

## **Summary of Studies Supporting USDA Product Licensure**

Establishment Name	Boehringer Ingelheim Animal Health USA Inc.
USDA Vet Biologics Establishment Number	124
Product Code	1288.4A
True Name	Bursal Disease-Marek's Disease Vaccine, Serotype 3, Live Virus
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	Infectious bursal disease virus, standard
Study Purpose	Efficacy against infectious bursal disease, standard
<b>Product Administration</b>	Subcutaneous
Study Animals	Chickens at day-of-age
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.
<b>USDA Approval Date</b>	January 21, 1999

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Study Type	Efficacy
Pertaining to	Marek's disease
<b>Study Purpose</b>	Efficacy against Marek's disease
<b>Product Administration</b>	Subcutaneous
Study Animals	Chickens at day-of-age
<b>Challenge Description</b>	GA-22 MDV
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.
<b>USDA Approval Date</b>	January 21, 1999

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Study Type	Safety
Pertaining to	ALL
Study Purpose	Demonstrate safety under field conditions
<b>Product Administration</b>	Subcutaneous
Study Animals	Chickens at day-of-age
<b>Challenge Description</b>	
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.
<b>USDA Approval Date</b>	September 14, 1999

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