Attachment 7

CERTIFICATE OF PHARMACEUTICAL MANUFACTURER IN THE FRAMEWORK OF THE AGREEMENT ON MUTUAL RECOGNITION RELATING TO CONFORMITY ASSESSMENT BETWEEN CANADA, ON THE ONE HAND, AND THE EEA EFTA STATES, ON THE OTHER

As requested by the

.....(*)

on ...../...../..... (date)

(reference:.....),

the Competent Authority of .....(**) confirms the following:

The company .....

whose legally registered address is:

.....

.....

.....

has been authorized, under Directive 75/319/EEC (Article 16) (EEA Agreement, Annex II, Chapter XIII, point 3) and Directive 81/851/EEC (Article 24) (EEA Agreement, Annex II, Chapter XIII, point 5) transposed in the national legislation of..... (**), under the authorisation reference number covering the following site(s) of manufacture (and contract testing laboratories, if any):

1.....

.....

2.....

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3.....

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