

How Can You Help?

The PHF, Inc. is committed to bringing increased awareness and research on the diagnosis and treatment of Hydrocephalus to the forefront of public attention, which is a priority for everyone touched by this condition.

There are several ways you or your organization can contribute to the PHF, Inc:

- *Become a Corporate Sponsor
*DONATE
- *Become a Member and help raise funds
*Join a local State Chapter

Proceeds raised benefit the PHF, Inc., a non-profit 501 (c) (3) charitable organization providing support, education and advocacy for children and families dealing with the complex issues of Hydrocephalus.

September is

'National Hydrocephalus Awareness Month'

Contact Us Today!

To donate or for more information:

(732) 634-1283

www.HydrocephalusKids.org
info@HydrocephalusKids.org



***"Funding A Cure
For
Hydrocephalus"***

About the PHF, Inc.

The mission of the PHF, Inc. is to educate the community by raising the level of awareness about this incurable brain condition.

The PHF, Inc. also provides support to the families, friends and children who are diagnosed with Hydrocephalus.

The PHF, Inc. advocates on behalf of the members of the Hydrocephalus community and works with policy makers at the State and Federal level to raise awareness and push for more research and support in the fight against Hydrocephalus.

The PHF, Inc. is a non-profit 501 (c) (3) charitable organization.

What is Hydrocephalus?

*Hydrocephalus is a lifelong condition, affecting over One Million Americans from newborns to seniors.

*An average of 40,000 shunt surgeries are performed each year in the United States.

*Americans spend in excess of \$1 BILLION per year to treat this condition.

*Lack of advancements in treatment have left many people with Hydrocephalus unable to lead full and productive lives.

*There is no known cure for this condition.



'Families First for Hydrocephalus The PHF, Inc. puts Families First for Hydrocephalus. The PHF, Inc. understands that our most important mission is supporting the persons and families affected by Hydrocephalus. Each member of our TEAM is either a person living with Hydrocephalus or has a Child with Hydrocephalus, giving us a unique level of understanding and experience that you can rely on during difficult and stressful times.

The PHF, Inc. and its ALL volunteer staff is available 24/7/365 to lend an ear, locate a local support group, refer a Doctor, Neurosurgeon or Hospital, and answer your questions and concerns. The PHF, Inc. is a valuable resource for information, knowledge, comfort and support. No one has to be alone. The PHF is here for you!