



Summary of Studies Supporting USDA Product Licensure

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| Establishment Name | Boehringer Ingelheim Animal Health USA Inc. |
| USDA Vet Biologics Establishment Number | 124 |
| Product Code | 1901.R1 |
| True Name | Rabies Vaccine, Live Canarypox Vector |
| Tradename(s) / Distributor or Subsidiary (if different from manufacturer) | |
| Date of Compilation Summary | May 17, 2019 |

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.

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| Study Type | Efficacy |
| Pertaining to | Rabies |
| Study Purpose | To evaluate efficacy against rabies twelve months post-vaccination to establish 1 year revaccination interval. |
| Product Administration | Subcutaneously (SQ) |
| 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 | March 13, 1997 |

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| Study Type | Safety |
| Pertaining to | ALL |
| Study Purpose | To evaluate safety under field conditions |
| Product Administration | Subcutaneously (SQ) |
| 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 | December 4, 1998 |