



MARCH
18
SATURDAY

SHAMROCKIN' FOR A CURE

TASTE OF SHAMROCKIN'
IN KIND SPONSORSHIP

SATURDAY, MARCH 18TH, 6:00 PM
AMERIS BANK AMPITHEATRE

shamrockinforacure.com

The mission of the Cystic Fibrosis Foundation is to cure cystic fibrosis and to provide all people with CF the opportunity to lead long, fulfilling lives by funding research and drug development, partnering with the CF community, and advancing high-quality, specialized care.



ABOUT SHAMROCKIN'



In 2002, the Baker's only son was diagnosed with cystic fibrosis. Soon after that, their second unborn son was also diagnosed with CF. Hopes of first football games, girlfriends, graduation and grandchildren turned to a future full of hospitals, mounting bills and a parent's worst nightmare. Fast forward to 2009 and ShamRockin' was born out of love and friendship. The idea of throwing a "Party with a Purpose" took off and has grown due to the blood, sweat and tears of hundreds of volunteers and supporters sharing the passion for a cure.

JOIN THE SHAMROCKIN' ARMY

ShamRockin' for a Cure provides a unique experience to give back to the community and have a great time while doing it. When you become a Taste of ShamRockin' in-kind sponsor, you become part of a family committed to curing CF and supporting the mission of the Cystic Fibrosis Foundation to cure cystic fibrosis and to provide all people with CF the opportunity to lead long, fulfilling lives by funding research and drug development, partnering with the CF community, and advancing high-quality, specialized care.

As a Taste of ShamRockin' Sponsor, your restaurant will have the opportunity to showcase your craft to guests through tasting experiences. Your commitment will include the following:

- Provide heavy hors d'oeuvres to showcase a signature sampling of your food (suggested 500 servings)
- Provide at least one chef and/or representative to serve; Up to 3 total restaurant representatives may attend to serve

Taste of ShamRockin' is an integral part of the ShamRockin' experience. With your support we are confident that one day—not one person will lose a child, sibling, parent, or friend to cystic fibrosis. You have an opportunity in your lifetime to be part of defeating this terrible disease.

BENEFITS OF JOINING TASTE OF SHAMROCKIN'



- Two (2) complimentary tickets, FMV \$104 to ShamRockin' in addition to servers representing your restaurant
- Restaurant logo/name in ShamRockin' for a Cure e-newsletter
- Restaurant logo/name and link to your website on ShamRockin' webpage
- Restaurant recognition on social media
- Networking with ShamRockin' for a Cure guests, North Atlanta's best chefs, and local business leaders
- Networking opportunities outside of ShamRockin' for a Cure event (FMV \$30)
- Inclusion on Taste of ShamRockin' event signage

RESTAURANT PARTICIPATION FORM

Restaurant Name

Chef's Name

Contact Name

Address

City

State

Zip

Email

Phone

INSURANCE POLICY INFORMATION

Insurance Contact Name

Insurance Contact Email

Insurance Contact Phone

I agree to participate as a Taste of ShamRockin' In-Kind Sponsor for ShamRockin' for a Cure on March 18, 2023.

I will have staff for my booth to serve samples or arrange with ShamRockin' committee an alternate server.

Signature_____ Date_____

Please provide a description below of what you intend on serving for our guest menu. This also helps us ensure a variety of tastings and an opportunity to highlight your item in promotional materials.

I'd like help: Setting Up Serving

I need access to electricity at the event for my tasting: Yes No

ShamRockin' will provide a banner for your restaurant space. Please let us know if you'd prefer to use yours.

For more information on Taste of ShamRockin, please contact:

Deb Riecke, debriecke@comcast.net, 770.695.3969 or Beth Culloty, mcullotyb@aol.com, 404.434.1886

Return completed form to Linda Blakely, CFF GA Chapter - georgia@cff.org or complete your form online at
shamrockinforacure.com/tasteofshamrockin

CFF Georgia Chapter | 57 Executive Park S, Ste 380 | Atlanta, GA 30329 | georgia@cff.org | 404.325.6973

The Cystic Fibrosis Foundation, a 501(c)(3) nonprofit organization, has unrestricted financial reserves of about 14 times its 2022 operating budget. These reserves are largely a result of the Foundation's successful venture philanthropy model, through which we have raised and invested hundreds of millions of dollars to help discover and develop breakthrough CF therapies. These funds and any future revenue from our model are reinvested into the CF Foundation's mission to cure cystic fibrosis and to provide all people with CF the opportunity to lead long, fulfilling lives. To obtain a copy of our latest Annual Report, visit <https://www.cff.org/about-us/annual-reports-and-financials>, email info@cff.org or call 1-800-FIGHT-CF.





Please Note

SPONSORSHIP DISCLAIMER:

- Events may be subject to change at any time, based on health concerns, including concerns based on guidance from the Centers for Disease Control and Prevention and local health officials. In these instances when events are moved to a virtual format, sponsorship benefits may be adjusted to suit the virtual format.
- Sponsors may be required to wear a mask, if they are interacting with event participants or manning a care and share table, based on venue guidelines, local municipality guidelines and/or the current COVID situation where the event is taking place.

IMPORTANT NOTE ON ATTENDANCE AT FOUNDATION EVENTS

To reduce the risk of COVID-19 the Foundation is taking steps to host safe events for our community. Please be advised that events may be subject to change at any time based on guidance from the Centers for Disease Control and Prevention and local health officials. To help minimize the risk of COVID-19, we ask that attendees at CF Foundation events follow these steps:

- Where mandated, practice physical distancing and maintain at least a safe 6-foot distance from persons outside of their household.
- Face masks are encouraged and should be worn in accordance with local guidelines. (Note, children under two years of age should not wear masks due to safety concerns and therefore should only attend outdoor CF Foundation events where they are able to distance and stay within their family group.)
- Follow basic infection, prevention and control practices by regularly washing hands with soap and water or with an alcohol-based hand gel, and by covering your cough or sneeze with a tissue or your inner elbow.
- Persons with CF should consult their physician before participation in any in-person event as they may be at an increased risk for severe illness from COVID-19.
- For indoor events, it is CFF policy to invite only one person with CF to be in-person at the event at a time. For outdoor events, people with CF should maintain a safe 6-foot distance from persons outside of their household at all times.

FOR YOUR SAFETY AND THE SAFETY OF OTHERS:

- For the health and wellbeing of the CF community, we strongly encourage that participants be fully vaccinated and have received the CDC recommended booster shot when joining us in-person for the event.
- Please be advised certain localities and venues may require you to be fully vaccinated and show proof of vaccination. We will communicate these requirements in event communications.
- We are counting on the fact that you appreciate the potential consequences of exposing members of the CF community to COVID-19 and will not attend an event if you are feeling ill or have been recently exposed to COVID-19.
- We cannot guarantee that all people in attendance at the event, including participants, sponsors, volunteers, vendors, or others, have been fully vaccinated.