A Guide for Health Professionals Working with Aboriginal Peoples

EXECUTIVE SUMMARY

This Policy Statement has been reviewed by the Aboriginal Health Issues Committee and approved by Executive and Council of the Society of Obstetricians and Gynaecologists of Canada.

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SUPPORTING ORGANIZATIONS
Assembly of First Nations
Canadian Institute of Child Health
Canadian Paediatric Society
College of Family Physicians of Canada
Congress of Aboriginal Peoples
Federation of Medical Women of Canada
Inuit Tapirisat of Canada
Metis National Council
National Indian and Inuit Community Health Representatives Organization
Pauktuutit Inuit Women's Association

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Executive Summary

Aboriginal health is a highly relevant concern to the health care community. Aboriginal patients, whether identified as Aboriginal or not, will be encountered by most Canadian health care practitioners at some point in their practice. Such encounters may range from the daily to the infrequent; from the urban specialist who may be required to communicate the management of a complex disease process to an Aboriginal patient flown in from a remote location, to the rural community health nurse who may be providing perinatal instruction to a young Aboriginal family, to the rural family physician who practices in or close to an Aboriginal community. In forging a balanced and positive health care practitioner-patient relationship, it is essential to understand the heritage context of the Aboriginal patient in order to negotiate chasms and find common ground.

History leaves a difficult contextual legacy. The adverse socioeconomic environment facing many Aboriginal
communities can be directly linked to the European appropriation of the traditional land of many Aboriginal nations: land that had been the basis of economic livelihood for Aboriginal communities prior to colonization. Additionally, the European colonists and their subsequent governments brought with them epidemics of disease, slavery, cultural suppression, imposed religious practices, family disruption and community relocation, and physical, emotional, and sexual violence. Although Canada is often referred to as the ideal place to live because of its quality of life, there are two realities in the nation. One is for most Canadians; the other is for ... Aboriginal peoples. The statistics on the first inhabitants of what is now Canada are much like the statistics for most indigenous populations who have undergone colonization: inadequate nutrition; substandard housing and sanitation; unemployment and poverty; discrimination and racism; violence; inappropriate or absent services; high rates of physical, social and emotional illness and injury; disability; and premature death.

One predictable consequence of these socioeconomic inequities is that Aboriginal peoples have an overall health status that falls well below that of other Canadians. Aboriginal life expectancy is approximately seven to eight years less than that of the average Canadian. Chronic diseases, post-neonatal mortality, accidental deaths, certain infectious diseases, and mental health problems continue to be disproportionately common among Aboriginal peoples. Although there have been some significant improvements in Aboriginal health over the past decades, the health care system has not yet been successful in rectifying this disparity.

The Royal Commission on Aboriginal Peoples (RCAP), whose purpose is to help restore justice to the relationship between Aboriginal and non-Aboriginal peoples in Canada by investigating the foundations of a fair and honourable relationship, has recognized the unacceptable level of health problems among Aboriginal peoples, the linkages of these health problems to larger social, economic, and emotional issues, and the inadequacy of the current health infrastructure to impact significant change. Its proposed new health strategy recommended long-term structural changes in government relations and economic development, as well as a reorganization of existing health and social service systems. Mainstream services were encouraged to "undertake a systematic assessment of their practices to see how they [could] improve their connections with Aboriginal peoples." This document is in keeping with previous recommendations by the Canadian Medical Association (CMA) which, in their submission to RCAP, recommended that the government of Canada "develop educational initiatives in cross-cultural awareness and other Aboriginal health issues for the Canadian population and in particular health care providers." In a later (1996) CMA sponsored workshop on Aboriginal women's health issues, the participants, comprised mainly of Aboriginal women, recommended that:

The Canadian Medical Association, the Native Physicians Association in Canada or both should develop, or facilitate the development of, a series of publications outlining the historical concepts and approaches to health of Aboriginal peoples. The publications should also include information on variability and multiculturalism among the First Nations and Aboriginal peoples of Canada— even within individual communities. These publications should be distributed to all health service providers to heighten awareness and sensitize them to the present state of Aboriginal health.

The Aboriginal Health Issues Committee of the Society of Obstetricians and Gynaecologists of Canada (SOGC), founded in 1994, is a multidisciplinary committee with Aboriginal and non-Aboriginal members, with representation from several Aboriginal organizations and backgrounds including First Nations, Inuit, and Métis. Its guiding principles, adopted at the first meeting, derive from the Keewatin Regional Health Board mission statement. The Committee additionally supports the recommendations and guiding principles in the CMA’s Bridging the Gap document.

This policy statement brings together concise information and specific recommendations regarding attitudes, knowledge base, and skills, to help the health professional build a more balanced two-way relationship with Aboriginal patients. Information and recommendations are grouped in four sections: sociodemographics, health definitions and issues, cross-cultural communication, and health resources. It is hoped that implementation of these recommendations and information will enhance understanding of Aboriginal cultures, approaches to health and healing, and worldviews.

A. METHODS
1. Social, cultural, and historical context

The creation of these evidence-based guidelines presented several challenges for the authors. For any such creation to be relevant, Aboriginal health issues should be framed in their social, cultural, and historical context. This adheres to RCAP findings regarding the causal connection between specific illnesses and "factors outside the boundaries of ordinary medicine—social, emotional, and economic conditions that in turn lead back to the corrupt, destabilizing, and demoralizing legacy of colonialism;" as well as the desire by Aboriginal peoples for their health issues to be understood holistically, summarized here by Henry Zoe of D’ogrib Treaty 11 Council:

For a person to be healthy [he or she] must be adequately fed, be educated, have access to medical facilities, have access to spiritual comfort, live in a warm and comfortable house with clean water and safe sewage disposal, be secure in cultural identity, have an opportunity to excel in a meaningful endeavour, and so on. These are not separate needs; they are all aspects of a whole.

The five volume RCAP report, the product of five years of extensive public hearings, community visits, expert consultation,
commissioned research studies, and literature review, is an important knowledge source for Aboriginal health needs.

2. Language, oral and written
Aboriginal cultures historically followed an “oral tradition” in which information was collected, conserved, and transmitted through carefully memorized verbal discourse often including stories and songs. Although there are examples of stories recorded through artwork prior to colonization, written information sources are of relatively recent European introduction into the Aboriginal community. For these reasons, a feeling of distrust regarding written information exists among some Aboriginal individuals, some of whom feel that important cultural information, such as traditional healing techniques, should only be transmitted orally and not be written down. Thus, at least some important information regarding Aboriginal health is unlikely to be found in a written format, and much data regarding Aboriginal peoples is not published in the standard accessible sources.

Additionally, lower rates of literacy and formal education achievement among Aboriginal peoples in Canada create systemic barriers to publication by Aboriginal authors; and those who do get published may decide that their information transmitted as a story, political organizational document, or other resource material is more consistent with their perspective than formal scientific literature. Using a comprehensive search strategy, LeMaster found only a limited number of published intervention studies in the health education literature which included “Native Americans.” However, the author had encountered elsewhere numerous unpublished newsletters and other commissioned documents from governmental and Aboriginal organizations, none of which could be found in a literature search, although some of them contained relevant data. LeMaster concluded that “it is very likely ... that many health education interventions are conducted by and for Native Americans, but results are not disseminated in the published literature.”

3. Language groups
Aboriginal peoples in Canada represent over 50 culturally distinct language groups. Although often grouped together by non-Aboriginals, it is important to remember that First Nations, Metis, and Inuit peoples each have a unique cultural, social, and historical context, and are represented by separate political organizations. Increasingly, specific and focussed health programming is being called for by different Aboriginal groups. For example, at the May 2000 Forum on Aboriginal Health Services and Issues, Okalik Eegeesiak, President of the Inuit Tapirisat of Canada, called for health programmes specifically for Inuit peoples, to be based on Inuit approaches to health issues. Another distinct set of health issues is raised by the historical exclusion of Metis people from health programmes administered by the federal government via the former Medical Services Branch. Although this policy statement presents one set of recommendations intended to apply generally, the specific and distinct health concerns of First Nations, Inuit, and Metis peoples are detailed in the text where possible.

4. Culturally appropriate, balanced format
Finally, the Committee wished to present the information and recommendations in a culturally appropriate, balanced format. A careful, systematic documentation of the enormous health problems facing Aboriginal peoples in Canada could easily lead to a document that, while factual, is bleak and depressing. The personal cultural perspective of one author was that “bad news” needs to be balanced with “good news.” This notion of “balance” is fundamental to Aboriginal concepts of health. The authors have therefore attempted to achieve a balanced presentation which communicates the inherent resources of Aboriginal individuals and communities at the same time as it identifies health issues.

5. Process
Just as this document attempts to bridge the perspectives of health professionals and Aboriginal clients, so the methodology has required a synthesis of the published, scientific knowledge base with a contextual, culturally-informed approach.

A framework for this policy statement was circulated to Committee members by the Committee chair in 1997. This framework was revised by Committee members who were then assigned different sections of the framework on which to work. Information on the different sections was brought back to the Committee and two Committee members organized and expanded upon this data to produce a poster presentation at the 1998 SOGC Annual Clinical Meeting (ACM) in Victoria.

A formal literature review was initiated in 1998, with a computerized Medline search conducted using the M eSH headings “Indians, North American or Eskimos” and “Health.” The search included all review articles, randomized clinical control trials, meta-analyses, and practice guidelines after 1966 inclusive. Ninety-five review articles, ten randomized clinical control trials, no meta-analyses, and one practice guideline were identified for a total of 106 articles, many from public health and social science sources. Papers were reviewed by a single researcher (JKS), and the article deemed relevant if it contained information about Aboriginal peoples (preferably in the Canadian context) and the topic was health (including sociodemographic determinants of health, health policy, and health education). Eighty-six of these articles were deemed relevant for review and all 86 articles were retrieved. Data was extracted and organized according to the four major sections of the framework: sociological context, health concerns, cross-cultural understanding, and Aboriginal health resources. Ancillary and unpublished references were recommended by members of the SOGC Aboriginal Health Issues Committee and the panel of expert reviewers.

A draft series of recommendations incorporating the previously adopted guiding principles as well as recommendations...
regarding Aboriginal health from other sources was produced and circulated to the Committee in early 1999. Members responded with their feedback and the recommendations were revised. Ongoing work by Committee members helped produce tools to educate health professionals in the areas of cross-cultural communication. A roleplaying session was piloted at the 1998 AGM of the Association of Professors of Obstetrics and Gynaecology and presented at the 1999 SOGC ACM in Ottawa. In the autumn of 1999, one of the Committee members was commissioned to finish the literature review and produce a draft document, which was circulated to Committee members and a panel of experts and then revised, before presentation for review by the SOGC Executive and Council.

RECOMMENDATIONS

A. SOCIOCULTURAL CONTEXT
1. Health professionals should have a basic understanding of the appropriate names with which to refer to the various groups of Aboriginal peoples in Canada.
2. Health professionals should have a basic understanding of the demographics of Aboriginal peoples in Canada.
3. Health professionals should familiarize themselves with the traditional geographic territories and language groups of Aboriginal peoples.
4. Health professionals should have a basic understanding of the disruptive impact of colonization on the health and well-being of Aboriginal peoples.
5. Health professionals should recognize that the current sociodemographic challenges facing many Aboriginal individuals and communities have a significant impact on health status.
6. Health professionals should recognize the need to provide health services for Aboriginal peoples as close to home as possible.
7. Health professionals should have a basic understanding of governmental obligations and policies regarding the health of Aboriginal peoples in Canada.
8. Health professionals should recognize the need to support Aboriginal individuals and communities in the process of self-determination.

B. HEALTH CONCERNS
1. Health professionals should appreciate holistic definitions of health as defined by Aboriginal peoples.
2. Health professionals should recognize that the degree of ill health in Aboriginal populations is unacceptable, and work with Aboriginal individuals and communities towards improved health outcomes.
3. Health professionals should recognize and respond to key areas of morbidity and mortality without stereotyping.

C. CROSS-CULTURAL UNDERSTANDING
1. Relationships between Aboriginal peoples and their health care providers should be based on a foundation of mutual respect.
2. Health professionals should recognize that the current health care system presents many gaps and barriers for Aboriginal individuals and communities seeking health care.
3. Health professionals should work proactively with Aboriginal individuals and communities to address these gaps and barriers.
4. Health professionals should work with Aboriginal individuals and communities to provide culturally appropriate health care.
5. Aboriginal peoples should receive treatment in their own languages, whenever possible.
6. Health care programmes and institutions providing service to significant numbers of Aboriginal peoples should have cultural interpreters and Aboriginal health advocates on staff.
7. Aboriginal peoples should have access to informed consent regarding their medical treatments.
8. Health services for Aboriginal peoples should recognize the importance of family and community roles and responsibilities when attempting to service Aboriginal individuals.
9. Health professionals should respect traditional medicines and work with Aboriginal healers to seek ways to integrate traditional and western medicine.
10. Health professionals should take advantage of workshops and other educational resources to become more sensitive to Aboriginal peoples.
11. Health professionals should get to know Aboriginal communities and the people in them.

D. ABORIGINAL HEALTH RESOURCES
1. Aboriginal communities and health professionals working with Aboriginal peoples should support the creation of community-directed health programmes and services for Aboriginal peoples.
2. Aboriginal communities and health professionals working with Aboriginal peoples should support the development of community-directed, participatory health research for Aboriginal peoples.
3. Aboriginal communities and health professionals working with Aboriginal peoples should recognize the need for preventative health programming in Aboriginal communities.
4. Aboriginal communities and health professionals working with Aboriginal peoples should encourage the education of Aboriginal health professionals committed to future work in Aboriginal communities.

REFERENCES


Front cover illustration: Jay Bell-Redbird, an Ojibwa artist from Wiikwemiking First Nation, created the logo on the cover for the SOGC Committee on Aboriginal Health Issues. The logo represents the diversity of Aboriginal peoples in Canada. The medicine wheel base, a symbol used by many indigenous communities, represents a paradigm of teaching and healing that promotes wholeness, harmony and balance. The four colours are significant in healing work and also represent the diversity of all humanity. The floral motif is very representative of beadwork designs done by women in the woodland and sub-arctic regions of Canada. The sash around the circle is a symbol of the Metis nation. The drum with a moon symbol is drawn in the style of the west coast peoples. The great white pine and the four roots of peace are important symbols of the Iroquois confederacy. The eagle, a sacred bird to many nations, represents freedom, honour, and respect. The Inuit woman with her baby in an amauti at the centre of the circle represents the importance of women as the bearers of life and the importance of children as the future hope of the people.