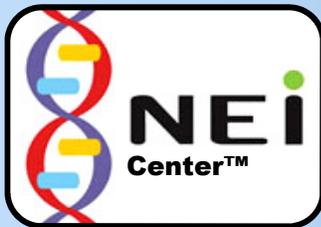


Most general practice physicians do not receive adequate training in medical schools for the complex neuro-endocrine-immune diseases. These diseases have not been embraced by any medical specialty because they cause dysfunctions in multiple body systems at the same time.

In addition to overall medical professional education of these diseases, we also work toward creating neuro-endocrine-immune disease specialty centers, called NEI Centers™.

Our goal is for NEI Centers™ to provide expert clinical care, research and social services access assistance.



Our Name

Our name is based on Pandora's Box in Greek mythology. When Pandora opened the box, disease and suffering spread into the world, but the last thing that flew out was hope. The butterfly represents the hope of patients being released from their cocoon of disease.

Our work brings hope to millions.

PANDORA Org is a 501(c) (3) nonprofit charity registered in Michigan. The full name is Patient Alliance for Neuro-endocrine-immune Disorders Organization for Research and Advocacy, Inc.



Our Advocates Extraordinaire™ program provides training so patients and their loved ones can effectively advocate to government officials and the news media.

Some of these individuals become PANDORA Org volunteers as ambassadors or chapter facilitators. Ambassadors represent us in local efforts or in national advocacy. Chapter facilitators lead a group in these efforts or lead a local patient support group.

Join us in improving patient quality of life by making a donation or becoming a PANDORA Org volunteer.

Call us at 231-421-8350.
Email us at info@pandoraorg.net
Donate or contact us through our website:



PANDORAorg.net

Mail check to: PANDORA Org
3625 Orchard View
Traverse City, MI 49684

Provided through an unrestricted grant from



Bringing Hope to Millions!

PANDORAorg.net

Serving those with neuro-endocrine-immune diseases (NEIDs), a.k.a. complex chronic diseases or chronic multi-system illnesses, including:

- Fibromyalgia Syndrome**
- Myalgic Encephalomyelitis / Chronic Fatigue Syndrome**
- Gulf War Illnesses**
- Chronic Tick-borne Infections**
- Environmental Illness**

Our vision:

Improving patient quality of life

Education

We give presentations to biology students about NEIDs. This plants a seed of knowledge and interest for those who may become nurses, physicians or researchers.

Through our Advocates Extraordinaire™ program, we train patients to be their own advocates when speaking to politicians and the news media.



We look for opportunities to arrange for physician education of NEIDs, such as this one in Alabama. As physicians become more knowledgeable of these diseases, they will diagnose patients earlier and provide better health care.



Awareness

Through public service announcements, brochures and motivating governments to recognize May 12 as NEIDs Awareness Day, we increase attention to and public knowledge of these diseases.

Advocacy

We speak out boldly at federal government advisory committee meetings on behalf of patient interests.



Patient Assistance

By arranging special outings, we help isolated patients keep an optimistic outlook.

When we can, we provide direct assistance to patients with electronics, gas cards or other items.



We have monthly support group meetings for social interaction, patient education and support.



We partnered with NeedyMeds to provide patients with low-cost medications.

Through our newsletter and social media, we keep patients informed of advocacy opportunities and research findings.

