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RETRACTION NOTE

Retraction note to: Prevalence of the C282Y and H63D mutations of the HFE hemochromatosis gene in Azerian major β-thalassemia and iron overload

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This article has been retracted at the request of the Editor-in-Chief per the Committee on Publication Ethics guidelines. The article shows evidence of irregularities in authorship during the submission process, there is strong reason to believe that the peer review process was compromised and the article contains patchwork plagiarism from a variety of sources.

The main sources are:
Hong Kong Med J 2000;6:153–8 | Number 2, June 2000

The C282Y mutation of the HFE gene is not found in Chinese haemochromatotic patients: multicentre retrospective study
WMS Tsui, PWY Lam, KC Lee, KF Ma, YK Chan, MWY Wong, SP Yip, CSC Wong, ASF Chow, STH Lo

European Journal of Immunogenetics
Volume 27, Issue 3, pages 129–134, June 2000
Iron-overload and genotypic expression of HFE mutations H63D/C282Y and transferrin receptor Hin6I and BanI polymorphism in German patients with hereditary haemochromatosis
R. Gottschalk, C. Seidl, S. Schilling, A. Braner, E. Seifried, D. Hoelzer and J. P. Kaltwasser
DOI: 10.1046/j.1365-2370.2000.00215.x

European Journal of Haematology
Vol 74 Issue 4
Prevalence of the H63D mutation of the HFE in north India: its presence does not cause iron overload in beta thalassemia trait
Gurjeewan Garewal, Reena Das, Jasmina Ahluwalia and R. K. Marwaha
DOI: 10.1111/j.1600-0609.2004.00390.x

Transfusion Clinique et Biologique
Volume 13, Issue 6, December 2006, Pages 353–357
Mutations du gène HFE chez des β-thalassémiques majeurs tunisiens et surcharge en fer
HFE gene mutations in Tunisian major β-Thalassemia and iron overload
F. Mellouli, W. El Borgi, H. Kaabi, E. Ben Hassen, R. Sassi, H. Hmida, G. Cherif, M. Maamar, B. Zouari, K. Boukef, M. Bejaoui, S. Hmida
DOI: 10.1016/j.tracli.2006.12.002

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Journal of Gastroenterology and Hepatology
Volume 28, Issue 7, pages 1087–1094, July 2013
Iron storage disease in Asia-Pacific populations: The importance of non-HFE mutations
Cameron J McDonald, Daniel F Wallace, Darrell H G Crawford, V Nathan Subramaniam
DOI: 10.1111/jgh.12222

Cold Spring Harb Perspect Med
2012;2:a011726
Pathophysiology and Clinical Manifestations of the β-Thalassemias
Arthur W. Nienhuis and David G. Nathan
DOI: 10.1101/cshperspect.a011726

As such the validity of the content of this article cannot be verified.