



# Monoclonal antibodies available in Canada

Barriers and vaccination remain the best means of preventing COVID-19 infection and complications. However, some patients with a hematological cancer are unlikely to have an effective immune response. Passive immunity through SARS-CoV-2 neutralizing antibodies is one way to provide additional protection to individuals with compromised immunity or those at high risk of COVID-19 complications.

## Evusheld (tixagevimab and cilgavimab)

### Authorized for

Pre-exposure prophylaxis of COVID-19 in people who have been recently exposed to SARS-CoV-2.

### Adults and adolescents (≥12 years old and weighing at least 40 kg)

Who are not currently infected with COVID-19 and have not recently been exposed to someone with COVID-19:

- Who have compromised immunity due to an underlying medical condition or immunosuppressive therapy and who are unlikely to have an effective immune response to vaccination OR
- For whom COVID-19 vaccination is not recommended.

### Activity against Omicron

Effective against most variants (Alpha, Beta, Delta, Gamma, Omicron BA.2).  
Decreased neutralizing activity against Omicron subvariants BA.1 and BA.1.1.

### Administration

300 mg: two separate and sequential 1.5-mL injections, preferably in each of the gluteal muscles or, alternatively, in the thigh.

\* Consider increasing the dose to 600 mg in regions where the BA.1 and BA.1.1 subvariants are present.

Monitoring: min. 30 minutes

Interaction with COVID-19 vaccines has not been studied and cannot be excluded. It is recommended to wait 14 days after the most recent vaccine dose.

### Manufacturer

AstraZeneca [https://pdf.hres.ca/dpd\\_pm/00065403.PDF](https://pdf.hres.ca/dpd_pm/00065403.PDF)



## Sotrovimab



**Authorized for** Treatment for a person with COVID-19 (confirmed by NAAT or antigen test) with mild or moderate symptoms for five days or less and not hospitalized due to COVID-19.

**Adults and adolescents (≥12 years old and weighing at least 40 kg)** Who are at high risk of the disease progressing to a form that requires hospitalization and/or results in death.

**Activity against Omicron** Effective against most variants (Alpha, Beta, Delta, Gamma, Omicron BA. 1 and BA1.1).  
Decreased neutralizing activity against Omicron subvariants BA.2 and BA.2.12.1, BA.4 and BA.5.

**Administration** 500 mg, administered as a single intravenous infusion over 60 minutes.  
Monitoring: one hour after the infusion.  
\* Consider increasing the dose to 1 g in regions where the BA.2, BA.2.12.1, BA.4 and BA.5 subvariants are present.

**Manufacturer** GlaxoSmithKline [https://pdf.hres.ca/dpd\\_pm/00062881.PDF](https://pdf.hres.ca/dpd_pm/00062881.PDF)



## Bamlanivimab



**Authorized for** Treatment for a person with COVID-19 (confirmed by NAAT or antigen test) with mild or moderate symptoms for 10 days or less and not hospitalized due to COVID-19.

**Adults and adolescents (≥12 years old and weighing at least 40 kg)** Who are at high risk of the disease progressing to a form that requires hospitalization and/or results in death.

**Activity against Omicron** Not very effective against most Omicron subvariants, so it is not used.

**Administration** 700 mg, administered as a single intravenous infusion over 60 minutes.  
Monitoring: min. one hour after the infusion.

**Manufacturer** Eli Lilly [https://pdf.hres.ca/dpd\\_pm/00060539.PDF](https://pdf.hres.ca/dpd_pm/00060539.PDF)

## Casirivimab and imdevimab

**Authorized for** Treatment for a person with COVID-19 (confirmed by NAAT or antigen test) with mild or moderate symptoms and not hospitalized due to COVID-19.

**Adults and adolescents (≥12 years old and weighing at least 40 kg)** Who are at high risk of the disease progressing to a form that requires hospitalization and/or results in death.

**Activity against Omicron** Not very effective against most Omicron subvariants, so it is not used.

**Administration** 1,200 mg casirivimab + 1,200 mg imdevimab, administered together in a single intravenous infusion over 60 minutes.  
Monitoring: min. one hour after the infusion.

**Manufacturer** Hoffmann-La Roche [https://pdf.hres.ca/dpd\\_pm/00066206.PDF](https://pdf.hres.ca/dpd_pm/00066206.PDF)

**References:** 1. Evusheld Product Monograph, last revised on April 14, 2022. 2. Sotrovimab Product Monograph, last revised on September 14, 2021. 3. Bamlanivimb Product Monograph, last revised on April 14, 2021. 4. Casirivimab and Imdevimab Product Monograph, last revised on June 7, 2022. 5. INESS <https://www.iness.qc.ca/covid-19/traitements-specifiques-a-la-covid-19.html>

## Outpatient antiviral treatments for COVID-19

Antivirals decrease the amount of virus in the body by blocking viral replication. This helps to reduce the severity of COVID-19 infection and shorten the duration of the illness.

### Paxlovid (nirmaltrelvir and ritonavir)

**Authorized for** Treatment for a person with COVID-19 (confirmed by NAAT or antigen test) with mild or moderate symptoms for five days or less and not hospitalized due to COVID-19.

**Adults** Who are at high risk of the disease progressing to a form that requires hospitalization and/or results in death.

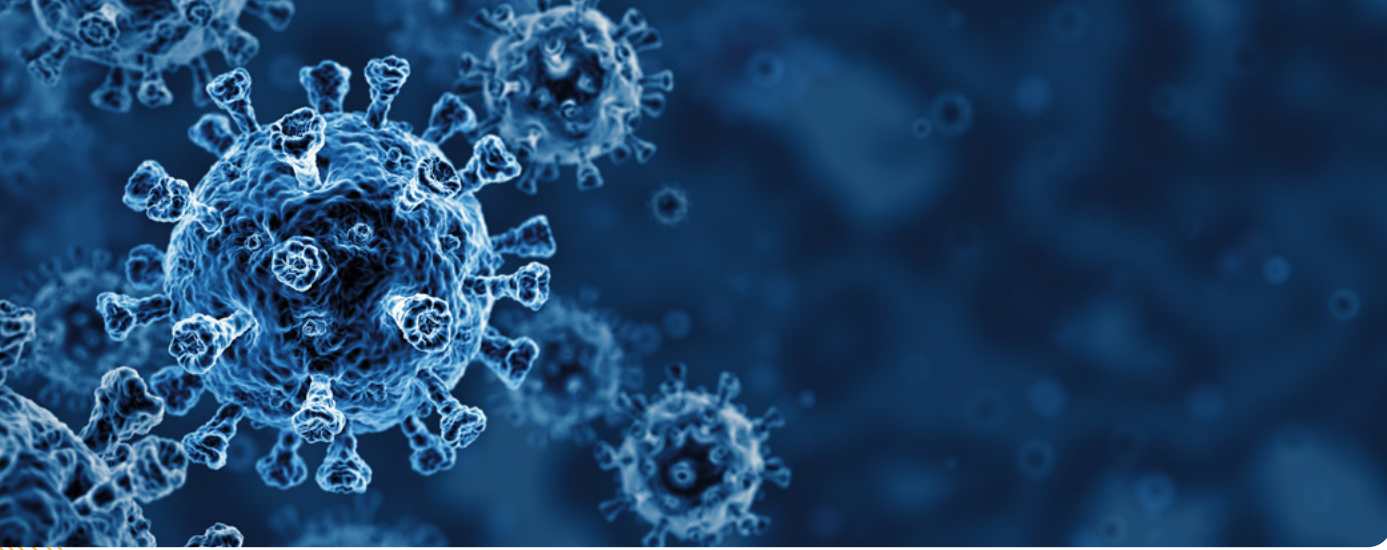
**Activity against Omicron** Effective against most variants, including Omicron.

**Administration** 300 mg nirmaltrelvir (two 150-mg tablets) and 100 mg ritonavir PO taken twice a day with or without food × five days.

\* Adjustment for kidney impairment eGFR ≥ 30 mL/min and < 60 mL/min = 150 mg of nirmaltrelvir and 100 mg of ritonavir twice a day × five days.

Monitor interactions with other drugs.

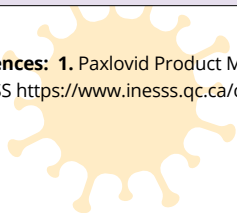
**Manufacturer** Pfizer [https://pdf.hres.ca/dpd\\_pm/00066256.PDF](https://pdf.hres.ca/dpd_pm/00066256.PDF)



## Veklury (remdesivir)

<b>Authorized for</b>	Treatment for a person with COVID-19 (confirmed by NAAT or antigen test) with mild or moderate symptoms for seven days or less and not hospitalized due to COVID-19.
<b>Adults and adolescents (≥12 years old and weighing at least 40 kg)</b>	Who are at high risk of the disease progressing to a form that requires hospitalization and/or results in death.
<b>Activity against Omicron</b>	Effective against most variants, including Omicron.
<b>Administration</b>	Remdesivir 200 mg IV D1 then 100 mg IV D2-3 over 30 to 120 minutes. Not to be administered if CrCl < 30 ml/min. Not to be administered if ALT ≥ 5 ULN.
<b>Manufacturer</b>	Gilead Sciences <a href="https://pdf.hres.ca/dpd_pm/00065539.PDF">https://pdf.hres.ca/dpd_pm/00065539.PDF</a>

**References:** 1. Paxlovid Product Monograph, last updated on June 13, 2021. 2. Veklury Product Monograph, last updated on April 22, 2022. 3. INESS <https://www.iness.qc.ca/covid-19/traitements-specifiques-a-la-covid-19.html>.



This publication was made possible thanks to the support of:

**PHARMACIE  
TORANI ET HADDAD**

**SP  
SpecPharma**  
TRAITEMENTS SPÉCIALISÉS

 LEUKEMIA &  
LYMPHOMA  
SOCIETY OF  
CANADA®

**Never hesitate to contact us, we're here to help!**

1 833 222-4884 • [info@bloodcancers.ca](mailto:info@bloodcancers.ca) • [bloodcancers.ca](http://bloodcancers.ca)