BC PHARMACARE Newsletter

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The PharmaCare Newsletter team works from the territory of the Lekwungen peoples, including the Songhees and Esquimalt Nations. Our gratitude extends to them, and all the Indigenous peoples on whose territories and lands we build relationships.

The PharmaCare Newsletter is published by the Pharmaceutical, Laboratory & Blood Services Division to provide information to British Columbia's health care providers.

The use of PharmaNet is not intended as a substitute for professional judgment. Information on PharmaNet is not exhaustive and cannot be relied upon as complete. The absence of a warning about a drug or drug combination is not an indication that the drug or drug combination is safe, appropriate or effective in any given patient. Health care professionals should confirm information obtained from PharmaNet, and ensure no additional relevant information exists, before making patient care decisions

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Q: Where can you find the updated table of all medications for osteoporosis that are approved in Canada?

HINT: The answer is in the May 2022 edition of PAD Refills. Don't forget to subscribe!

Special release – June 17, 2022: Guidance for B.C. pharmacies during the specialized infant formula shortage

To conserve hypoallergenic infant formula during the <u>global shortage</u>, the BC Ministry of Health issues this guidance for pharmacies for its sale and distribution:

- Keep specialized formulas behind the counter amino acid-based (AAF) and extensively hydrolyzed formulas
 (EHF)
- Sell specialized formulas in limited quantities 7-10 day supply. Use judgement if clients face barriers getting to a pharmacy. For example, sell in larger quantities to people in remote communities
- Sell specialized formulas only to families with clear medical need. See decision tree
- Consider purchase limits on regular formula. The shortage could put pressure on regular formula supply
- Post notices indicating that specialized formula is behind the counter and being carefully distributed

This decision tree helps providers guide families to choose the best formula for each family:

• Use of Specialized Infant Formula During a Shortage – decision tree

These measures are to *conserve supply*. No formula supply issues exist currently in B.C. More stock of both specialized formula types have been routed to B.C.

See the ministry's information bulletin B.C. takes steps to protect supply of specialized infant formulas.

Background on formula shortage

Due to a global shortage of infant formulas for infants with food allergies and certain medical conditions, Canada is experiencing an acute shortage of:

- Extensively hydrolyzed formula (EHF) for moderate food allergies, and
- Amino acid formulas (AAF) for complex or severe food allergies

To mitigate shortages, Health Canada has implemented an <u>interim policy</u> allowing the temporary importation and sale of foreign-labelled product.

The BC Ministry of Health is working with Health Canada to address concerns about supply.

Health Canada:

- Expects the shortage of hypoallergic formula to last until the fall
- Is monitoring for equitable distribution of the limited supply across the country
- May allocate supplies to be used in emergency situations
- Is importing product from countries with similar manufacturing practices and standards

The temporary shortage was caused by Abbott's interim closure of its formula manufacturing plant in Michigan, and voluntary recall of certain formulas produced there. The facility reopened on June 4; however, production was halted again this week due to local storms and flooding. As production resumes, specialized infant formulas will gradually become more available in the coming months.

We will share further information as soon as possible.

Resources

For healthcare providers:

- Use of Specialized Infant Formula During a Shortage decision tree
- List of imported products, with printable labels Health Canada
- NEW <u>Hypoallergenic infant formula shortage Information for pharmacy professionals</u> Canadian Pharmacists Association
- Interim policy on importation Health Canada
- Algorithm (French only) to help health professionals choose alternatives to specialized formulas Quebec's Ministry of Health and Social Services
- Health Link BC. Call 811 for dietitian support with formula decisions

For patients:

- <u>Information for families during the shortage of formula for infants with food allergies</u> (web page with downloadable PDF version) Health Canada
- Fact sheet for patients Health Canada
- B.C. takes steps to protect supply of specialized infant formulas
- Health Link BC. Call 811 for dietitian support with formula decisions

Special release – June 13, 2022

In transition: COVID-19 antiviral support line for clinicians

June 24, 2022 will be the last active date for the COVID-19 antiviral support line for clinicians. The support line is staffed by pharmacists and assists with questions related to prescribing and dispensing Paxlovid.

- Monday to Friday, 8:30 am to 4:30 pm (until June 24, 2022)
- 1-866-604-5924

Starting June 27, 2022, questions about prescribing and dispensing Paxlovid should be directed to the Ministry of Health's phone line:

- Monday to Friday, 8:30 am to 4:30 pm (starting June 27, 2022)
- 1-844-915-5005

New Paxlovid™ assessment fee and increase to follow-up fee

As of June 13, 2022, pharmacists dispensing nirmatrelvir/ritonavir (Paxlovid™) can claim a temporary \$30 clinical assessment fee (PAX-A) in addition to the standard dispensing fee. Also on June 13, 2022, the Paxlovid follow-up fee (PAX-F) will increase from \$15 to \$25.

Paxlovid Assessment fee (PAX-A)

The new fee takes into account Paxlovid's complexity, the potential for drug-drug interactions (DDIs), and the extra time needed for clinical assessments. B.C. dispensing data to June 5, 2022 indicates that more than 30% of dispenses in B.C. have a DDI management plan, and more than 12% require renal dose adjustments.

Claims for the temporary \$30 PAX-A fee (PIN: 66128340) must be submitted on the same day as the Paxlovid claim, from the same pharmacy.

The PAX-A fee can be claimed following an assessment of the Paxlovid prescription. This must include a DDI check with two independent sources, even if no drug therapy problem is identified.

If a pharmacist determines it is inappropriate or unsafe to dispense Paxlovid to a patient with a valid prescription, they may claim a <u>refusal to fill fee</u>, but **not the PAX-A fee**.

The end date of the temporary fee will be announced in a future PharmaCare Newsletter.

Required activities:

- Assess the Paxlovid prescription for completeness and appropriateness
- Include *DDI* or *Renal* in the SIG field for the Paxlovid dispense, as needed
- Communicate with the prescriber as needed to resolve drug therapy problems, including any DDIs and recommended management plans
- Keep records as usual.

- An optional form in Appendix C in the <u>Dispensing Paxlovid and monitoring adverse drug events: A guide</u>
 <u>for B.C. pharmacists</u> guide may be used to document DDIs and associated management plans but is not required.
- Submit the PAX-A claim the same day as the Paxlovid claim using
 PIN 66128340

Procedure:

- 1. Enter the PAX-A PIN 66128340 in the DIN/PIN field
- 2. If necessary, enter \$0 for drug cost, mark-up and fee
- 3. Enter 1 in the QTY field
- 4. In the PRACT ID Ref field, enter P1 College of Pharmacists of BC
- 5. In the PRACT ID field, enter your College ID
- 6. At start of SIG, enter the 10-digit phone number of the pharmacy where the service took place

Paxlovid follow-up fee (PAX-F) increase

As of June 13, 2022, the PAX-F fee (PIN: 66128313) will increase from \$15 to \$25.

Follow-up should be conducted with the patient or caregiver 6 to 10 days after dispensing Paxlovid.

Required activities:

- All relevant information must be recorded in the SIG of the PAX-F claim using the codes outlined in Appendix A
 of the Dispensing Paxlovid and monitoring adverse drug events: A guide for B.C. pharmacists
- Moderate and severe ADEs must be recorded in the Adverse Reaction field in PharmaNet
- Follow-up and additional clinical information may be documented separately, following usual pharmacy
 procedures, systems, or forms already in place. A form in Appendix B in the <u>Dispensing Paxlovid and monitoring</u>
 adverse drug events: A guide for B.C. pharmacists may be used as a follow-up tool, but is not required

To date:

- Pharmacists have followed up on approximately 64% of eligible Paxlovid dispenses
- Of those follow-ups:
 - More than half (54%) of patients experienced at least 1 ADE
 - Over 2,700 ADEs have been identified, with 76% classified as Mild
 - 89% of patients completed all 5 days of therapy

Reminder:

^{*}Only one fee can be claimed per treatment course

Like other prescription drugs, Paxlovid must be dispensed in accordance with the College of Pharmacists of BC's Community Standards of Practice. When dispensing Paxlovid, pharmacists should ensure they have the knowledge, skills and abilities to do so. Review the prescription for completeness and appropriateness, and review the patient's PharmaNet profile for drug therapy problems, therapeutic duplications and any other potential problems.

Resources:

Dispensing Paxlovid and monitoring adverse drug events: A guide for B.C. pharmacists

Special release - June 10, 2022: Shortage of infant formula

The temporary closure of a large U.S. manufacturing plant has affected global imports of infant formula, especially hypoallergenic formula. There are two types of hypoallergenic infant formula:

- Extensively hydrolyzed formula (EHF) for babies with moderate food allergies
- Amino acid formulas (AAF) for babies with complex or severe food allergies

To mitigate shortages, Canada has implemented an <u>interim policy</u>. Despite the interim policy, there remains very limited incoming supply of both EHF and AAF to Canada. Shortages are expected to last until mid-June and to affect community/retail supply.

Pharmacists are encouraged to refer families with questions to Health Canada's <u>Information for families during the shortage of formula for infants with food allergies</u>.

The Ministry will continue to monitor the situation and may provide additional guidance in the near future.

Manual payments

COVID-19 vaccination – weekend premium

As in PharmaCare added a \$4.00 premium to the COVID-19 vaccination administration fee, for vaccines administered on weekend only from December 11, 2021 to March 27, 2022. (Note: "weekend" refers to Saturday, Sundays and provincial statutory holidays.) PharmaCare has assessed the eligibility of the COVID-19 vaccine claims administered on weekends and has calculated the lump-sum payment to the eligible pharmacies for these claims.

Payments for eligible weekend premiums will be included in the weekly payment on June 13, 2022. These payments will appear on the Pharmacy Remittance Advice Form under the Adjustment Code "7–Manual Payment."

Rapid antigen tests - April 2022 payment

As in <u>PharmaCare Newsletter 22-004</u>, effective April 11, 2022, pharmacies would be paid \$75.00 per case distributed of the Rapid Response® COVID-19 antigen test (RAT) kit.

PharmaCare has calculated the total monthly fees owed to each pharmacy at the rate of \$75.00 per case recorded with PIN 66128325. The April 2022 payment will be included in the weekly payment on June 20, 2022. The payment will appear on the Pharmacy Remittance Advice Form under the Adjustment Code "7—Manual Payment."

Pharmacies are encouraged to order more RAT kits to maintain sufficient stock levels.

Temporary coverage for Nicorette lozenges in short supply

Nicorette lozenges RX 2 mg and 4 mg (NPNs 80053099 and 80053100), packaged for BC PharmaCare nicotine replacement therapies, are in short supply.

The over-the-counter (OTC) products (same NPNs) will be temporarily covered as eligible smoking cessation products until the RX versions are available. The OTC products aren't experiencing any supply challenges.

Tixagevimab/cilgavimab (Evusheld™) for prevention of COVID-19

Tixagevimab/cilgavimab (Evusheld™) is a long-acting monoclonal antibody, approved by Health Canada for prevention of COVID-19 in immunocompromised individuals.

On May 25, 2022, the BC Centre for Disease Control (BCCDC) released the <u>Clinical Practice Guide for the use of</u>
<u>Evusheld™ (PDF)</u>. Given the lack of evidence demonstrating clear benefit and potential risk of cardiac serious adverse events, Evusheld is not routinely recommended to all immunocompromised patients.

If used on a case-by-case basis, Evusheld should only be prescribed after careful consideration of theoretical benefit and risk and limited to patients who are:

- Severely immunocompromised (categorized as Clinically Extremely Vulnerable Group 1), AND
- Who have no known cardiovascular disease, AND
- Who have additional risk factors or exceptional circumstances that correlate with an extremely high risk of poor outcomes from COVID-19.

Currently, Evusheld is dispensed through the provincial Product Distribution Centre only.

Note: Evusheld is not used to treat COVID-19 infection.

See BCCDC documents for further details:

- Health Care Provider Information (PDF, 446KB)
- Patient Information (PDF, 347KB)

Advance notice: elbasvir-grazoprevir (Zepatier®) delisting

PharmaCare will delist elbasvir-grazoprevir (Zepatier®) 50 mg/100 mg tablet on **August 1, 2022**, because of the manufacturer's decision to remove it from market.

In April 2022, Merck Canada Inc. announced it was ceasing the sale of Zepatier, after evaluating clinical use and the availability of alternative therapies. The decision was not related to any quality or safety issues, the company stated.

Zepatier was listed as a limited coverage drug on March 1, 2017, as part of a coverage expansion of direct-acting antivirals for chronic hepatitis C (CHC).

Zepatier can still be sold—and will still be covered by PharmaCare—until **July 31, 2022**, by which the manufacturer estimates remaining inventory will be depleted.

The <u>Adults with Chronic Hepatitis C information sheet (PDF, 129KB)</u> will be updated on August 1, to reflect Zepatier's new non-benefit status.

Dispensing EDRDs and billing

Pharmacy billing overview

Expensive drugs for rare diseases (EDRDs) in B.C. are considered PharmaCare non-benefits (i.e., not eligible for PharmaCare coverage or by Special Authority request). However, in exceptional cases, coverage may be requested by a treating physician through the EDRD process. Requests are assessed for approval on a case-by-case, last resort basis. Under the EDRD process, the Ministry of Health (the Ministry) is the payer of last resort.

At this time, only Trikafta® is billed through PharmaNet. Payments for all other EDRDs are coordinated with the Provincial Health Services Authority (PHSA) Provincial Specialized Programs (PSP), as outlined below.

Pharmacy billing procedures for EDRD patients

Trikafta patients with no third-party coverage

Claims for patients on Trikafta who do not require coordination of benefits (i.e., those who are fully funded through the Ministry) can be billed directly through PharmaNet.

Trikafta patients with partial third-party coverage

Common partial third-party coverage may include patients with a copay, an annual maximum and/or a lifetime maximum. At this time, pharmacy software does not permit PharmaCare to be designated as the secondary payer when adjudicating claims.

To process Trikafta claims for patients with partial third-party coverage:

- 1. The pharmacy must set up an account with PHSA for manual invoicing. Email PSP@phsa.ca with the subject "Trikafta: New Pharmacy Account".
 - a. If your pharmacy already has an account set up with PHSA for billing of other EDRD drugs, follow the usual process.
- 2. Trikafta claims must be adjudicated:
 - a. through PharmaCare as the first payer, noting that you will receive a "DIN/Drug is a non-benefit" message, and
 - b. the patient's third-party plan as the second payer.
- 3. The remaining co-pay, after third-party reimbursement, must be manually invoiced to PHSA PSP at monthly intervals.
- 4. For patients with third-party annual or lifetime maximums, inform PSP@PHSA.ca once this maximum has been reached.

Claims for all other EDRD therapies (non-Trikafta patients)

If applicable, the patient's third-party plan must be billed first. Any remaining copay must be invoiced to PHSA.

- If your pharmacy already has an account set up with PHSA for billing of other EDRD drugs, follow the usual process.
- If your pharmacy does not have an account with PHSA, email PSP@phsa.ca with the subject "EDRD: New Pharmacy Account".

Refill process

Patients who obtain medications funded through the EDRD program must contact their respective pharmacies for a medication refill:

- The routine shipping of drugs (or automatic dispensing) to patients is not permitted.
- Requesting refills is the responsibility of the patient.
 - Patients are advised to provide enough time for the pharmacy to order medication refills (recommended 2 weeks notice)

Frequency of dispensing

The Ministry requires pharmacies to dispense EDRD medications in quantities of 30 days or less.

- In some circumstances, the dispensing pharmacy and/or patient may request from the Ministry an alternate dispensation interval on a temporary or permanent basis.
 - Email <u>PSP@phsa.ca</u> in the event a patient needs to exceed this maximum. Requests will be reviewed on a case-by-case basis.

Reminders

COVID-19 single-entry claims: don't enter PIN or DIN

Pharmacists are reminded not to submit COVID-19 vaccination claims in PharmaNet under the PIN or DIN. Such claims may be subject to recovery and pharmacies may be contacted to reverse the claims.

As of <u>April 1, 2022</u>, pharmacies only need to record COVID-19 vaccinations once ("single entry"), in the ImmsBC application. The entered information will automatically be transmitted to PharmaNet.



Methadone interaction fees are consistently higher than other clinical service fees. Injection fees have a growth rate of 30.7% over the last 10 years.

Find more stats like this in 2020/2021 PharmaCare Trends.

Regular benefits

Effective May 31, 2022, the following products are listed as PharmaCare regular benefits:

Drug name	propylthiouracil (Halycil™)		
Date effective	May 31, 2022		
Indication	In conjunctionThe control of	of thyrotoxicosis prior t	hasten recovery while awaiting the effects of
DIN	02521059	Strength and form	50 mg tablet
Covered under lans	Fair PharmaCare, B, (C, F, W	

Drug name	biosynthetic human insulin (Entuzity™ KwikPen®)		
Date effective	May 31, 2022		
Indication	diabetes mellitus		
DIN	02466864	Strength and form	500 U/mL pre-filled vial
Covered under lans	Fair PharmaCare, B, C, F, W, P		

Limited coverage benefits

As of May 17, 2022, the following product is covered as a limited coverage benefit under the DIN below:

Drug name	trientine (MAR-trientine)		
Date effective	May 17, 2022		
Indication	Wilson's disease		
DIN	02504855	Strength and form	250 mg capsule
Covered under plans	Fair PharmaCare, B, C, F, W		

Non-benefits

As of May 10, 2022, PharmaCare has determined the following products will not be covered per the DINs below:

Drug name	pegfilgrastim (Ziextenzo®)	
Date effective	May 10, 2022	
DIN	02497395	

Drug name	pegfilgrastim (Nyvepria™)
Date effective	May 10, 2022
DINs	02506238

Your Voice: Patient input needed for drug decisions

The knowledge and experience of patients, caregivers and patient groups are integral to **B.C.'s drug review process**.

The Ministry depends on pharmacies and practitioners to help connect patients and their caregivers with opportunities to give input. If you have a patient currently taking one of the drugs under review or who has the condition the new drug treats, please encourage them to visit http://www.gov.bc.ca/BCyourvoice.

Currently, input is needed for the following:

DRUG	anifrolumab (Saphnelo™)
INDICATION	systemic lupus erythematosus (SLE)
INPUT WINDOW	May 25 to June 22, 2022

DRUG	faricimab (TBC)	
INDICATION	neovascular (wet) age-related	
	macular degeneration (nAMD)	
INPUT WINDOW	May 25 to June 22, 2022	

DRUG	pitolisant hydrochloride (Wakix®)	
INDICATION narcolepsy		
INPUT WINDOW	May 25 to June 22, 2022	

DRUG	dapagliflozin (Forxiga®)
INDICATION	chronic kidney disease (CKD)
INPUT WINDOW	May 25 to June 22, 2022
DRUG	cariprazine (Vraylar®)
INDICATION	bipolar disorder
INPUT WINDOW	May 25 to June 22, 2022

FNHA Partnership series: Coming Together for Wellness

This article is part of a 10-article series by the Ministry of Health and the First Nations Health Authority (FNHA) to increase awareness of First Nations issues and build cultural humility, and as a result, safety in B.C.'s health system. The series began in the PharmaCare Newsletter, edition 21-010.

Article #8: Accessing the first BGTS fill

<u>Article 6</u> of this series indicated that FNHA can add a Certificate of Training in Blood Glucose Monitoring for its clients who live with diabetes. This article explores the topic in greater detail.

PharmaCare covers blood glucose test strips (BGTS) for eligible B.C. residents who have completed blood glucose monitoring training at an accredited diabetes education centre (DEC) or a designated primary care network (PCN). A Certificate of Training needs to be in the client's PharmaNet profile in order for a pharmacy to receive payment for a BGTS claim.

Historical and geographical context

FNHA functions as a DEC for clients who can't access training from the usual DEC or a PCN. Distance has been a barrier to access training and education. This is one example of the health disparities faced by First Nations in B.C. due to the lasting effects of structural racism and colonization existing in the healthcare system and broader determinants of health.

<u>Article 2</u> of the series highlighted how colonization and forced dislocation of First Nations disrupted access to healthy, traditional foods and food practices. First Nations living in rural and remote regions have inequitable access to quality,

healthy and affordable food and medicines. Furthermore, intergenerational trauma stemming from experiences in Indian hospitals and residential schools led to mistrust and avoidance of the healthcare system. These combined factors have contributed to an increased risk of type 2 diabetes and related complications in First Nations people, showing <a href="https://diabetes.ncbi.nlm.

FNHA's DEC program continues to evolve as many First Nations communities have holistic diabetes programming focused on managing diabetes through nutrition, activity, medications, and emotions. There are well-developed programs like the mobile diabetes clinics run by Seabird Island Band and Carrier Sekani <a href="mailto:Fam



The Carrier Sekani Family Services clinic team provides mobile diabetes prevention and intervention education, check-ups and testing in communities across northern British Columbia.

How FNHA clients can get the first BGTS fill

The first fill of BGTS can be pre-activated for Plan W beneficiaries by calling First Nations Health Benefits at 1-855-550-5454.

Healthcare providers, diabetes educators, health directors, and health advocates may also call Health Benefits on behalf of a client.

Pharmacy support

Pharmacists can support FNHA clients in the following ways:

- Support access for an FNHA client who visits the pharmacy for BGTS for the first time and has no coverage in
 place. The pharmacist should contact Health Benefits and consider using the
 FNHA Transitional Payment Request (TPR) (PDF, 208KB) form to provide the client immediate access.
- Encourage ongoing learning about diabetes and counsel on how to monitor their blood sugar levels with purpose (PDF, 205KB).
- Seek opportunities to strengthen care-team relationships and collaboration with diabetes care providers at local First Nations Health Service organizations.

Next in the Coming Together for Wellness Series: Next Steps in Indigenous Healthcare in British Columbia

Suggested reading

• First Nations Population Health and Wellness Agenda (p. 92-97) (PDF, 22.7MB)