Dear Director:

RE: [Product name/generic name, strength, and dosage form (the “Product”) manufactured by <name of manufacturer> (“the Manufacturer”)].

For the purpose of this letter, the following terms have the following meanings:

“DIDFA” means the Drug Interchangeability and Dispensing Fee Act, R.S.O.1990, c.P.23;

“ODBA” means the Ontario Drug Benefit Act, R.S.O. 1990, c.O.10;

“Person” means one of the persons listed in subsection 12.1(1) of the DIDFA and subsection 11.5(1) of the ODBA;

“Rebate” means a rebate as defined under subsection 12.1(14) of the DIDFA and subsection 11.5(15) of the ODBA.

The Manufacturer hereby represents and warrants that the Manufacturer has not provided a Rebate to any Person with respect to the Product contrary to the ODBA and/or DIDFA since Health Canada approved the Product for sale in Canada.
[Signature]

[Name and Title of Senior Company Official]

I have authority to bind the Manufacturer