



## e-form

# Application for the Transfer of a Marketing Authorisation Holder during the Authorisation Procedure – (National Phase)



Please note that submission must be sent electronically (CD/DVD or through the Common European Submission Portal – CESP).

This application enables a company to take over responsibility from another company (different legal entity) where the application is in identical terms to the existing marketing authorisation.

DOCUMENTS WHICH HAVE TO BE SUBMITTED WITH THE APPLICATION FORM FOR A MARKETING AUTHORISATION TRANSFER FOR AN AUTHORISED PRODUCT.

Details of documents to be submitted : (Kindly indicate if submitted)

Legal document for Transfer (application form).

Proof of payment (as per current legal notice on fees).

Proof of establishment of the new Marketing Authorisation Holder (from official sources).

SmPC (Word format) /Package Leaflet & Labelling bearing the new:

- Marketing Authorisation Holder Name
- Address
- Marketing Authorisation Number for Malta

Type IB variation (Type IB C.I.g.z) has been submitted and other relevant variations pertaining to the transfer (e.g. change in name of the medicinal product affected due to transfer).

Pharmacovigilance system master file (PSMF) : (Kindly indicate if submitted )

PSMF declaration

Required date for Change of ownership of the existing (as proposed in the Marketing Authorisation application) Marketing Authorisation Holder to the new proposed holder.

Date:

The earliest transfer date of the Marketing Authorisation Holder is the date of receipt of the application by the Medicines Authority.

#### 1 DETAILS OF THE PROPOSED MARKETING AUTHORISATION HOLDER AFTER TRANSFER

Name and address:

Proposed trading style (if applicable):

Name and address of the applicant acting on behalf of the proposed Marketing Authorisation Holder, if different:

#### 2 DETAILS OF THE COMPANY CURRENTLY PROPOSED AS THE MARKETING AUTHORISATION HOLDER IN THE MARKETING AUTHORISATION APPLICATION IN PROCESS.

Name and address:

#### 3. DETAILS OF THE PRODUCT

Current MA Number:

Name of product:

Name of active substance(s):

Pharmaceutical form:

Strength(s):

STATEMENT TO BE SIGNED BY THE COMPANY CURRENTLY PROPOSED AS THE MARKETING AUTHRISATION HOLDER.

This dossier includes all of the data in support of the original application together with all correspondence with the Medicines Authority concerning the product and all pharmacovigilance data both before and after the issue of the original MA.

3. I acknowledge our responsibilities in the event of any adverse reaction or quality defect associated with any remaining product bearing our name, address and MA number.

4. I acknowledge our responsibilities in the event of the necessity to recall from the market any remaining product bearing our name, address and MA number.

REASON FOR TRANSFER APPLICATION:

1. I hereby notify the Medicines Authority that

is to be transferred to

2. I confirm that the entire dossier for the product has been transferred to

Signed:

Date:

Status of signatory:

Telephone No:

Fax No:

Email address:

STATEMENT TO BE SIGNED BY THE PROPOSED MARKETING AUTHORISATION HOLDER AFTER TRANSFER.

REASON FOR TRANSFER APPLICATION:

1. I will have the sole responsibility for the product including obtaining approval for any changes subsequent to the grant of this product information.

2. I have received the entire dossier for

from

This dossier includes all of the data in support of the original product authorisation application together with all correspondence with the Medicines Authority concerning the product and all Pharmacovigilance data both before and after the issue of the original product authorisation.

3. I have been assured by the current MA holder/applicant that, apart from the change of name and address of the product authorisation holder and the product authorisation number, the dossier on which the transfer is based is identical in every respect to that submitted by the original holder.

Signed:

Date:

Status of signatory:

Telephone No:

Fax No:

Email address: