

Summary of Studies Supporting USDA Product Licensure

Establishment Name	Boehringer Ingelheim Animal Health USA Inc.
USDA Vet Biologics Establishment Number	124
Product Code	1151.26
True Name	Bovine Rhinotracheitis-Virus Diarrhea Vaccine, Modified Live Virus
Tradename(s) / Distributor or Subsidiary (if different from manufacturer)	Boehringer Ingelheim (Canada) Ltd. Pyramid 2 + Type II BVD - Boehringer Ingelheim (Canada) Ltd. Pyramid 3 - Boehringer Ingelheim Vetmedica, Inc. Pyramid 3 - No distributor specified
Date of Compilation Summary	April 08, 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	Bovine Virus Diarrhea (BVD)
Study Purpose	Demonstration of efficacy against BVD Type 1 clinical disease,
	leukopenia, and viremia
Product Administration	
Study Animals	Bovine
Challenge Description	BVD1b NY-1 strain
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	June 14, 1994

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Study Type	Efficacy
Pertaining to	Bovine Virus Diarrhea (BVD)
Study Purpose	Demonstration of efficacy against persistent infection of calves
	with BVD Type 1
Product Administration	Pregnant heifers or cows prior to breeding
Study Animals	Bovine
Challenge Description	BVD1b 97B1415 strain
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 10, 2003

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Study Type	Efficacy									
Pertaining to	Bovine Virus Dian	rhea (BVD)								
Study Purpose	Demonstration of	efficacy against	BVD Type 1 (re	espiratory						
	disease) 217 days after vaccination									
Product Administration	One dose, subcutaneously									
Study Animals	46 bovine calves (23 vaccinates and 23 controls), 25 to 37 days									
	of age	of age								
Challenge Description	Challenged with BVD Type 1b, Isolate CA0401186A, 217 days									
	(7 months) after v	(7 months) after vaccination								
Interval observed after	Calves were obser	Calves were observed and blood was collected for 14 days after								
challenge	challenge to evalu	ate viremia and	leukopenia.							
Results	Results of the stud	ly are summariz	ed as follows:							
	Blood was evaluate leukopenia (at least pre-challenge base	st one white blo	od cell count belo	/						
		Viremia	Leukopenia							
	Vaccinates	0/23 (0%)	6/23 (26%)							
	Controls 22/23 (96%) 21/23 (91%)									
	See tables on the following pages for data.									
USDA Approval Date	July 14, 2014									

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Viremia in Vaccinates (23 bovine)

Animal						Da	ys Po	ost-C	Chall	enge					Overall
ID	1	2	3	4	5	6	7	8	9	10	11	12	13	14	Result
21	-	-	-	-	-	-	-	-	-	-	-	ı	-	-	-
32	-	-	-	-	-	-	-	-	-	-	-	1	-	-	-
34	-	ı	ı	-	-	-	-	ı	-	-	-	ı	-	-	-
37	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
41	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
43	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
48	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
49	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
51	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
57	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
61	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
64	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
67	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
68	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
73	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
82	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
86	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
87	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
94	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
97	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
102	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
104	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
108	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-

^{+ =} positive for virus (highlighted yellow)

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^{- =} negative for virus

Viremia in Controls (23 bovine)

Animal						Da	ys Po	st-C	halle	nge					Overall
ID	1	2	3	4	5	6	7	8	9	10	11	12	13	14	Result
31	-	-	-	-	-	+	+	-	+	-	-	-	-	-	+
33	-	-	-	-	-	+	-	-	-	ı	1	-	-	-	+
36	-	+	+	+	-	+	-	-	-	-	-	-	-	-	+
38	-	-	-	+	+	+	+	-	+	-	-	-	-	-	+
44	-	-	-	+	+	+	-	-	-	-	-	-	-	-	+
47	-	-	-	+	+	-	-	-	-	-	-	-	-	-	+
50	-	-	-	-	-	+	+	-	-	-	-	-	-	-	+
53	-	-	-	-	+	+	+	+	+	-	-	-	-	-	+
56	-	-	-	+	+	+	+	-	-	-	-	-	-	-	+
58	-	-	-	+	+	+	+	-	-	-	-	-	-	-	+
59	-	-	-	+	+	+	+	+	+	-	-	-	-	-	+
62	-	-	-	-	+	-	+	-	-	-	-	-	-	-	+
66	-	-	-	-	+	+	-	+	-	-	-	-	-	-	+
69	-	-	-	-	+	-	-	-	-	-	-	-	-	-	+
70	-	-	-	-	+	+	+	-	-	-	-	-	-	-	+
74	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
85	-	-	-	-	+	+	-	-	-	-	-	-	-	-	+
89	-	-	-	-	+	+	+	-	-	-	-	-	-	-	+
92	-	-	-	-	-	+	+	-	-	-	-	-	-	-	+
98	-	-	-	-	+	+	+	-	-	-	-	-	-	-	+
101	-	-	-	-	-	+	+	+	-	-	-	-	-	-	+
106	-	-	-	-	+	-	-	-	-	-	-	-	-	-	+
107	-	-	-	-	+	+	+	-	-	-	-	-	-	-	+

^{+ =} positive for virus (highlighted yellow)

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^{- =} negative for virus

Leukopenia in Vaccinates (23 bovine)

				Wi	nite Blo	od Ce	ll Cour	ıt per e	ach Da	y Post	-Challe	enge				Overall
ID	Base -line	1	2	3	4	5	6	7	8	9	10	11	12	13	14	Result
21	13.5	10.4	11.3	11.6	11.9	11.3	12.4	12.2	11.6	12.4	9.6	9.6	9.3	9.7	10.4	-
32	8.4	8.1	7.8	7.9	NA	9.9	8.9	9.7	8.9	7.1	6.6	8	8.4	10.7	12	-
34	8.9	8.7	9	9.3	9.5	8.5	8.9	9	8.6	8	8.2	4.7	5.1	6.9	8.6	+
37	13.9	12.4	12.7	12.6	11.9	11.6	11	11.4	9.4	9.3	9	9.9	10.1	10.1	9.9	-
41	8.1	8.3	7.8	9.1	8.3	7.5	7.3	7.4	7.5	5.4	5.3	5.1	6.2	7.8	8.1	-
43	8.0	7.7	8	8.8	8.1	8.6	8	8.9	7.8	8.2	6	6.9	8.1	7.4	6.8	-
48	6.2	6.6	6.8	7	6.7	6.5	6.5	6.8	7.2	3.1	3.8	5	5.6	5.8	6.1	+
49	11.1	10.6	10.9	11.1	11.8	10.5	9.4	10.1	10.4	10.5	8	8.8	8.6	10.9	11.9	-
51	8.0	8.5	9.4	10.8	10.4	11.3	9.7	10.1	10.3	8.7	5.5	6.7	7.4	7.5	8.2	-
57	8.5	9.4	9.6	9.1	9.7	8.8	9.4	9.1	9	8.8	8.4	7	8	7.5	7.3	-
61	7	7	6.8	7.9	4.6	3.9	4.5	6.6	7	6.8	6.3	8.2	8.8	8	7.9	+
64	8.1	6.6	6.1	6.6	7.2	6.2	6.9	7.2	7.1	7.4	7.1	6.8	7.8	9.5	9	-
67	6.2	5.8	5.9	6.6	6.7	6.3	3.5	4.7	5	5.1	5.4	6.4	6.4	8.2	6.1	+
68	7.6	8.2	8.5	8.7	8.7	7	5.1	5.6	6.3	6.6	7.2	7.9	8.1	9.3	9.4	-
73	8.0	8	7.9	8	8.1	7.8	7.7	6.1	5.7	5.8	5.9	6.1	7.3	7.4	9.2	-
82	11.6	11.1	11.2	17.2	12.1	11.6	11.6	11	11.5	11.4	5.7	8.1	9.6	11.5	10.8	+
86	6.6	7.2	7.1	6.8	6.8	5.9	6.3	6.1	5.8	6.7	4.2	4.6	5.9	5.9	7.2	-
87	9.2	9.4	9.9	10	7.4	8.3	7.5	10.4	8.8	8.6	9.7	9.1	8.9	9.1	9.6	-
94	11.6	11.2	11.4	10.6	10.1	9.3	9.3	9.4	9.9	10.2	8.3	8.6	9.4	11.4	11.8	-
97	6.4	6.8	7	8.1	8.3	7.7	7.4	7.9	8.3	7.5	4.8	5.3	5.7	6.6	5.8	-
102	8.7	8.3	8.8	8.7	8.9	8.2	8.2	8.5	8	6	3.8	5.5	6.3	7.1	8.1	+
104	8.3	8.4	8.6	9.2	8.9	7.9	8.1	8.9	8.5	6.9	5.3	6.7	6.7	9.9	7.9	-
108	8.1	8.1	9	7.7	8.4	11.4	7.8	9.3	11.9	10.3	10.3	8	12.7	10.6	10.2	-

White Blood Cell Count:

- Baseline is the average white blood cell (WBC) counts from 3 consecutive days, prior to challenge
- Numerical values = white blood cell count in K/μL
- Highlighted yellow = positive for leukopenia, meaning a 40% or more reduction in the total WBC count compared to baseline on that day
- NA = not applicable / no measurement on that day

Overall Result:

- + = positive for leukopenia at least one day (highlighted yellow)
- = negative for leukopenia on every day

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Leukopenia in Controls (23 bovine)

				WI	nite Blo	od Ce	ll Cour	t per e	ach Da	y Post	-Challe	enge				Overall
ID	Base -line	1	2	3	4	5	6	7	8	9	10	11	12	13	14	Result
31	7.8	7.7	7.3	3.8	NA	4.5	3.7	3.9	6.1	10.6	7	5.9	5.5	5.9	3.2	+
33	11.3	10.5	11.9	6.5	7	7.3	5.7	6	10.3	7.9	6.8	7	7.4	8.1	7.7	+
36	8.9	9.2	8.8	5.1	6.4	5.7	5.3	5.2	6.6	7.5	6.3	5.7	5.9	5.9	6	+
38	6.5	5.9	7	4.6	4.6	3.9	3.6	3.4	7	7.3	6.1	5.4	5.3	5.6	5	+
44	10.3	10.2	10	4.1	6.2	5.8	4.3	7.3	12.7	8	5.4	5.7	6.3	5.5	5.5	+
47	7.9	8.1	7.6	4.8	5.9	5.8	4.5	5.3	7.9	6.5	6.2	5.8	5.6	5.6	6	+
50	8.2	8.9	8.9	5.9	4.6	6	5.1	6.3	10.4	8.9	6.3	5.1	5.1	5.7	5.6	+
53	7.5	8.3	8.5	3.9	4.5	4.3	3.1	3.4	3.6	7.5	6.1	4.4	5.6	5.8	5.5	+
56	9.8	7.6	9.1	3.9	6	4.5	4.4	5.3	12.2	9.8	5.4	5.2	6	5.5	6.3	+
58	8.3	8.7	9.8	5.7	6.3	7.1	5.1	5.2	8.2	8.4	6.1	6.4	7.4	6.2	6.5	-
59	10.1	11.1	12.2	8.5	7.3	8.7	7.6	7.8	9.4	12.7	13	9.9	9.5	9.9	21.6	-
62	9.5	10.2	9.8	4.9	5.5	5.1	4.2	3.9	7.5	10.7	8.6	7	6.1	6.2	5.5	+
66	6.7	8.5	8	4.5	6	5.2	4.5	5.8	7	5.8	4.8	4.6	5.1	5.4	3.9	+
69	11.1	12.1	11.4	5.1	8.1	8.8	7.2	9.4	11.3	10.5	7.8	7.4	7.1	7.4	7.7	+
70	10.7	10.1	9	4.4	6	5.8	4.4	4.5	7.5	10.8	8.5	7.4	5.6	6.4	5.8	+
74	5.8	6.2	6.4	4.2	4.7	4.2	3.3	3	4	7	5.2	4.4	4.6	4.1	4.5	+
85	8.6	8	7.8	5.7	6.8	5.4	4.5	4.3	5	6.1	6.3	7.4	7.2	7.6	7.4	+
89	8.9	8.4	8.5	3.9	5.8	6.1	4.6	4.1	9.1	8.6	5.7	4.7	5.2	5.1	5.7	+
92	7.8	7.4	5.9	4.4	4.6	3.8	3.2	3.6	4.1	4.6	3.6	3.8	3.8	4.5	4.4	+
98	7.9	7.4	8	4.5	5.9	4.2	4.1	4.1	6.5	7.3	6.5	5.8	6.1	5.9	5.8	+
101	9.9	9.3	10.5	8.5	7.6	7.2	5.6	5.5	5.3	13.4	9.6	7.2	6.5	7	5.4	+
106	7.1	7.2	7.3	4.4	5.3	4.6	3.7	5.2	7.6	5.3	5.1	6.4	5.6	6.3	6.9	+
107	10.7	10.5	10.1	6.7	7.9	7	5.5	5.7	7.8	8.9	8.7	7.5	7.1	7.6	8.8	+

White Blood Cell Count:

- Baseline is the average white blood cell (WBC) counts from 3 consecutive days, prior to challenge
- Numerical values = white blood cell count in K/μL
- Highlighted yellow = positive for leukopenia, meaning a 40% or more reduction in the total WBC count compared to baseline on that day
- NA = not applicable / no measurement on that day

Overall Result:

- + = positive for leukopenia at least one day (highlighted yellow)
- = negative for leukopenia on every day

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Study Type	Efficacy
Pertaining to	Bovine Virus Diarrhea (BVD)
Study Purpose	Demonstration of efficacy against persistent infection of calves
	with BVD Type 2
Product Administration	Pregnant heifers or cows prior to breeding
Study Animals	Bovine
Challenge Description	BVD2a 96B2222 strain
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 10, 2003

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Study Type	Efficacy
Pertaining to	Bovine Virus Diarrhea (BVD)
Study Purpose	Demonstration of efficacy against BVD Type 2clinical disease,
	leukopenia, and viremia
Product Administration	
Study Animals	Bovine
Challenge Description	BVD2a IAF 103, BT-4A-2 strain
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	August 4, 2003

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Study Type	Efficacy
Pertaining to	Infectious Bovine Rhinotracheitis (IBR)
Study Purpose	Demonstration of efficacy against IBR (respiratory disease)
Product Administration	
Study Animals	Bovine
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	September 9, 1994

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Study Type	Safety
Pertaining to	All fractions
Study Purpose	To demonstrate safety under field conditions
Product Administration	
Study Animals	Bovine
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 10, 2002

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Study Type	Safety
Pertaining to	All fractions
Study Purpose	To demonstrate safety under field conditions
Product Administration	
Study Animals	Bovine: pregnant cows and calves nursing pregnant cows provided
-	the cows were vaccinated pre-breeding
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 6, 2005

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