APPLICATION FOR WAIVER
Section 29(3) § of the Medicines Act (395/87)

The grounds for the waiver shall be given on the appendix.

Under section 29 of the Medicines Act, marketing authorisation or registration lapses, i.e. ceases to be valid if an authorised or registered product has not been placed on the market within three years of the granting of the authorisation or registration or if an authorised or registered product has been absent from the market for a period of three consecutive years.

For reasons relating to the health of humans or for other particular reasons The Finnish Medicines Agency may upon application from the marketing authorisation or registration holder decide that the marketing authorisation or registration shall not lapse.

Product type	
☐ Human medicinal product	☐ Registered product

Product data

Marketing authorisation number or registration number	Procedure number (mutual recognition or decentralised procedure)
Date of issue of marketing authorisation or registration	
Name of medicinal product	
Strength	
Pharmaceutical form	
Name of marketing authorisation or registration holder	

Contact information for applicant

Name of company
Surname
Given name
Address
Telephone
Email address

Other items of note

Signature

Date	Signature

Please return the signed form and appendix electronically or to:

Finnish Medicines Agency
P.O. Box 55
FI-00034 FIMEA, FINLAND

APPENDIX
Grounds for continuing validity of Marketing authorisation or Registration

Marketing authorisation number or registration number

Lääkealan turvallisuus- ja kehittämiskeskus | Säkerhets- och utvecklingscentret för läkemedelsområdet | Finnish Medicines Agency
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