



## Examining the Genetic Basis of Gender-Specific Neurodegenerative Disorders and How It Contributes to their Development

Jposh Karsh\*

*Department of Pharmaceutics, The State University of New Jersey, United States of America*

### DESCRIPTION

Gender-specific neurodegenerative diseases are illnesses that affect different genders differently. Examples include Alzheimer's disease, Parkinson's disease, and Amyotrophic Lateral Sclerosis (ALS). While there are many factors that can contribute to the development of these diseases, genetics play an important role. Research has shown that certain genetic markers can make certain individuals more susceptible to developing a gender-specific neurodegenerative disorder than others. Genetic risk factors for gender-specific diseases such as Alzheimer's and Parkinson's are largely determined by family history. Individuals with a parent or grandparent who had the disorder may be at increased risk for developing it themselves. Additionally, some genetic mutations have been linked to an increased risk for specific disorders. For example, mutations in the apolipoprotein E gene have been associated with an increased risk of developing Alzheimer's disease in women over 65 years old. In addition to genetic risk factors, environmental factors can also play a role in contributing to the development of a gender-specific neurodegenerative disorder. These could include lifestyle choices as well as exposure to toxins and pollutants over time. It is important to note that no single factor is solely responsible for the development of these diseases; rather, it is usually a combination of factors that can lead to their onset.

Neurodegenerative disorders have long been known to affect certain genders more than others. Recent studies have focused on exploring the genetic basis of these gender-specific neurodegenerative disorders in order to better understand their cause and development. This finding suggests that individuals who are already at higher risk for cancer may also be at an increased risk for developing neurodegenerative diseases. Additionally, Alzheimer's disease is more common among women than men after age 65 due to life expectancy differences between genders and some researchers speculate that estrogen levels could be a contributing factor as well. Research into the genetics and hormones behind gender-specific neurodegenerative diseases helps scientists identify potential biomarkers and treatments

for these conditions. Ultimately, this helps us better understand the underlying causes of these disorders so we can work towards improving patient outcomes.

Research has shown that gender plays a powerful role in the onset and progression of neurodegenerative disorders. While some diseases, such as Parkinson's disease, are known to affect men more than women, other disorders have differing onset ages and course of progression depending on gender. According to a paper published in neuromolecular medicine, there are several different genetic pathways that may be involved in the development of gender-specific neurodegenerative disorders. One study looked at Huntington's disease, which is known to disproportionately affect men. Researchers found that while both genders had similar levels of the mutant huntingtin gene implicated in Huntington's disease, male patients had much higher concentrations of the protein itself which leads to the development of the disorder. They concluded that there may be additional genetic components unique to each gender playing an important role in disease development and progression. In addition to Huntington's disease, researchers have also studied Amyotrophic Lateral Sclerosis (ALS) which is often referred to as Lou Gehrig's disease. This condition is more common among men than women, and recent studies suggest that genetic factors are likely at play. One particular study found that certain gene variants were more commonly associated with men who had ALS compared with men without ALS. It was also found that one variant was only present among female patients with ALS, suggesting a possible female-specific association with this gene variant and (Amyotrophic Lateral Sclerosis) ALS. These findings indicate that there are complex genetic pathways underlying gender-specific neurodegenerative disorders. Further research will help uncover why certain genes are associated with specific diseases in one gender but not another, as well as what other risk factors contribute to their development. With this new knowledge, medical professionals can develop more targeted treatments for each individual patient based on their unique risk profile.

**Correspondence to:** Jposh Karsh, Department of Psychology, University of Wisconsin-Madison, Madison, United States of America; E-mail: karsh.lu.ck.jposh@email.com

**Received:** 09-Oct-2023, Manuscript No. JPP-23-23459; **Editor assigned:** 12-Oct-2023, PreQC No. JPP-23-23459 (PQ); **Reviewed:** 26-Oct-2023, QC No. JPP-23-23459; **Revised:** 11-Jul-2024, Manuscript No. JPP-23-23459 (R); **Published:** 18-Jul-2024, DOI: 10.35248/2153-0645.24.15.116

**Citation:** Karsh J (2024) Examining the Genetic Basis of Gender-Specific Neurodegenerative Disorders and How It Contributes to Their Development. *J Pharmacogenom Pharmacoproteomics*. 15:116.

**Copyright:** © 2024 Karsh J. This is an open access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

It is becoming increasingly evident that gender-specific neurodegenerative disorders, such as Alzheimer's disease and Parkinson's disease, have a strong genetic component. As we learn more about the how genetics impacts individuals'

susceptibility to these conditions, we can begin to develop treatments that take this factor into account. The connection between genetics and gender-specific neurodegenerative diseases has been established for some time.