



**AU-PANVAC
Laboratories**



Quality Control of PPR & FMD Vaccines

Dr Charles BODJO, Ag Director AU-PANVAC

AU-PANVAC's Mission & Mandates

 **MISSION:** “To promote the use of **GOOD QUALITY VACCINES** and **DIAGNOSTIC REAGENTS** for the control, eradication and surveillance of animal diseases in Africa.”

 MANDATES

- 1 **INDEPENDENT QUALITY CONTROL** of vet. vaccines.
- 2 Development and Production of **DIAGNOSTICS**
- 3 **TRANSFER OF VACCINE PRODUCTION TECHNOLOGIES**
- 4 **STANDARDIZATION** of production and **HARMONIZATION** of the quality control techniques in Africa
- 5 **CAPACITY BUILDING** to veterinary laboratories.
- 6 **AUDITING AND CERTIFICATION of Vaccine Manufacturing Facilities** *in collaboration with National Regulatory Authorities in Africa.*



International Recognitions of AU-PANVAC



World Organisation
for Animal Health
Founded as OIE

- **Collaborating Center For Quality Control of Veterinary Vaccine** (WOAH Gen. Assembly Resolution 32, Paris, May 2013)



Food and Agriculture
Organization of the
United Nations

- **Reference Centre for Technical Assistance in Quality Control of Veterinary Vaccine** (11th May, FAO Rome, 2015)



GF-TADs
GLOBAL FRAMEWORK FOR THE
PROGRESSIVE CONTROL OF
TRANSBOUNDARY ANIMAL DISEASES



Food and Agriculture
Organization of the
United Nations



World Organisation
for Animal Health
Founded as OIE

- **Rinderpest Holding Facility (RHF) for Africa** (among five other RHF's: US, UK, France, China & Japan)



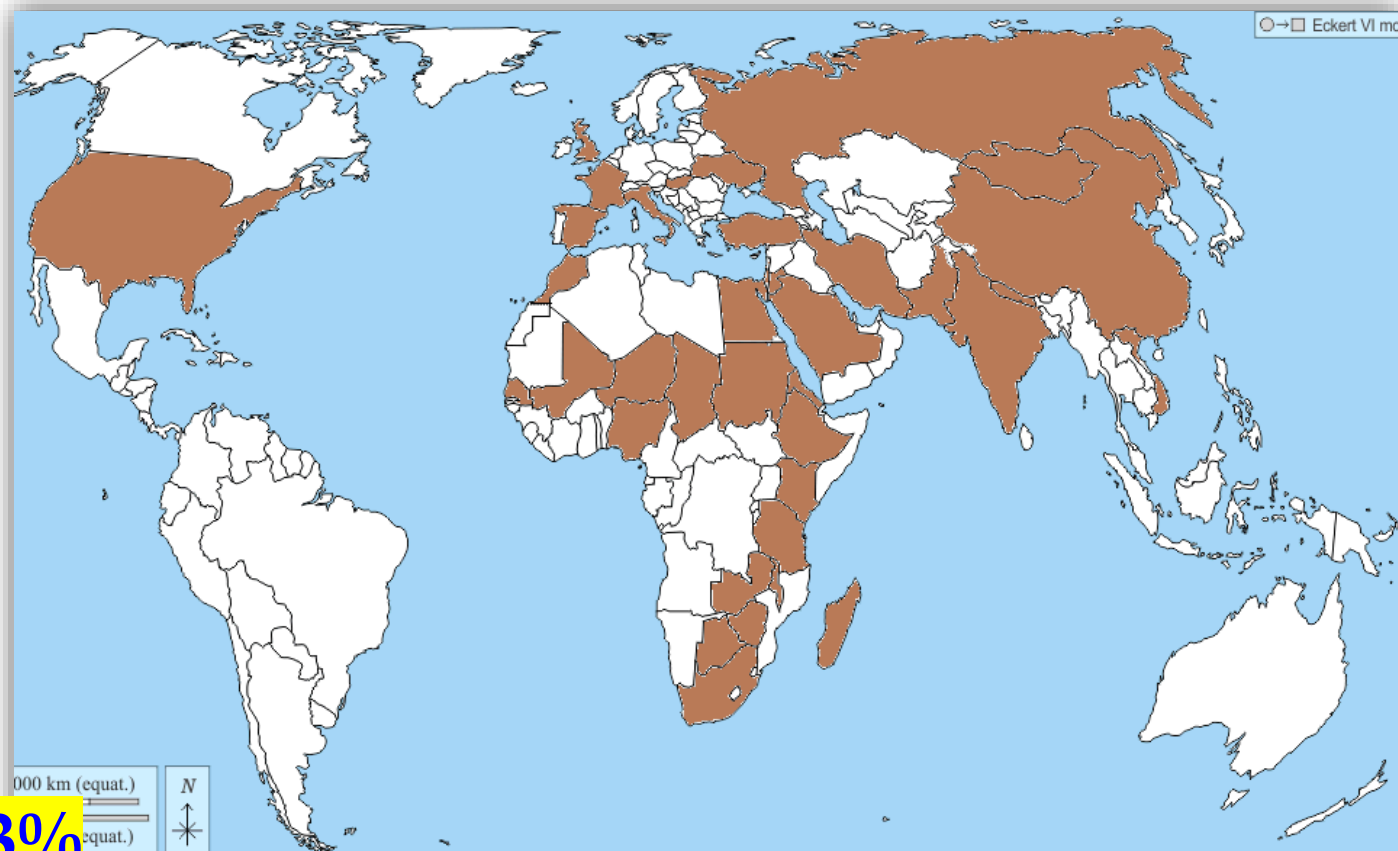
AU-PANVAC
ETHIOPIA.



Vaccine Quality Control Activities

❑ African Manufacturers (22): 67%

- Egypt (**VSRVI, MEVAC**) & Morocco (**MCI & BIOPHARMA**),
- Senegal (**ISRA**), Mali (**LCV**), Niger (**LABOCEL**) & Nigeria (**NVRI**)
- Cameroun (**LANAVET**), Chad (**IREDA**)
- Ethiopia (**NVI**) & Kenya (**KEVEVAPI, INNOVA**), Tanzania (**TVI, HESTER Africa**)
- Botswana (**BVI**), South Africa (**OBP, MSD**), Madagascar (**IMVAVET**), Malawi (**CTTBD**), Zambia (**CVRI**), Zimbabwe (**CVL**)



❑ Manufacturers out of Africa (23): 33%

France, Spain, UK, Hungary (**Boehringer, CEVA, ZOETIS, HIPRA, Vaxxinova**), Kazakhstan (**BIOTRON**), Pakistan, Russia (**ARRIAH**), Turkey (**DOLVET, VETAL**), Jordan (**JOVAC**), Iran (**VIRA Vaccine, RAZIVSRI**), India (**Brilliant Biopharma, Biomed Indian Immunologicals, Ventri**), Ukraine (**Biotestlab**), USA (**Zoetis, CEVA**), Vietnam (**AVAC**), China (**SINOPHARM ANIMAL HEALTH CORPORATION, SHAN DONG LVDU, TECON BIOLOGY JOINT STOCK COMPANY**), Saudi Arabia (**IBRIZ**)

Vaccine Quality Control Activities...

- More than **50 types of Vet. Vaccines** (for all animal species) tested
- **≈ 300-400 batches** received annually

≈ 2 Billion doses of vaccines to protect all animal species

- Required 4 Weeks Test/Batch
- Publication of batches passed QC tests on the AU-PANVAC website:

WWW.AUPANVAC.ORG



Current PPR vaccines

- ❑ PPRV Nigeria 75/1 Strain (*Diallo et al. 1989; Diallo 2004*): Lineage II, **Most widely used in the world (ONLY currently used in AFRICA)**
- ❑ PPRV Sungri/96 (*Singh et al. 2009; Saravanan et al. 2010*): Lineage IV, **used only in India**
- ❑ Duration of immunity: **immunity for at least 3 years.**
- ❑ Vaccine produced in freeze-dried form,
 - Heat sensitive.
 - Need to **maintain cold chain** to guarantee vaccine efficacy for field use.

Peste des Petits Ruminants Vaccine (Nigerian Strain 75/1) Confers Protection for at Least 3 Years in Sheep and Goats

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Received July 2014

Abstract

The present study reports the duration of immunity and protective efficacy of Peste des Petits Ruminants (PPR) vaccine (Nigerian strain 75/1) in sheep and goats. A total of 105 sheep and goats were divided into three groups A, B and C. Group A received normal recommended dose (1.0 ml) of PPR vaccine, group B received half dose (0.5 ml) of PPR vaccine and group C was kept as unvaccinated control group in contact with vaccinated animals. The post vaccination dynamics of antibodies against PPR virus was studied. It was found that significant antibody titres persisted for 3 years post vaccination in sheep and goats vaccinated with either full dose or half dose of PPR vaccine. The challenge protection studies were carried out in experimental animals at 24 and 36 month post vaccination. The vaccinates withstood challenge and were found completely resistant clinically and virologically to virulent PPR virus for 24 and 36 months post vaccination. The unvaccinated control animals developed typical clinical signs of PPR and the challenged virus was detected in ocular, nasal and oral secretions of these animals. This study demonstrated that a single immunization with PPR vaccine conferred solid protection in sheep and goats for 3 years.

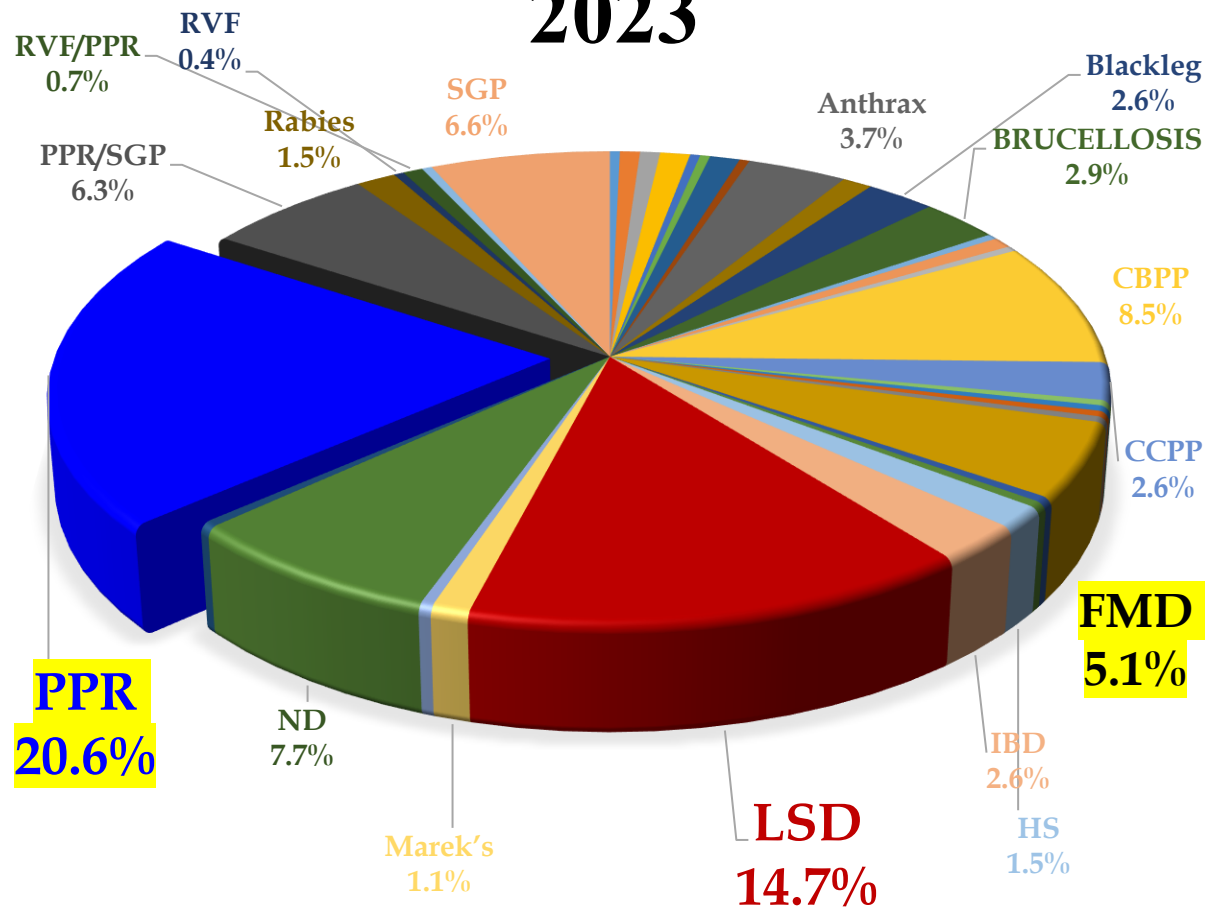
Keywords

PPR Vaccine, Small Ruminants

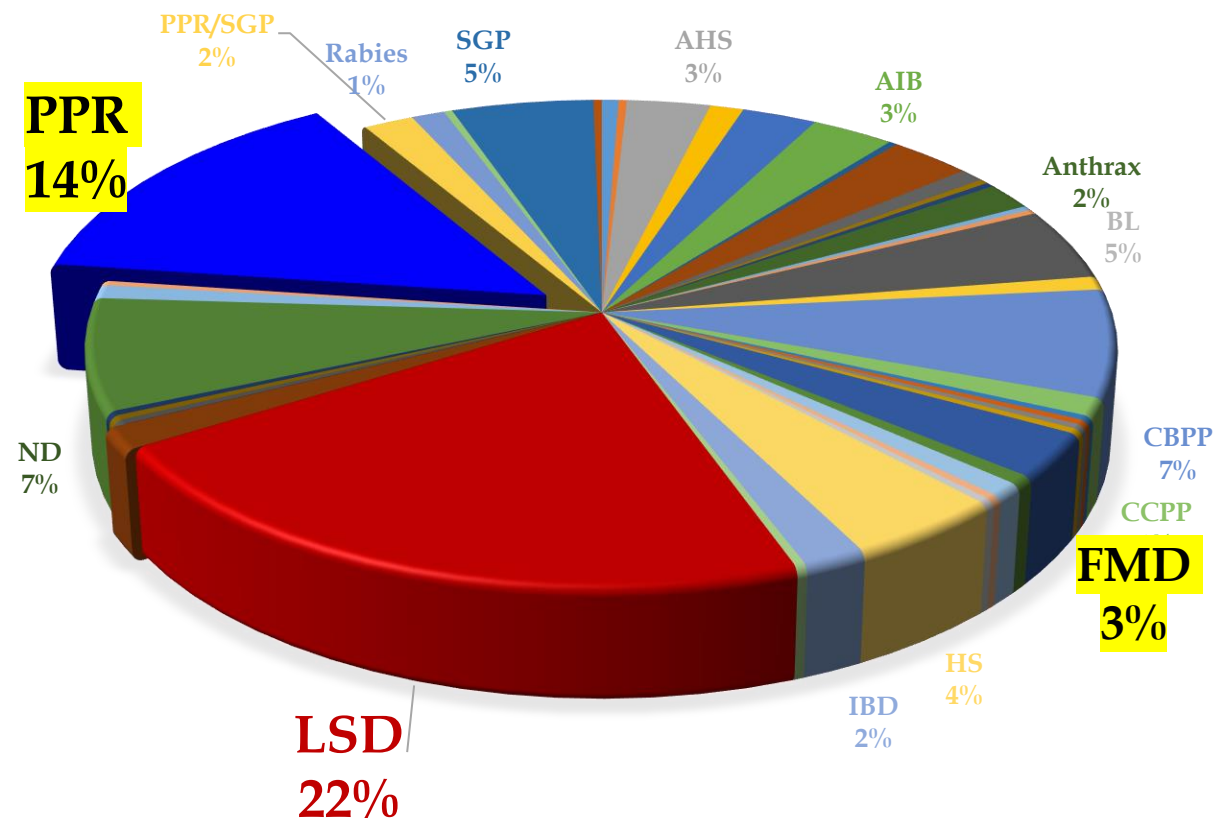


Proportion of PPR Vaccines

2023



2024



PPR vaccine was the major vaccine tested throughout the years, however this seems to be declined in favor of LSD vaccines



PPR Vaccines Quality Control Pass Rate

Year	Batches received	Pass Rate (%)
2020	61	96.7
2021	78	98.7
2022	88	84.1
2023	43	80
2024	70	93.3



❑ In December 2017, FAO and WOAHP PPR-Secretariat, with the support of the Global Alliance for Livestock Veterinary Medicines (GALVmed), organised a workshop on PPR thermotolerant vaccines at the FAO Headquarters in Rome:



- Define criteria for evaluation of PPR Thermotolerant vaccine
- Proposal to use an average temperature of **40°C** to evaluate thermotolerant PPR vaccines



Standard for PPR Thermotolerant Vaccine

❑ Define Criteria/Standards for Testing PPR Thermotolerant Vaccine

- (1) **Titer $\geq 2.5 \log_{10}$** (TCID₅₀/ml) after incubation **40°C/Day-5**
- (2) **Titer Loss** after post exposure should be **$\leq 1 \log_{10}$**

❑ For comparison with other viral vaccines:

- *Rindertest Thermostable vaccine*: **Titer $\geq 2.5 \log_{10}$** (TCID₅₀/ml) after incubation **45°C/14 Days** & **Titer loss $\leq 1.6 \log_{10}$**
- *Measles-Mumps vaccine*: **Titer $\geq 3 \log_{10}$** (TCID₅₀/ml) after incubation **37°C/7Days** & **Titer loss $\leq 1 \log_{10}$**





Article

Peste des Petits Ruminants Vaccine: Criteria for Assessing Its Thermotolerance

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<https://doi.org/10.3390/v17091151>



Criteria for Testing Thermotolerant PPR Vaccine Submitted to the WOAHA Biological Commission

- ❑ Presentation on the criteria and the Standard Operating Procedure (SOP) for testing thermotolerant PPR vaccines was made on Sept. 2025 to WOAHA Biological Commission.
- ❑ The WOAHA Biological Commission agreed to:
 - Incorporate the criteria for assessing thermotolerant of PPR vaccines in the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, Chapter 3.8.8 (Section C.2.2.4 or C.2.3.3 on batch release).
 - Detailed step-by-step SOP for testing thermotolerant PPR vaccines should be prepared and maintained through available platforms such as the [WOAHA PPR Reference Laboratory Network website](#)."



Development of PPR Diagnostics



PPR-bELISA (*Archives of Virology, 2018*)

Archives of Virology

<https://doi.org/10.1007/s00705-018-3782-1>

ORIGINAL ARTICLE



Development and validation of an epitope-blocking ELISA using an anti-haemagglutinin monoclonal antibody for specific detection of antibodies in sheep and goat sera directed against peste des petits ruminants virus

Sanne Charles Bodjo¹ · Jean-de-Dieu Baziki¹ · Nick Nwankpa¹ · Ethel Chitsungo¹ · Yao Mathurin Koffi² · Emmanuel Couacy-Hymann² · Mariame Diop³ · Daniel Gizaw⁴ · Idris Badri Adam Tajelser⁵ · Mamadou Lelenta⁶ · Adama Diallo⁷ · Karim Tounkara⁸

Received: 29 June 2017 / Accepted: 7 February 2018
© Springer-Verlag GmbH Austria, part of Springer Nature 2018

Assay Performances:

Specificity: 99%

Sensitivity: 93.74%



Evaluation ELISAs Cross reactivity against CDV Antibodies

Sera from Dogs (*immunized with CDV vaccine*) tested with ID Screen® PPR Competition (IDVET) and PPR-bELISA (PANVAC):

	ID Screen® PPR Competition (IDvet)	PPR-bELISA (PANVAC)
% Positive	12/63	0/63
% Doubtful	7/63	0/63
Negative	44/63	63/63



Goats Immunized with CDV Vaccine

Sera from Goats immunized with CDV vaccine tested with PPR-cELISA (IDVET) and PPR-bELISA (PANVAC)

□ Day 0

SAMPLE TYPE	ID Screen cELISA (% PI)	PANVAC bELISA (% PI)
DO-691	72.47 (Neg)	-0.37 (Neg)
DO-692	57.73 (Neg)	6.41 (Neg)
DO-685	60.15 (Neg)	-7.44 (Neg)
DO-686	51.40 (Neg)	-11.24 (Neg)
DO-696	71.18 (Neg)	0.90 (Neg)

□ Day 31

SAMPLE TYPE	ID Screen cELISA (% PI)	PANVAC bELISA (% PI)
D31-691	67.38 (Neg)	-7.05 (Neg)
D31-692	68.27 (Neg)	-5.34 (Neg)
D31-685	38.59 (Pos)	-0.02 (Neg)
D31-686	25.58 (Pos)	9.19 (Neg)
D31-696	68.27 (Neg)	6.32 (Neg)



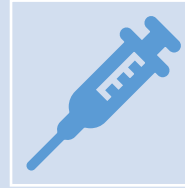
False positives results can occur with ID Screen® PPR Competition (IDvet)



Quality Control of FMD Vaccines & Challenges



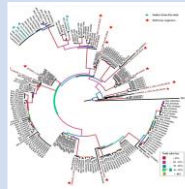
FMD is an important constraint to livestock with impact on global trade.



Vaccination is a critical component of FMD control



7 serotypes of FMDV (O, A, C, SAT1, SAT2, SAT3, and Asia1) and multiple subtypes are currently recognized



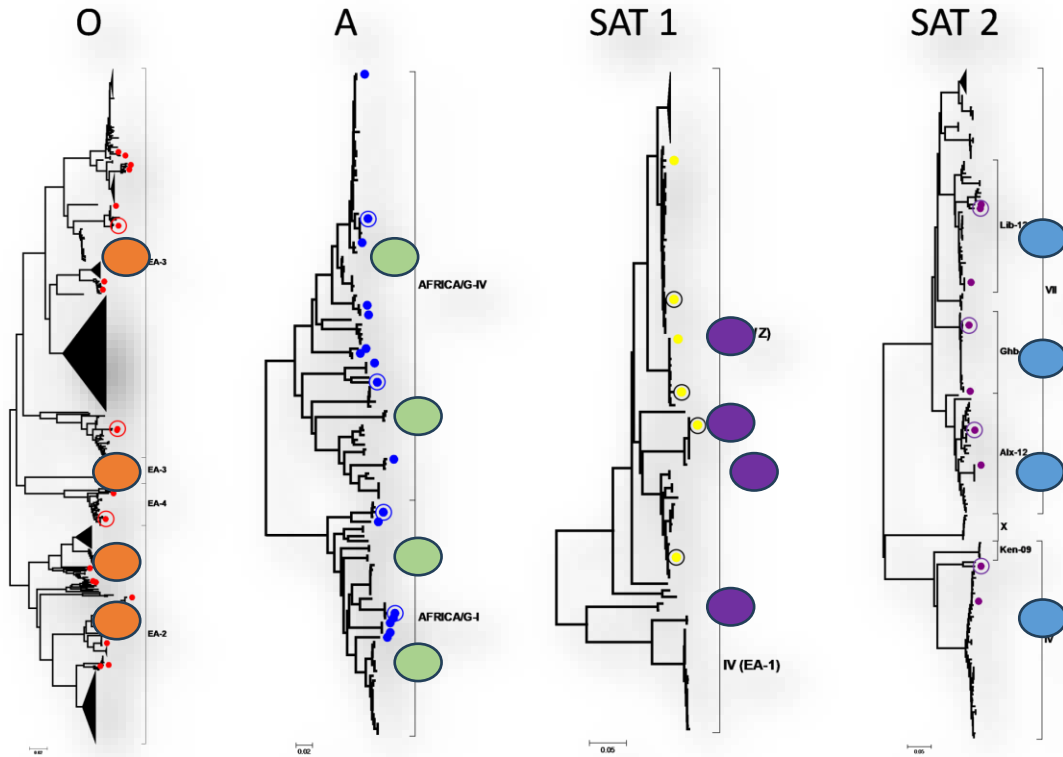
Constant evolving of antigenic diversity of FMD virus require that vaccine strains **BE MATCHED** with field strains.



Development of FMDV Reference Panel for Vaccine Quality Control



Reference Panels for Quality Control of FMD



Selection of a panel of **16 FMD Viruses** (**O**, **A**, **SAT1** & **SAT2**) among O = 709; A = 138; SAT 1 = 143; SAT 2 = 190 from the repository WRLFMD covering the genetic diversity circulating in **Eastern African Region (10 countries)** and used for VNT



East African Region FMD Virus Panel

16 VIRUSES SELECTED

Serotype A	Serotype O	Serotype SAT 1	Serotype SAT 2
SUD/9/2018	ETH/4/2015	TAN/22/2013	ETH/16/2015
ETH/2/2018	ETH/9/2019	KEN/10/2013	KEN/19/2017
UGA/28/2019	ETH/30/2016	TAN/27/2012	EGY/1/2018
ETH/19/2019	KEN/4/2018	TAN/22/2014	ETH/11/2018



Immunogenicity Trials on the Reference Panel

❑ Sera from vaccinated animals with 13 batches of commercial FMD vaccines supplying of quadrivalent FMD vaccine O, A, SAT 1, and SAT 2.

❑ Coloured boxes show sera from vaccinated animal with FMD vaccines that induced ≥ 1.5 log₁₀ antibody responses in $\geq 60\%$ of cattle

❑ Grey colour indicates positive responses at day 0 results were not provided to confirm the validity of the trial).



Results showed variability in vaccine potency and some batches failed to provide adequate cross-protection

Vaccine Batch	Trial Location ^a	Serotypes Included	Time between manufacture and use (months)	No. of Cattle Vaccinated (and of unvaccinated controls where present)	Score ^c			
					O	A	SAT 1	SAT 2
1	TPI (F)	O, A, SAT 1, SAT 2	5	5				
2	TPI (F)	O, A, SAT 1, SAT 2	8	10				
2b	TPI (F)	O, A, SAT 1, SAT 2	8	5				
3	AU-P (E)	O, A, SAT 1, SAT 2	6	10(2) ^b				
3b	AU-P (E)	O, A, SAT 1, SAT 2	6					
4	AU-P (E)	O, A, SAT 1, SAT 2	6	10(2) ^b				
4b	AU-P (E)	O, A, SAT 1, SAT 2	6					
5	AU-P (E)	O, A, SAT 1, SAT 2	10	10(2) ^b				
5b	AU-P (E)	O, A, SAT 1, SAT 2	10					
6	Other (F)	O, A, SAT 1, SAT 2	ND ^d	5				
7	Other (F)	O, A, SAT 1, SAT 2	ND	5				
8	Other (F)	O, A, SAT 1, SAT 2	ND	5				
9	Producer (E)	O, A, SAT 1, SAT 2	13	5 ^b				
9b	Producer (E)	O, A, SAT 1, SAT 2	13					
10	Producer (E)	O, A, SAT 1, SAT 2	2	6				
11	TPI (F)	O, A, SAT 2	17	5				
12	Other (F)	O, A, SAT 2	6	5 ^b				
12b	Other (F)	O, A, SAT 2	6					
13	TPI (F)	SAT 1, SAT 2	2	10				
13b	TPI (F)	SAT 1, SAT 2	2	5				



Publications on East African FMD Virus Panel

Publication 1 : 2021

Rev. Sci. Tech. Off. Int. Epiz., 40 (1)

A.B. Ludi, V. Mioulet, L. Bakkali Kassimi, D.J. Lefebvre, K. De Clercq,
E. Chitsungo, N. Nwankpa, W. Vosloo, D.J. Paton & D.P. King

Selection and use of reference panels: a case study highlighting
current gaps in the materials available for foot and mouth disease 239

*Sélection et utilisation des panels de référence : à partir de l'exemple
de la fièvre aphteuse, étude soulignant les lacunes actuelles en la matière
(résumé)..... 247*

*Selección y uso de paneles de referencia: estudio de las carencias
de los paneles disponibles actualmente a partir del ejemplo de la fiebre
aftosa (resumen)..... 248*

<https://doi.org/10.20506/rst.40.1.3221>

Publication 2: 2025

npj | vaccines

Published in partnership with the Sealy Institute for Vaccine Sciences

Article



<https://doi.org/10.1038/s41541-025-01128-7>

An antigen panel to assess the regional relevance of foot and mouth disease vaccines

Check for updates

David J. Paton¹ ✉, Ginette Wilsden¹, Clare FJ Browning¹, Efrem A. Foglia², Antonello Di Nardo¹,
Nick J. Knowles¹, Jemma Wadsworth¹, Simon Gubbins¹, Ethel Chitsungo³,
Cisse Rahamatou Moustapha Boukary², Gelagay Ayelet², Charles S. Bodjo³, Nick Nwankpa³,
Emiliana Brocchi², Santina Grazioli², Anna Ludi¹ & Donald P. King¹

<https://doi.org/10.1038/s41541-025-01128-7>



Future Direction

- ❑ Developing similar approach of FMD virus panel for:



Northern



Western



Central



Southern

- ❑ Exploration of guinea pigs as an alternative model for vaccine testing.



AU-PANVAC New Facility Complex



- Fully supported by **US - DTRA**



Enhance Biosafety and Biosecurity measures



Expand Laboratory operation
Training & capacity

- Groundbreaking Ceremony held on **21st Feb. 2024** & presided by H.E Excellency Moussa Faki Mahamat, Chairperson of AUC



Construction AU-PANVAC New Lab. Complex

❑ Commencement of Construction work March 2026



Construction site visit by His Excellency Moses Vilakati, Commissioner for Agriculture, Rural Development, Blue economy, and Sustainable Environment (ARBE)

❑ Construction completion expected in 2028





AU-PANVAC!

ADDING VALUE TO ANIMAL HEALTH AND HUMAN LIVES!!

Thank you

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