

Healthcare-associated infections surveillance report

Carbapenemase-Producing Organisms (CPOs) Update

December 2019

Highlights for Q2 of 2019/20 (June 28 – September 19, 2019)

- 41 patients were newly confirmed with CPO
- 46 new cases of carbapenemase genes were identified, including 5 patients each having 2 or more different genes
- NDM were the most common carbapenemase genes identified, accounting for 56.5% (26/46)
- Surveillance information was reported for 38 out of 46 new cases, 2 of which were reported at community healthcare settings
- 14 of the reported cases had healthcare encounters outside Canada and 10 cases reported travel without healthcare encounters.

What are carbapenemase-producing organisms (CPOs)?

Carbapenems are a class of antibiotics usually reserved to treat serious infections, and often considered one of the antimicrobial treatments of last resort. Over the last decade, some bacteria have developed resistance to carbapenems by producing an enzyme (carbapenemase) that breaks down the structure of these antibiotics and makes them ineffective for treatment. These antibiotic-resistant bacteria are called carbapenemase-producing organisms (CPOs). The most common carbapenemases in Canada include NDM, KPC, and OXA-48.

Why are CPOs considered important?

CPOs are an important emerging threat to healthcare settings and the community. First, these organisms are often resistant to multiple classes of antimicrobials, substantially limiting treatment options. Second, infections caused by these organisms are associated with high mortality rates, up to 50% in some studies. Third, many carbapenem resistance genes can be transmitted from one species of bacteria to another, potentially facilitating widespread resistance. Fourth, since Enterobacteriaceae are a common cause of infections, carbapenem resistance in these organisms could have far-reaching impact. Finally, outbreaks of CPOs are more difficult and costly to contain.

How are CPOs spread?

People can carry CPOs without having any symptoms of illness (this is called colonization), but they can still pass the germs to other people. CPOs usually spread person-to-person through direct contact with infected or colonized people, or by contaminated surfaces. This can happen in both community and healthcare settings. Without proper precautions, CPOs can spread easily from person-to-person in hospitals, especially in countries where CPOs are endemic.

How can the spread of CPOs be prevented?

Good hand hygiene by both healthcare providers and patients, such as washing hands often with soap and water or using an alcohol-based hand sanitizer, is a simple and effective way to prevent the spread of CPOs. The public should avoid unnecessary access to health care in endemic countries. In healthcare settings, identifying CPO cases and placing colonized or infected patients on contact precautions, using medical devices and antimicrobials wisely, and carefully cleaning and disinfecting rooms as well as medical equipment can significantly reduce the risk of CPO transmission.

How can CPOs be treated?

If a person is colonized with CPO, they do not need to be treated with antibiotics. If a person has an infection with CPO, the antibiotics that will work against it are limited, but some options are still available. In addition, some infections may be treatable with other therapies, such as draining the infection.

Tracking CPOs in BC

The first CPO case in British Columbia (BC) was identified in 2008 from a traveller returning from an endemic country where the patient had received medical procedures. Since then, the health authorities (HA), BC Center for Disease Control's Public Health Laboratory (PHL), the Provincial Infection Control Network of BC (PICNet), and the BC Ministry of Health have been working collaboratively to identify and monitor CPOs in the province.

A mandatory CPO surveillance program was established in BC's acute care facilities in July 2014. CPO-suspect isolates are required to be submitted to PHL for molecular testing and genotyping analysis. If the CPO is identified for the first time or identified with a gene encoding a new carbapenemase among inpatients, it is considered a new case of CPO and is to be reported to PICNet, who is responsible for publicly reporting the data. CPO was further designated a reportable condition in BC by the Provincial Health Officer on December 22, 2016. Under the revised provincial surveillance protocol for CPO, endorsed by the Provincial Communicable Diseases Policy Advisory Committee of BC, all newly identified cases of CPO in any health care setting (both acute care and community care) are to be reported to PICNet as of December 19, 2017.

Summary of CPO cases for Q2 of 2019/20

CPOs have been identified among patients in both acute care and community care settings, but remain uncommon in the majority of hospitals and communities. This quarterly report summarizes cases of CPOs newly identified by PHL and surveillance information for new cases reported to PICNet during fiscal quarter 2 of 2019/20 (Q2: June 28 – September 19, 2019).

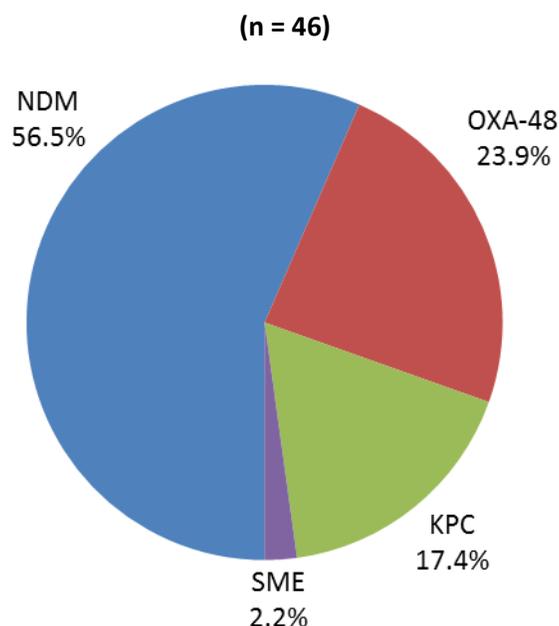
Of the isolates submitted to PHL during Q2, 44 isolates obtained from 41 patients were newly confirmed with CPO. Among them, 46 new cases of carbapenemase genes were identified, including four patients who each had two different carbapenemase genes and one patient who had three different carbapenemase genes – each gene identified for the first time in a given patient is considered a new case of CPO.

Of the 46 genes newly identified, 26 were NDM (accounting for 56.5%), 11 were OXA-48 (23.9%), 8 were KPC (17.4%), and 1 was SME (2.2% each) (Figure 1).

Surveillance information was collected and reported to PICNet for 38 out of 46 new cases (Table 1) – 36 cases (94.7%) were identified in acute care facilities and two cases (5.3%) were identified in community healthcare settings. Of the 36 cases in acute care facilities, 21 cases (55.3% of total reported cases) were identified in Fraser Health, 14 cases (36.8%) were identified in Vancouver Coastal Health, and 1 case (2.6%) was identified in Interior Health.

The surveillance information collected includes risk factors that may have contributed to CPO acquisition in the prior 12 months, including travel or healthcare encounters outside Canada (e.g. overnight hospitalization, medical or surgical procedures, etc.); close contact with a known CPO patient or the patient's environment; healthcare encounter in BC; and transfer from or stay in a care unit which was under investigation for CPO transmission. Among 38 reported cases, 14 cases (36.8%) reported healthcare exposure outside Canada. Another 10 cases reported travel without healthcare encounter. In addition, 24 cases (63.2%) were associated with other risk factors listed in the provincial surveillance protocol¹. Four cases (10.5%) reported no risk factors listed in the provincial surveillance protocol.

¹ These risk categories are not mutually exclusive – patients reporting healthcare exposure outside Canada may also be identified with other risk factors listed in the provincial surveillance protocol.

Figure 1. Distribution of carbapenemase genes newly identified in BC, Q2 of 2019/20 (June 28 – September 19, 2019)**Table 1. Number of new cases of CPO reported in BC by healthcare setting, Q2 of 2019/20** (June 27 – September 19, 2019)* (n = 38)

Healthcare setting	NDM	OXA-48	KPC	IMP	Total
Acute care facilities	22	7	7	0	36
<i>Interior Health</i>	0	1	0	0	1
<i>Fraser Health</i>	15	3	3	0	21
<i>Vancouver Coastal Health</i>	7	3	4	0	14
<i>Island Health</i>	0	0	0	0	0
<i>Northern Health</i>	0	0	0	0	0
<i>Provincial Health Services Authority</i>	0	0	0	0	0
Community healthcare settings	2	0	0	0	2
Subtotal in Q1 2019/20	24	7	7	0	38
Total in 2019/20	60	28	12	1	101

* based on the date of specimen collection from which a carbapenemase-encoding gene was first identified from the patient.

For more information about CPOs and the provincial surveillance program, please visit the PICNet website at <https://www.picnet.ca/surveillance/cpo>.