



Sackville Kinsmen Funspiel  
for Cystic Fibrosis  
Lakeshore Curling Club  
Saturday February 25, 2023



## DONATION REQUEST

Dear Community Friend:

The Kinsmen Club of Sackville is currently organizing its **20<sup>th</sup> ANNUAL FUNSPIEL FOR CYSTIC FIBROSIS** at the Lakeshore Curling Club, Sackville Sports Stadium on **Saturday, February 25, 2023**. All funds raised go directly to Cystic Fibrosis Canada to support their research programs. Through this annual event we have raised over \$110,000 for Cystic Fibrosis (CF).

Your business can contribute to this worthy cause through the following participation opportunities:

- Event Sponsorship – further details on request
- Prizes for participants to thank them for their effort in raising funds
- Auction items
- Promotional items to be given as door prizes

Any donation will be greatly appreciated, and we will ensure your company receives exposure on our prize table and registration table at the event. One of our members will be able to pick up any items that you can donate. Community minded organizations like yours enable us to make a difference for people living with cystic fibrosis.

If your company is interested in becoming more involved in this project, please contact us for information on becoming a Silver, Gold or Platinum Sponsor.

We also welcome you to enter a team in this fun event. We are always seeking new teams to help spread the word and perhaps try out curling for the first time. All are welcome - new and experienced curlers – we have instructors on hand and a beginner's clinic for those who have never curled before.

If you have any questions, please feel welcome to contact us:

**Gerry Archibald (902) 497-6562 or [garchibald38@gmail.com](mailto:garchibald38@gmail.com)**

**Kinsmen Club of Sackville website: [www.sackvillekinsmen.ca](http://www.sackvillekinsmen.ca)**

CF is a fatal, inherited disease that causes progressive damage to the respiratory and digestive systems. Since 1964, Kinsmen and Kinettes have raised more than \$50 million in support of cystic fibrosis research and treatment programs. This support has helped Cystic Fibrosis Canada (CFC) to fund research discoveries that are making a difference in the lives of young Canadians living with CF.

When CFC was established in 1960, most children with CF did not live long enough to attend kindergarten; today, half of all Canadians with CF are expected to live into their 40s and beyond. Although CF remains a devastating disease, this progress is a tremendously encouraging sign that, together, we are charting a course which may lead to a cure, or an effective control.