

## Summary of Studies Supporting USDA Product Licensure

Establishment Name	Diamond Animal Health, Inc.			
USDA Vet Biologics Establishment Number	213			
Product Code	1101.20			
True Name	Bovine Rhinotracheitis Vaccine, Modified Live Virus			
Tradename(s) / Distributor or Subsidiary (if different from manufacturer)	Titanium IBR - Elanco Animal Health Titanium IBR - Elanco US, Inc Diamond Animal Health, Inc. Titanium IBR - Elanco, Division Eli Lilly Canada, Inc Diamond Animal Health, Inc. Titanium IBR - No distributor specified			
Date of Compilation Summary	August 16, 2021			

## 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.

Study Type	Efficacy				
	5				
Pertaining to	Infectious Bovine Rhinotracheitis (IBR) Virus				
Study Purpose	To demonstrate effectiveness against disease caused by infectious				
	bovine rhinotracheitis virus.				
Product Administration	1 dose to calves 6-8 months of age.				
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	July 13, 1998				

Study Type	Safety			
Pertaining to	ALL			
Study Purpose	Demonstrate safety of product under typical use conditions			
<b>Product Administration</b>	One dose			
Study Animals	Bovine			
Challenge Description	NA			
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	July 13, 1998			

Study Type	Safety						
Pertaining to	ALL						
Study Purpose	To demonstrate safety in pregnant cows and calves nursing						
· I	pregnant cows						
Product	Two doses administered subcutaneously to heifers and cows, 1						
Administration	dose prior to breeding and 1 dose during pregnancy at different						
	trimesters. Heifers and cows were confirmed to be pregnant at						
	administration during pregnancy.						
Study Animals	Heifers and cows-separate groups vaccinated during each trimester.						
v	Similar sized groups in each trimester were maintained as controls.						
Challenge Description	NA						
Interval observed after	Heifers and cows observed from pre-breeding vaccination through						
challenge	birth of calves. Nursing calves observed through 4 weeks of age.						
Results	Summary of Calving Rates						
			d/Total deliveries	)			
			,	, ,			
	Trimester	Vacc	inates	Controls			
	1	200/2	208 (96%)	205/213 (96%)			
	2		313 (96%)	293/308 (95%)			
	3		205 (94%)	195/208 (94%)			
	Total		726 (96%)	693/729 (95%)			
	First Trimester (<	93 day	vs of gestation)				
	<u>`</u>		Vaccinate	Control			
	Enrolled		209	213			
	Excluded (not re	Excluded (not related		0			
	to vaccination)						
	Aborted or stillborn		6	5			
			2	3			
	Second Trimester (94-187 days of gestation)						
		Vaccinate					
	Enrolled Excluded (not related		315	310			
			2	2			
	to vaccination)						
	Aborted or stillborn		7	9			
	Died at or after birth		4	6*			
	*one death was fr	om a s	et of twins; the ot	her was normal			
	Third Trimester (188-250 days of gestation)						
			Vaccinate	Control			
	Enrolled		205	208			
	Aborted or stillborn Died at or after birth		9	9*			
			3	4**			
	*one stillborn was from a set of twins; the other was normal						
	**one dead was from a set of twins; the other was normal						

USDA Approval Date	March 5, 2013