



Benefitting The Cystic Fibrosis Foundation
Monday, October 10, 2019
Walk of Fame Park | Nashville, TN
Registration 1 p.m. | Tournament 2 p.m.



	Number of Teams & Tickets (2 players per team)	Customized Board Set* Includes logo on boards	Company name displayed during event	Emcee Recognition	Social Media Recognition	Partner for the Cure Wall (plaque displayed for one year)
\$10,000 Scoreboard Sponsor \$8,480 tax deductible	4 teams 20 audience tickets	2 used during event	X Logo displayed	X	X	X
\$7,500 Beverage Sponsor \$6,740 tax deductible	2 teams 10 audience tickets	1 used during event	X	X	X	X
\$7,500 Food Sponsor \$6,740 tax deductible	2 teams 10 audience tickets	1 used during event	X	X	X	X
\$5,000 Championship Board Sponsor \$4,320 tax deductible	2 teams 8 audience tickets	1 used during final round	X	X	X	X
\$3,000 Final Four Board Sponsor \$2,320 tax deductible	2 teams 8 audience tickets	1 used during semi-final round	X	X	X	X
\$2,500 Board Sponsor \$1,900 tax deductible	2 teams 6 audience tickets	1 used during event	X	X	X	X
\$1,500 Volunteer Tent Sponsor \$1,180 tax deductible	1 team 6 audience tickets			X		X
\$1,000 Elite Team Sponsor \$760 tax deductible	1 team 4 audience tickets					
\$300 Team Entry \$220 tax deductible	1 team					

*Not available after September 28, 2019

Can't sponsor this year? Form a Corporate Team! Encouraging team participation among employees is a wonderful way to support your employees and cystic fibrosis. Unite your employees in the common goal of saving lives, strengthen morale, promote team building, and ultimately help find a cure for cystic fibrosis.



[A Look at who we are supporting](#)

Mission Statement:

The mission of the Cystic Fibrosis Foundation is to cure cystic fibrosis and to provide all people with the disease the opportunity to live full, productive lives by funding research and development, promoting individualized treatment and ensuring access to high-quality, specialized care.



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 40s.

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.





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Company Name _____

Contact Name _____

Contact Title _____

Mailing Address _____

City _____ State _____ Zip _____

Phone # _____

Email _____ Website _____

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

- | | |
|---|---|
| <input type="checkbox"/> Scoreboard Sponsor \$10,000 (\$8,480 tax deductible) | <input type="checkbox"/> Board Sponsor \$2,500 (\$1,900 tax deductible) |
| <input type="checkbox"/> Beverage Sponsor \$7,500 (\$6,740 tax deductible) | <input type="checkbox"/> Volunteer Tent \$1,500 (\$1,180 tax deductible) |
| <input type="checkbox"/> Food Sponsor \$7,500 (\$6,740 tax deductible) | <input type="checkbox"/> Elite Team Entry \$1,000 (\$760 tax deductible) |
| <input type="checkbox"/> Championship Board \$5,000 (\$4,320 tax deductible) | <input type="checkbox"/> Team Entry \$300 (\$220 tax deductible) |
| <input type="checkbox"/> Final Four Board \$3,000 (\$2,320 tax deductible) | <input type="checkbox"/> Audience Member Entry \$50 (\$10 tax deductible) |
- \$50 x number of tickets _____ = \$ _____

Sponsorship Payment:

Check enclosed

Payable to Cystic Fibrosis Foundation. Mail to CFF, 4538 Trousdale Dr. Nashville, TN 37204.

Please invoice

Thank you for your support!

Bob Jackson | Board Co-Chair | The Crichton Group

3011 Armory Drive | Suite 250 | Nashville, TN 37204

o: 615.986.6223 | bjackson@thecrichtongroup.com | thecrichtongroup.com

The Cystic Fibrosis Foundation has unrestricted financial reserves of about 10 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 lifelong 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 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.