# **PATIENT ENGAGEMENT IN RESEARCH PLAN**

Patients, survivors, thrivers, caregivers and families (collectively, called "patients") make sure people affected by cancer shape the way that research funding and funding decisions are made.





### WHY ENGAGE?

"Involving patients in the review process of research grants brings a perspective that many researchers are not familiar with. The sense of participating in an important program was rewarding. I found the experience invaluable."

Angus, Patient/Survivor/Caregiver (PSC) Reviewer & Advisory Council on Research (ACOR) member

## **WAYS TO ENGAGE**

#### Lead

We will support patient leadership. Last year CCS welcomed patient members to our Advisory Council on Research (ACOR).

#### **Co-Create**

We will identify issues and co-develop solutions with patient partners. Patient partners engage as Program Advisors and Patient / Survivor / Caregiver Reviewers in all of our current research competitions.

#### Consult

We will seek input and advice from patient partners. For instance, patient partners co-developed our website, reviewed and were featured in our 2022 Research Impact Report.

#### Inform

We will share information in multiple ways. This includes research competition news and results and patient partner experiences in emails, on our website, by video and other ways.





#### COMPENSATION

We created a Patient Partner Compensation Policy designed to facilitate patient participation and to support equity, diversity and inclusion. We offer compensation to all of our patient partners in accordance with our policy.



#### **CONTACT US**

To learn more about being a patient partner and engaging with patient partners, visit our website: cancer.ca/ENpatientengagement. Have questions or comments? Contact: research@cancer.ca.