**Model for manufacturers of a**

**MARKETING INFORMATION FORM for use in the United Kingdom**

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 **United Kingdom.**

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| Addressee:  | Control Records Office, NIBSC, Blanche Lane, South Mimms, Potters Bar, EN6 3QG, United Kingdom (UK) t: +44 1707 641000 e: marketinginformationform@mhra.gov.uk   |
|   |   |
| Trade name:  | ‘*Trade name of the product in the UK’*  |
| Batch number(s) appearing on the market package:  | *‘Batch number of the product in the UK’*  |
| Other batch identification numbers associated with this batch1:  | *‘Filling bulk number, final lot number and packaging lot number’*  |
| Number of containers to be marketed in the UK:  |   |
| EU Market authorisation number:  | *‘MA number used for EU certification’*  |
| UK Market authorisation number:  | *‘MA number in the UK’*  |
| Name and address of marketing authorisation holder:  | *‘MA holder in the UK’*  |
| Date of start of period of validity:  |   |
| Date of expiry in the UK:  |   |
| Intended date of marketing (dd/mm/yyyy):  |   |
|   |   |
| OMCL where certificate was issued:  |   |
| Official batch release certificate number:  |   |

I hereby declare that:

-this batch is in compliance with the above marketing authorisations 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.

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

 Sufficient detail should be given to allow clear traceability back to the level of the final bulk.