

RABISIN displays consistent high potent glycoprotein G level as tested by ELISA test at release

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INTRODUCTION

- Importance of using high-quality rabies vaccines to achieve the goal of Zero by 30.
- **First** in vitro ELISA test able to quantify the rabies glycoprotein G antigen in RABISIN formulated vaccine was developed as an alternative to NIH potency test in 2015¹.
- ELISA titration results of recent batches of RABISIN produced in 2023 are presented

METHODS

- Glycoprotein G is the primary antigen that stimulates virus-neutralizing antibody production (immunity) against rabies virus infection.
- Quantity of glycoprotein G of RABISIN® measured to ensure the highest quality in terms of immunogenicity
- Key advantages
 - ✓ specificity
 - ✓ robustness



A versatile *in vitro* ELISA test for quantification and quality testing of infectious, inactivated and formulated rabies virus used in veterinary monovalent or combination vaccine

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In vivo Potency test
NIH

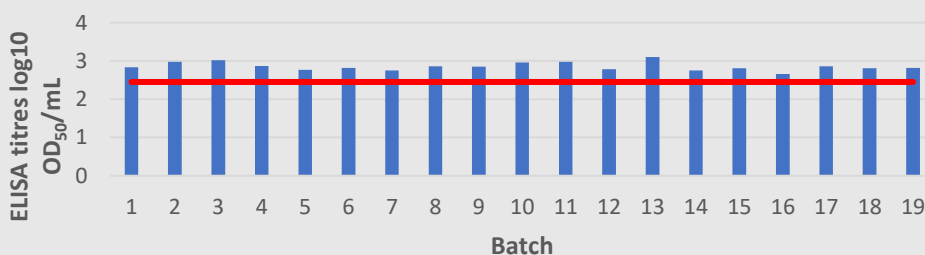


In Vitro Potency

- 19 batches of RABISIN® produced in 2023 tested by ELISA

RESULTS

Potency measured by Glycoprotein G level



- Best alternative to NIH:
 - ✓ Animal welfare
 - ✓ More discriminant to confirm batch-to-batch consistency,
 - ✓ Safety benefit for laboratory staff
 - ✓ Lead-time to market significantly reduced

CONCLUSION

High potent glycoprotein G quantity for RABISIN and batch to batch consistency

RABISIN® is a registered trademark of Boehringer Ingelheim in France and elsewhere

References:

1. C. Sigollot-Claude *et al.* A versatile in vitro ELISA test for quantification and quality testing of infectious, inactivated and formulated rabies virus used in veterinary monovalent or combination vaccine, *Vaccine*, Volume 33, Issue 32, 2015, Pages 3843-3849,