progressive fatigue, frequent fainting and pallor for three months, no obvious blood loss. The physical examination showed moderate to marked pallor without jaundice and just palpable splenomegaly.

The blood tested: Hb 3.8 g%, Hct 12.4 %, MCV 50.4 fl, MCH 15.4 pg, MCHC 30.5 g%, RDW 32.3 %, WBC 8,400/mm³, platelet 526,000/mm³, ferritin 5.0 ng/ml, serum iron 33 mcg/dl (normal 35-165), TIBC 123 mcg/dl (normal 200-360), cholesterol 113 mg%, albumin 3.5 g%, globulin 3.0 g%, total bilirubin 0.3 mg%, direct bilirubin 0.1 mg%, alkaline phosphatase 72 U/L, SGOT 14 U/L, SGPT 11 U/L, creatinine 0.97 mg%, serum erythropoietin > 200 mIU/ml (normal 2.60-34.0).

The gastroscopy revealed minimal Mallory-Weiss tear 2 points while the colonoscopy revealed unremarkable study.

The hemoglobin analysis using the high-performance liquid chromatography (HPLC) method (BioRad®): A₂A, Hb A₂ 2.1 %, Hb F 0 %.

She was diagnosed as having severe iron deficiency anemia and continuously treated with the iron tablets and folic acid for 2 months. The blood was tested again: Hb 7.7 g%, Hct 26.1 %, MCV 69.3 fl, MCH 20.5 pg, MCHC 29.6 g%, RDW 20.7 %, WBC 9,100/mm³, platelet 306,000/mm³, reticulocyte 6.0 %, ferritin 6.2 ng/ml, serum iron 50 mcg/dl, transferrin 267.9 mg/dl, Hb analysis using the HPLC method: A₂AH Bart, Hb F 0.8 %, Hb A₂ 1.7 %

Besides iron deficiency anemia, the diagnosis of Hb H disease was added and the treatment with iron tablets was still going on. After 5 months of treatment: Hb 8.8 g%, Hct 31.2 %, WBC 6,100/mm³, MCV

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The only one clue to suspect that our patient should not have pure IDA is the just palpable splenomegaly which can be found around 20.5% of Hb H disease patients [12]. The MCV and MCH among IDA and beta thalassemia trait is significantly increased to 5.8+0.78 % after the iron treatment [16]. And those of Hb H disease are 43.9-75.1 fl and 15.1-28.0 pg, respectively [12]. Both are mostly overlap between two entities. The MCV and MCH during IDA in our case were 50.4 fl and 15.4 pg which appeared closely similar to those of Hb H disease but less than those of the IDA. However, it could not be simply concluded because the Hb concentration in our patient was 3.8 g%, less than 6.2 g% of the IDA group and there is the correlation between the Hb concentration and the MCV in IDA patients, viz., the lower Hb concentration the lower MCV [18].

**Conclusion**

A 50-year-old Thai woman presented with marked microcytic anemia due to iron deficiency. After the iron treatment, hemoglobin H band could be detected. Therefore, the hemoglobin analysis should be delayed in cases with suspected thalassemia during having severe iron deficiency anemia until the iron fortification could be completed.

**References**


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