Gorodnichy v. CBSA (FPSLREB 566-02-48305)
Exhibit G-19

Exhibit sheet description:

New evidence submitted on October 22, 2026 – 2nd Book of documents. Document at page 102 (Portal that grievor created with colleagues to allow people to submit side effects).

Source documents and page ranges:

566-02-48305_Grievor_Book_of_Documents-NEW_02_HC_CHANGES_TO_C19V_SIDE_EFFECT_REPORTING-AND-ETHICAL_PROBLEMS.pdf:

PDF pages 102-104

Reader guidance:

Source: https://open-canada.github.io/vitals/report-side-effect, linked from https://open-canada.github.io/vitals

Evidence:

This page was created by R4GC community led by Gorodnichy to provide a link to the online vaccine side-effect reporting form. Later it was found that Health Canada removed this link immediately before

COVID-19 vaccination began.

The page also provides guidance on how to report vaccine side-effects using the example of Gorodnichy's family member, who got shingles after Moderna. It also provides links to the reports that document severe side effects, including deaths, from COVID-19 vaccines. One of these documents was released by Pfizer under the court order on 17 November 2021:

CUMULATIVE ANALYSIS OF POST-AUTHORIZATION ADVERSE EVENT REPORTS OF

PF-07302048 (BNT162B2) RECEIVED THROUGH 28-FEB-2021. Pfizer,

https://phmpt.org/wpcontent/uploads/2021/11/5.3.6-postmarketing-experience.pdf listed as 5.3.6 postmarketing experience.pdf at https://phmpt.org/pfizer-16-plus-documents

vitals

Official Canada Vital (Deaths) Statistics, analyzed using Data Science

View On GitHub

This project is maintained by open-canada

Reporting COVID-19 vaccines side effects

Background

Reporting of serious (or life threatening) side effects of vaccines is mandatory in Canada. However, reporting of non-serious (or not life threatening) side effects in Canada is voluntary. Many doctors may not have time to do it, nor they provide guidance to their patients on how they can do it.

For this reason many non-serious side effects remain unreported. Many doctors will not do it, and consumers do not know how to do it. In fact, many Canadians may not even know that they can do report vaccine side-effect themselves, i.e. without a medical doctor. There are no instructions on the Health Canada portal on how to do it. The link to the Health Canada Voluntary Side Effect Reporting portal may also be rather difficult to find.

This page is created to address this gap and to help Canadians to report any vaccine (or any drug) side-effect, as a consumer. Health Canada encourages everyone to do so. – It is thanks to such efforts that vaccines (and drugs) can be improved for the benefits of all.

Once you know how to do it, it should not take more than fifteen minutes to do it. This is how you do it. An Example is provided here.

Submitting Voluntary Side Effect Report via Health Canada portal

Step 1: Go to the Health Canada Voluntary Side Effect Reporting portal : https://hpr-rps.hres.ca/side-effects-reporting-form.php?form=voluntary

Step 2: Submit electronic form there as shown in this Example.

- There are three places there that you need to know how to fill out:
  - in Section B: Patient ID write your initials;
  - in Section D: Product DIN and Product Name - paste the vaccine names (and batch/lot number, if available), as printed on your receipt.

- Make sure you provide a valid email address, which will be used to send you to further instructions on how ontinue side-effect report process.

- If you still have problems submitting the form (i.e., clicking on button “SUBMIT” does not lead you anywhere),
  call Heath Canada Voluntary Reporting support line: Tel. 866-234-2345, if you leave a message. They should respond within 24 hours. If they did not, please call again.

Step 3: A few (2-5) days later you should receive email from *Canada Vigilance (HC/SC)* *canadavigilance@hc-sc.gc.ca* (with Subject: “Correspondence from the Marketed Health Products Directorate - Adverse Reaction Number 0009xxxxx), which will provide you with the Adverse Reaction Report (AER) number and which will ask you to submit additional details though their portal.

If you don’t receive email from them within 5 business days, please check you Spam folder and, if it is not there, please call them at Tel. 866-234-2345 and leave a message.

As they note in their email: *“Posting of reports in the online database remains 3 months behind the date of reporting. This delay accounts for the necessary activities involved in report processing”*

Additional resources:

The following additional resources are provided to help you prepare your side-effect report. There you can check if your symptoms are already reported in national adverse event reporting systems, and if so, how common they are, for which type of vaccine, and which vaccine batch (a.k.a. lot, which is the number printed on your vaccination receipt)

566-02-48305

- Recalls and safety alerts, Government of Canada https://recalls-rappels.canada.ca/: for [Moderna], for [Pfizer], for [Astrazeneca])

- 'Reported side effects following COVID-19 vaccination in Canada', Canadian COVID-19 vaccination safety report, Public Health Agency of Canada, https://health-infobase.canada.ca/covid-19/vaccine-safety/

- US Government Vaccine Adverse Event Reports (VAERS): https://vaers.hhs.gov. You can also check if your vaccine batch (lot) is among the "bad ones", according to VAERS data, here,

- Comprehensive world-wide COVID-19 vaccine reports repository

- CUMULATIVE ANALYSIS OF POST-AUTHORIZATION ADVERSE EVENT REPORTS OF PF-07302048 (BNT162B2) RECEIVED THROUGH 28-FEB-2021. Prepared by Pfizer, Released to Public Health and Medical Professionals for Transparency (https://phmpt.org) under the court order on 17 November 2021: https://phmpt.org/wp-content/uploads/2021/11/5.3.6-postmarketing-experience.pdf

The authoritative source for COVID-19 information is Canada.ca/coronavirus.

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