



2019 SPONSORSHIP OPPORTUNITIES

Saratoga Springs CF Cycle for Life
Sunday, September 22, 2019
Saratoga Polo Association – Saratoga Springs, NY

Northeastern New York Chapter
518.453.3583 – ne-ny@cff.org

The **MISSION** of the Cystic Fibrosis Foundation is to **CURE CYSTIC FIBROSIS** and to provide all people with the disease the **OPPORTUNITY TO LEAD FULL, PRODUCTIVE LIVES** by funding **RESEARCH AND DRUG DEVELOPMENT**, promoting **INDIVIDUALIZED TREATMENT** and ensuring **ACCESS TO HIGH-QUALITY, SPECIALIZED CARE**.



THE CYSTIC FIBROSIS FOUNDATION,

leading the way

ABOUT THE CYSTIC FIBROSIS FOUNDATION

Founded in 1955, the Cystic Fibrosis Foundation is the world's leader in the search for a cure for cystic fibrosis. The Foundation was started by parents desperate to save their children's lives. Their relentless and impassioned determination to prolong life has resulted in tremendous strides over the past 60 years in accelerating innovative research and drug development, as well as advancing care and advocacy. Virtually every approved cystic fibrosis drug therapy available now was made possible because of the Foundation and its supporters. Still, we believe no one should have to die at a young age. **We will not rest until we have a cure for all people living with CF.**

ABOUT CYSTIC FIBROSIS

Cystic fibrosis is a rare, genetic, life-shortening disease that affects every organ in the body and makes breathing difficult. Some people with the disease say it's like breathing through a narrow straw. In people with CF, a defective gene causes a thick buildup of mucus in the lungs, pancreas and other organs. In the lungs, the mucus clogs the airways and traps bacteria, leading to life-threatening lung infections. Sixty years ago, most children did not live long enough to attend elementary school, but thanks to Foundation-based research and care, the median survival age of people with CF is now into the 40's.

While people with CF are living longer than in the past, we still lose precious young lives every day.

WHY SUPPORT THE FOUNDATION?

Many people with CF are living long enough to realize their dreams of attending college, pursuing careers, getting married, and having children. This is due in large part to the work of the Foundation and the amazing CF community. We helped discover the gene that causes CF, created a state-of-the-art model for CF care, and have funded groundbreaking research. But we're not done. We are working every day to build on this incredible momentum, and we won't stop until there is a cure for *all* people living with CF.



Despite tremendous progress, we are not yet done. Our story will not be finished until we have achieved normal life spans and have a cure for 100 percent of people with CF.

Research Advances

Drugs that treat the genetic cause of CF are rapidly becoming available to more individuals as a result of the Foundation's efforts. There are now two FDA-approved therapies that treat the basic defect in cystic fibrosis for more than half of the population. Perhaps most exciting, more and more people will be helped by these treatments in coming years. In fact, by 2026, we expect that close to 95 percent of people with CF will benefit from similar drugs.

Better Today's

In addition to adding tomorrows by pursuing innovative treatments, we're also committed to helping people with CF live the best lives that they can today. Along with new therapies and a one-time cure, we're still steadfast in our efforts to develop treatments that address the symptoms of CF. We are continuing to invest in CF care by increasing our support for the 120 centers in our innovative care network, with a focus on adult care and mental health. And we're actively focused on lung transplant initiatives for the many adults with CF who need them. We are enabling efforts by and for people with CF. And we're working diligently in the policy arena to ensure that people with CF have access to the care they need. In addition, CF Foundation *Compass*, exists to help people with CF and their families with insurance, financial, legal and other complex issues they are facing.

A Cure for All

But we don't just want to treat CF. We want to end CF – for everyone, including those with rare or nonsense mutations. To bring that vision to life, we are focused on a very high-tech and experimental process called gene editing. Gene editing will remove the genetic mutation that causes CF and replace it with a normal sequence of DNA. Research into using gene editing to cure CF is already underway. Although there are many hurdles, with time and continued effort, we believe we will see a permanent, one-time cure in our lifetimes.

We need your involvement now as much as ever before to reach that goal.

CF CYCLE FOR LIFE

sponsorship

CF Cycle for Life is the Cystic Fibrosis Foundation's premier cycling event – a ride with a purpose. Each year, thousands of riders join forces in the fight against CF, pedaling along scenic routes and across multiple distances in a once-in-a-lifetime experience. Whether you're a first-time participant or a professional cyclist, you can turn your passion into progress and inspire change through what you love.

Cystic Fibrosis Cycle for Life provides a great opportunity for companies and groups of friends and family members to come together and ride in support of a worthy cause. Those who participate are given the opportunity to display their community involvement and raising funds to provide all people with cystic fibrosis the opportunity to lead full, productive lives.



We invite you to be a part of this year's event and join us in riding to new distances by supporting the Cystic Fibrosis Foundation and CF Cycle for Life as an event sponsor.



CHAPTER-WIDE SPONSORSHIP LEVELS AND BENEFITS

	Presenting \$15,000	Premier \$10,000	Gold Spokes \$7,500	Silver Spokes \$5,000	Why I Ride \$2,500	Start/ Finish \$2,000	Rest Stop \$1,000	Mile Marker \$500
PRE- AND POST-EVENT RECOGNITION								
Event presented by company name (limited to one sponsor only)	X							
Company logo on the front of the event brochure	X							
Recognition in chapter's event press release	X							
Company logo on cycle guide	X							
Company logo on event fact sheet	X	X						
Company logo on poster	X	X	X	X	X			
Company logo on brochure in sponsorship recognition section (back of brochure)	X	X	X	X	X	X		
DAY-OF-EVENT RECOGNITION								
Corporate spokesperson to speak during program	X	X						
Company logo on commemorative event shirt	X	X	X	X				
Company-provided banner displayed at main stage area	X	X	X	X				
Company-provided banner displayed at Why I Ride wall					X			
Company-provided banner displayed at start/finish line						X		
Company-provided banner at rest stop							X	
Company logo on mile marker	All	3	2	1				1
Company table to distribute company information at events	X	X	X	X	X			
WEB/SOCIAL MEDIA								
Logo placement on event website	X	X						
Company logo on chapter event e-communications	X	X	X	X	X			



2019 Saratoga Springs CF Cycle for Life

Company Name _____

Contact Name _____

Contact Title _____

Mailing Address _____

City _____ State _____ Zip _____

Phone# _____ Fax# _____

Email _____ Web Address _____

I would like to become a sponsor at the following level:

Presenting Level \$15,000*

Why I Ride Level \$2,500*

Premier Level \$10,000*

Start/Finish Level \$2,000*

Gold Spokes Level \$7,500*

Rest Stop Level \$1,000*

Silver Spokes Level \$5,000*

Mile Marker Level \$500*

I cannot sponsor but want to make a general donation to the event: _____

Sponsorship Payment:

Check enclosed (payable to the Cystic Fibrosis Foundation) Please invoice Please call for payment: _____

Credit Card Payment: Number: _____ Expiration Date: _____

Name on Card: _____ Total amount to charge: _____

Signature: _____

This signature authorizes the Cystic Fibrosis Foundation to charge the credit card number above the stated and agreed upon amount. Credit card information will be securely destroyed immediately after processing.

Northeastern New York Chapter
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Please email or mail completed form to E: ne-ny@cff.org P: 518.459.2903

For questions, please contact: Jillian Farrell – E: jfarrell@cff.org P: 518.453.3583

The Cystic Fibrosis Foundation has unrestricted financial reserves of about 13 times its 2018 budgeted expenses following a one-time royalty sale in 2014. These funds, along with the public's continuing support, are needed to help accelerate our efforts to pursue a cure for this fatal disease, fund development of new therapies, and help all people with CF live full, productive lives. To obtain a copy of our latest Annual Report, visit <https://www.cff.org/About-Us/Reports-and-Financials/>, email info@cff.org or call 1-800-FIGHT-CF.

Important Note on Attendance at Foundation Events: To reduce the risk of getting and spreading germs at CF Foundation-sponsored events, we ask that everyone follow basic best practices by regularly cleaning your hands with soap and water or with an alcohol-based hand gel, covering your cough or sneeze with a tissue or your inner elbow and maintaining a safe 6-foot distance from anyone with a cold or infection. Medical evidence shows that germs may spread among people with CF through direct and indirect contact as well as through droplets that travel short distances when a person coughs or sneezes. These germs can lead to worsening symptoms and speed decline in lung function. To further help reduce the risk of cross-infection, the Foundation's attendance policy recommends that all people with CF maintain a safe 6-foot distance from each other at all times while attending an outdoor Foundation-sponsored event.