

Blooming Rose Foundation



Hope

Faith



Keeping positive is not hard when surrounded by those who believe.

Finding your faith

For many of us, the key to feeling positive and optimistic for our child(ren)'s future is connecting with other CF parents and CFers, and utilizing the support system that we have created.

All of our consultants want to help, please don't hesitate to contact any or all of them. We have different perspectives, but we all share a common vision of a positive journey filled with a hopeful future for all of our families.

Positive Parents

Kathleen Quinn East (MT)

c: 406.599.0139

e: kat@bloomingrosefoundation.org

Kate Gettel (AZ)

c: 520.250.0889

e: kgettel@gmail.com

Carin Stein (CA)

c: 858.245.1787

e: carinnirac@hotmail.com

Jill Hansen-Twardoski (MT)

c: 406.396.7279

e: j_twardoski@hotmail.com

Positive CFers:

Marianne Phillips Bertucci (PA)

(bi-lateral transplant recipient)

mbertucci43@yahoo.com

Sharlie Ross Kaltenbach (CA)

c: 619.750.7527

e: sharlie@larsenglobal.com

Josh Mogren (MN)

e: welcometojoshland@gmail.com

Somer Love (UT)

c: 801.971.2912

e: somermlove@yahoo.com

Emily Schaller (MI)

c: 734.341.5867

e: emily@letsrockcf.org

Ronnie Sharpe (AZ)

c: 602.315.5133

e: ronnie@cysticlifef.org

*Realize your child has more
than hope, they have a
future.*

What is Cystic Fibrosis?

CF brings about strength in young children that you can hardly believe is possible, it creates heroes and warriors.

CF empowers parents to become advocates for their children, never taking the backseat, never forgetting the importance of each day and each decision that is made.

CF creates a support system between parents of children with CF that you otherwise would never experience.

CF is a disease that makes us all stronger and gives our children the experiences to be our greatest teachers on this earth.

CF is something that we will fight and WIN!!

Belief

In honor of Maylie and her spirit for life we have started the Blooming Rose Foundation, a nonprofit organization. It is designed to assist families that have been given a new diagnosis of Cystic Fibrosis. We would like to share with other families what we have learned about the latest research, medications, and promising future. We hope to help other children with Cystic Fibrosis thrive when surrounded by those with positive outlooks based on knowledge and hope. We have faith in the future of every individual with Cystic Fibrosis as the cure rapidly approaches!!

Websites/Blogs:

www.runsickboyrun.com

www.cysticlifef.org

www.welcometojoshland.blogspot.com

www.cfri.org

www.cff.org

www.cfvoice.com

www.lovetobreathe.blogspot.com

<http://www.esiason.org/>

www.letsrockcf.org

<http://maryq.usana.com/>