Make the Bonnell Foundation part of your Estate planning

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FOR IMMEDIATE RELEASE

HELP FUND A CURE FOR CYSTIC FIBROSIS BY PURCHASING A 2011-12 *PORTRAITS OF CYSTIC FIBROSIS* CALENDAR

(September 28, 2010) The 2011-12 *Portraits of Cystic Fibrosis* calendars are now available. The black and white calendars feature the beautiful faces of people living with Cystic Fibrosis (CF), from toddlers to adulthood. The calendar made its debut in 2003 and continues to be a tribute to the people living with Cystic Fibrosis. Since its inception, $46,000 has been donated to fund CF research through the sale of the calendars. "The Portraits of Cystic Fibrosis calendar is dedicated to everyone with CF fighting to stay healthy. We all have to do our part to help find a cure for cystic fibrosis, so that one day no CF parent will experience the pain and loss of losing a child to this disease" says Laura Bonnell, Founder of The Bonnell Foundation: Living with Cystic Fibrosis.

Cystic Fibrosis is a genetic disease that causes chronic and fatal lung infections and interferes with digestion. The life expectancy for a child born with CF is their mid-30's. CF affects approximately 30,000 people in the United States. There is no cure.

Calendars can be purchased by visiting The Bonnell Foundation: Living with Cystic Fibrosis web site www.roadmaptoCF.org, Click on the order form. Calendars are $15 and checks can be made payable to The Bonnell Foundation. Proceeds from the calendars, purchased through The Bonnell Foundation: Living with Cystic Fibrosis help support research at the University of Michigan Health System, Children's Hospital of Michigan, The CF Foundation and Michigan Pulmonary Disease Community, Inc.

The Bonnell Foundation: Living with Cystic Fibrosis (501(C)3 pending 70908G) was created in 2010 to help parents of CF families, fund research for CF, and offer scholarships (for lung transplants and/or insurance costs) to persons struggling with CF.

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