Name of Journal: *World Journal of Hepatology*

Manuscript NO: 63116

Manuscript Type: MINIREVIEWS

Wilson's disease: revisiting an old friend

Wilson's disease: revisiting an old friend

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Wilson disease | Nature Reviews Disease Primers

https://www.nature.com/articles/s41572-018-0018-3

Sep 06, 2018 · **Wilson disease** (WD) is a potentially treatable, inherited disorder of copper metabolism that is characterized by the pathological accumulation ...

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Author: Julie M Button, Shefah A Qazi, Josep... **Publish Year:** 2020

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Non-invasive cellular and molecular imaging techniques are emerging as a multidisciplinary field that offers promise in understanding the components, processes, dynamics and therapies of **disease** at a molecular level. Magnetic resonance imaging (MRI) is an attractive technique due to the absence of r ...

Cited by: 153 **Author:** Dipanjan Pan, Shelton D. Caruthers, Ang...
Publish Year: 2011

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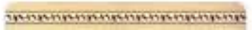

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Trientine dihydrochloride for Wilson's disease ...

https://www.nps.org.au/.../articles/trientine-dihydrochloride-for-wilsons-disease

Apr 26, 2021 · **Wilson's disease** is an autosomal recessive disorder. This genetic defect affects the transport of copper. Free copper is toxic so it causes cell damage as it accumulates, firstly in the liver and then in the brain. The **disease** is therefore also known as hepatolenticular degeneration. The treatment ...

A century for progress in the diagnosis of Wilson disease ...

https://www.sciencedirect.com/science/article/pii/S0946672X14001217

Oct 01, 2014 · The diagnosis of **Wilson disease** has evolved from the original description of a neurological syndrome by **Wilson** and other contemporaries at the turn of the 20th century to where we recognize that there is a spectrum of clinical liver and neuropsychiatric **disease** diagnosed by a combination of clinical and biochemical tests and more recently by molecular genetic analysis.

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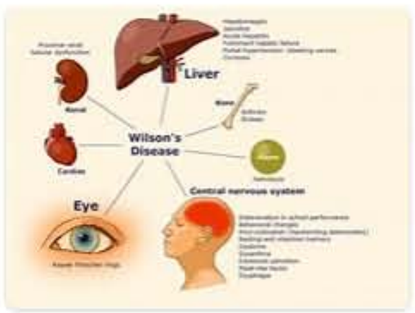

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The Diagnostic Approach to Wilson Disease - ScienceDirect

https://www.sciencedirect.com/science/article/pii/B9780128110775000086

Jan 01, 2019 · The diagnosis of **Wilson disease** (WD) should be excluded in any patient with unexplained

Wilson's Disease

Medical Condition

A genetic disorder causing excessive copper accumulation in the liver, brain and other organs.

- Rare (Fewer than 200,000 cases per year in US)
- Not known to be contagious
- No vaccine available
- Requires lab test or imaging
- Treatable by a medical professional
- Can last several months

Caused by inheritance through parents Symptoms include fatigue, abdominal pain, uncontrolled movements or stiffness. Treatment involves medications and surgery to remove copper deposition.

Treatments

Treatment involves medications and surgery to remove copper deposition.