Marketing Information Form (MIF)

Notification of the intention to market a batch of a medicinal product which is subjected to Swiss Official Control Authority Batch Release (OCABR) 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	Immunological medicinal product for venterinary use
Name of applicant (Marketing Authorization Holder in Switzerland)	
Trade name of the product	
Marketing authorization number	
Name and address of Manufacturer (galenic production) *(In case the Manufacturer is located outside the EU/EEA please use the form for request for OCABR/OBPR instead of MIF)*	
Bulk lot number *(for veterinary medicinal products only)*	
Filling lot number	
Packaging lot number *(Batch number appearing on market package)*	
Batch number of diluents (*where appropriate, for veterinary medicinal products only*)	
Type of container	
Number of doses* / Volume per container (**not applicable for blood products for human use*)	/
Date of start of period of validity	
Expiry date	
OMCL which performed EU batch release	
Official EU batch release certificate number	
Type of submitted certificate *(Applicable to veterinary medicinal products only)*	OBPR

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	