**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](mailto: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.