

**OFFICIAL CONTROL AUTHORITY  
BATCH RELEASE  
OF  
IMMUNOLOGICAL VETERINARY  
MEDICINAL PRODUCTS**

**2012**

**RABIES VACCINE FOR FOXES (LIVE)**

<b>Document title</b>	<b>Official control authority batch release of rabies vaccine for foxes (live)</b>
<b>Legislative basis</b>	<b>Council Directive 2001/82/EC amended by Directive 2004/28/EC</b>
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<b>Custodian Organisation</b>	<b>The present document was elaborated by the EDQM through the OMCL network and is finalised under PA/PH/OMCL (04) 5 DEF CORR</b>

## **OFFICIAL CONTROL AUTHORITY BATCH RELEASE OF RABIES VACCINE FOR FOXES (LIVE)**

*OMCLs performing batch release on this product should receive a completed signed protocol from the MAH (model template available separately on the EDQM website ([www.edqm.eu](http://www.edqm.eu))) and the required samples.*

*The licensing authority provides the OMCL with all necessary data from the quality part of the dossier such as relevant Pharmacopoeia monographs, list of tests to be performed on each batch and the SOPs as presented in the dossier.*

### **1 INTRODUCTION**

Official Control Authority Batch Release of immunological products for veterinary use is performed within the framework of Article 82 of Directive 2001/82/EC as amended by Directive 2004/28/EC and following the current guideline on EC administrative procedure for official control authority batch release of IVMPs.

The Ph Eur monograph 0746 is relevant for this product.

### **2 SAMPLING AND TESTS TO BE PERFORMED BY THE OFFICIAL CONTROL LABORATORY**

The following samples should be supplied to the Official Medicines Control Laboratory performing batch release:

At least 20 containers of each final lot (blisters and baits).

The Control Laboratory should perform the following tests:

- Appearance
- Virus titre