

All I Want for Christmas is a...Cure

To Benefit the Cystic Fibrosis Foundation



All I Want for Christmas is a Cure! This year the Cystic Fibrosis Foundation is hosting its fourth annual Holiday Party on Saturday, November 21, 2009. This event will feature a festive evening of Christmas carols, holiday cheer, and other fabulous surprises. The evening will also include upscale live and silent auctions that will be the talk of the town! A variety of items will be highlighted, and an item from your business would be a wonderful addition!

The CF Foundation proudly displays all auction items, incorporating promotional materials of the donors. Your contribution is fully tax-deductible, and the CF Foundation will acknowledge your generosity in the event's program book. With over 400 professionals at this joyous event, we offer great product exposure in front of a lively crowd.

Cystic fibrosis (CF) is a genetic disease affecting 30,000 children and adults in the United States alone; including Tommy and Nico pictured above. The respiratory system and the digestive tract are both compromised by CF, making involuntary activities, like breathing and eating, extraordinarily difficult for these patients on a daily basis. Scientific research is at its peak in this field and nearly ninety cents of every dollar raised by the CF Foundation is donated directly towards finding a cure.

I hope that you will join Tommy and Nico's family in this fight by making a donation to the Cystic Fibrosis Foundation. If you require more information about our Foundation and the work that we do, please contact Elizabeth Cravedi at (312) 236-4491 or ecravedi@cff.org.

On behalf of the children, their families and the CF Foundation, I thank you in advance for your consideration and support.

Sincerely,

Elizabeth Cravedi