University of San Diego Digital USD

At Risk for Huntington's Disease

Department of History

9-24-2005

Dr. Martha Nance to do live chat on HD

Kenneth P. Serbin University of San Diego

Follow this and additional works at: https://digital.sandiego.edu/huntingtons

Part of the Nervous System Diseases Commons

Digital USD Citation

Serbin, Kenneth P., "Dr. Martha Nance to do live chat on HD" (2005). *At Risk for Huntington's Disease*. 15. https://digital.sandiego.edu/huntingtons/15

This Blog Post is brought to you for free and open access by the Department of History at Digital USD. It has been accepted for inclusion in At Risk for Huntington's Disease by an authorized administrator of Digital USD. For more information, please contact digital@sandiego.edu.

More

At Risk for Huntington's Disease

HD is a genetically caused brain disorder that causes uncontrollable bodily movements and robs people's ability to walk, talk, eat, and think. The final result is a slow, ugly death. Children of parents with HD have a 50-50 chance of inheriting the disease. There is no cure or treatment.

Blog Archive

- ▶ 2021 (12)
- 2020 (16)
- ▶ 2019 (19)
- 2018 (16)
- ▶ 2017 (14)
- ▶ 2016 (13)
- ▶ 2015 (24)
- ▶ 2014 (24)
- ▶ 2013 (30)
- ▶ 2012 (26)
- ▶ 2011 (33)
- ▶ 2010 (26)
- ▶ 2009 (21)
- ▶ 2008 (7)
- ▶ 2007 (7)
- 2006 (4)
- ▼ 2005 (17)
 - December (2)
 - ▼ September (2)
 - Dr. Martha Nance to do live chat on HD An uncertain journey along the genetic trail
 - ▶ July (1)
 - ▶ May (1)
 - ► April (1)
 - ▶ March (4)
 - ► February (3)
 - ► January (3)

About Me

View my complete profile

HD Links

Huntington's Disease Society of America International Huntington Association Huntington's Disease Drug Works SATURDAY, SEPTEMBER 24, 2005

Dr. Martha Nance to do live chat on HD

- Monday, Sept. 26, at 1 p.m. ET
- Outlook: Huntington's Disease
 - Incurable Disease Looms Before Those With Gene
- Dr. Martha Nance
 - Struthers Parkinson's Center and the University of MinnesotaMonday,
 - September 26, 2005; 1:00 PM

Today there are thousands of Americans who have had themselves tested for the gene for Huntington's Disease and know they will get it. They've seen a parent suffer with it or die from it. But there's still no cure. As genetic testing advances, more and more Americans could find themselves in similar positions, dreading an all-too-familiar fate. In an article in Sunday's Outlook section, Dr. Martha Nance, a Minneapolis neurologist who runs a Parkinson's Disease clinic and who is doing research on treatments for Huntington's, discusses the moving tale of one person who has the gene and the race to find a way to treat people like him.Dr. Martha Nance will be online Monday, Sept. 26, at 1 p.m. ET to discuss her Sunday Outlook article on Huntington's Disease.

Click on this link for more information: <u>http://www.washingtonpost.com/wp-</u>

dyn/content/discussion/2005/09/23/DI2005092300953.html

Posted by Gene Veritas at 11:29 AM 💽 M 🖪 🔚 😭

1 comment:

Evan Light said...
"Gene",



As a caregiver for a fiance in the mid-stages of HD, I greatly appreciate your writing. I'm attempting to pull together whatever HD information that I can via RSS/Atom feeds. I don't suppose that you'd mind turning the Atom on for your site?

Thanks, Shepherd <u>8:13 AM, October 30, 2005</u>

Post a Comment

Newer Post

Home

Older Post

curehd.blogspot.com/2005/09/dr-martha-nance-to-do-live-chat-on-hd.html

Subscribe to: Post Comments (Atom)

Huntington's Disease Lighthouse Hereditary Disease Foundation Huntington's Disease Advocacy Center Thomas Cellini Huntington's Foundation HDSA Orange County (CA) Affiliate HD Free with PGD! Stanford HOPES Earth Source CoQ10, Inc.

HD Blogs and Individuals

Chris Furbee: Huntingtons Dance Angela F.: Surviving Huntington's? Heather's Huntington's Disease Page