Marketing Information Form

Notification of the intention to market a batch of a medicinal product which is subjected to Swiss Official Control Authority Batch Release according to EU/EEA Guidelines and the MRA Switzerland – EC, Annex 1, Chapter 15. For detailed information refer to *Official Control Authority Batch Release*.

Type of medicinal product	Medicinal product derived from human blood or plasma for human use
Name of applicant (Marketing Authorization Holder)	
Trade name of the product	
Marketing authorization number	
Manufacturer	
Batch number appearing on market package (Packaging Lot number)	
Other batch numbers (Filling Lot number)	
Batch number of diluent (*where appropriate, only for veterinary medicinal products*)	
Type of container	
Number of doses* / Volume container (**not applicable for blood products for human use*)	
Date of start of period of validity *(only applicable to human medicinal products)*	
Expiry date	
EU OMCL performing batch release	
Official EU batch release certificate number	
Type of submitted certificate *(only applicable to veterinary medicinal products)*	n/a

I hereby declare that:
- this batch is in compliance with the above marketing authorization and the relevant European Pharmacopoeia monographs
- this batch is the batch referred to in the accompanying batch release certificate

For human medicinal products: A copy of the official EU batch release certificate is attached
For veterinary medicinal products: A copy of the OBPR/OCABR certificate and the manufacturer's protocol are attached

Name of qualified person	
Date of issue	
Signature of qualified person	