



Rett Syndrome has been reversed in the lab and could be one of the first childhood neurological diseases to be cured. We are hopeful Rett research could also lead the way to treatments for millions of others suffering from these types of disorders.

*Aleksandra Djukic, MD, PhD.
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Associate Professor of Neurology and Pediatrics
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From the moment I heard about GP2C I knew they were special. The support, the inspiration, the knowledge they have and share has been amazing. I have made wonderful friends who I know will always be there for my daughter and our family. My journey as a parent to a child with Rett Syndrome was forever changed and made easier just by being a small part of this wonderful organization's family.

Dawn - Rett Syndrome parent

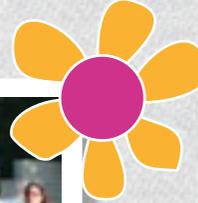
I am most impressed by the way Girl Power 2 Cure has responded to the need to give Rett children powerful access to learning and experiences long held unattainable. Girl Power 2 Cure stays on the cutting edge. This is what makes the charity unique and outstanding.

Donor

Visit us online:

Resources, materials and activities for:

- Rett families and their daughters
- Girls wanting to help a cause for girls
- Therapists and Educators
- Service Groups



Get involved!

Visit our website to learn about supporting GP2C as a donor, volunteer, event host, or community outreach organizer! We need YOUR POWER to help girls battling Rett Syndrome!

Girl Power 2 Cure, Inc.

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Amelia Island, FL 32034
904.277.2628

our websites:

gp2c.org rettgirl.org
rettuniversity.org

GIRL POWER 2 CURE, INC IS A 501(c)(3) NONPROFIT ORGANIZATION
TAX ID 20-8011668



Growing hope for girls with Rett Syndrome

FIND HER. HELP HER. CURE HER.



We educate and raise awareness.

Because girls with Rett Syndrome have the right to be heard.

Through webinars, events, and awareness campaigns, we have inspired a movement of thousands, and need thousands more.

We support families.

Because they are the heroes, tirelessly caring for their daughters, sisters and granddaughters.

We open our arms to every family, meet them where they are and provide them with the resources and support they need to fight for their daughters and for a cure. From helpful product reviews and hands-on training to fundraising efforts, we walk with them on the journey that many of us have made ourselves.

We fund research.

Because the world needs these girls.

They are brave, they are bright, and they are beautiful. We cannot rest knowing Rett Syndrome is curable. We are determined to see the girls and women who are suffering live better lives. Through a strong partnership with the Rett Syndrome Research Trust, we fund research that is crucial to developing treatments and a cure.

We inspire action.

Because change starts with a simple belief - that girls losing the ability to walk, speak or use their hands for a lifetime is unacceptable.

Our flower is our brand, the cornerstone, always in bloom with hope and positive energy ready to grow wherever there is inspiration to fight for our cause.

www.gp2c.org

about Rett Syndrome

- Debilitating neurological disorder that primarily affects females.
- The leading genetic cause of severe neurological impairment in girls - most cannot speak, walk or use their hands.
- As prevalent as Cystic Fibrosis, ALS and Huntington's.
- Another little girl is born with Rett Syndrome every 90 minutes.
- It is random and can strike any family, anywhere.
- There are close to half a million girls and women suffering worldwide.

“For us, change starts with a simple belief - that girls losing the ability to walk, speak or use their hands for a lifetime is unacceptable.”



#gp2c #powertheflower #rettsyndrome #untilshecan

