

DEDICATED *to*
REVERSING *the*
IRREVERSIBLE



the Foundation for Peripheral Neuropathy

Since 2007, *the* Foundation for Peripheral Neuropathy has focused on **improving the lives of those affected by peripheral neuropathy** through awareness, education, advocacy and research to advance the discovery of new therapies and cures.

Why we *advocate* for progress

Patients can display symptoms of pain or numbness, and have balance and mobility issues. We want to improve their lives by alleviating, or better yet eliminating, these issues with better treatments and cures.

How we *support* patients

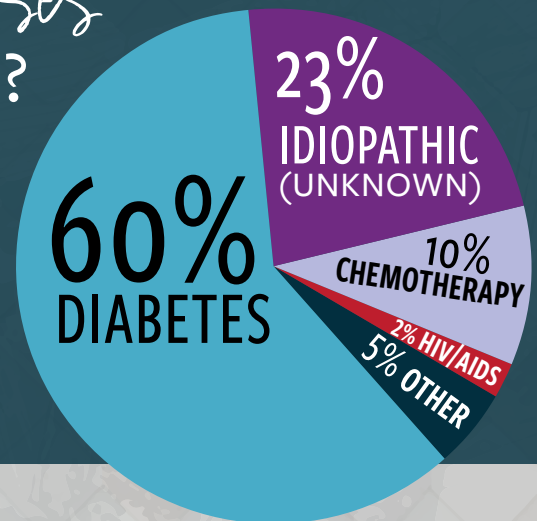
- » Providing information for patients, their families and healthcare providers
- » Funding scientific research and advocating for government funding to accelerate treatments and cures
- » Raising awareness of peripheral neuropathy and its causes

What is peripheral neuropathy?

Peripheral neuropathy (PN) occurs when the peripheral nerves are damaged or destroyed and can't send messages from the brain to other parts of the body. A neurological disorder, rather than a disease, PN has hundreds of known causes, and endless *unknown* causes.



What *causes* PN?



OUR

Purpose

Our work focuses on peripheral neuropathy awareness, education, advocacy and research.

We hope to continuously **identify more treatments** and, one day, **discover cures** for peripheral neuropathy (PN).

RESEARCH + ADVOCACY

FPN funds research through awards and grants, and provides research assistance through our biobank, the Peripheral Neuropathy Research Registry (PNRR). We also advance research through our advocacy work, successfully lobbying for expanded federal funding for peripheral neuropathy research.



AWARENESS

We help the world realize the prevalence of the condition through partnerships, advocacy, awareness campaigns and educational programs. An estimated 30 million people in the U.S., and millions more worldwide, struggle with PN.

GET INVOLVED:
FoundationForPN.org/aware



EDUCATION

Our ever-expanding library of educational resources highlights medical advances, research, and tips for living well with PN, providing information for patients, their families and healthcare providers.

CHECK OUT OUR RESOURCE LIBRARY:
FoundationForPN.org/library

FIND SUPPORT GROUP INFORMATION:
FoundationForPn.org/groups



OUR

Impact

\$5 MILLION
Funded over \$5 million in research

2000+

With a statistically significant cohort of 2000+ patient enrollments in the PNRR, more research studies are planned to answer the difficult questions surrounding PN

250K

Provided hundreds of webinars, newsletters, and other educational materials to 250,000 patients across the globe

100+

Partnered with a growing network of 100+ research scientists and medical experts

\$13 MILLION

Successfully expanded federal funding to include peripheral neuropathy in the Department of Defense's Peer Reviewed Medical Research Program, providing \$13 million in PN funding in the initial two years

20+

published research papers and investigations facilitated utilizing the PNRR

OUR NEWEST PROGRAM:

Two Clinical Research Training Scholarships in peripheral neuropathy awarded in collaboration with the American Brain Foundation and the American Academy of Neurology to encourage young researchers to study peripheral neuropathy

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Support THE CAUSE

- » Learn more at **FoundationForPN.org**
- » Follow, like and share our content on social media
- » Attend a free upcoming webinar FoundationForPN.org/webinars
- » Donate online today at FoundationForPN.org/donate

the Foundation for Peripheral Neuropathy is a 501(c)(3) non-profit organization; all donations are tax deductible.



IRREVERSIBLE REVERSING the DEDICATED to

Gold
Transparency
2023

Candid.

the **Foundation for
Peripheral Neuropathy**

is governed by a board of directors and scientific advisory board, who together help us maintain a comprehensive view of the field. Learn more at **FoundationForPN.org/leadership**.

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