

FOOT AND MOUTH DISEASE - VERY COMPLEX AND COSTLY DISEASE



)isease

- Caused by FMDV
- Picornavirus
- Genus Aphthovirus
- Vesicular disease of cloven hoofed animals
- Seven global FMDV pools
- Multiple serotypes within a pool
- Multiple strains within serotype
- Each pool requires a tailored, multivalent vaccine



Highly politicized diseaseVaccination programs

- often under Government control
- Largest impact on trade of any animal disease
- Animal welfare issues
- Impact on farmers and producers
- Significant economic burden
- Severe impact of incursion for "FMD-Free" Countries



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Regional production

- Produced from virulent, wild-type strains
- Risk of release from manufacturing sites
- Do not protect across serotypes, limited cross-protection within a serotype
- NSP-depletion to accommodate "DIVA" compatibility
- Not 100% due to residual NSP's present in vaccines
- Protect from clinical signs
- Do not fully prevent fever, viremia, shedding, or persistence



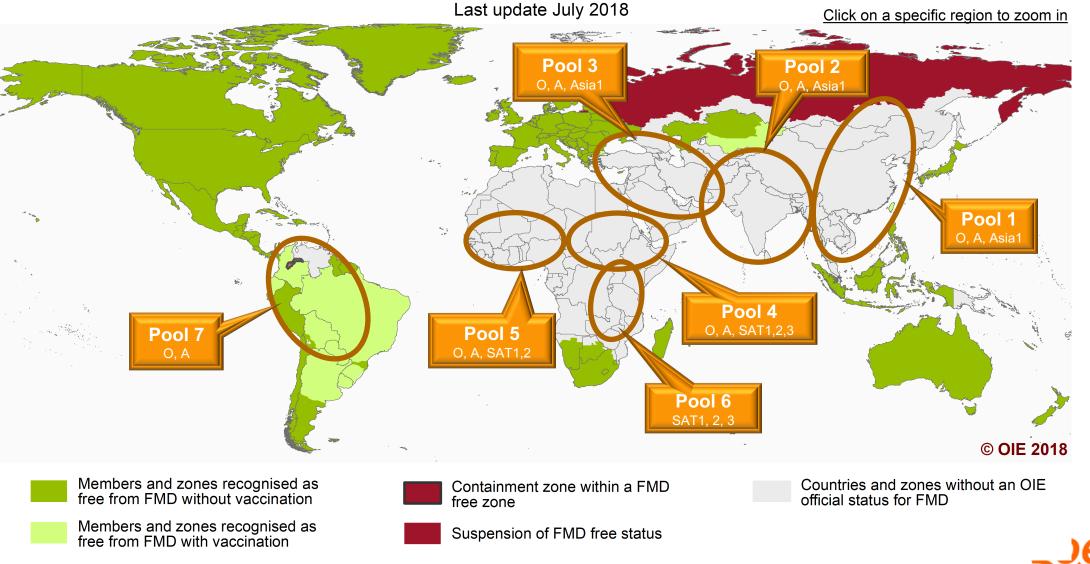
Safe vaccine platform

- Prevention of fever, viremia, shedding, and persistence
- Rapid onset of immunity
- Long duration of immunity
- Full DIVA compatibility
- Ability to rapidly address new strains



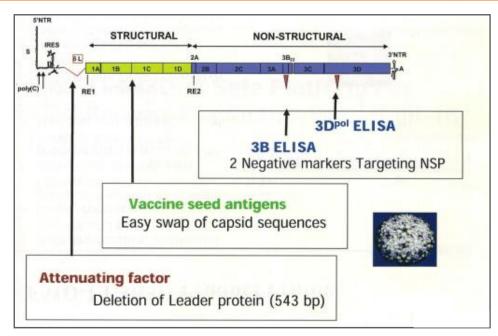
WORLD-WIDE DISTRIBUTION OF FMD

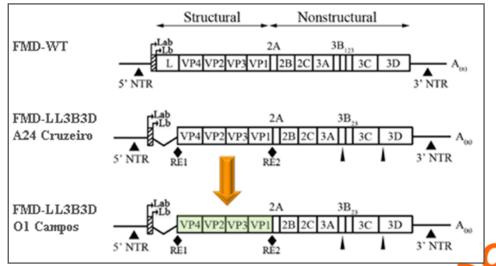
OIE Members' official FMD status map



DISRUPTIVE FMD VACCINE - LL3B3D PLATFORM

- Collaborative effort between the USDA-ARS and Zoetis
- Cassette construction
 - Allows rapid insertion of capsid coding regions from emerging strains
- Double BEI inactivated and formulated with a proprietary oil-based adjuvant
- Safe and easy production technology
 - FMD-LL3B3D A24 Cruzeiro platform backbone lacks Lpro region
 - Attenuated in cattle and pig
 - Uses the same production systems as current inactivated FMDV vaccines
- Non-transmissible (cattle & swine)
- Fully DIVA compatible
 - Two independent and stable negative markers
 - Genetically altered key epitopes in 3B and 3D NSPs
- High potency
 - Potent immune responses to inserted capsid proteins of target strain
 - Proprietary adjuvant increases antibody and cellular immune responses





SAFETY OF THE LIVE FMD LL3B3D VACCINE STRAINS - CATTLE AND SWINE

FMD-LL3B3D platform viruses are incapable of replicating to detectable levels in cattle or swine





Most susceptible species

Construct	Inoculation Route	# Animals
FMD-LL3B3D-A24 Cruzeiro	Intralingual (7x10 ⁶)	2
FMD-LL3B3D-A24 Cruzeiro	Aerosol (1x10 ⁶ to 3x10 ⁶)	3
FMD-LL3B3D-A24 Cruzeiro	Aerosol and Contact / (1x10 ⁶)	9

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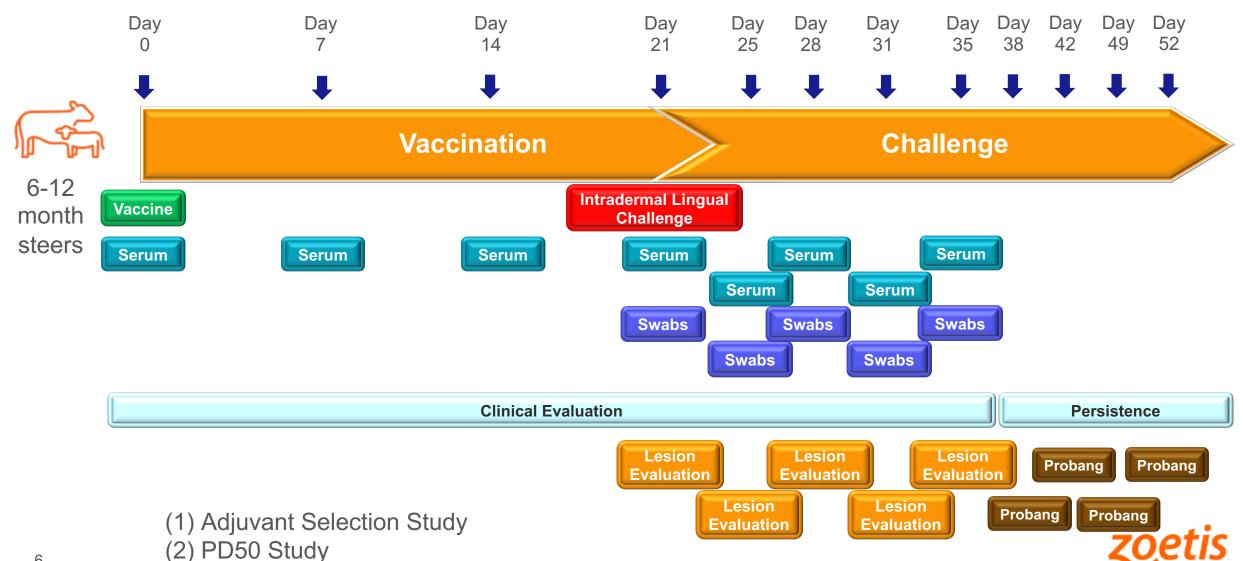
- √ No clinical disease
- √ No viral shedding
- √ No fever spike
- ✓ No contact transmission
- ✓ Very limited if any immune response

Construct	Inoculation Route	# Animals
FMD-LL3B3D-A24 Cruzeiro	Heelbulb and Contact (1x10 ⁵)	4
FMD-LL3B3D-Asia1 Shamir	Heelbulb and Contact (1x10 ⁶)	5
FMD-LL3B3D-A Turkey 06	Heelbulb and Contact (1x10 ⁶)	5
FMD-LL3B3D-O1 Campos	Heelbulb and Contact (1x10 ⁶)	4
FMD-LL3B3D-A Argentina 2001	Heelbulb and Contact (2x10 ⁶)	4
FMD-LL3B3D-C3 Indaial	Heelbulb and Contact (2.8x10 ⁶)	4



> Safety study results supported the US Select Agent Exclusion

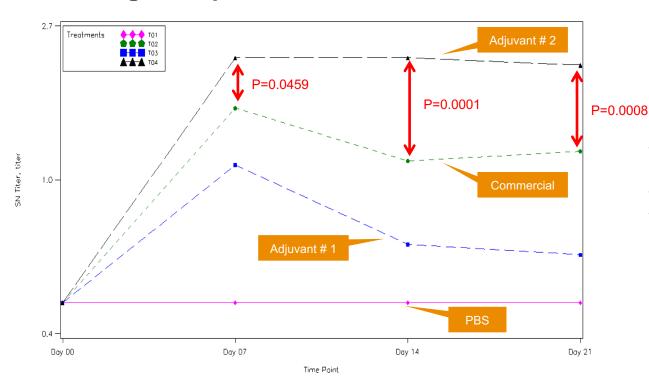
SUMMARY OF FMD-LL3B3D A24 CRUZEIRO CATTLE EFFICACY STUDY DESIGNS



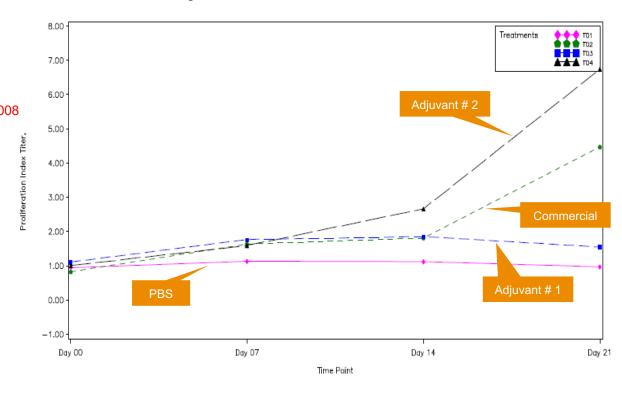
(1) FMD-LL3B3D A24 CRUZEIRO ADJUVANT SELECTION STUDY—SEROLOGY/CMI

Robust antibody and cellular responses with adjuvant # 2

A: Serologic Responses



B: Cellular Responses

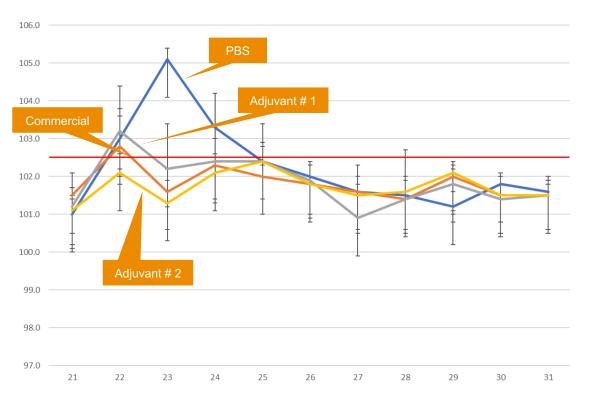




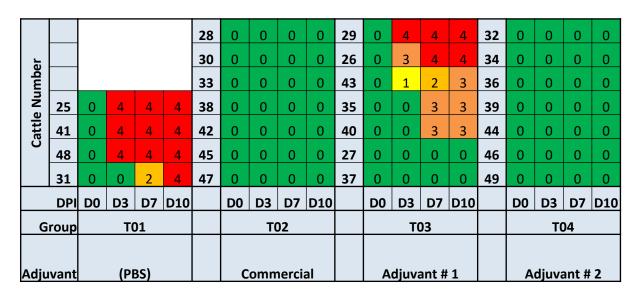
(1) FMD-LL3B3D A24 CRUZEIRO ADJUVANT SELECTION STUDY CLINICAL

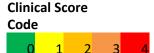
Prevention of clinical signs of FMD with adjuvant # 2 upon challenge

A: Temperature (°F)



B: Clinical Signs





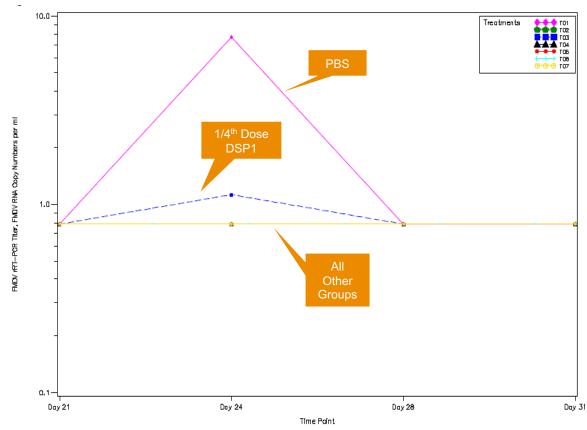


(2) FMD-LL3B3D A24 CRUZEIRO VACCINE IN ADJUVANT # 2 – PD50 STUDY



• Full dose prevented viremia, clinical signs, & persistent infection

A: Viremia



B: Clinical Signs and Persistence

		Clinical Signs			Log10 RNA Copies (Probang)			Virus Isolation (Probang)					
		Day of Study											
Group	Animal	20	23	27	30	38	42	49	52	38	42	49	52
PBS	R14-84	0	2	4	4	4.29	4.72	0	3.83	Positive	Positive	Positive	Positive
	R14-85	0	4	4	4	4.26	6.01	5.14	4.7	Positive	Positive	Positive	Positive
	R14-86	0	1	4	4	0	3.62	0	0	Negative	Negative	Negative	Negative
	R14-87	0	4	4	4	0	0	0	0	Negative	Negative	Negative	Negative
Full Dose	R14-72	0	No	No	No	0	0	0	0		Negative		
	R14-73	0	No	No	No	0	0	0	0		Negative		
DSP1	R14-74	0	No	No	No	0	0	0	0	Negative	Negative	Negative	Negative
DOFT	R14-75	0	No	No	No	0	0	0	0	Negative	Negative	Negative	Negative
1/4th Dose	R14-76	0	No	No	No	4.98	4.68	0	0	Positive	Positive	Negative	Positive
	R14-77	0	1	1	1	5.52	3.43	0	0	Positive	Positive	Negative	Negative
DSP1	R14-78	0	No	No	No	0	4.35	0	5.3	Positive	Positive	Positive	Positive
DOI 1	R14-79	0	No	No	No	0	0	0	0	Negative	Negative	Negative	Negative
1/16th	R14-80	0	No	No	No	0	0	4.88	4.59	Positive	Negative	Positive	Positive
Dose	R14-81	0	No	No	No	5.08	4.01	3.98	4.65	Positive	Positive	Positive	Positive
DSP1	R14-82	0	No	No	No	0	4.47	6.12	4.32	Positive	Positive	Positive	Positive
DOI 1	R14-83	0	No	No	No	0	0	0	0	Positive		Positive	
Full	R14-60	0	No	No	No	0	0	0	0		Negative		
Dose	R14-61	0	No	No	No	0	0	0	0	Negative	Negative	Negative	Negative
DSP2	R14-62	0	No	No	No	4.75	0	0	0	Negative	Negative	Negative	Negative
DO: 2	R14-63	0	No	No	No	0	0	0	0	Negative	Negative	Negative	Negative
1/4th	R14-64	0	No	No	No	0	0	0	0	Negative	Negative	Negative	Negative
1/4th Dose DSP2 1/16th Dose DSP2	R14-65	0	No	No	No	4.1	4.11	0	3.39	Positive	Positive	Positive	Positive
	R14-66	0	No	No	No	0	0	0	0	Negative	Negative	Negative	Negative
	R14-67	0	No	No	No	4.14	5.08	5.18	4.82		Positive		
	R14-68	0	No	No	No	0	0	0	0		Negative		
	R14-69	0	No	No	No	0	0	0	0	Negative	Negative	Negative	Negative
	R14-70	0	No	No	No	0	0	0	0	Negative	Negative	Negative	Negative
	R14-71	0	No	No	No	5.34	5.46	4.49	3.7	Positive	Positive	Positive	Positive

Total abnormal vesicle sco

0

2



EFFICACY OF THE FMD-LL3B3D A24 CRUZEIRO IN CATTLE

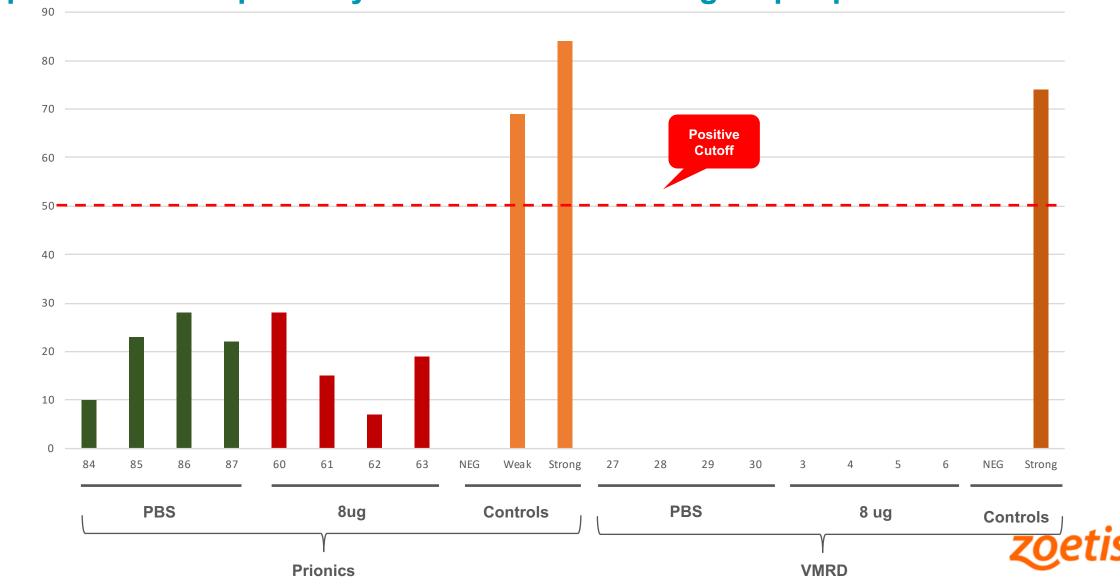
FMD-LL3B3D A24 Cruzeiro vaccine prevented FMD lesions at 1/16th dose

- Vaccines formulated with inactivated FMD-LL3B3D A24 Cruzeiro and a proprietary Zoetis oil-based adjuvant
- Clinical outcomes
 - Higher serum antibody titers compared to conventional vaccine
 - More robust CMI response compared to conventional vaccine
 - Complete prevention of FMD lesions
 - Even with 1/16 dose (0.5 μg antigen)
 - Full dose (8 μg) vaccine prevented persistent infection (DSP # 1 and # 2)
 - Prevention of fever
 - Prevention of viremia (DSP # 2)
 - Significant reduction in shedding (data not shown)



DIVA COMPATIBILITY WITH CURRENT COMMERCIAL ASSAYS - (PrioCHECK AND VMRD)

Complete DIVA compatibility due to absence of target epitope



% Inhibition

LL3B3D PLATFORM VACCINE - CONCLUSIONS

Inactivated FMD-LL3B3D platform vaccine offers protection of livestock from the threat of natural or intended FMD outbreaks with full DIVA compatibility

- ✓ Prevents FMD lesions and fever even after 1 vaccination in cattle
- ✓ Prevents FMD persistent infection
- ✓ Allows differentiation of infected from vaccinated animals (DIVA)
- ✓ Safe *in vitro* and *in vivo*: produced without the need for virulent FMDV
- ✓ No need for adaptation of field strains to cell culture
- ✓ Affords a rapid response capability



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