

Cure Finders



Noah Colbert, Age 3

Noah is our inspiration for raising funds to cure cystic fibrosis. As an active three year old, Noah loves riding his scooter, cheering for his beloved Denver Broncos, and playing pirates with his big sister, Madeleine. However, Noah has cystic fibrosis, an inherited life-threatening, chronic disease that affects the lungs and digestive system of about 70,000 children and adults worldwide.

Every day, Noah must dedicate several hours to keeping himself healthy. He does chest physiotherapy twice a day to keep mucus from settling in his lungs. He takes enzymes and special vitamins before every meal to help him digest fats and gain weight, and he uses a variety of nasal sprays to help him breathe.

We have seen much progress in cystic fibrosis research and treatments since Noah was born in 2005, yet we continue to support the Cystic Fibrosis Foundation in their mission to find a cure for this disease. Our hope is that people living with cystic fibrosis won't have to spend several hours a day on therapies and endure numerous surgeries to keep them alive. Our hope is that Noah will have to take only a couple of pills a day to stay healthy. Our hope is that Noah will live to be a grandfather.

Thank you for being **CureFinders** - your contribution is immeasurable. Your efforts are giving people hope, health, and longer lives!

The Colbert Family
Jim, Libby, Madeleine (age 5) & Noah (age 3)



...adding tomorrows every day.

For more information, contact:
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