



Summary of Studies Supporting USDA Product Licensure

Establishment Name	Biomune Company
USDA Vet Biologics Establishment Number	368
Product Code	1231.1J
True Name	Bronchitis Vaccine, Georgia Type, Live Virus
Tradename(s) / Distributor or Subsidiary (if different from manufacturer)	CEVAC IBRON - No distributor specified CEVAC IBRON GA L - No distributor specified
Date of Compilation Summary	November 09, 2017

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						
Pertaining to	Infectious Bronchitis Virus (IBV)						
Study Purpose	Pivotal efficacy against IBV, Georgia 13 Type						
Product Administration	One dose administered by the coarse spray route						
Study Animals	30 chickens per treatment group vaccinated at day of age						
Challenge Description	Heterologous IBV Georgia 13 Type administered at 28 days post vaccination						
Interval observed after challenge	Daily observation for 5 days post challenge; IBV evaluated in the target tissue day 5 post challenge						
Results	<p>A chicken was considered affected by the challenge (positive) if IBV was recovered from the target tissue.</p> <p>The study fulfilled 9 CFR 113.327(c)</p> <table border="1" data-bbox="580 936 1329 1088"> <thead> <tr> <th>Treatment</th> <th>Number protected/Total</th> </tr> </thead> <tbody> <tr> <td>Vaccinated, challenged</td> <td>26/30</td> </tr> <tr> <td>Placebo-vaccinated, challenged control</td> <td>0/30</td> </tr> </tbody> </table> <p>Raw data are shown on attached page.</p>	Treatment	Number protected/Total	Vaccinated, challenged	26/30	Placebo-vaccinated, challenged control	0/30
Treatment	Number protected/Total						
Vaccinated, challenged	26/30						
Placebo-vaccinated, challenged control	0/30						
USDA Approval Date	July 22, 2015						

Vaccinate ID	Virus Recovery	Control ID	Virus Recovery
1	Neg	1	Pos
2	Pos	2	Pos
3	Neg	3	Pos
4	Neg	4	Pos
5	Neg	5	Pos
6	Neg	6	Pos
7	Neg	7	Pos
8	Neg	8	Pos
9	Neg	9	Pos
10	Neg	10	Pos
11	Neg	11	Pos
12	Neg	12	Pos
13	Neg	13	Pos
14	Neg	14	Pos
15	Pos	15	Pos
16	Neg	16	Pos
17	Neg	17	Pos
18	Neg	18	Pos
19	Neg	19	Pos
20	Neg	20	Pos
21	Neg	21	Pos
22	Pos	22	Pos
23	Neg	23	Pos
24	Neg	24	Pos
25	Neg	25	Pos
26	Neg	26	Pos
27	Neg	27	Pos
28	Neg	28	Pos
29	Neg	29	Pos
30	Pos	30	Pos

Study Type	Efficacy						
Pertaining to	Infectious Bronchitis Virus (IBV)						
Study Purpose	Pivotal efficacy against IBV, Georgia 08 Type infection						
Product Administration	One dose administered by the coarse spray route						
Study Animals	30 chickens per treatment group vaccinated at day of age						
Challenge Description	Homologous IBV Georgia 08 administered at 26 days post vaccination						
Interval observed after challenge	Daily observation for 5 days post challenge; IBV evaluated in the target tissue day 5 post challenge						
Results	<p>A chicken was considered affected by the challenge (positive) if IBV was recovered from the target tissue.</p> <p>The study fulfilled 9 CFR 113.327(c).</p> <table border="1" data-bbox="579 936 1329 1088"> <thead> <tr> <th>Treatment</th> <th>Number protected/Total</th> </tr> </thead> <tbody> <tr> <td>Vaccinated, challenged</td> <td>27/30</td> </tr> <tr> <td>Placebo-vaccinated, challenged control</td> <td>1/30</td> </tr> </tbody> </table> <p>Raw data are shown on attached page.</p>	Treatment	Number protected/Total	Vaccinated, challenged	27/30	Placebo-vaccinated, challenged control	1/30
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Vaccinated, challenged	27/30						
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USDA Approval Date	July 22, 2015						

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1	Neg	1	Pos
2	Neg	2	Pos
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5	Neg	5	Pos
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7	Pos	7	Pos
8	Neg	8	Pos
9	Neg	9	Pos
10	Neg	10	Pos
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22	Neg	22	Pos
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25	Neg	25	Pos
26	Neg	26	Pos
27	Neg	27	Pos
28	Neg	28	Pos
29	Neg	29	Pos
30	Neg	30	Pos

Study Type	Efficacy						
Pertaining to	Infectious Bronchitis Virus (IBV)						
Study Purpose	Pivotal efficacy against IBV, DMV/1639/11 type infection						
Product Administration	One dose administered by the gel spray route (gel droplet by oral administration)						
Study Animals	30 chickens per treatment group vaccinated at day of age						
Challenge Description	Heterologous IBV DMV/1639/11 administered at 28 days post vaccination						
Interval observed after challenge	Daily observation for 5 days post challenge; IBV evaluated in the target tissue day 5 post challenge.						
Results	<p>A chicken was considered affected by the challenge (positive) if IBV was recovered from the target tissue.</p> <p>The study fulfilled 9 CFR 113.327(c).</p> <table border="1" data-bbox="579 972 1329 1126"> <thead> <tr> <th>Treatment</th> <th>Number protected/Total</th> </tr> </thead> <tbody> <tr> <td>Vaccinated, challenged</td> <td>30/30</td> </tr> <tr> <td>Placebo-vaccinated, challenged control</td> <td>0/30</td> </tr> </tbody> </table> <p>Raw data are shown on attached page.</p>	Treatment	Number protected/Total	Vaccinated, challenged	30/30	Placebo-vaccinated, challenged control	0/30
Treatment	Number protected/Total						
Vaccinated, challenged	30/30						
Placebo-vaccinated, challenged control	0/30						
USDA Approval Date	December 21, 2015						

Vaccinate ID	Virus Recovery	Control ID	Virus Recovery
1	Neg	1	Pos
2	Neg	2	Pos
3	Neg	3	Pos
4	Neg	4	Pos
5	Neg	5	Pos
6	Neg	6	Pos
7	Neg	7	Pos
8	Neg	8	Pos
9	Neg	9	Pos
10	Neg	10	Pos
11	Neg	11	Pos
12	Neg	12	Pos
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22	Neg	22	Pos
23	Neg	23	Pos
24	Neg	24	Pos
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26	Neg	26	Pos
27	Neg	27	Pos
28	Neg	28	Neg
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Study Type	Safety																																			
Pertaining to	ALL																																			
Study Purpose	Demonstrate safety of product under typical use conditions.																																			
Product Administration	One dose administered via the coarse-spray application.																																			
Study Animals	Broiler chickens at day-of-age. 107,800 were vaccinated with product vaccine and 107,800 were kept as controls. Animal were observed daily for mortality through 14 days after vaccination.																																			
Challenge Description	Not applicable																																			
Interval observed after challenge	Not applicable																																			
Results	<table border="1"> <thead> <tr> <th>Location</th> <th>Treatment</th> <th>Total Placed</th> <th>14 Day Mortality</th> <th>% Mortality</th> </tr> </thead> <tbody> <tr> <td>1</td> <td>Product Vaccine</td> <td>24,100</td> <td>611</td> <td>2.54%</td> </tr> <tr> <td>1</td> <td>Control</td> <td>24,100</td> <td>560</td> <td>2.32%</td> </tr> <tr> <td>2</td> <td>Product Vaccine</td> <td>38,600</td> <td>946</td> <td>2.45%</td> </tr> <tr> <td>2</td> <td>Control</td> <td>38,600</td> <td>515</td> <td>1.33%</td> </tr> <tr> <td>3</td> <td>Product Vaccine</td> <td>45,100</td> <td>483</td> <td>1.07%</td> </tr> <tr> <td>3</td> <td>Control</td> <td>45,100</td> <td>542</td> <td>1.20%</td> </tr> </tbody> </table> <p>No adverse reactions attributable to the vaccine were recorded.</p>	Location	Treatment	Total Placed	14 Day Mortality	% Mortality	1	Product Vaccine	24,100	611	2.54%	1	Control	24,100	560	2.32%	2	Product Vaccine	38,600	946	2.45%	2	Control	38,600	515	1.33%	3	Product Vaccine	45,100	483	1.07%	3	Control	45,100	542	1.20%
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Study Type	Safety																																			
Pertaining to	ALL																																			
Study Purpose	Demonstrate safety of product under typical use conditions.																																			
Product Administration	One dose administered via the gel droplet application (oral route).																																			
Study Animals	Broiler chickens at day-of-age. 44,000 were vaccinated with product vaccine and 44,000 were kept as controls. Animals were observed daily for mortality through 21 days after vaccination.																																			
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