

“. . .With the help of other FPIES families, we are coping, handling and even growing stronger as a family because of it” - An FPIES mom

The FPIES Foundation, a 501(c)(3) non-profit, was founded by parents of children with FPIES. The Foundation is dedicated to overcoming the challenges of FPIES by offering tools for education, support and advocacy to empower families and the medical community.

Awareness is Action!
Our shoes may be small but our steps will be big!

Support us at:
www.TheFPIESFoundation.org

FPIES (Food Protein-Induced Enterocolitis Syndrome) is a rare type of food allergy affecting the gastrointestinal tract. Reactions can require emergency care and be triggered by any food.

FPIES is not well-known, even within the medical community, and awareness is desperately needed to facilitate earlier diagnosis and improved symptom management for affected children. Together, we can raise awareness and help to improve the quality of life for affected children and their families! **You can help:**

- Create an awareness event with materials provided by The FPIES Foundation
- Volunteer to provide educational materials to parent groups or pediatric offices
- Reach out to affected families in your community and create a support group

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