



NOTIFICATION OF CANCELLING A MARKETING AUTHORISATION/ REGISTRATION

The marketing authorisation or registration holder or a person or company authorised by the holder should submit a written notification before the end of the authorisation year if the authorisation is to be cancelled and the annual fee left unpaid.

The marketing authorisation/registration is to be cancelled starting from the following date

Year	Month	Day
Reason for cancellation		
Type of medicinal product/preparation		
Human medicine	Veterinary medicine	Herbal medicinal product
Parallel imported product		

Product details

Marketing authorisation number/Registration number
Name of the medicinal product
Strength
Pharmaceutical form
Marketing authorisation holder/Registration holder

Contact details for the sender

Company name	
Surname	First name
Address	
Phone	
E-mail address	
Other comments	
If the product to be cancelled has Finland as the RMS, is the change of RMS pending? Yes No, Marketing Authorisation will be cancelled in all CMS countries	

If the medicinal product is for human use, the reasons for withdrawal are the ones given in the Medicines Act, section 27, third paragraph?

Yes No

If you answer yes, this notification has to be accompanied with two separate reports in English (cover letter and notification report table 'Notification of withdrawn products') and both of them have to be sent to both Fimea and EMA (withdrawnproducts@ema.europa.eu).

Date and signature

Date	Signature
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NB. A single form can be used for the cancellation of several marketing authorisations, provided that all the marketing authorisations listed in the form have the same cancellation date.

The signed form should be submitted to registry@fimea.fi or by CESP-submission.