

# grand chefs experience

## CYSTIC FIBROSIS FOUNDATION

### Restaurant Participation Form



**Saturday, March 2, 2024**

Field Museum | 1400 S Lake Shore Drive, Chicago, IL

6:00pm - 12:00am | [www.cffgrandchefs.com](http://www.cffgrandchefs.com)

Contact Hannah Klinnert Sowa at [hklinnert@cff.org](mailto:hklinnert@cff.org) or Olivia Coady at [ocoady@cff.org](mailto:ocoady@cff.org) for more information.



# Driven by a Dream

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.



Gene & Georgetti, 2023

## About the Cystic Fibrosis Foundation

We are driven by a dream that one day – every person with cystic fibrosis will have the chance to live a long, healthy life. Recognized globally, the Cystic Fibrosis Foundation has led the way in the fight against cystic fibrosis, fueling extraordinary medical and scientific progress.

The life expectancy of someone born with CF has doubled in the last 30 years. Despite this progress, many people with CF do not benefit from existing therapies. Our vision is a cure for every person with cystic fibrosis and a life free from the burden of this disease. We will not leave anyone behind. You are an important part of our progress. Together, let's make CF stand for Cure Found.



**CYSTIC FIBROSIS  
FOUNDATION®**

## About Cystic Fibrosis

In people with cystic fibrosis, a defective gene causes a thick buildup of mucus in the lungs, pancreas and other organs. In the lungs, mucus clogs the airways and traps bacteria, leading to infections, extensive lung damage and respiratory failure. Many people with CF do not benefit from existing therapies, either because their disease is too advanced or because their specific genetic mutations will not respond.

Those who can take current therapies continue to face challenging complications--leading to hospitalizations, missed school and work, and significant interruptions in their daily routine. And, many children and adults with CF still face the sobering prospect of a shortened life span.



# Meet Kathy and Kent Duncan

Board members and Grampion Chairs, Kathy and Kent Duncan have been actively fighting to make CF stand for Cure Foundation ever since they learned their grandson, Luca, had cystic fibrosis. As pillars in our Chicago CF community, their vision is clear: a cure for ALL living and impacted by cystic fibrosis. Advances in genetic-based therapies have progressed significantly, but our job is not finished.

“As members of the cystic fibrosis community, we have found a second family of compassionate, loving people. We have felt an incredible amount of support, leaning on these friendships that were formed through understanding. From the Foundation, from our family and our friends, from doctors, from other grandparents, from all of you. It truly takes a village and there is no amount of thanks that will represent our gratitude for all that are in this fight with us.”



[Learn more about the Duncans' CF experience!](#)

## A CURE FOR ALL.

Winning this fight means working harder and faster. Genetic therapies – our best hope for curing cystic fibrosis – are more complex than anything we have ever done. Progressing a genetic therapy could cost the Foundation 10 times more than the development of a novel therapy a decade ago. The Foundation is leading the way in applying this emerging science to CF.

In addition to CF, there are more than 1,000 diseases caused by this type of genetic mutation, virtually all of which have eluded researchers seeking FDA-approved therapies. While our focus is on cutting-edge research for CF, our hope is that future genetic-based therapies could also benefit many others living with genetic diseases. You are an important part of our progress. Corporate donations help accelerate scientific advancements.

Our vision is a **CURE** for **100%** of people living with cystic fibrosis.

**We will not leave anyone behind.**

# 1 IN 31

AMERICANS  
ARE SYMPTOMLESS  
CARRIERS OF THE  
DEFECTIVE CF GENE

# 56

MEDIAN PREDICTED AGE FOR  
SOMEONE BORN WITH CF — MORE  
THAN DOUBLED SINCE THE START  
OF THE CF FOUNDATION

# 0

CURES EXIST FOR  
CYSTIC FIBROSIS



Bar Siena, 2023



## The Grand Chefs Experience

In 2023, The Grand Chefs Experience raised over **\$560,000** in support of the CF Foundation. We look forward to the one-of-a-kind tasting once again, welcoming over **600 supporters** to an exclusive evening of custom cocktails and signature hors d'oeuvres.

The evening includes a spirited live program featuring our honored CF Fighter and the opportunity to support scientific advancements during our bid for a cure and dynamic live auction.

Throughout the evening, food will be abundant, libations will be flowing, and your support will make us one step closer to a CURE.



Adalina, 2023







# 2024 Grand Chefs Experience **Participation Opportunities**

## **Grand Chefs Tasting 6:00PM - 8:00PM**

- Prepare small bites for 600 guests (approximately 300 portions)

## **Nightcap Bartender 9:30PM - 12:00AM**

- Select a donated spirit from Diageo and prepare a signature cocktail for 700 guests (approximately 350 portions)
- Provide necessary juices and mixers

## **Nightcap Pastry Chef 9:30PM - 12:00AM**

- Prepare a small pastry/dessert for 700 guests (approximately 350 portions)

## **Nightcap 'Late-Night' Chef 9:30PM - 12:00AM**

- Prepare a small savory dish for 700 guests (approximately 350 portions)





## You Will Receive:

- Chef/bartender & restaurant name on event materials
- Social media (Facebook & Instagram) posts featuring your participation
- One 6' table for presentation and one 6' table for backup along with black cloths
- Sign with name, logo, and marker to write the dish/drink being served
- Electrical outlet(s) upon request
- Disposables upon request
- Ice upon request
- Opportunity to interact with active CF Foundation supporters, food lovers and corporate supporters from throughout Chicagoland

## We Ask You To:

- Prepare a dish/drink for the appropriate number of guests
- Provide all food, necessary equipment, and decor
- Provide staff to help serve and interact with guests
- Have all items loaded at the Field Museum by the appropriate time
- Donate an item to be used in the silent or live auction
- **Provide a Certificate of Liability Insurance**
- **Provide necessary paperwork as required by The Field Museum (see page 8)**
- Provide head shot, bio, hi-res logo and quote about why you are participating in the Grand Chefs Experience.



# 2024 Grand Chefs Experience Commitment Form



Chef/Bartender Name: \_\_\_\_\_

Restaurant/Bar: \_\_\_\_\_

Phone: \_\_\_\_\_ Email: \_\_\_\_\_

Restaurant Social: \_\_\_\_\_ Chef/Bartender Social: \_\_\_\_\_

Public Relations Contact:

Name: \_\_\_\_\_ Agency: \_\_\_\_\_

Phone: \_\_\_\_\_ Email: \_\_\_\_\_

I hereby consent to the publication, exhibition, reproduction or other release, public or private, by the Cystic Fibrosis Foundation or any of its chapters, officers, or other representatives (including the release of any assignee, transferee, affiliated entity or license) of my name and story, photograph, recording or other likeness (collectively, "Media"), as it relates to the 2024 Grand Chefs Experience

If I cannot be present that night, for any reason, I will let the Cystic Fibrosis Foundation know who my substitute will be. The Foundation cannot be responsible for any items left behind by vendors.

I will provide a Certificate of Liability Insurance and a COI Listing the Field Museum of Natural History as additionally insured.

## **I would like to Participate in the Following:**

- ☐ Chefs Tasting (6:00PM-8:00PM)
- ☐ Nightcap Pastries (9:30PM-12:00AM)
- ☐ Nightcap Bartender (9:30PM-12:00AM)
- ☐ Nightcap Late Night Bites (9:30PM - 12:00AM)

I am excited to donate the following to the auction:

- |  |  |  |
|--|--|--|
| <input type="checkbox"/> Dinner for Four | <input type="checkbox"/> Includes Pairings     | <input type="checkbox"/> Cooking Class   |
| <input type="checkbox"/> Dinner for Six  | <input type="checkbox"/> Includes Kitchen Tour | <input type="checkbox"/> Cocktail Class  |
| <input type="checkbox"/> Dinner for Ten  | <input type="checkbox"/> In-Home Experience    | <input type="checkbox"/> Contact me about contributing a Live Auction Experience |

Fair Market Value: \$\_\_\_\_\_

Restrictions: \_\_\_\_\_

To redeem, please have the winner contact: \_\_\_\_\_

## **Questions?**

Hannah Klinnert Sowa at [hklinnert@cff.org](mailto:hklinnert@cff.org) and 872-265-1008 or Olivia Coady at [ocoady@cff.org](mailto:ocoady@cff.org) and 872-265-1012.

The Cystic Fibrosis Foundation, a 501(c)(3) nonprofit organization, has unrestricted financial reserves of about 10 times its 2023 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](mailto:info@cff.org) or call 1-800-FIGHT-CF.



# INSURANCE COVERAGE REQUIRED

The Cystic Fibrosis Foundation requires proof of insurance to be provided before the event.

The Field Museum requires each Food Vendor to carry the following insurance coverages during the Event:

- Commercial General Liability Insurance with limits of not less than \$2,000,000 per occurrence and \$5,000,000 in the aggregate, including Coverage A, bodily injury and property damage liability and property damage liability, and Coverage B, personal and advertising injury liability and broad form contractual liability coverage, premise operations liability inclusive of broad form contractual liability, products and completed operations, independent contractors, and medical payments. Such liability limits can be satisfied through any combination of primary and excess insurance policies. Coverage shall be no less than the coverage provided under Insurance Services Office Form No. CG 00 01 12 07 or the most recently approved State of Illinois version of this form.
- Workers' Compensation and Employer Liability Insurance as prescribed by applicable law covering all employees who are to provide a service under this Agreement, and Employers Liability coverage with limits of not less than \$1,000,000 each accident or illness.
- Business Automobile Liability Insurance, with limits of not less than \$1,000,000 per occurrence for bodily injury and property damage arising out of owned, non-owned, and hired vehicles coverage Symbol "1".
- Additional Insured. The Museum and the Chicago Park District shall be named as an additional insured on a primary, non-contributory basis on the Commercial General Liability and Business Automobile policies.

The Field Museum requires all restaurants provide names of attendees in advance.

Names of each individual need to be provided in advance of the event for security purposes.

All information needs to be provided to the Cystic Fibrosis Foundation by February 16, 2024.

## IN KIND GUIDELINES

### Important Note on Attendance at Foundation Events

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 COVID19.
- 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.

### CFF Restricted Logo Use Acknowledgement

This Restricted Logo Use Sponsor Acknowledgement ("Acknowledgement") between the Cystic Fibrosis Foundation ("CFF") and the Sponsor, who accepts this Acknowledgement ("Sponsor"). CFF grants to Sponsor a revocable, nonexclusive license to use the Logo (as defined below) in connection with the event to which the Sponsor is contributing ("Event"). Sponsor shall not use the Logo in any manner except as expressly set forth herein from the date of this signed Acknowledgement to up to the Event date. Sponsor agrees to use the Logo only, in connection with the Event and further agrees not to use the Logo in connection with any other sponsors whose logo or brand are deemed to be offensive, defamatory or vulgar. All references to the Logo shall mean the logo associated with the Event. Sponsor agrees not to use the CFF's Logo in a manner inconsistent with proper trademark use, or in any manner that tarnishes the name or reputation of the Cystic Fibrosis Foundation. Sponsor agrees to comply with any requirements established by CFF concerning the style, design, display and use of the Logo with every use of the Logo. Sponsor agrees to send CFF the website link or other relevant materials upon completion of adding the Logo to any promotional materials or third-party sites as part of the sponsorship. Sponsor shall also comply with requests from CFF for additional information, documents, or specimens concerning its use of the Logo. Sponsor agrees to modify or terminate any use of the Logo within 10 days of notice by CFF that, in its sole discretion, such use is not approved. Sponsor agrees that ownership of the Logo and the goodwill relating thereto shall remain vested in CFF both during the period of this license and thereafter. Participant agrees never to challenge, encourage a third-party challenge, or support any challenge to CFF's ownership of the Logo. Sponsor agrees to promptly inform CFF of the use of any marks similar to the Logo and any potential infringements of CFF's Logo that come to Sponsor's attention. Sponsor represents and warrants that it shall not use the Logo in any way that, directly or indirectly, raises any revenue for Sponsor. Sponsor and CFF agree that this license shall be royalty-free. Nothing herein shall be construed as an endorsement by CFF of the Sponsor or the Sponsor's business or activities. The Sponsor agrees and accepts this Acknowledgement upon providing CFF with a sponsorship payment.