Improving access to lung cancer treatment in northern Canada: the role of oral molecularly targeted agents

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Canadians with cancer who live in the North experience a number of barriers to accessing diagnostic and treatment services. Accumulating evidence suggests that the travel burden experienced by cancer patients in remote and rural communities may influence their treatment choices. Some Canadian Inuit living in the Arctic, who must travel over 2000 km for cancer therapy, are reportedly forgoing treatment altogether.

Better understanding of the molecular biology of cancer and technologic advances in the ability to measure gene and protein abnormalities have recently led to the development of agents that target specific molecular pathways responsible for cancer growth and proliferation. The efficacy of these molecularly targeted therapies is largely governed by the importance of the targeted tumour pathway, which is increasingly determined by specific molecular biomarkers. For example, in the first-line treatment setting of advanced non–small-cell lung cancer, the oral epidermal growth factor receptor inhibitor gefitinib has been shown to improve progression-free survival, relative to chemotherapy, for some patients but not others.

In molecularly defined subsets of patients, molecularly targeted agents are becoming the standard of care across many tumour sites, resulting in substantial improvements in progression-free survival and overall survival, particularly for patients with advanced forms of cancer not amenable to surgery. Although some molecularly targeted agents are delivered intravenously with chemotherapy, to enhance cytotoxicity, a number of these agents are taken as daily oral therapies.

For northern residents, the introduction of oral molecularly targeted agents, which can be delivered in the community and require less frequent monitoring and specialist reassessment than chemotherapy (every 6–8 wk v. 3 wk), is highly relevant. Furthermore, whereas patients receiving chemotheraphy may experience lengthy or delayed toxic effects, necessitating access to 24-hour emergency health services, most oral agents have a short pharmacokinetic half-life, which allows for rapid reversal of treatment-induced toxic effects (e.g., rash, diarrhea, anorexia) on drug withdrawal. Oral therapies may therefore be a viable option in isolated northern communities.

For Canadian Inuit, the potential relevance of targeted cancer therapy is underscored by the availability of several targeted oral agents for some of the most common cancers affecting Arctic populations. For example, in Nunavut, lung cancer represents 39% of all cancer diagnoses, compared with 13% nationally. High rates of lung cancer among the Inuit are largely attributable to their high smoking rate (> 60%), which is the focus of many public health promotion and cancer prevention initiatives in Nunavut. Of note, four oral molecularly targeted agents are currently approved by Health Canada for the treatment of advanced non–small-cell lung cancer in patients harbouring either epidermal growth factor receptor mutations (three agents) or translocations of the anaplastic lymphoma kinase gene (one agent), and one or more of these agents are funded by provincial and territorial drug benefit plans or federal programs (e.g., Non-Insured Health Benefits Program) in most jurisdictions in Canada. The influence of these oral molecularly targeted agents on clinical outcomes will ultimately depend on the molecular characteristics of lung cancer among the Inuit, which are currently unknown. Given their probable Asian ancestry and the high prevalence of epidermal growth factor receptor mutations in Asian populations with non–small-cell lung cancer, targeted therapy with these agents may represent an important treatment opportunity for the Inuit.

### Key points

- Orally administered molecularly targeted agents represent an important treatment opportunity for patients with lung cancer who reside in remote northern communities.
- Appropriate infrastructure is required to support the testing and early identification of patient subgroups eligible for targeted therapy.
- Enhanced oncology training for rural health care professionals and stronger linkages to regional cancer centres must be developed and maintained.
- More research is required to establish the incidence of molecularly defined oncogenic drivers in northern populations.

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Although some of the earlier molecularly targeted agents in oncology were approved without mandated biomarkers, it has become apparent that in many instances genetic profiling and mutational analysis are essential for their targeted, cost-effective delivery. As such, biomarker identification will likely be a requirement for regulatory approval of molecularly targeted agents in the future. Unfortunately, in Canada, the infrastructure to support molecular screening and early identification of patients eligible for targeted therapy is unevenly distributed across the provinces, and in some cases may depend on external financial subsidies. An ethical problem arises when approval of targeted therapies is limited to molecularly defined subsets, in the absence of uniform access to molecular testing and mechanisms to allow for the early identification of eligible patients. In the Canadian context, it is imperative that the Inuit and other northern residents be included in the debate on this issue, given the unique challenges they face in accessing cancer treatments.

The survival gains achieved with targeted therapy for cancer in certain molecularly defined subsets have been astounding. Despite the challenges that may exist, national standards for molecular testing following a cancer diagnosis, as have been successfully implemented in other countries, should be developed in Canada. Screening and molecular testing for oral targeted therapies are particularly relevant to cancer care for Canadians in the North, who stand to reap substantial benefit from cancer treatment in the community. Concurrently, strategies to monitor patients must be developed to ensure the safe and effective delivery of personalized therapies in rural settings. In this context, enhanced oncology training for rural health care professionals and improved patient education will be important to avoid and manage acute treatment-induced toxic effects, particularly given the higher incidence of such effects (e.g., rash, diarrhea) reported with some of the newer, more potent second-generation targeted therapies. In addition, stronger linkages with regional cancer centres must be developed and maintained using existing technologies such as telehealth, to avoid unnecessary travel and to provide the necessary support for health care providers in the North.

Canadians should have access to modern cancer therapies, regardless of where they live. With the introduction of orally administered targeted agents, there may be an opportunity to increase the proportion of northern patients getting access to effective anticancer therapies. Targeted therapy may also improve their quality of life, by reducing the travel burden associated with treatment and enabling treatment in the community. Finally, although molecularly targeted therapies are more expensive than most chemotherapy, the increased cost may be offset by a decrease in hospital admissions and a reduction in federally and provincially reimbursed travel and lodging, as a result of fewer trips to urban cancer treatment centres. With the right support mechanisms and molecular screening infrastructure, targeted therapy can be an important component of better cancer care for northern residents.

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