

## Appendix: Patient Group Conflict of Interest Declaration

1. Did you receive help from outside your patient group to complete this submission? If yes, please detail the help and who provided it.

No

2. Did you receive help from outside your patient group to collect or analyze data used in this submission? If yes, please detail the help and who provided it.

No

3. List any companies or organizations that have provided your group with financial payment over the past two years AND who may have direct or indirect interest in the drug under review.

Company
Abbvie Corp
AMGEN Canada Inc
Astrazeneca Canada
Bayer Inc.
BioCanRX
Boehringer Ingelheim Ltd.
Bristol Myers Squibb Canada
Cdn Partnership Against Cancer Corp
Coalition Priorite Cancer au Quebec
Eli Lilly Canada Inc
E-Z em Canada Inc.
Ferring Pharma
GlaxoSmithKline
Hoffmann-La Roche
Innovative Medicines Canada
Janssen Inc
Merck Canada Inc
Novartis Pharma Canada
Pfizer Canada Inc.
Taiho Pharma Canada

I hereby certify that I have the authority to disclose all relevant information with respect to any matter involving this patient group with a company, organization, or entity that may place this patient group in a real, potential, or perceived conflict of interest situation.

Name: Barry D. Stein  
Position: President  
Patient Group: Colorectal Cancer Canada  
Date: March 22, 2019

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To maintain the objectivity and credibility of the CADTH CDR and pCODR programs, all participants in the drug review processes must disclose any real, potential, or perceived conflicts of interest. This Patient Group Conflict of Interest Declaration is required for participation. Declarations made do not negate or preclude the use of the patient group input. CADTH may contact your group with further questions, as needed.

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Company
Bayer Inc.

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Name: Diana Arajs  
Position: Chair  
Patient Group: Sarcoma Cancer Foundation of Canada  
Date: March 11, 2019

## Patient Input Template for CADTH CDR and pCODR

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Note that this same information can be used for our submissions for:

Abiraterone (Zytiga) for prostate cancer

Lenvatinib (Lenvima) for Hepatocellular Carcinoma

Larotrectinib

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We contracted an independent contractor to interview patients who were taking Lenvatinib for hepatocellular carcinoma for a qualitative submission.

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The consultant collected and analyzed the interviews with patients in order to prepare the quantitative submission for hepatocellular carcinoma.

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Company
Eisai
Bayer
Janssen

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Name: Jackie Manthorne  
Position: President & CEO  
Patient Group: Canadian Cancer Survivor Network  
Date: March 29, 2019

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Company
Bayer

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Name: Christina Sit  
Position: Program Manager  
Patient Group: Lung Cancer Canada  
Date: March 11, 2019

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CCSN was responsible for submitting the larotrectinib patient input to pCODR. I was one member of a large group involved. I reviewed the questionnaire and conducted one patient interview. I do not have knowledge outside of that involvement.

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Please see above

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Company
none

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Name: Antonia Palmer

Position: Co-Founder

Patient Group: Neuroblastoma Canada

Date: May 30, 2019

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Company
N/A

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Name: Sarai Porretta

Position: Administrator

Patient Group: Ontario Parents Advocating for Children with Cancer (OPACC)

Date: May 17, 2019