

Mind Cases in Movement Disorders

Scenario 45: Keith

1. Huntington's chorea, specifically juvenile-onset Huntington's disease. Differential diagnosis, symptoms are usually mistaken for Parkinson's disease. Degenerative and hereditary nature of this condition help to distinguish it from Parkinson's. Huntington's involves the basal ganglia.
2. Marked by an abnormally high amount of Huntingtin protein, and basal ganglia damage. It is autosomal dominant (one copy of the gene from one parent will result in the disorder manifesting) and symptoms usually appear around the age of 35. In Keith's case he is exhibiting the juvenile form, and the earlier the onset typically the more severe and quick the decline in quality of life.
3. Anticholinergics help, because in Huntington's there is an oversupply of dopamine (anticholinergics block dopamine action) - the opposite of Parkinson's.
4. The abnormal hormone levels and raised by this disease and the effects it causes on the basal ganglia fully explain his symptoms. But because of his family history he had a higher percentage of showing symptoms of the disease at an early age.
5. There is no treatment to stop or reverse the disease. However there are medications that slow and help make the symptoms more bearable to live with. They say those that exercise more regularly fare far better than those who don't.

Reference:

What is Huntington's Disease? by J. Robbins | Updated September 26, 2018. from <https://www.mayoclinic.org>

Mini Cases In Movement Disorders Answer Key

Emilie Sanchez



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