

Summary of Studies Supporting USDA Product Licensure

Establishment Name	Boehringer Ingelheim Animal Health USA Inc.
USDA Vet Biologics Establishment Number	124
Product Code	8201.03
True Name	Clostridium Perfringens Types C & D Toxoid
Tradename(s) / Distributor or Subsidiary (if different from manufacturer)	Caliber 3 - No distributor specified
Date of Compilation Summary	April 03, 2020

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	Clostridium perfringens Type C
Study Purpose	Demonstration of efficacy against Clostridium perfringens Type C
Product Administration	
Study Animals	Bovine and Ovine
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	November 23, 1998

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Study Type	Efficacy
Pertaining to	Clostridium perfringens Type D
Study Purpose	Demonstration of efficacy against Clostridium perfringens Type D
Product Administration	
Study Animals	Bovine and Ovine
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.
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Safety
All fractions
To demonstrate safety under field conditions
Study data are not available.

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Study Type	Safety
Pertaining to	All fractions
Study Purpose	To demonstrate safety under field conditions
Product Administration	
Study Animals	Bovine (cattle) including pregnant and lactating cattle
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 10, 1998

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