

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

Establishment Name	Boehringer Ingelheim Animal Health USA Inc.
USDA Vet Biologics Establishment Number	124
Product Code	7423.00
True Name	Clostridium Chauvoei-Septicum-Novyi-Sordellii-Perfringens Types C & D-Haemophilus Somnus Bacterin-Toxoid
Tradename(s) / Distributor or Subsidiary (if different from manufacturer)	Bar Vac 7/Somnus - No distributor specified Fermicon 7/Somnugen - Boehringer Ingelheim (Canada) Ltd.
Date of Compilation Summary	September 04, 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.

124 7423.00 Page 1 of 9

Study Type	Efficacy
Pertaining to	Clostridium perfringens Type D
<b>Study Purpose</b>	Demonstration of efficacy against Clostridium perfringens Type D
<b>Product Administration</b>	
Study Animals	
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>	October 13, 1981

124 7423.00 Page 2 of 9

Study Type	Efficacy
Pertaining to	Clostridium chauvoei
Study Purpose	Demonstration of efficacy against Clostridium chauvoei
<b>Product Administration</b>	
Study Animals	
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>	October 13, 1981

124 7423.00 Page 3 of 9

Study Type	Efficacy
Pertaining to	Clostridium novyi
Study Purpose	Demonstration of efficacy against Clostridium novyi
<b>Product Administration</b>	
Study Animals	
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>	October 13, 1981

124 7423.00 Page 4 of 9

Study Type	Efficacy
Pertaining to	Clostridium perfringens Type C
<b>Study Purpose</b>	Demonstration of efficacy against Clostridium perfringens Type C
<b>Product Administration</b>	
Study Animals	
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>	October 13, 1981

124 7423.00 Page 5 of 9

Study Type	Efficacy
Pertaining to	Clostridium septicum
Study Purpose	Demonstration of efficacy against Clostridium septicum
<b>Product Administration</b>	
Study Animals	
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>	October 13, 1981

124 7423.00 Page 6 of 9

Study Type	Efficacy
Pertaining to	Clostridium sordellii
Study Purpose	Demonstration of efficacy against Clostridium sordellii
<b>Product Administration</b>	
Study Animals	
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>	October 13, 1981

124 7423.00 Page 7 of 9

Study Type	Efficacy
Pertaining to	Haemophilus somnus
Study Purpose	Demonstration of efficacy against Haemophilus somnus
<b>Product Administration</b>	
Study Animals	Bovine
<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>	May 5, 1981

124 7423.00 Page 8 of 9

Study Type	Safety
Pertaining to	All fractions
Study Purpose	To demonstrate safety under field conditions
<b>Product Administration</b>	
Study Animals	
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. Study data, however, are no longer available.
<b>USDA Approval Date</b>	October 13, 1981

124 7423.00 Page 9 of 9