

Recommendations: COVID-19 Vaccine Booster Doses for Adolescents

Published: March 8, 2022

Overview

On January 28, 2021, Canada's National Advisory Committee on Immunization (NACI) released recommendations for the off-label use of booster COVID-19 vaccine doses for adolescents 12-17 years of age at high risk of severe outcomes from COVID-19 due to medical and/or social risk factors.¹ Following this release, the Ministry of Health (MOH) requested the Ontario Immunization Advisory Committee (OIAC) to review NACI's guidance and provide recommendations on the use of booster doses (i.e., an additional dose following the completion of a primary vaccine series) for this age group in Ontario.

At the time of this request, select adolescent groups had already been made eligible for a booster dose by the MOH, including healthcare workers ≥ 16 years of age, Indigenous adolescents ≥ 12 years of age, and consistent with one of the specific NACI recommendations, adolescents ≥ 12 years of age with medical comorbidities; as such, these adolescent groups were out of scope for OIAC recommendations.

The OIAC met on February 9, 2022 to review and discuss current Ontario epidemiology, vaccine effectiveness and safety, and equity considerations to inform adolescent COVID-19 vaccine booster dose recommendations in Ontario. This document provides a summary of the evidence, considerations, and the OIAC's recommendation.

Recommendation

All adolescents 12 to 17 years of age be made eligible for a booster dose of COVID-19 vaccine, using the Pfizer-BioNTech (30 mcg per dose) vaccine at the NACI recommended interval of at least six months (168 days) after completion of the primary COVID-19 vaccine series.

Background

On May 5, 2021, Health Canada expanded the age indication of a primary series of the Pfizer-BioNTech COVID-19 vaccine to 12 years of age and older,² from the previous indication of 16 years of age and older made on December 9, 2020.³ In the context of increasing vaccine supply, Ontario expanded eligibility to all individuals ≥ 12 years of age on May 23, 2021.⁴ As of February 13, 2022, 82.9% of adolescents 12-17 years of age in Ontario had completed a two-dose primary series of COVID-19 vaccine.⁵ Despite high provincial vaccine coverage among adolescents, vaccine coverage varies across Ontario Marginalization Index dimensions quintiles.⁶ For example, lower vaccine coverage has been observed among adolescents (and other age groups) residing in neighbourhoods with high material deprivation as compared to those residing in less deprived neighbourhoods.⁶ As of mid-February 2022, approximately 80% of adolescents who have received two doses are at least 168 days from their last dose.⁵

Booster doses of Pfizer-BioNTech (30 mcg) COVID-19 vaccine were authorized by Health Canada on November 9, 2021 for those ≥ 18 years of age,⁷ but have not yet been authorized for use in those 12-17 years of age. On January 28, 2022 NACI issued off-label recommendations for booster doses of COVID-19 vaccines in adolescents 12 to 17 years of age who may be at higher risk of severe outcomes from COVID-19 infection.¹ This included individuals experiencing biological and/or social risk factors that may intersect, and systemic barriers to accessing health care. NACI recommended that "a booster dose of an mRNA COVID-19 vaccine may be offered ≥ 6 months after completion of a primary COVID-19 vaccine series to adolescents 12 to 17 years of age:

- a) with an underlying medical condition at high risk of severe illness due to COVID-19 (including those who are immunocompromised and who received a three-dose primary series)
- b) who are residents of congregate living settings (e.g., shelters, group homes, quarters for migrant workers, correctional facilities)
- c) who belong to racialized and/or marginalized communities disproportionately affected by COVID-19".¹

In addition to the groups NACI identified based on increased risk of severity, the OIAC deliberated on making booster recommendations for all adolescents in light of the increased transmissibility of the Omicron variant, the anticipated relaxation of public health measures in the coming months, and the goals of the COVID-19 immunization program. The program goals include minimizing serious illness, preserving health system capacity, and reducing transmission to protect high risk populations.⁸

Following the release of NACI's recommendations, several provinces and territories updated their guidance on booster vaccine eligibility for adolescents, with some jurisdictions following NACI's recommendations and others expanding eligibility to all adolescents 12-17 years of age.

Evidence Summary and Considerations

The following summary provides an overview of the evidence reviewed and considerations discussed by the OIAC.

Vaccine Effectiveness

- In the pre-Omicron era, mRNA vaccines demonstrated a high degree of effectiveness against infection with SARS-CoV-2 and symptomatic COVID-19 in adolescents.⁹⁻¹³ Vaccines have also been shown to be highly effective (i.e., >90%) in reducing the risk of serious outcomes in adolescents, including hospitalization and intensive care unit admission,^{12,14,15} as well as multisystem inflammatory syndrome (MIS-C) due to COVID-19.^{16,17}
- Vaccine effectiveness (VE) has been shown to decrease over time. When Delta was the dominant variant, VE against symptomatic infection declined as time since vaccination increased.^{12,18,19} At the time of the OIAC meeting no data on VE, as estimated through studies using the test-negative design, had been published for adolescents in the Omicron era. However, VE against symptomatic Omicron infection has been demonstrated to be lower and to wane faster as compared to VE against Delta in adults.^{20,21} VE against symptomatic Omicron infection has been estimated to be restored to approximately 50-75% in the first three months following a booster dose of mRNA vaccine.^{20,21} Although it is not direct evidence of VE, significant waning of antibodies to both wild-type virus and the Omicron variant have been described in adolescents six months following a second dose COVID-19 mRNA vaccine.²²
- While effectiveness against severe disease appears to be better maintained relative to symptomatic infection, VE against Omicron hospitalizations is lower than against Delta hospitalizations in adults,^{20,21} and this trend may be applicable to adolescents as well. For example, in New York State, surveillance data demonstrated a decreasing protective effect of two doses of vaccine against hospitalization during a time when Omicron became the dominant circulating variant. Unadjusted VE (as estimated by comparing rates of hospitalization by vaccination status) against hospitalization in those 12-17 years of age decreased from 95% in early December 2021 to 75% in early January 2022.²³ However, if hospitalizations with an incidental positive COVID-19 test result were included in the analyses, this would likely reduce VE estimates.
- The absolute risk of COVID-19 hospitalization in adolescents 12-17 years of age is lower relative to older age groups.^{5,24} Despite hospitalization being a rare outcome of COVID-19 among adolescents, Ontario surveillance data have demonstrated the impact of vaccination on adolescent hospitalizations. As of January 30, 2022, the rate of hospitalization among confirmed cases of COVID-19 in Ontario in the last 60 days was approximately four times greater in unvaccinated adolescents 12-17 years of age (0.24 per 100,000 person-days, n=21) compared to those who had completed their primary series (0.06 per 100,000 person-days, n=29).²⁴ Although young adults (i.e., 18-29 year-olds) are also at a low risk of hospitalization relative to older age groups, Ontario surveillance data has demonstrated that providing a booster dose further reduces the risk of hospitalization beyond the risk reduction observed following the primary series. In individuals 18-29 years of age, persons with a booster dose had the lowest hospitalization rate (0.06 per 100,000 person-days) compared to those with only a primary series (0.11 per 100,000 person-days) or those who were unvaccinated (0.44 per 100,000 person-days).²⁴ Similar trends of a reduced risk of hospitalization with increasing number of doses have also been observed in Alberta and the United States, including for young adult age groups.^{15,25}

Vaccine Safety

- Reports of myocarditis or pericarditis following the Pfizer-BioNTech COVID-19 vaccine have been identified through post-marketing vaccine safety surveillance. This surveillance has found consistent trends in higher reporting rates in adolescents and young adults as compared to older adults, males as compared to females (across ages), and after a second dose as compared to a first dose. In Ontario, reporting rates of myocarditis or pericarditis following the Pfizer-BioNTech COVID-19 vaccine have been highest in those 12-17 years of age, although this outcome is still rare with <0.01% individuals in this age group who received a second dose experiencing myocarditis or pericarditis.²⁶ In addition, SARS-CoV-2 infection is known to be associated with a risk of myocarditis.
- There is currently uncertainty regarding the rate of myocarditis or pericarditis following a booster dose and how it may compare to rates observed following first and second doses.^{27,28} In Ontario, few events of myocarditis or pericarditis have been reported in the passive vaccine safety surveillance system following a booster dose, including in those 18-24 years of age.²⁹ In Israel, where a three-month interval between second and booster doses have been used, rates following a booster dose of Pfizer-BioNTech were lower than after dose two but higher than after dose one across age groups, including adolescents.²⁷ A longer interval between doses of the primary series has been associated with a decreased risk of myocarditis or pericarditis.³⁰ However, the impact of the interval between second and booster doses on the risk of myocarditis or pericarditis is unclear but might follow a similar trend to that observed for the primary series. If a longer interval is associated with a reduction in risk of myocarditis or pericarditis, this would support using a longer interval (i.e., six-month interval) between the primary series and a booster dose, and may have additional benefits such as durability of the immune response.

Equity

- Over the course of the pandemic, rates of COVID-19 cases have been higher in neighbourhoods with higher neighbourhood diversity and material deprivation.³¹ The overall (i.e., all ages) rate of COVID-19 hospitalizations has also exhibited a similar trend, with the highest age-standardized rates of hospitalization observed in the most diverse and most deprived neighbourhoods. In addition to these area-level measures, the collection of race-based data in Ontario has demonstrated the risk of COVID-19 infection, hospitalization and death at the individual level has been highly inequitable and racialized.³²
- A targeted approach to booster dose eligibility that focuses only on those adolescents identified by NACI may inadvertently exclude adolescents who experience intersecting risk factors and barriers to accessing health care. Depending on definitions for booster dose eligibility and/or approaches to program implementation, this strategy may not be comprehensive of all adolescents at increased risk of severity. Further, a targeted approach may result in stigmatization of populations identified as being eligible. A universal approach to booster eligibility may reduce this potential stigmatization and increase trust in booster doses among adolescents and their families. A universal approach also offers the opportunity to roll out tailored strategies within universal eligibility in order to reach those most at risk of severe disease. It may also allow for a focus on access, as opposed to defining eligibility, with implementation efforts focused on the higher priority groups as identified by NACI, and supported by local epidemiology and community needs.

Additional Considerations

- While the protection of booster doses against symptomatic infection and transmission may be modest and the duration of this protection is unknown, a booster dose may help with the preservation of in-person learning, and the ability to attend work and participate in recreational activities for adolescents. Further benefits may also extend beyond the individual, including to the contacts of this age group, such as young children who are not yet eligible for vaccination as well as older adults who are at increased risk of severe outcomes.
- Completion of the primary COVID-19 vaccine series should continue to be promoted and prioritized in this age group, given the important impact of two doses at reducing hospitalization as compared to the incremental benefit of a booster dose.²⁴ However, and as outlined by NACI, there are select groups of adolescents who may benefit in particular from a booster dose due to their increased risk of severe outcomes from COVID-19. The additional benefit from a third dose may be of increasing importance for these groups if other layers of protection are reduced as public health measures are relaxed.
- Given the large number of infections that have occurred during the Omicron wave, including among adolescents, there are considerations related to the timing of booster doses for those infected following their primary series. NACI has recommended that these individuals should wait to receive their dose until three months after symptom onset or positive test, as long as six months have also passed since completion of the primary series.³³
- The OIAC acknowledges that many low- and middle-income countries have not had sufficient supply of COVID-19 vaccines to adequately protect individuals at highest risk of severe disease (e.g., older adults, adults with comorbidities) and that much work needs to be done to improve access to COVID-19 vaccines in less resourced jurisdictions.

References

1. Wong E MJ, Ismail S, Stirling R, Zafack J, Forbes N, et al; Public Health Agency of Canada; National Advisory Committee on Immunization. Rapid response: guidance on the use of booster COVID-19 vaccine doses in adolescents 12-17 years of age [Internet]. Ottawa, ON: Government of Canada; 2022 [cited 2022 Feb 14]. Available from: <https://www.canada.ca/content/dam/phac-aspc/documents/services/immunization/national-advisory-committee-on-immunization-naci/guidance-use-booster-covid-19-vaccines-adolescents-12-17-years-age.pdf>
2. Health Canada. Health Canada authorizes use of the Pfizer-BioNTech COVID-19 vaccine in children 12 to 15 years of age [Internet]. Ottawa, ON: Government of Canada; 2021 [cited 2022 Feb 4]. Available from: <https://www.canada.ca/en/health-canada/news/2021/05/health-canada-authorizes-use-of-the-pfizer-biontech-covid-19-vaccine-in-children-12-to-15-years-of-age.html>
3. Health Canada. Health Canada authorizes first COVID-19 vaccine [Internet]. Ottawa, ON: Government of Canada; 2020 [updated 2020 Dec 9; cited 2022 Feb 23]. Available from: <https://www.canada.ca/en/health-canada/news/2020/12/health-canada-authorizes-first-covid-19-vaccine0.html>
4. Government of Ontario. COVID-19 vaccine booking expanding to youth 12+ ahead of schedule [Internet]. Toronto, ON: Government of Ontario; 2021 [cited 2022 Feb 15]. Available from: <https://news.ontario.ca/en/release/1000185/covid-19-vaccine-booking-expanding-to-youth-12-ahead-of-schedule>
5. Ontario Agency for Health Protection and Promotion (Public Health Ontario). COVID-19 data tool [Internet]. Toronto, ON: Queen's Printer for Ontario; 2021 [cited 2022 Feb 15]. Available from: <https://www.publichealthontario.ca/en/Data-and-Analysis/Infectious-Disease/COVID-19-Data-Surveillance/COVID-19-Data-Tool?tab=summary>
6. Ontario Agency for Health Protection and Promotion (Public Health Ontario). COVID-19 vaccination coverage in Ontario by neighbourhood diversity and material deprivation: December 14, 2020 to December 5, 2021 [Internet]. Toronto, ON: Queen's Printer for Ontario; 2021 [cited 2022 Feb 15]. Available from: https://www.publichealthontario.ca/-/media/Documents/nCoV/epi/covid-19-immunization-diversity-deprivation-epi-summary.pdf?sc_lang=en
7. Health Canada. Health Canada authorizes the use of the Pfizer-BioNTech Comirnaty COVID-19 vaccine as a booster shot [Internet]. Ottawa, ON: Government of Canada; 2021 [cited 2022 Feb 17]. Available from: <https://www.canada.ca/en/health-canada/news/2021/11/health-canada-authorizes-the-use-of-the-pfizer-biontech-comirnaty-covid-19-vaccine-as-a-booster-shot.html>
8. Public Health Agency of Canada. Statement from the Council of Chief Medical Officers of Health (CCMOH): COVID-19 vaccination and the use of COVID-19 vaccine boosters [Internet]. Ottawa, ON: Government of Canada; 2022 [updated 2022 Feb 4; cited 2022 Feb 4]. Available from: <https://www.canada.ca/en/public-health/news/2021/11/statement-from-the-council-of-chief-medical-officers-of-health-ccmoh-covid-19-vaccination-and-the-use-of-covid-19-vaccine-boosters.html>
9. Glatman-Freedman A, Bromberg M, Dichtiar R, Hershkovitz Y, Keinan-Boker L. The BNT162b2 vaccine effectiveness against new COVID-19 cases and complications of breakthrough cases: a nation-wide retrospective longitudinal multiple cohort analysis using individualised data. *eBioMedicine*. 2021;72:103574. Available from: <https://dx.doi.org/10.1016/j.ebiom.2021.103574>

10. June Choe Y, Yi S, Hwang I, Kim J, Park YJ, Cho E, et al. Safety and effectiveness of BNT162b2 mRNA Covid-19 vaccine in adolescents. *Vaccine*. 2022;40(5):691-4. Available from: <https://dx.doi.org/10.1016/j.vaccine.2021.12.044>
11. Lutrick K, Rivers P, Yoo YM, Grant L, Hollister J, Jovel K, et al. Interim estimate of vaccine effectiveness of BNT162b2 (Pfizer-BioNTech) vaccine in preventing SARS-CoV-2 infection among adolescents aged 12-17 years - Arizona, July-December 2021. *MMWR Morb Mortal Wkly Rep*. 2021;70(5152):1761-5. Available from: <https://dx.doi.org/10.15585/mmwr.mm705152a2>
12. Powell AA, Kirsebom F, Stowe J, McOwat K, Saliba V, Ramsay ME, et al. Adolescent vaccination with BNT162b2 (Comirnaty, Pfizer-BioNTech) vaccine and effectiveness of the first dose against COVID-19: national test-negative case-control study, England. *medRxiv* 21267408 [Preprint]. 2021 Dec 11 [cited 2022 Feb 25]. Available from: <https://doi.org/10.1101/2021.12.10.21267408>
13. Reis BY, Barda N, Leshchinsky M, Kepten E, Hernan MA, Lipsitch M, et al. Effectiveness of BNT162b2 vaccine against Delta variant in adolescents. *N Engl J Med*. 2021;385(7):2101-3. Available from: <https://dx.doi.org/10.1056/NEJMc2114290>
14. Olson SM, Newhams MM, Halasa NB, Price AM, Boom JA, Sahni LC, et al. Effectiveness of BNT162b2 vaccine against critical Covid-19 in adolescents. *N Engl J Med*. 2022;386(8):713-23. Available from: <https://dx.doi.org/10.1056/NEJMoa2117995>
15. Centers for Disease Control and Prevention. Rates of laboratory-confirmed COVID-19 hospitalizations by vaccination status [Internet]. Atlanta, GA: Centers for Disease Control and Prevention; 2022 [2022 Feb 15]. Available from: <https://covid.cdc.gov/covid-data-tracker/#covidnet-hospitalizations-vaccination>
16. Levy M, Recher M, Hubert H, Javouhey E, Flechelles O, Leteurtre S, et al. Multisystem inflammatory syndrome in children by COVID-19 vaccination status of adolescents in France. *JAMA*. 2022;327(3):281-3. Available from: <https://dx.doi.org/10.1001/jama.2021.23262>
17. Zambrano LD, Newhams MM, Olson SM, Halasa NB, Price AM, Boom JA, et al. Effectiveness of BNT162b2 (Pfizer-BioNTech) mRNA vaccination against multisystem inflammatory syndrome in children among persons aged 12-18 years - United States, July-December 2021. *MMWR Morb Mortal Wkly Rep*. 2022;71(2):52-8. Available from: <https://dx.doi.org/10.15585/mmwr.mm7102e1>
18. Centers for Disease Control and Prevention. Updates to the evidence to recommendation framework: Pfizer-BioNTech vaccine booster doses in 12–15 year olds [Internet]. Atlanta, GA: Centers for Disease Control and Prevention; 2022 [updated 2022 Feb 2; cited 2022 Jan 21]. Available from: https://www.cdc.gov/vaccines/acip/meetings/downloads/slides-2022-01-05/06_COVID_Oliver_2022-01-05.pdf
19. Prunas O, Weinberger DM, Pitzer VE, Gazit S, Patalon T. Waning effectiveness of the BNT162b2 vaccine against infection in adolescents. *medRxiv* 22268776 [Preprint]. 2022 Jan 5 [cited 2022 Feb 25]. Available from: <https://dx.doi.org/10.1101/2022.01.04.22268776>
20. Buchan SA, Chung H, Brown KA, Austin PC, Fell DB, Gubbay JB, et al. Effectiveness of COVID-19 vaccines against Omicron or Delta symptomatic infection and severe outcomes. *medRxiv* 21268565 [Preprint]. 2022 Jan 28 [cited 2022 Feb 25]. Available from: <https://dx.doi.org/10.1101/2021.12.30.21268565>

21. UK. Health Security Agency. COVID-19 vaccine surveillance report Week 6 [Internet]. London: Crown Copyright; 2022 [cited 2022 Feb 25]. Available from: https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/1054071/vaccine-surveillance-report-week-6.pdf
22. Burns MD, Bartsch YC, Boribong BP, Loisel M, Davis JP, Lima R, et al. Durability and cross-reactivity of SARS-CoV-2 mRNA vaccine in adolescent children. medRxiv 22268617 [Preprint]. 2022 Jan 10 [cited 2022 Feb 25]. Available from: <https://doi.org/10.1101/2022.01.05.22268617>
23. New York State. Department of Health. Special report: pediatric COVID-19 update: January 14, 2022 [Internet]. New York, NY: New York State; 2022 [cited 2022 Feb 15]. Available from: https://www.health.ny.gov/press/releases/2022/docs/pediatric_covid-19_hospitalization_report_2021-01-14.pdf
24. Ontario Agency for Health Protection and Promotion (Public Health Ontario). Confirmed cases of COVID-19 following vaccination in Ontario: December 14, 2020 to January 30, 2022 [Internet]. Toronto, ON: Queen's Printer for Ontario; 2022 [cited 2022 Feb 15]. Available from: https://www.publichealthontario.ca/-/media/Documents/nCoV/epi/covid-19-epi-confirmed-cases-post-vaccination.pdf?sc_lang=en
25. Government of Alberta. COVID-19 Alberta statistics [Internet]. Calgary, AB: Government of Alberta; 2022 [cited 2022 Feb 15]. Available from: <https://www.alberta.ca/stats/covid-19-alberta-statistics.htm#vaccine-outcomes>
26. Ontario Agency for Health Protection and Promotion (Public Health Ontario). Myocarditis and pericarditis following vaccination with COVID-19 mRNA vaccines in Ontario: December 13, 2020 to November 21, 2021 [Internet]. Toronto, ON: Queen's Printer for Ontario; 2022 [cited 2022 Feb 25]. Available from: https://www.publichealthontario.ca/-/media/Documents/nCoV/epi/covid-19-myocarditis-pericarditis-vaccines-epi.pdf?sc_lang=en
27. Israel. Ministry of Health, Division of Epidemiology. [Vaccine safety] [Webinar]. Jerusalem: State of Israel; 2022 [presented 2022 Jan 17; cited 2022 Feb 15]. Available from: https://www.gov.il/BlobFolder/reports/vaccine-efficacy-safety-follow-up-committee/he/files_publications_corona_vaccine-safety-17012022.pdf
28. Patone M, Mei XW, Handunnetthi L, Dixon S, Zaccardi F, Shankar-Hari M, et al. Risk of myocarditis following sequential COVID-19 vaccinations by age and sex. medRxiv 21268276 [Preprint]. 2021 Dec 25 [cited 2022 Feb 25]. Available from: <https://dx.doi.org/10.1101/2021.12.23.21268276>
29. Ontario Agency for Health Protection and Promotion (Public Health Ontario). Weekly summary: adverse events following immunization (AEFIs) for COVID-19 in Ontario: December 13, 2020 to February 6, 2022 [Internet]. Toronto, ON: Queen's Printer for Ontario; 2022 [cited 2022 Feb 15]. Available from: <https://www.publichealthontario.ca/-/media/documents/ncov/epi/covid-19-aeFi-report.pdf?la=en>
30. Buchan SA, Seo CY, Johnson C, Alley S, Kwong JC, Nasreen S, et al. Epidemiology of myocarditis and pericarditis following mRNA vaccines in Ontario, Canada: by vaccine product, schedule and interval. medRxiv 21267156 [Preprint]. 2021 Dec 5 [cited 2022 Feb 25]. Available from: <https://dx.doi.org/10.1101/2021.12.02.21267156>

31. Ontario Agency for Health Protection and Promotion (Public Health Ontario). Weekly epidemiologic summary: COVID-19 in Ontario – focus on January 30, 2022 to February 5, 2022 [Internet]. Toronto, ON: Queen’s Printer for Ontario; 2022 [cited 2022 Feb 15]. Available from: https://www.publichealthontario.ca/-/media/Documents/nCoV/epi/covid-19-weekly-epi-summary-report.pdf?sc_lang=en
32. McKenzie K DS, Peterson S. Tracking COVID-19 through race-based data [Internet]. Toronto, ON: Ontario Health; 2021 [cited 2022 Feb 25]. Available from: <https://www.ontariohealth.ca/our-work/equity-inclusion-diversity-and-anti-racism/report-covid-19-race-based-data#:~:text=This%20report%20examines%20race%2Dbased,infection%20compared%20with%20white%20Ontarians>
33. Forbes N, Krishnan R, Ismail S, Salvadori M, Warshawsky, B, Young K, et al; Public Health Agency of Canada, National Advisory Committee on Immunization. Rapid response: updated guidance on COVID-19 vaccination timing for individuals previously infected with SARS-CoV-2 [Internet]. Ottawa, ON: Government of Canada; 2022 [cited 2022 Mar 3]. Available from: <https://www.canada.ca/content/dam/phac-aspc/documents/services/immunization/national-advisory-committee-on-immunization-naci/naci-rapid-response-updated-guidance-covid-19-vaccination-timing-individuals-previously-infected-sars-cov-2.pdf>

About the Ontario Immunization Advisory Committee

The OIAC is a multidisciplinary scientific advisory body that provides evidence-based advice to Public Health Ontario on vaccines and immunization matters including vaccine program implementation in Ontario, priority populations and clinical guidance. The focus of the OIAC's work is on publicly-funded vaccines and immunization programs in Ontario, including COVID-19 and those under consideration for new programming.

For more information about the OIAC and its members contact secretariat@oahpp.ca.

Acknowledgements

The statement was prepared by the OIAC Secretariat on behalf of the OIAC. The OIAC also acknowledges the contribution of PHO staff within Health Protection, Health Promotion, Chronic Disease and Injury Prevention (Health Equity Team), Communications Services and Library Services.

OIAC Members

Dr. Jessica Hopkins, co-chair
Chief Health Protection and
Emergency Preparedness Officer
Public Health Ontario

Dr. Jeffrey Pernica, co-chair
Head, Division of Infectious Disease
Department of Pediatrics
McMaster University

Dr. Juthaporn Cowan
Associate Scientist
The Ottawa Hospital Research Institute

Dr. Vinita Dubey
Associate Medical Officer of Health
Toronto Public Health

Dr. Julie Emili
Associate Medical Officer of Health
Region of Waterloo

Dr. Mariam Hanna
Pediatric Allergist and Clinical Immunologist

Susie Jin
Pharmacist

Dr. Allison McGeer
Professor, Laboratory Medicine and Pathobiology
Dalla Lana School of Public Health
University of Toronto

Dr. Justin Presseau
Scientist
The Ottawa Hospital Research Institute

Dr. Maurianne Reade
Family Physician; Associate Professor
Northern Ontario School of Medicine

Richard San Cartier
Clinical Team Lead
N'Mninoeyaa Aboriginal Health Access Centre

Fairleigh Seaton
Director, Infectious Disease Prevention
and Environmental Health
Kingston, Frontenac and Lennox & Addington
Public Health

Dr. Wendy Whittle
Maternal Fetal Medicine Specialist
Mount Sinai Hospital

OIAC Ex-Officio Members

Tara Harris
Manager
Immunization and Emergency Preparedness
Public Health Ontario

Robert Lerch
Director (Acting)
Health Protection and Surveillance Policy and
Programs Branch
Ministry of Health

Dr. Daniel Warshafsky
Associate Chief Medical Officer of
Health (Acting)
Office of Chief Medical Officer of Health,
Public Health, Ministry of Health

Dr. Sarah Wilson
Public Health Physician and Medical
Epidemiologist
Public Health Ontario

Citation

Ontario Agency for Health Protection and Promotion (Public Health Ontario). Recommendations: COVID-19 vaccine booster doses for adolescents. Toronto, ON: Queen's Printer for Ontario; 2022.

Disclaimer

This document was prepared by the OIAC for Public Health Ontario (PHO). The OIAC provides evidence-based advice to PHO on vaccines and immunization matters. The OIAC's work is guided by the evidence available at the time this document was prepared. The application and use of this document is the responsibility of the user. PHO assumes no liability resulting from any such application or use. This document may be reproduced without permission for non-commercial purposes only and provided that appropriate credit is given to PHO. No changes may be made to this document without prior and expressed written permission from PHO.

Questions about the information in this document can be sent to secretariat@oahpp.ca.

Public Health Ontario

Public Health Ontario is an agency of the Government of Ontario dedicated to protecting and promoting the health of all Ontarians and reducing inequities in health. Public Health Ontario links public health practitioners, front-line health workers and researchers to the best scientific intelligence and knowledge from around the world.

For more information about PHO, visit publichealthontario.ca

© Queen's Printer for Ontario, 2022

