Model for manufacturers of a
MARKETING INFORMATION FORM

Notification of the intention to market a batch of an immunological medicinal product, which has a marketing authorisation, or medicinal product derived from human blood or plasma, which has a marketing authorisation, in the following EC/EEA member state .....

Addressee:	*'Name and address of specified contact person(s) in the member state/EU where the batch of product is to be marketed'*

Trade name:	'*Trade name of the product in the member state/EU where the batch of product is to be marketed'*
Batch number(s) appearing on the market package:	*'Batch number of the product as in the member state/EU where the batch of product is to be marketed '*
Other batch identification numbers associated with this batch[1]:	*'Filling bulk number, final lot number and packaging lot number'*
Number of containers to be marketed in the member state:	
Market authorisation number:	*'MA number in the member state/EU where the batch of product is to be marketed'*
Name and address of marketing authorisation holder :	'*MA holder in the member state/EU where the batch of product is to be marketed'*
Date of start of period of validity:	
Expiry date in the member state where the batch is to be marketed:	
Intended date of marketing:	

OMCL performing batch release:	
Official batch release certificate number:	

I hereby declare that:

- this batch is in compliance with the above marketing authorisation and the relevant European Pharmacopoeia monographs ;
- this batch is the batch referred to in the accompanying batch release certificate.

A copy of the batch release certificate is attached.

Signature of qualified person:	
Name of qualified person:	
Date of issue:	

[1] Sufficient detail should be given to allow clear traceability back to the level of the final bulk