



Please join us for the 14th Annual Redbone@Large Nantucket Slam September 10-12, 2017 Nantucket, MA

To register, make a donation or for additional information please visit <https://nantucketslam2017.events.cff.org>

To share the event with friends & family, view 2016 tournament photos and event information visit us on FB



The Nantucket Slam is an all-inclusive two-day competitive catch-and-release fishing tournament designed to please the recreational and avid angler alike.

Anglers are invited with or without their own boats and will be supplied with all necessary tackle and equipment. To accommodate various boat sizes and styles of fishing, three categories are offered: Fly, Spin/Plug and Trolling.

This is a catch and release tournament consisting of two - six anglers per boat. Prizes will be awarded for the largest and most fish per species as well as Division Champion, First Runner Up and Grand Champion.

Accommodations, Marina & Dining

Jared Coffin House a stately three-story mansion in the heart of Nantucket Cottages at the Boat Basin reminiscent of a seaside village with an in-town location. Nantucket Boat Basin world-renowned 240-slip marina.



Sunday Evening: Welcome Reception Nantucket Prime Monday Evening: NE Lobster-bake and auction at the White Elephant Monday & Tuesday: Morning breakfast Tuesday Awards Reception: Featuring Spankey's Famous Raw Bar

On behalf of the Cystic Fibrosis Foundation and the 2017 Nantucket Slam Committee, please join us in September to "Catch the Cure" for cystic fibrosis.

Table listing names of participants and organizers: Kevin Barrett, Peter Bishop, Bob Coughlin, Ralph Hardy, Christina Martin, Billy O'Connor, Christy Bassett Baker, Jim Boushell, Cameron Gammill, Rob Hardy, Jay Miller, Jim Sperzel, George Bassett, Jr., Mike Callahan, Christian Giardini, Lynne Burchell Heyer, Paula Perini Miller, Jeff Bilezikian, Alex Clark, Matt Gutwill, Jeff Heyer, Doug Nelson.



SPACE IS LIMITED. TO ENSURE YOUR LODGING AND BOAT PREFERENCES ARE MET, PLEASE REGISTER ASAP.

For additional information or to register please call Karen Kelly (800) 966-0444 or kkelly@cff.org

The Mission of the Cystic Fibrosis Foundation (CFF) 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. We will not rest until we find a cure for all people with cystic fibrosis.

The Cystic Fibrosis Foundation Is...Hope in Action.

The CFF is the world's leader in the search for a cure. We fund more CF research than any other organization, nearly every CF drug today was made possible by Foundation support.

We are a nonprofit donor-supported organization dedicated to attacking CF from every angle. Our focus is to support the development of new drugs to fight the disease, improve quality of life for those with CF and ultimately to find a cure. The Foundation's drug development model has been recognized by Harvard Business School and by publications such as Forbes, The New Yorker and Bloomberg Businessweek. The CFF is one of the most efficient organizations of its kind and is an accredited charity of the Better Business Bureau's Wise Giving Alliance.

OUTSTANDING CORPORATE SUPPORTER



Compass is a personalized service to help people and families affected by CF with insurance, financial, legal and other issues you are facing. It's free, confidential and available to everyone.



Click here to visit our website To become a teamMate and volunteer with the CFF MA/RI Chapter please contact Liz Feigen at efeigen@cff.org or (800) 966-0444

Photography and Video Notice:

When you attend this Cystic Fibrosis Foundation event, you enter an area where photography, audio and video recording may occur. By entering the event premises, you consent to such recording media and its release, publication, exhibition, broadcasting or reproduction to be used for news, web casts, promotional purposes, telecasts, advertising, and inclusion on websites. You further waive all rights that you may have to any claims in connection with any exhibition, streaming, web casting, televising, or other publication of these materials, regardless of purpose or sponsorship of such exhibiting, broadcasting, web casting or other publication.

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 inviting only one person with CF to attend the indoor portion of a Foundation-sponsored event at a specific time. For the outdoor portion, the Foundation recommends that all people with CF maintain a safe 6-foot distance from each other at all times.

The Cystic Fibrosis Foundation has unrestricted financial reserves of about 12 times its budgeted annual 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, develop lifesaving new therapies and help all people with CF live full, productive lives. To request a copy of our Strategic Plan, email info@cff.org or call 800 FIGHT-CF.