

Summary of Studies Supporting USDA Product Licensure

Establishment Name	Intervet Inc.
USDA Vet Biologics Establishment Number	165A
Product Code	4875.A1
True Name	Encephalomyelitis-Influenza Vaccine, Eastern & Western & Venezuelan, Killed Virus, Tetanus Toxoid
Tradename(s) / Distributor or Subsidiary (if different from manufacturer)	Encevac TC-4 + VEE - Merck Animal Health
Date of Compilation Summary	February 08, 2018

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	Clostridum tetani
Study Purpose	Demonstrate efficacy against <i>C. tetani</i>
Product Administration	One dose administered subcutaneously.
Study Animals	Ten guinea pigs (5 females and 5 males, 450-550g)
Challenge Description	Not applicable
Interval observed after	Six weeks after vaccination, guinea pigs were bled for serological
challenge	testing.
Results	Efficacy of <i>C. tetani</i> was demonstrated in laboratory animals according to 9CFR 113.114(c). Satisfactory result is an antitoxin titer of at least 2.0 A.U. per mL for the serum pool. Pooled Guinea Pig Antitoxin titer (A.U./mL) 2.082
USDA Approval Date	June 15, 2010

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Study Type	Efficacy			
Pertaining to	Eastern Equine E	Encephalomyelitis (EEE)	
Study Purpose	Demonstrate effi	cacy against EEE		
Product Administration	Two doses admir	nistered intramuscu	larly 3 weeks ap	oart.
Study Animals	Twelve guinea pi	igs, 10 vaccinates a	nd 2 controls, ea	ach 300-500g
Challenge Description	Not applicable			
Interval observed after	14 days post 2nd	vaccination, guine	a pigs were bled	for
challenge	serological testin	g.		
Results	according to 9CF Satisfactory test	was demonstrated in FR 113.207(b). The result is a Virus New 20 vaccinates (2 nd states).	utralization Tite	r of ≥ 1:40 in
	Treatment		Test	
	group	Results	Disposition	
	Vaccinates	$17/20 \ge 1:40$	Satisfactory	
	Controls	2/2 <1:4	Ballstactory	
USDA Approval Date	June 15, 2010		·	

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Study Type	Efficacy			
Pertaining to	Venezuelan Equi	ne Encephalomyel	itis (VEE)	
Study Purpose	Demonstrate effic	cacy against VEE		
Product Administration	Two doses admir	nistered intramuscu	ılarly 3 weeks ap	oart.
Study Animals	Twelve guinea pi	igs, 10 vaccinates a	and 2 controls, ea	ach 300-500g
Challenge Description	Not applicable			
Interval observed after	14 days post 2nd	vaccination, guine	a pigs were bled	for serological
challenge	testing per SAM	110.		
Results	according to 9CF	result is a Virus Ne	,	
	Treatment		Test	
	group	Results	Disposition	
	Vaccinates	$10/10 \ge 1:4$	Satisfactory	
	Controls	2/2 <1:4]
USDA Approval Date	June 15, 2010			

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Study Type	Efficacy			
Pertaining to	Western Equine	Encephalomyelitis	(WEE)	
Study Purpose	Demonstrate effi-	cacy against WEE		
Product Administration	Two doses admir	nistered intramuscu	ılarly 3 weeks ap	art.
Study Animals	Twelve guinea pi	igs, 10 vaccinates a	and 2 controls, ea	ach 300-500g
Challenge Description	Not applicable			
Interval observed after	14 days post 2nd	vaccination, guine	a pigs were bled	
challenge	for serological te	sting.		
Results	according to 9CF	result is a Virus Ne	,	
	Treatment		Test	
	group	Results	Disposition	
	Vaccinates	$9/10 \ge 1:40$	Satisfactory	
	Controls	2/2 <1:4	Butisfactory	
USDA Approval Date	June 15, 2010			

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Study Type	Efficacy
Pertaining to	Equine Influenza Virus (EIV)
Study Purpose	To demonstrate efficacy of updated EIV strains FL/13 and RI/07
Product Administration	
C4	
Study Animals	
Challenge Description	
Interval observed after	
challenge	
Results	This product class allows the manufacturer to update micro- organisms in this vaccine under expedited procedures to respond to emerging needs. Abbreviated data to support influenza strain updates to the product composition were evaluated by USDA- APHIS and found to be acceptable based on regulations and policies at the time of approval. Full vaccination-challenge studies may not have been required for these updates.
USDA Approval Date	March 8, 2016

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Study Type	Efficacy			
Pertaining to	Equine Influer	nza Virus (E	IV)	
Study Purpose	To demonstrat	te efficacy a	gainst EIV six	months after
	vaccination.			
Product Administration	Two doses ada	ministered ir	ntramuscularly	(IM) three weeks apart.
Study Animals	18 vaccinate a	nd 7 control	horses were u	sed at 6 months of
	age.			
Challenge Description			d with EIV stra	
				econd vaccination.
Interval observed after				oost-challenge for
challenge		Nasal swab	s were collecte	ed daily for virus
	isolation.			
Results		-		f any clinical sign at
	•	_		od (nasal discharge,
	coughing, resp	oiration, tem	perature >102.	5°F).
			Ţ	
		# of	Presence of	
	Group	Animals	clinical sign	S
	Vaccinates	18	14	
	Controls	7	7	
		- 1		considered negative for
	_	• •	_	vabs were virus
	negative, other	rwise it was	positive.	
		,, <u> </u>		
		# of	Virus	
	Group	Animals	Isolation	
	Vaccinates	18	12	
	Controls	7	7	
	Raw data show		ed pages.	
USDA Approval Date	August 8, 200	5		

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Rectal body temperatures of horses vaccinated with vaccine 111103 on days post-challenge with virulent EIV KY99. Table 1.

					-	Rody tem	temperatures	(ab)	on days	post chi	challenge	with EIV	V KY99:				
HOTSE	Group	-	0	-	2	1	7	2	9		8	6	10	11	12	13	14
		100.3	100.0	9.66	99.9	101.5	101.1	100.0	8.66	100.2	99.1	100.2	98.1	100.7	6.66	1001	100.6
1		2 66	100.0	99.7	Iσ	99.5	8.66	6.86	89.3	100.0	99.1	8.66	99.2	100.8	9.66	100.3	100.1
4		8.86	99.1	6.66	100.1	98.6	8.66	98.9	99.3	98.9	97.8	99.5	99.2	100.5	6.66	101.7	100.0
7		8.66	100.9	100.6	101.9	99.8	8.66	6.66	100.0	99.2	99.1	100.3	99.2	100.8	63.6	100.2	99.7
16		99.1	100.1	99.3	IN	100.5	101.5	1001	98.9	98.8	98.5	98.3	98.8	1001	99.4	100.0	100.6
0		99.7	100.6	99.9	10	93.6	100.2	99.4	100.0	99.7	99.5	100.0	98.6	100.0	100.6	100.9	100.2
20		100.3	100.5	9.66	99.7	99.7	100.0	99.3	99.2	99.2	100.5	8.66	100.0	100.5	100.9	100.2	100.3
30		101.2	100.1	100.5	105.4	102.6	104.4	102.1	100.9	9.66	100.4	101.5	100.7	102.4	101.3	102.1	101.4
3		98.8	100.8	100.0	100.4	99.3	100.5	9.66	7.66	99.7	98.3	93.6	100.0	101.2	100.0	100.4	102.7
33	Vaccinates	99.2	100.0	99.4	104.6	100.0	100.0	98.7	100.0	100.8	98.6	99.0	99.3	100.6	9.66	100.9	100.6
34		8.66	100.2	99.7	103.6	100.0	101.3	102.1	102.7	102.1	101.5	102.9	102.0	100.4	1001	8.66	99.4
35		99.2	99.7	99.1	9	99.5	100.6	100.4	7.66	8.86	98.5	1001	1001	101.6	100.2	100.3	100.5
36		99.7	9.66	8.66	100.5	100.0	100.5	9.66	99.4	100.4	98.6	99.5	0.66	100.6	100.7	66.66	100.4
000		99.7	100.4	100.0	100.5	100.3	100.5	8.86	100.0	99.4	98.7	100.8	9.66	100.5	1001	100.9	100.0
90		99.4	100.4	99.9	102.2	6.66	101.6	1001	1001	99.1	99.1	99.7	99.5	100.3	99.2	66.66	100.4
200		99.2	100.0	99.4	102:6	99.7	7.66	99.4	9.66	69.7	99.3	99.66	98.8	100.2	99.0	1001	100.2
43		99.7	100.4	8.66	99.4	8.66	1001	98.6	8.66	98.7	99.0	99.4	9.66	9.66	99.9	93.6	99.9
47		100.2	100.2	100.1	101.6	99.2	1001	99.9	100.5	99.66	99.3	8.86	99.5	99.3	99.7	99.5	100.6
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α		0.66	66	98.8	101.9	100.8	103.0	99.1	100.7	100.2	6.86	66.66	99.3	100.4	1001	100.3	99.6
-		100.4	100.5	1001	101.9	6.66	8.66	100.9	1001	99.2	100.6	101.3	99.5	101.0	6.66		100.3
2		99.5	7.66	100.3	103.6	102.4	102.8	99.5	99.3	100.4	99.5	99.2	98.8	101.0	1001	100.7	99.8
26	Controls	100.1	100.6	99.5	4	103.6	102.2	103.1	104.2	103.2	102.3	105.4	99.5	8.66	8.66	7.66	100.1
8		99.2	100.0	99.0	104.7	103.7	104.0	104.6	104.7	103.0	103.6	103.3	101.6	101.2	99.5	99.2	99.2
44		99.1	6.66	8.66	103.2	102.6	104.4	97.2	8.66	100.4	104.6	103.8	102.6	102.9	100.6	100.0	99.4

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Coughing observations of horses vaccinated with vaccinated 111103 on days postchallenge with virulent EIV KY99. Table 2.

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KY99:	12	N	N	N	Z	z	z	N	z	z	N	Z	N	z	z	N	N	Z	z	1000	z	N	z	Z	z	z	z	
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N=no coughing , C=coughing 1 time during the observation period, C, C=coughing 2 or during the observation period

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Nasal discharge observations of horses vaccinated with vaccine 111103 on days postchallenge with virulent EIV KY99. Table 3.

Horse			Nasal		discharge	1	Serva	observations	å	days I	ost-c	post-challenge		with E	EIV KY	KY99:	
No.	Group	-1	0	l-	7	1	4	2	9		8	6	10	11	12	13	14
-		z	Z	z	z	z	z	Œ	z	z	Σ	z	Z	N	SM	N	z
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4		z	z	z	z	z	z	z	z	z	z	N	N	Z	z	z	z
14		z	z	z	z	z	z	z	z	Z	Z	SM	CM	SM	z	N	z
16		z	z	z	z	z	z	z	z	z	N	N	z	z	z	N	z
19		z	z	z	z	z	z	z	z	z	N	N	Z	z	z	z	z
29		z	z	z	z	z	z	z	SM	N	SM	z	Z	z	N	N	z
30		z	z	z	z	z	SM	z	SM	SM	SM	z	z	z	N	SM	N
32		z	z	z	z	z	z	z	z	z	N	N	Z	N	z	z	N
33	Vaccinates	z	z	z	z	z	z	z	Z	N	Z	z	Z	N	N	z	z
34		z	z	z	z	z	z	SM	SM	N	N	SM	z	SM	N	N	N
32		z	z	z	z	z	z	z	z	z	N	N	N	z	z	SM	N
36		z	z	z	z	N	z	z	z	z	z	z	N	N	N .	z	Z
39		z	z	z	z	z	z	z	z	z	Z	z	Z	Z	N	Z	SM
40		z	z	z	z	z	z	z	z	z	SM	N	N	z	z	z	N
41		z	z	z	z	z	z	z	z	z	z	N	N	N	z	z	N
		z	z	z	z	z	SM	z	z	N	SM	z	z	N	N	N	SM
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13		z	z	z	z	z	SM	z	SM	SM	SM	SM	Z	CM	N	N	N
25	Controls	z	z	z	z	z	z	CM	SM	CM	CM	CM	CM	N	N	SM	z
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	_							L									
		3			300	7	400	0000	0.00	J. 40	SM.	tani saht	1 m11/	in in in	- Lug		

(score of 0), S-copious serous discharge (score of 1), SM-slight mucopurulent discharge (score of 2), CM=copious mucopurulent discharge (score of 4) N=normal

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Abnormal respiration and depression observations of horses vaccinated with vaccine 111103 on days post-challenge with virulent EIV KY99. Table 4.

N N N N N N N N N N N N N N N N N N N	Group	7	0	-	0	Observations	tions	on days	ys post	t-chal	-challenge	with 1	EIV KY	KY99:	12	13	14
N		Z	z	z	z	z	z	z	z	z	N	z	z	z	N	z	z
N		z	z	z	z	z	z	N	z	z	N	Z	Z	z	N	N	Z
The control of the co		Z	z	z	z	z	z	N	N	z	N	N	Z	z	z	z	z
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lethargy or inappetence. (score of 1) Observations were scored as not observed (score of 0) and observed N=normal, Ab=abnormal respiration of >36 per minute, D=depression,

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Virus isolation from nasal swabs from horses vaccinated with vaccine 111103 on days post-challenge with virulent EIV KY99. Table 5.

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Study Type	Safety			
Pertaining to	ALL			
Study Purpose	To demonstrate safety under field conditions			
Product Administration	298 horses received 2 doses intramuscularly 3 to 4 weeks apart for			
	primary immunization. 254 horses received 1 dose			
	intramuscularly.			
Study Animals	552 horses of various ages, breeds and sex in 5 different states.			
	177 horses were 4-months of age or younger at the time of the			
	initial vaccination.			
Challenge Description	Not applicable			
Interval observed after	Horses were observed immediately following vaccination and then			
challenge	daily for 3 days post-vaccination			
Results	Doses are reported due to difference in vaccination schedule.			
	Score	# of Cases	% of Total]
	0	820	96.47	
	1	25	2.94	
	2	3	0.35	
	3	2	0.24	
	4	0	0	
	5	0	0	
	Total # of Doses administered = 850			
	Score Overview:			
	0 – No reaction			
	1 – Localized swelling at or near the injection site, which is not			
	visible; detectable only by palpation. Not clinically significant.			
	2 – Localized visible swelling at or near the injection site. Not			
	painful.			
	3 – Localized visible swelling at or near the injection site. Raised,			
	circumscribed and painful.			
	4 – Visible diffused swelling involving a substantial area around the injection site. Very painful and hot. Horse is stiff and/or			
	reluctant to move.			
	5 – Generalized or systemic reaction, including anaphylaxis or			
	elevated temperature.			
USDA Approval Date	February 8, 2006			

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