



## CERTIFICATE OF ANALYSIS

Product: **Biocan R vakcina A.U.V.**

Batch No.: **40 56 20 B** 1 dose

Expiry: **04. 06. 2015**

Date of production: **05. 06. 2013**

### Results of analysis:

Tests	Specification	Results
Appearance	Slightly pink solution, which might content fine sediment.	conforms
Content of the original package	min. 1 ml, 5 ml, 10 ml, 20 ml	1.04 ml
Sterility	Vaccine should be sterile.	sterile
Safety	The product is satisfactory if no systemic or local abnormal reaction is observed during the testing period.	conforms
Potency	The vaccine must contain min. 2 IU/ml.	5.4 IU/ml
Identity	In the serum of vaccinated animals must be specific antibody levels $\geq 0.5$ IU/ml.	1.21 IU/ml
Inactivation	The vaccine must not contain a vivid rabies virus.	conforms
pH	7.2 – 8.2	7.80
Aluminium oxide	0.15 – 0.30 %	0.204 %
Merthiolate	0.0085 – 0.0115 %	0.0096 %
Air tightness	No colour change.	conforms

I certify that this product has been tested in the laboratories of Bioveta a.s. and has been found to meet specification requirements. Preparation is suitable with control regulations of the valid PN No. 233.



Mgr. Hana Nežvalová  
HEAD of QC, QP  
03/10/2013



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