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Wilson's disease: A review of what we have learned

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Abstract

Wilson's disease, which results from the defective ATP7B protein product, is characterized by impaired copper metabolism and its clinical consequences vary from an asymptomatic state to fulminant hepatic failure, chronic liver disease with or without cirrhosis, neurological, and psychiatric manifestations. A high grade of suspicion is warranted to not miss cases of Wilson's disease, especially less florid cases with only mild elevation of transaminases, or isolated neuropsychiatric involvement. Screening in first and second

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www.genome.gov/27532725
Wilson disease causes a person'.s body to store too much of the mineral copper. Many foods contain copper, and it is important for people to have a small....

Wilson's disease - Wikipedia, the free encyclopedia
en.wikipedia.org/?title=Wilson's_disease
Symptoms usually appear between the ages of 6 and 20 years, but cases in much older people have been described. Wilson'.s disease occurs in 1 to 4 per....

Wilson (Wilson's) Disease Inheritance - Wilson Disease Association
www.wilsonsdisease.org/wilson-disease/wilsondisease-inheritance.php
Most patients have no family history of Wilson disease. ... members to learn if they could be affected but do not yet have symptoms, to learn they are carriers, or to....

Wilson Disease - Canadian Liver Foundation
www.liver.ca/liver-disease/types/wilsons-disease.aspx
Wilson disease is a hereditary disease in which excessive amounts of copper accumulate in the body, mainly in ... In order to have the disease, a person must get two defective genes, one from each parent. ... What else can we learn about it?

Canadian Liver Foundation | Wilson Disease
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www.niddk.nih.gov/health...**diseases**/**wilson-disease**/.../facts.asp... 翻译此页

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