Summary of Studies Supporting USDA Product Licensure

Establishment Name	Intervet Inc.
USDA Vet Biologics Establishment Number	165A
Product Code	1905.20
True Name	Rabies Vaccine, Killed Virus
Tradename(s) / Distributor or Subsidiary (if different from manufacturer)	Prorab-1 - Merck Animal Health Prorab-1 - No distributor specified
Date of Compilation Summary	December 23, 2021

Disclaimer: Do not use the following studies to compare one product to another. Slight differences in study design and execution can render the comparisons meaningless.

Study Type	Efficacy
Pertaining to	Rabies virus
Study Purpose	To demonstrate effectiveness and a 2-year duration of immunity against rabies.
Product Administration	Intramuscular (IM) and Subcutaneous (SC)
Study Animals	Dogs
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. No data are published because this study was submitted to USDA-APHIS prior to January 1, 2007, and APHIS only requires publication of data submitted after that date.
USDA Approval Date	July 22, 1993

Study Type	Efficacy
Pertaining to	Rabies virus
Study Purpose	To demonstrate effectiveness and a 1-year duration of immunity against rabies.
Product Administration	Intramuscular (IM)
Study Animals	Cats
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. No data are published because this study was submitted to USDA-APHIS prior to January 1, 2007, and APHIS only requires publication of data submitted after that date.
USDA Approval Date	July 22, 1993

Study Type	Safety
Pertaining to	ALL
Study Purpose	Demonstrate safety under typical field conditions.
Product Administration	One dose administered subcutaneously (SC) or intramuscularly (IM)
Study Animals	Dogs
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. No data are published because this study was submitted to USDA-APHIS prior to January 1, 2007, and APHIS only requires publication of data submitted after that date.
USDA Approval Date	February 25, 1994

Study Type	Safety
Pertaining to	ALL
Study Purpose	Demonstrate safety under typical field conditions.
Product Administration	One dose administered subcutaneously (SC)
Study Animals	Cats
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. No data are published because this study was submitted to USDA-APHIS prior to January 1, 2007, and APHIS only requires publication of data submitted after that date.
USDA Approval Date	February 25, 1994

Study Type	Safety
Pertaining to	ALL
Study Purpose	Demonstrate safety under typical field conditions.
Product Administration	One dose administered intramuscularly (IM)
Study Animals	Cats
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. No data are published because this study was submitted to USDA-APHIS prior to January 1, 2007, and APHIS only requires publication of data submitted after that date.
USDA Approval Date	February 25, 1994