

Supplement Article

Crohn's and Colitis Canada's 2021 Impact of COVID-19 and Inflammatory Bowel Disease in Canada: COVID-19 Vaccines—Biology, Current Evidence and Recommendations

Sanjay K. Murthy, MD, MSc¹, M. Ellen Kuenzig, PhD^{2,3,6}, Joseph W. Windsor, PhD^{4,6}, Jean-Eric Ghia, PhD⁵, Anne M. Griffiths, MD^{2,3,6,7}, Remo Panaccione, MD⁸, Cynthia H. Seow, MBBS(Hons), MSc^{4,9}, Eric I. Benchimol, MD, PhD^{2,3,6,7}, Charles N. Bernstein, MD^{10,11}, Alain Bitton, MD¹², James Guoxian Huang, MBBS², Jennifer L. Jones, MD, MSc¹³, Kate Lee, MBA, PhD¹⁴, Gilaad G. Kaplan, MD, MPH^{4,9}, Mariam S. Mukhtar, MD¹⁵, Parul Tandon, DO¹⁶, Laura E. Targownik, MD, MSc¹⁶, Deanna L Gibson, PhD¹⁷

¹The Ottawa Hospital IBD Centre, Department of Medicine, University of Ottawa, Ottawa, Ontario, Canada; ²SickKids Inflammatory Bowel Disease Centre, Division of Gastroenterology, Hepatology and Nutrition, The Hospital for Sick Children, Toronto, Ontario, Canada; ³Child Health Evaluative Sciences, SickKids Research Institute, Toronto, Ontario, Canada; ⁴Department of Medicine, University of Calgary, Calgary, Alberta, Canada; ⁵Department of Immunology & Internal Medicine section of Gastroenterology, Max Rady College of Medicine, Rady Faculty of Health Sciences, University of Manitoba and University of Manitoba Inflammatory Bowel Disease Clinical and Research Centre, Winnipeg, Manitoba, Canada; ⁶ICES, Toronto, Ontario, Canada; ⁷Department of Paediatrics and Institute of Health Policy, Management and Evaluation, University of Toronto, Toronto, Ontario, Canada; ⁸Division of Gastroenterology and Hepatology, Department of Medicine, University of Calgary, Calgary, Alberta, Canada; ⁹Department of Community Health Sciences, University of Calgary, Calgary, Alberta, Canada; ¹⁰Department of Internal Medicine, Max Rady College of Medicine, Rady Faculty of Health Sciences, University of Manitoba, Winnipeg, Manitoba, Canada; ¹¹University of Manitoba IBD Clinical and Research Centre, Winnipeg, Manitoba, Canada; ¹²Department of Medicine, McGill University Health Centre, McGill University, Quebec, Canada; ¹³Department of Medicine, Dalhousie University, Halifax, Nova Scotia, Canada; ¹⁴Crohn's and Colitis Canada, Toronto, Ontario, Canada; ¹⁵Department of Internal Medicine, King Abdulaziz University Hospital, Jeddah, Saudi Arabia; ¹⁶Division of Gastroenterology and Hepatology, Mount Sinai Hospital, University of Toronto, Toronto, Ontario, Canada; ¹⁷Department of Biology, Faculty of Science; Department of Medicine, Faculty of Medicine, The University of British Columbia, Okanagan campus, Kelowna, British Columbia, Canada.

Correspondence: Sanjay K. Murthy, MD, MSc, FRCPC, The Ottawa Hospital IBD Centre, Department of Medicine, University of Ottawa, 501 Smyth Road, Ottawa, ON, K1H 8L6, Canada, e-mail: smurthy@toh.ca

ABSTRACT

The COVID-19 pandemic has ushered a globally focused vaccine development program that produced multiple successful vaccines within a year. Four SARS-CoV-2 vaccines have been approved for use in Canada, using two different technologies, all of which have shown excellent efficacy in reducing the rate of symptomatic COVID-19 infection and 100% efficacy in preventing death from COVID-19. People with inflammatory bowel disease (IBD), like many others with immune-mediated chronic diseases, were excluded from the pivotal trials of these vaccines, leading to early hesitancy by regulatory bodies to endorse administering the vaccines to these groups. However, recent data has shown that the adverse event rate to SARS-CoV-2 vaccine among people with IBD is similar to the general population. Early data has further shown that people with IBD are capable of mounting a robust immune response to SARS-CoV-2 vaccines, particularly following a second dose, whereas the response to the first dose is blunted in those receiving anti-TNF therapy or conventional immunosuppressants (azathioprine, 6-mercaptopurine, methotrexate). Based on these data and evidence from previous vaccine

Received: June 25, 2021; Accepted: August 14, 2021.

© The Author(s) 2021. Published by Oxford University Press on behalf of the Canadian Association of Gastroenterology. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted reuse, distribution, and reproduction in any medium, provided the original work is properly cited.

Key Messages

- Although people with IBD were excluded from the SARS-CoV-2 vaccine trials, real-world evidence suggests that these vaccines are safe and elicit robust immune responses and protect against SARS-CoV-2 infection following two doses of mRNA vaccine.
- The effectiveness of the first dose of vaccine (for a two-dose vaccine) is limited (reduced immunogenicity) for people with IBD receiving anti-TNF therapy or conventional immunosuppressants (e.g., prednisone, azathioprine, 6-mercaptopurine, methotrexate).
- National and international expert panels recommend that all people with IBD receive a complete series SARS-CoV-2 vaccine at the earliest opportunity, irrespective of vaccine type, disease status, or treatment.

programs among people with IBD, multiple national and international expert panels have recommended that individuals with IBD receive complete vaccination against SARS-CoV-2 as soon as possible.

Keywords: COVID-19; Crohn's disease; Inflammatory bowel disease; SARS-CoV-2; Ulcerative colitis; Vaccination; Vaccine

Introduction

In response to the COVID-19 pandemic, international pharmaceutical companies have rapidly developed, tested and produced highly effective vaccines against the Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) virus. More than 200 COVID-19 vaccine candidates are under development or in clinical trials, using both traditional (inactivated or live attenuated vaccines) and newer (recombinant protein vaccines, vectored vaccines and RNA and DNA vaccines) technologies (1,2). Four vaccines having now been approved for use in Canada ([COVID-19 Vaccines: Authorized vaccines: Canada.ca](https://www.canada.ca/en/health-canada/services/covid-19/vaccines/authorized-vaccines.html)).

Many individuals with immune-mediated chronic diseases, including all persons with inflammatory bowel disease (IBD), were excluded from the clinical trials that ultimately led to the approval of these vaccines. Thus, there was initially some uncertainty regarding efficacy and safety of these vaccines in people with IBD. This article reviews the immunology, mechanisms and evidence for currently available (as of writing: May 25, 2021) COVID-19 vaccines and discusses the potential risks and benefits of people with IBD being vaccinated.

IMMUNITY AND COVID-19: NATURAL AND VACCINE IMMUNITY

Natural COVID-19 infection often results in exaggerated and dysfunctional innate and adaptive immune responses (3,4). Inflammasome activation and increases in inflammatory monocytes and neutrophils lead to an overexuberant cytokine storm of IL-1 β , IL-6 and TNF- α , exacerbating severe vascular and respiratory insults (5). Indeed, late-stage pathology in COVID-19 is driven primarily by host immune responses to SARS-CoV-2 (4). Furthermore, evidence reveals T follicular helper cell differentiation may be faulty in individuals who develop lethal COVID-19 infection, given the lack of mature germinal centers and resultant impairment

in the somatic hypermutation process required for class switching to IgG in these individuals (6). Conversely, some studies have found that convalescent patients have circulating T follicular helper memory cells, which are positively correlated with spike protein-specific neutralizing IgG, IgM and IgA antibodies (7). While neutralizing antibodies will afford protection against SARS-CoV-2, a misguided humoral response contributes to some of the immunopathology of COVID-19, including autoantibody production (8). These autoantibodies not only impair immune function and exacerbate COVID-19 severity but also have long-lasting potential for systemic autoimmune diseases post-infection. Improved understanding of the natural immune response will be instrumental in developing an effective vaccine and other therapeutic strategies.

Controlling COVID-19 requires effective and safe vaccines that provide disease-attenuating immunity, including robust T-cell and B-cell memory responses with the development of germinal centers and high-affinity neutralizing antibodies against COVID-19 and any variants. As variants of concern circulate in the population, several types of vaccines may become important in achieving herd immunity (when 70–95% of the population is immune) (9). Vaccine-induced protective immunity is safer and likely to be more effective than natural immunity. There is no evidence to suggest that T follicular helper cells or memory B cells responses would be blunted relative to natural infection or that vaccination would produce the plethora of maladaptive immune and autoimmune reactions seen with natural COVID-19 infection (e.g., cytokine storm or multisystem inflammatory syndrome in children). While some early research suggested immunity might be short-lived to SARS-CoV-2 infection, recent studies have shown that humoral response is long-lived and antigen-specific, including to the spike protein (10), providing good rationale for using the spike protein as a vaccine candidate—as is the case with all currently approved SARS-CoV-2 vaccines in Canada.

CURRENTLY APPROVED SARS-COV-2 VACCINES: MECHANISMS, EFFICACY AND SAFETY

Four SARS-CoV-2 vaccines have received Health Canada authorization for use in persons aged 18 years or older; one of these (Pfizer-BioNTech) has recently received approval for use in persons aged 12 and older (*COVID-19 Vaccines: Authorized vaccines - Canada.ca*): 1) Pfizer-BioNTech BNT162b2 mRNA vaccine (Pfizer, Inc., NY; BioNTech SE, Mainz, Germany); 2) NIH-Moderna mRNA-1273 vaccine (Moderna, Inc., Cambridge); 3) AstraZeneca ChAdOx1 nCoV-19 adenovirus vector vaccine (AstraZeneca plc, Cambridge, England) and 4) Janssen Ad26.COV2.S adenovirus vector vaccine (Janssen Pharmaceuticals, Beerse, Belgium). Large clinical trials have demonstrated these vaccines to be highly efficacious and safe (11–14). Mild side effects, such as injection site reactions (muscle pain, redness and swelling), fatigue, malaise, headaches, joint pains, low-grade fevers and chills were common in all trials. Serious adverse events were rare, and no deaths attributable to the vaccine were encountered in any trials. Importantly, the approved vaccines do not demonstrate the potential for virus activation or integration into the human genome.

MESSENGER RNA (mRNA) VACCINES

The Pfizer-BioNTech vaccine (approved by Health Canada December 9, 2020) and the NIH-Moderna vaccine (approved by Health Canada December 23, 2020) use a lipid nanoparticle delivery system to transport modified SARS-CoV-2 genetic material (messenger ribonucleic acid [mRNA]) encoding the virus' spike protein to host cells (11,12). The viral mRNA enters host cells and uses its translational machinery to produce copies of the spike protein, which then embed into the host's cell membranes, prompting an adaptive immune response resulting in memory T and B lymphocytes producing neutralizing antibodies to the spike protein. These lymphocytes recognize and fight future SARS-CoV-2 infection faster and more effectively than their precursors. Both vaccines are administered as two intramuscular injections 21 (Pfizer) or 28 (Moderna) days apart. To vaccinate a maximum number of individuals amid a vaccine shortage, the National Advisory Council on Immunization (NACI) has recommended that the doses may be administered up to 16 weeks apart, based on suggestion of high-level protection against COVID-19 beyond 14 days following the first dose of either of these vaccines (11,12,15).

A large multinational RCT in 43,548 persons, 16 years of age or older who were healthy or had stable chronic disease (excluding immune-mediated inflammatory diseases) demonstrated 95.3% efficacy of the Pfizer-BioNTech vaccine in reducing symptomatic COVID-19 infections at least seven days after the second dose and 90% efficacy in reducing severe disease leading to hospitalization or death (one versus nine

severe COVID-19 cases in vaccinated versus unvaccinated individuals) (11). A similarly designed RCT conducted in the US in 30,420 persons aged 18 years or older who were healthy or had stable chronic disease demonstrated 94.1% efficacy of the NIH-Moderna vaccine at least 14 days after the second dose and 100% efficacy in reducing severe disease leading to hospitalization or death (12). Recent data have shown these vaccines to be more than 90% effective out to six months without any serious safety concerns (16).

ADENOVIRUS VECTOR VACCINES

The AstraZeneca vaccine (approved by Health Canada February 26, 2021) and the Janssen Ad26.COV2.S vaccine (approved by Health Canada on March 5, 2021) use a non-replicating adenovirus vector to deliver SARS-CoV-2 DNA encoding the spike protein to human cells, which uses the host machinery to produce viral spike protein (13). The AstraZeneca vaccine is administered as two intramuscular injections between four and 12 weeks apart, while the Janssen vaccine is administered as a single intramuscular injection.

A pooled interim analysis of 11,636 healthy adults randomized to either active vaccine or control across four RCTs held in the UK, Brazil and South Africa showed 66.7% efficacy of the AstraZeneca ChAdOx1 nCoV-19 vaccine in preventing symptomatic COVID-19 infection at least 14 days after the second dose and 100% efficacy in preventing hospitalization and/or death from SARS-CoV-2 (13). Further exploratory analyses have shown that vaccine efficacy between day 22 and day 90 following a single standard dose was 76.0%; effectiveness ≥ 14 days after the second dose among those who received a second dose of vaccine ≥ 12 weeks after the first dose was 81.3%. In an RCT of 44,325 healthy adults, the Janssen vaccine was 66.3% effective at preventing laboratory-confirmed symptomatic COVID-19 infection 14 days post-vaccination, 93.1% effective at preventing hospitalization for severe COVID-19 and 100% effective at preventing death due to COVID-19 (17).

VACCINE-INDUCED IMMUNE THROMBOTIC THROMBOCYTOPENIA (VITT)

Recent rare reports of cerebral and systemic venous thromboembolic events (VTEs), associated with thrombocytopenia and hemorrhage with both the AstraZeneca and Janssen vaccines have raised concerns regarding the risk-benefit ratio of these vaccines, particularly among younger individuals who are at very low risk of dying from COVID-19 (18–20). Termed vaccine-induced immune thrombotic thrombocytopenia (VITT), the VTE that develop with VITT have been often associated with life-threatening cerebral VTE (21,22). While the rate of VITT with the AstraZeneca vaccine was initially suggested to be about 1 in 250, persons (18), more recent data have suggested that the rate may be much higher (1 in 60,000

persons, 0.0017%) (20). The risk is expected to be similar for people with IBD as the mechanism of VTE development in VITT differs from traditional VTE (22).

In light of the risk of VITT, NACI has issued a statement that mRNA vaccines are preferred over non-mRNA vaccines. These facts have also prompted several provincial health authorities to suspend administration of the AstraZeneca vaccine as the first dose until more data are available and to consider administering an mRNA vaccine for the second dose among those who received the AstraZeneca vaccine for the first dose (20,23). These policy changes have also come in the face of increasing supply of mRNA vaccines, diminishing supply of AstraZeneca vaccine and decreasing rates of COVID-19 in society.

Importantly, the background risk of VTE in the Canadian population is roughly 0.1% (24), and the risk of VTE among individuals hospitalized with COVID-19 is as high as 15% (25). Considering the adverse impacts of COVID-19, the benefits of these vaccines in preventing symptomatic and severe COVID-19 may still outweigh the potential risks for individuals who have risk factors for acquiring COVID-19 or for having an adverse outcome with COVID-19, such as those who are elderly, have multimorbidity, or live in a region with high SARS-CoV-2 transmission rates (particularly of novel variant strains), and especially in regions where mRNA vaccines are in short supply (18).

VACCINE PRODUCTION AND DISTRIBUTION IN CANADA

For many years, Canada was a leader in vaccine production (26). While plants in Toronto and Montreal have continued producing vaccines, they are not equipped to manufacture the COVID-19 vaccines, resulting in the Canadian government seeking vaccines from the U.S., Europe and India. (*Canadian Health Policy*, December 2020. ISSN 2562-492). However, this led to a lag in Canadian vaccine procurement and vaccination efforts when Moderna and Pfizer reduced vaccine production to build capacity for larger-scale production. Further, transport and storage logistics of existing vaccines coming from Europe and the US to a central repository in Canada (and later to regional distribution centres and provincial or territory points of care) has been challenging for some of the COVID-19 vaccines, particularly those requiring storage at -20°C to -80°C .

The Canadian government plans to increase the domestic production capability of COVID-19 vaccines (27). First, a new facility will be created for the National Research Council in Montreal, where ten million Novavax vaccines will be made, pending Health Canada approval (as of writing, the Novavax SARS-CoV-2 vaccine [NVX-CoV2373] is completing Phase 3 trials with healthy adults in the United Kingdom and is still pending approval by Health Canada [application made on January 29, 2021]; Novavax is a recombinant nanoparticle vaccine utilizing the spike glycoprotein of the SARS-CoV-2 virus).

Additionally, companies that have demonstrated past success in manufacturing capabilities have received massive government investment. For example, VIDO-interact in Saskatoon has been allocated upwards of \$45 million to build a manufacturing plant, and Medicago has been allocated upwards of \$170 million to expand its manufacturing capacity, in addition to a specific order of 76 million doses of its vaccine, pending Health Canada approval.

SARS-COV-2 VACCINE RECOMMENDATIONS FOR PEOPLE WITH IBD

People with chronic immune-mediated diseases such as IBD were excluded from the trials that evaluated the currently approved SARS-CoV-2 vaccines; therefore, concerns have been raised by healthcare providers and the IBD community regarding the safety and efficacy of SARS-CoV-2 vaccines for people with IBD, as well as the influence of drug therapy on vaccine immunogenicity and safety. Importantly, as none of the approved vaccines are live attenuated vaccines, there is no reason to suspect that individuals with IBD receiving immunosuppressive therapy would be at increased risk of virus reactivation. Moreover, multiple observational studies have now been published on outcomes of SARS-CoV-2 vaccination in people with IBD with no cause for concern indicated by any.

In a study of 246 people with IBD who received a SARS-CoV-2 vaccine, the overall rate of adverse events was similar to the general population, while biologic therapy was associated with fewer adverse events, possibly due to blunting of an aggressive immune response (28). A study of 1500 individuals with IBD in the UK (CLARITY-IBD) reported that infliximab treatment was associated with a less robust immune response to the first dose of the Pfizer or AstraZeneca vaccine as compared to vedolizumab. Moreover, individuals receiving concomitant azathioprine or methotrexate had reduced seroconversion with both infliximab and vedolizumab (29). Similarly, an earlier UK study of 7000 people with IBD reported that serological responses to SARS-CoV-2 infection were attenuated in individuals receiving infliximab relative to those receiving vedolizumab and further blunted by concomitant azathioprine or methotrexate (30). Overall, these findings are consistent with observations of reduced immunogenicity to other vaccines in people with IBD receiving anti-TNF therapy (alone or in combination) and traditional immunosuppressive therapies, such as azathioprine, 6 mercaptopurine, methotrexate and higher doses of corticosteroids (prednisone ≥ 20 mg per day), including those for pneumococcus (31–34), influenza viruses (35–39), hepatitis B virus (40), hepatitis A virus (41) and herpes zoster virus (42). Immune responses to conventional vaccines do not appear to be impacted by vedolizumab (influenza and hepatitis B) (43) or ustekinumab (influenza, pneumococcal, tetanus) (44,45).

Importantly, in the CLARITY-IBD study, seroconversion was robust following the second dose of vaccine and in individuals who received a dose of vaccine following recovery from COVID-19 infection. Furthermore, a national US cohort study of nearly 15,000 predominantly white males with IBD receiving a wide spectrum of medications in the Veterans Health Administration System reported that the risk of infection with SARS-CoV-2 was 1.34% in those who were unvaccinated and 0.11% among those who were at least seven days from their second dose of mRNA vaccine (80.4% vaccine effectiveness) (46).

Canadian (47), European (48) and international (49) organizations recommend that people with IBD be vaccinated against SARS-CoV-2 at the earliest opportunity, irrespective of vaccine type, disease status or treatment and without interruption of scheduled therapy. Crohn's and Colitis Canada further recommends that persons with IBD receive a scheduled second dose of vaccine three to four weeks following the first dose to boost immunity, rather than extending the interval for the second dose by up to 16 weeks as for the general population. As individuals with IBD receiving immunosuppressive medications may be at higher risk of contracting severe COVID-19 than the general public (50,51), vaccination is particularly relevant in this group. NACI now "preferentially recommends that a complete two-dose vaccine series with an mRNA COVID-19 vaccine (Pfizer-BioNTech, Moderna) should be offered to individuals in the authorized age group, including those who are immunosuppressed, have an autoimmune condition, are pregnant or are breastfeeding." Furthermore, it recommends that "with the increase of COVID-19 vaccine supply in Canada, second doses should be offered as soon as possible, with priority given to those at highest risk of severe illness and death from COVID-19 disease" (*NACI updated COVID-19 vaccine statement, May 28, 2021: Summary—Canada.ca*).

Regardless of vaccination status, individuals with IBD should remain vigilant with practicing recommended public health measures to prevent SARS-CoV-2 infection and transmission. Given the impaired serological responses to SARS-CoV-2 among people on immunosuppressive treatment, serological testing and virus surveillance should be considered to detect suboptimal vaccine responses, infection persistence and viral evolution to inform public health policy.

Vaccine recommendations for special populations with IBD (children and adolescents, pregnant people and seniors) are provided in articles dedicated to those populations (Benchimol, this volume; Bernstein, this volume).

CONCLUSION

The collective efforts of scientists, physicians, politicians and industry have led to the fastest ever vaccine development program against any infectious disease known to mankind.

Moreover, efficacy and safety of the approved vaccines have been extremely high. While SARS-CoV-2 variants are on the rise across the globe, we are closer to achieving herd immunity with the development and mass rollout of these efficacious vaccines. Despite people with IBD being excluded from the pivotal COVID-19 vaccine trials, emerging evidence and past evidence from other vaccination programs suggest that these vaccines should be similarly safe and effective in those with IBD. As such, the overwhelming recommendation by all societies is that individuals with IBD should receive any of the available vaccines at the earliest opportunity, without delay or interruption of their IBD treatment. Future research is needed to determine if vaccine efficacy wanes among those receiving anti-TNF therapy or conventional immunosuppressive agents and if a timely booster dose is warranted to increase protection. It is important that people with IBD are not excluded from existing vaccination programs and that they do their part to procure timely vaccination to contribute to the ultimate goal of achieving herd immunity.

Funding

This supplement is sponsored by Crohn's and Colitis Canada and a Canadian Institutes of Health Research (CIHR) COVID-19 Rapid Research Funding Opportunity (Funding Reference Number – VR5 172684). Crohn's and Colitis Canada received partial funding support from Pfizer Canada, AbbVie Corporation (Canada), and Takeda Canada Inc. after completing the draft of the impact of COVID-19 and IBD report. Only Crohn's and Colitis Canada was involved in the research, writing, and conclusions of this report. The other sponsors had no role in the development or conclusions of this report.

CONFLICT OF INTEREST

Eric Benchimol has acted as a legal consultant for Hoffman-LaRoche Limited and Peabody & Arnold LLP for matters unrelated to medications used to treat inflammatory bowel disease and has received honoraria from McKesson Canada. He is Chair of the Scientific and Medical Advisory Council of Crohn's and Colitis Canada. Charles Bernstein is supported in part by the Bingham Chair in Gastroenterology. He is on Advisory Boards for AbbVie Canada, Amgen Canada, Bristol Myers Squibb, Janssen Canada, Pfizer Canada, Roche Canada, Sandoz Canada, Takeda Canada. He is a Consultant for Mylan Pharmaceuticals and Takeda. He has received educational grants from AbbVie Canada, Pfizer Canada, Takeda Canada, Janssen Canada. He is on the speaker's panel for AbbVie Canada, Janssen Canada, Pfizer Canada, Takeda Canada and Medtronic Canada. Received research funding from AbbVie Canada, Pfizer Canada, Sandoz Canada. Alain Bitton has participated in advisory boards with AbbVie, Janssen, Pfizer, Takeda, Hoffman-LaRoche, Amgen. He has received research support from AbbVie. He has received educational support from Fresenius Kabi, Takeda. Jean-Eric Ghia is a scientific analyst for the Canadian Broadcasting Corporation / Radio-Canada and acted as a scientific director for La Liberté. Anne Griffiths holds the Northbridge Financial Corporation Chair in IBD, has been a

consultant for AbbVie, Amgen, Bristol Myers Squibb, Janssen, Lilly, Merck, Pfizer, has received speaker fees from AbbVie, Janssen, Nestle and investigator-initiated research support from AbbVie. Jennifer Jones has received honoraria for speaking and consulting for AbbVie, Janssen, Pfizer, Shire and Takeda.

Gilad Kaplan has received honoraria for speaking or consultancy from AbbVie, Janssen, Pfizer and Takeda. He has received research support from Ferring, Janssen, AbbVie, GlaxoSmith Kline, Merck and Shire. He has been a consultant for Gilead. He shares ownership of a patent: TREATMENT OF INFLAMMATORY DISORDERS, AUTOIMMUNE DISEASE and PBC. UTI Limited Partnership, assignee. Patent WO2019046959A1. PCT/CA2018/051098. 7 Sept. 2018. Remo Panaccione reports consultant work for: AbbVie, AGI Therapeutics, Alba Therapeutics, Amgen, Astellas, Athersys, Atlantic Healthcare, BioBalance, Boehringer-Ingelheim, Bristol-Myers Squibb, Celgene, CoMentis, Cosmo Technologies, Coronado Biosciences, Cytokine Pharmasciences, Eisai Medical Research, Elan, EnGene, Eli Lilly, Enteromedics, Ferring, Flexion Therapeutics, Genentech, Genzyme, Gilead, Given Imaging, GlaxoSmithKline, Human Genome Sciences, Ironwood, Janssen, Merck & Co., Merck Research Laboratories, Merck Serono, Nisshin Kyorin, Novo Nordisk, NPS Pharmaceuticals, Optimer, Orexigen, PDL Biopharma, Pfizer, Procter and Gamble, Santarus, Shire Pharmaceuticals, Sigmoid Pharma, Sirtris (a GSK company), Sandoz, S.L.A. Pharma (UK), Targacept, Teva, Therakos, Tillotts, TxCell SA, Speaker's fees for AbbVie, Amgen, Celgene, Ferring, Janssen, Merck, Novartis, Pfizer, Prometheus, Sandoz, Shire, Takeda Advisory board attendance for AbbVie, Abbott, Allergan, Amgen, Biogen Idec, Eisai, Ferring, Genentech, Janssen, Merck, Shire, Elan, GlaxoSmithKline, Hospira, Pfizer, Bristol-Myers Squibb, Takeda, Cubist, Celgene, Salix, Roche. Research/educational support from AbbVie, Ferring, Janssen, Shire Takeda. Cynthia Seow has been on advisory boards for Janssen, AbbVie, Takeda, Ferring, Shire, Pfizer, Sandoz, Pharmascience and a speaker for Janssen, AbbVie, Takeda, Ferring, Shire, Pfizer, Pharmascience. Laura Targownik has received research funding from AbbVie Canada, Takeda Canada, Sandoz Canada, Amgen Canada, Gilead Canada, Roche Canada and Pfizer Canada and has been on Advisory Boards for Janssen Canada, AbbVie Canada, Takeda Canada, Pfizer Canada, Merck Canada, Roche Canada, Sandoz Canada and Amgen Canada. None: Deanna Gibson, Ellen Kuenzig, Kate Lee, Sanjay Murthy, James Guoxian Huang, Mariam S. Mukhtar, Parul Tandon, Joseph Windsor.

Supplement sponsorship. This supplemental issue was produced with support from Pfizer Canada. Pfizer Canada had no role in the creation of the Impact of COVID-19 & IBD Report, nor did it influence its contents. This supplemental issue was also produced with support from Crohn's and Colitis Canada and a Canadian Institutes of Health Research (CIHR) COVID-19 Rapid Research Funding Opportunity grant (Funding Reference Number – VR5 172684).

References

- Krammer F. SARS-CoV-2 vaccines in development. *Nature* 2020;586(7830):516–27.
- Rolak S, Hayney MS, Farraye FA, et al. What gastroenterologists should know about COVID-19 vaccines. *Clin Gastroenterol Hepatol* 2021;19(4):657–61.
- Vabret N, Britton GJ, Gruber C, et al.; Sinai Immunology Review Project. Immunology of COVID-19: Current state of the science. *Immunity* 2020;52(6):910–41.
- Lucas C, Wong P, Klein J, et al.; Yale IMPACT Team. Longitudinal analyses reveal immunological misfiring in severe COVID-19. *Nature* 2020;584(7821):463–9.
- Jeyanathan M, Afkhami S, Smail F, et al. Immunological considerations for COVID-19 vaccine strategies. *Nat Rev Immunol* 2020;20(10):615–32.
- Kaneko N, Kuo HH, Boucay J, et al.; Massachusetts Consortium on Pathogen Readiness Specimen Working Group. Loss of Bcl-6-expressing T follicular helper cells and germinal centers in COVID-19. *Cell* 2020;183(1):143–57.e13.
- Zhang J, Wu Q, Liu Z, et al. Spike-specific circulating T follicular helper cell and cross-neutralizing antibody responses in COVID-19-convalescent individuals. *Nat Microbiol* 2021;6(1):51–8.
- Wang EY, Mao T, Klein J, et al.; Yale IMPACT Team. Diverse functional autoantibodies in patients with COVID-19. *Nature* 2021;595(7866):283–8.
- Aschwanden C. The false promise of herd immunity for COVID-19. *Nature* 2020;587(7832):26–8.
- Turner JS, O'Halloran JA, Kalaidina E, et al. SARS-CoV-2 mRNA vaccines induce persistent human germinal centre responses. *Nature* 2021;596(7870):109–13.
- Polack FP, Thomas SJ, Kitchin N, et al.; C4591001 Clinical Trial Group. Safety and efficacy of the BNT162b2 mRNA Covid-19 vaccine. *N Engl J Med* 2020;383(27):2603–15.
- Baden LR, El Sahly HM, Essink B, et al.; COVE Study Group. Efficacy and safety of the mRNA-1273 SARS-CoV-2 vaccine. *N Engl J Med* 2021;384(5):403–16.
- Voysey M, Clemens SAC, Madhi SA, et al.; Oxford COVID Vaccine Trial Group. Safety and efficacy of the ChAdOx1 nCoV-19 vaccine (AZD1222) against SARS-CoV-2: An interim analysis of four randomised controlled trials in Brazil, South Africa, and the UK. *Lancet* 2021;397(10269):99–111.
- National Institutes of Health Press Release. Janssen Investigational COVID-19 Vaccine: interim analysis of phase 3 clinical data released. 2021. <https://www.nih.gov/news-events/news-releases/janssen-investigational-covid-19-vaccine-interim-analysis-phase-3-clinical-data-released>. January 29, 2021. (Accessed May 5, 2021).
- Dagan N, Barda N, Kepten E, et al. BNT162b2 mRNA Covid-19 vaccine in a nationwide mass vaccination setting. *N Engl J Med* 2021;384(15):1412–23.
- Doria-Rose N, Suthar MS, Makowski M, et al.; mRNA-1273 Study Group. Antibody persistence through 6 months after the second dose of mRNA-1273 vaccine for Covid-19. *N Engl J Med* 2021;384(23):2259–61.
- Sara E, Oliver JWG, Heather Scobie, et al. Center for Disease Control and Prevention, Morbidity and Mortality Weekly Report. The Advisory Committee on Immunization Practices' interim recommendation for use of Janssen COVID-19 vaccine — United States, February 2021. https://www.cdc.gov/mmwr/volumes/70/wr/mm7009e4.htm?_cid=mm7009e4_w. (Accessed May 5, 2021).
- CNN. EU agency finds AstraZeneca vaccine can cause rare blood clots, as UK advises other shots for under-30s. 2021. <https://www.cnn.com/2021/04/07/health/astrazeneca-coronavirus-vaccine-europe-uk-ema-intl/index.html>. (Accessed May 5, 2021).
- Mahase E. Covid-19: Unusual blood clots are “very rare side effect” of Janssen vaccine. *BMJ* 2021;373:n1046.
- CBC News. Future of AstraZeneca COVID-19 vaccine in question in Canada over blood clots, supply issues. 2021. <https://www.cbc.ca/news/health/astrazeneca-vaccine-paused-canada-blood-clot-vitt-1.6022821>. (Accessed May 5, 2021).
- Pottegård A, Lund LC, Karlstad Ø, et al. Arterial events, venous thromboembolism, thrombocytopenia, and bleeding after vaccination with Oxford-AstraZeneca ChAdOx1-S in Denmark and Norway: Population based cohort study. *BMJ* 2021;373:n1114.
- Greinacher A, Thiele T, Warkentin TE, et al. Thrombotic thrombocytopenia after ChAdOx1 nCoV-19 vaccination. *N Engl J Med* 2021;384(22):2092–101.
- CTV News. Canada could be mixing and matching COVID-19 vaccines by the summer. 2021. <https://www.ctvnews.ca/health/coronavirus/canada-could-be-mixing-and-matching-covid-19-vaccines-by-the-summer-tam-1.5425807>. May 12, 2021. (Accessed May 5, 2021).
- Payne JG, Tagalakis V, Wu C, et al.; CanVECTOR Network. Current estimates of the incidence of acute venous thromboembolic disease in Canada: A meta-analysis. *Thromb Res* 2021;197:8–12.
- Tan BK, Mainbourg S, Friggeri A, et al. Arterial and venous thromboembolism in COVID-19: a study-level meta-analysis. *Thorax* 2021;thoraxjnl-2020-215383; Online ahead of print. doi: 10.1136/thoraxjnl-2020-215383. PMID: 33622981; PMCID: PMC7907632.
- Christopher J. Ruddy. Royal Society of Canada: Canada's vaccine legacy: influenza, polio & COVID-19 vaccine(s). RSC COVID-19 Series, Publication #39; September 24, 2020. 2020. <https://rsc-src.ca/en/voices/canada%E2%80%99s-vaccine-legacy-influenza-polio-covid-19-vaccines>. (Accessed May 5, 2021).

27. Government of Canada. Background – Government of Canada investments in COVID-19 vaccines, therapeutics and biomanufacturing capacity. 2021. <https://www.canada.ca/en/innovation-science-economic-development/news/2021/02/backgrounder--government-of-canada-investments-in-covid-19-vaccines-and-biomanufacturing-capacity.html>. (Accessed May 5, 2021).
28. Botwin GJ, Li D, Figueiredo J, Cheng S, Braun J, McGovern DPB, Melmed GY. Adverse events after SARS-CoV-2 mRNA vaccination among Patients with inflammatory bowel disease. *Am J Gastroenterol* 2021; Online ahead of print doi: [10.14309/ajg.000000000001342](https://doi.org/10.14309/ajg.000000000001342). PMID: 34047304.
29. Kennedy NA, Lin S, Goodhand JR, et al. Infliximab is associated with attenuated immunogenicity to BNT162b2 and ChAdOx1 nCoV-19 SARS-CoV-2 vaccines in patients with IBD. *Gut* 2021;gutjnl-2021-324789; Online ahead of print. doi: [10.1136/gutjnl-2021-324789](https://doi.org/10.1136/gutjnl-2021-324789). PMID: 33903149; PMCID: PMC8076631.
30. Kennedy NA, Goodhand JR, Bewshea C, et al.; Contributors to the CLARITY IBD study. Anti-SARS-CoV-2 antibody responses are attenuated in patients with IBD treated with infliximab. *Gut* 2021;70(5):865–75.
31. Fiorino G, Peyrin-Biroulet L, Naccarato P, et al. Effects of immunosuppression on immune response to pneumococcal vaccine in inflammatory bowel disease: A prospective study. *Inflamm Bowel Dis* 2012;18(6):1042–7.
32. Banaszekiewicz A, Targońska B, Kowalska-Duplaga K, et al. Immunogenicity of 13-valent pneumococcal conjugate vaccine in pediatric patients with inflammatory bowel disease. *Inflamm Bowel Dis* 2015;21(7):1607–14.
33. Melmed GY, Agarwal N, Frenck RW, et al. Immunosuppression impairs response to pneumococcal polysaccharide vaccination in patients with inflammatory bowel disease. *Am J Gastroenterol* 2010;105(1):148–54.
34. Lee CK, Kim HS, Ye BD, et al.; Korean Association for the Study of Intestinal Diseases (KASID) Study. Patients with Crohn's disease on anti-tumor necrosis factor therapy are at significant risk of inadequate response to the 23-valent pneumococcal polysaccharide vaccine. *J Crohns Colitis* 2014;8(5):384–91.
35. Cullen G, Bader C, Korzenik JR, et al. Serological response to the 2009 H1N1 influenza vaccination in patients with inflammatory bowel disease. *Gut* 2012;61(3):385–91.
36. Lu Y, Jacobson DL, Ashworth LA, et al. Immune response to influenza vaccine in children with inflammatory bowel disease. *Am J Gastroenterol* 2009;104(2):444–53.
37. deBruyn J, Fonseca K, Ghosh S, et al. Immunogenicity of influenza vaccine for patients with inflammatory bowel disease on maintenance infliximab therapy: A randomized trial. *Inflamm Bowel Dis* 2016;22(3):638–47.
38. Mamula P, Markowitz JE, Piccoli DA, et al. Immune response to influenza vaccine in pediatric patients with inflammatory bowel disease. *Clin Gastroenterol Hepatol* 2007;5(7):851–6.
39. Hagihara Y, Ohfuji S, Watanabe K, et al. Infliximab and/or immunomodulators inhibit immune responses to trivalent influenza vaccination in adults with inflammatory bowel disease. *J Crohns Colitis* 2014;8(3):223–33.
40. Pratt PK Jr, David N, Weber HC, et al. Antibody response to hepatitis B virus vaccine is impaired in patients with inflammatory bowel disease on infliximab therapy. *Inflamm Bowel Dis* 2018;24(2):380–6.
41. Park SH, Yang SK, Park SK, et al. Efficacy of hepatitis A vaccination and factors impacting on seroconversion in patients with inflammatory bowel disease. *Inflamm Bowel Dis* 2014;20(1):69–74.
42. Wasan SK, Zullo S, Berg A, et al. Herpes Zoster vaccine response in inflammatory bowel disease patients on low-dose immunosuppression. *Inflamm Bowel Dis* 2016;22(6):1391–6.
43. Wyant T, Leach T, Sankoh S, et al. Vedolizumab affects antibody responses to immunisation selectively in the gastrointestinal tract: randomised controlled trial results. *Gut* 2015;64(1):77–83.
44. Doornekamp L, Goetgebuer RL, Schmitz KS, et al. High immunogenicity to influenza vaccination in Crohn's disease patients treated with ustekinumab. *Vaccines (Basel, Switzerland)* 2020;8(3):455.
45. Brodmerkel C, Wadman E, Langley RG, et al. Immune response to pneumococcus and tetanus toxoid in patients with moderate-to-severe psoriasis following long-term ustekinumab use. *J Drugs Dermatol* 2013;12(10):1122–9.
46. Khan N, Mahmud N. Effectiveness of SARS-CoV-2 vaccination in a veterans affairs cohort of patients with inflammatory bowel disease with diverse exposure to immunosuppressive medications. *Gastroenterology* 2021;S0016-5085(21)03066-3; Online ahead of print. doi: [10.1053/j.gastro.2021.05.044](https://doi.org/10.1053/j.gastro.2021.05.044). PMID: 34048782; PMCID: PMC8146263.
47. Tse F, Moayyedi P, Waschke KA, et al. COVID-19 vaccination in patients with inflammatory bowel disease: Communiqué from the Canadian Association of Gastroenterology. *J Can Assoc Gastroenterol* 2021;4(1):49.
48. Alexander JL, Moran GW, Gaya DR, et al.; Inflammatory Bowel Disease section of the British Society of Gastroenterology and the the Inflammatory Bowel Disease Clinical Research Group. SARS-CoV-2 vaccination for patients with inflammatory bowel disease: A British Society of Gastroenterology Inflammatory Bowel Disease section and IBD Clinical Research Group position statement. *Lancet Gastroenterol Hepatol* 2021;6(3):218–24.
49. Siegel CA, Melmed GY, McGovern DP, et al.; International Organization for the Study of Inflammatory Bowel Disease (IOIBD); International Organization for the Study of Inflammatory Bowel Diseases (IOIBD). SARS-CoV-2 vaccination for patients with inflammatory bowel diseases: Recommendations from an international consensus meeting. *Gut* 2021;70(4):635–40.
50. Brenner EJ, Ungaro RC, Geary RB, et al. Corticosteroids, but not TNF antagonists, are associated with adverse COVID-19 outcomes in patients with inflammatory bowel diseases: Results from an international registry. *Gastroenterology* 2020;159(2):481–491.e3.
51. Ungaro RC, Brenner EJ, Geary RB, et al. Effect of IBD medications on COVID-19 outcomes: Results from an international registry. *Gut* 2021;70(4):725–32.