
**Council for Trade-Related Aspects
of Intellectual Property Rights**

Original: English/
French

**NOTIFICATION UNDER PARAGRAPH 2(C) OF THE DECISION OF 30 AUGUST 2003
ON THE IMPLEMENTATION OF PARAGRAPH 6 OF THE DOHA DECLARATION
ON THE TRIPS AGREEMENT AND PUBLIC HEALTH**

CANADA

The following notification has been received from the delegation of Canada on 4 October 2007 for circulation to the Council for TRIPS.

In accordance with paragraph 2(c) of the WTO General Council Decision of 30 August 2003 on the Implementation of Paragraph 6 of the Doha Declaration on the TRIPS Agreement and Public Health, attached* ¹ is the authorization granted on 19 September 2007 by Canada's Commissioner of Patents to Apotex, Inc, pursuant to section 21.04 of the *Patent Act*.² Rwanda had previously filed a related notification, dated 17 July 2007, pursuant to paragraph 2(a) of the same Decision of the WTO General Council (IP/N/9/RWA/1).

Pursuant to paragraph 2(c) and 2(b)(iii) of the Decision of 30 August 2003 on the Implementation of paragraph 6 of the Doha Declaration on the TRIPS Agreement and Public Health, the information on the shipment (quantities and distinguishing features) will be posted on the licensee's website at: www.apotex.com/apotriavir/abouttriavir.asp.

* In English only.

¹ Canada's Commissioner of Patents authorization can also be found at: http://strategis.gc.ca/sc_mrksv/cipo/new/CAMR_Authorization.pdf.

² The WTO General Council Decision was implemented in Canada by an *Act to amend the Patent Act and the Food and Drugs Act (The Jean Chrétien Pledge to Africa Act)*, of 2004. The text of Canada's Patent Act may be found at http://laws.justice.gc.ca/en/showdoc/cs/P-4/bo-ga:s_21_01/en#anchorbo-ga:s_21_01.

ANNEX

AUTHORIZATION UNDER SECTION 21.04 OF THE PATENT ACT

In the matter of application for authorization number CAMR-1 by Apotex, Inc. for export to Rwanda of the following pharmaceutical product:

(a) if the pharmaceutical product is a drug as set out in section 2 of the *Food and Drugs Act*: A fixed dose combination tablet of lamivudine (150mg) + nevirapine (200mg) + zidovudine (300mg), as provided in Schedule 1 to the *Patent Act*

(b) if the pharmaceutical product is a medical device:

1. I hereby authorize Apotex, Inc. whose postal address is

150 Signet Drive
Toronto, Ontario
M9L 1T9

to make, construct and use, the patented inventions identified in patent numbers 2,311,988; 2,070,230; 2,068,790; 2,286,126; 2,216,634; 2,105,487; 2,059,263; 2,009,637 and 2,030,056 solely for purposes directly related to the manufacture of the above-mentioned pharmaceutical product, and to sell it for export to the above-mentioned country or WTO Member.

2. The quantity of the pharmaceutical product authorized to be manufactured by this authorization is 15,600,000 tablets.

3. In accordance with section 21.09 of the Act, this authorization is valid for a period of two years, beginning on the date shown below.

Granted at Gatineau, Quebec, the 19th day of September, 2007-10-05

Mary Carman
Commissioner of Patents
