

# WOAH Reference Laboratory Reports Activities 2022

## Activities in 2022

This report has been submitted : 15 février 2023 15:57

### Laboratory Information

<b>Name of disease (or topic) for which you are a designated WOA Reference Laboratory:</b>	Rift Valley fever
<b>Address of laboratory:</b>	CIRAD, UMR ASTRE, TA A15/E, Campus International de Baillarguet, 34398 MONTPELLIER Cedex 5
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<b>Name (including Title) of Head of Laboratory (Responsible Official):</b>	VACHIERY Nathalie
<b>Name (including Title and Position) of WOA Reference Expert:</b>	CETRE-SOSSAH Catherine
<b>Which of the following defines your laboratory? Check all that apply:</b>	Research agency

### TOR1: DIAGNOSTIC METHODS

1. Did your laboratory perform diagnostic tests for the specified disease/topic for purposes such as disease diagnosis, screening of animals for export, surveillance, etc.? (Not for quality control, proficiency testing or staff training)

Yes

Diagnostic Test	Indicated in WOA Manual (Yes/No)	Total number of test performed last year	
Indirect diagnostic tests		Nationally	Internationally
ELISA IgG	Yes	47	120
ELISA IgM	Yes		82
Direct diagnostic tests		Nationally	Internationally
RTqPCR	Yes		459
Partial sequencing	Yes		

			4
Full sequencing	Yes		1

## TOR2: REFERENCE MATERIAL

2. Did your laboratory produce or supply imported standard reference reagents officially recognised by WOA?H?

No

3. Did your laboratory supply standard reference reagents (nonWOAH-approved) and/or other diagnostic reagents to WOA?H Members?

Yes

TYPE OF REAGENT AVAILABLE	RELATED DIAGNOSTIC TEST	PRODUCED/ PROVIDE	AMOUNT SUPPLIED NATIONALLY (ML, MG)	AMOUNT SUPPLIED INTERNATIONALLY (ML, MG)	NO. OF RECIPIENT WOA?H MEMBER COUNTRIES	COUNTRY OF RECIPIENTS
Standard RVF positive goat serum	ELISA	produced		1 ml	10	Africa

4. Did your laboratory produce vaccines?

Not applicable

5. Did your laboratory supply vaccines to WOA?H Members?

Not applicable

## TOR3: NEW PROCEDURES

6. Did your laboratory develop new diagnostic methods for the designated pathogen or disease?

No

7. Did your laboratory validate diagnostic methods according to WOA?H Standards for the designated pathogen or disease?

No

8. Did your laboratory develop new vaccines for the designated pathogen or disease?

No

9. Did your laboratory validate vaccines according to WOA?H Standards for the designated pathogen or disease?

No

## TOR4: DIAGNOSTIC TESTING FACILITIES

10. Did your laboratory carry out diagnostic testing for other WOA?H Members?

Yes

NAME OF WOA?H MEMBER COUNTRY SEEKING ASSISTANCE	DATE	WHICH DIAGNOSTIC TEST USED	NO. SAMPLES RECEIVED FOR PROVISION OF DIAGNOSTIC SUPPORT	NO. SAMPLES RECEIVED FOR PROVISION OF CONFIRMATORY DIAGNOSES
GABON	2022-01-06	RTqPCR	434	
GABON	2022-11-04	ELISA IgG	114	
SOUTH AFRICA	2022-01-06	ELISA IgM	6	

11. Did your laboratory provide expert advice in technical consultancies on the request of an WOA Member?

Yes

NAME OF THE WOA MEMBER COUNTRY RECEIVING A TECHNICAL CONSULTANCY	PURPOSE	HOW THE ADVICE WAS PROVIDED
SUDAN	RVF training	Email Exchanges
BURUNDI	RVF vaccine production and training offers	Email Exchanges
CHAD	Possibility of collaboration with IRED for capacity building in animal disease diagnostic	Email exchanges

## **TOR5: COLLABORATIVE SCIENTIFIC AND TECHNICAL STUDIES**

12. Did your laboratory participate in international scientific studies in collaboration with WOA Members other than the own?

Yes

Title of the study	Duration	PURPOSE OF THE STUDY	PARTNERS (INSTITUTIONS)	WOAH MEMBER COUNTRIES INVOLVED OTHER THAN YOUR COUNTRY
Study of the mechanisms of neuropathology associated with Rift Valley Fever virus infection Rift Valley fever virus (NEURORIFT)	3 years	Study of neuropathology associated with Rift Valley Fever virus infection with humans and ruminants RVFV strains from Mayotte and Mauritania respectively	INSERM, UMR PCCEI, Montpellier, France	FRANCE
Impact of genetic diversity of Rift Valley fever virus on its replicative capacity	3 years	Investigate the genetic diversity of Rift Valley fever virus in order to adapt prevention and control tools	INRAE, UMR IVPC, Lyon, France	FRANCE
Epidemiological and socio-economic status of Rift Valley fever (RVF) in Burundi	3 years	Investigate the socio-economic status of Rift Valley fever (RVF) in Burundi	LNV, Bujumbura, Burundi	BURUNDI
WOAH twinning LNERV-CIRAD	2 years	2 years Support LNERV to become regional ref lab for RVF	ISRA-LNERV	SENEGAL
WOAH twinning LCV-CIRAD	2 years	Support LCV to become regional ref lab for RVF	LCV	MALI
RFOROA One health	1 year	Diagnostic support IRED, Smithsonian Institute	IRED, Smithsonian Institute	CHAD

## **TOR6: EPIZOOLOGICAL DATA**

14. Did your Laboratory collect epidemiological data relevant to international disease control?

Yes

IF THE ANSWER IS YES, PLEASE PROVIDE DETAILS OF THE DATA COLLECTED:

- 1) development of a rapid Diagnostic test to detect 2)RVF genetic diversity and evolution of RVFV genotype in West Africa;3) quality of RVFV sequence data published

15. Did your laboratory disseminate epidemiological data that had been processed and analysed?

Yes

IF THE ANSWER IS YES, PLEASE PROVIDE DETAILS OF THE DATA COLLECTED:

- 1) development of a rapid Diagnostic test to detect 2)RVF genetic diversity and evolution of RVFV genotype in West Africa;3) quality of RVFV sequence data published

16. What method of dissemination of information is most often used by your laboratory? (Indicate in the appropriate box the number by category and list the details in the box)

a) Articles published in peer-reviewed journals:

[22-01] Barry Y, Elbara A, Bollahi MA, Ould El Mamy AB, Fall M, Beyit AB, Khayar MS, Demba BA, Limine Haki ML, Faye O, Plee L, Bonbon E, Doumbia B, Arsevska E, Cetre-Sossah C. 2022. Rift Valley fever, Mauritania, 2020: Lessons from a one health approach. *One Health* 15. 100413. <https://doi.org/10.1016/j.onehlt.2022.100413>

[22-02] Seck I, Lo MM, Fall AG, Diop M, Ciss M, Cêtre-Sossah CB, et al. 2022. Identification of drivers of Rift Valley fever after the 2013–14 outbreak in Senegal using serological data in small ruminants. *PLoS Negl Trop Dis* 16(2): e0010024. <https://doi.org/10.1371/journal.pntd.0010024>

b) International conferences:

12th International ESVV Congress, 20-23 September, 2022, Ghent, Belgium. Impact of genetic diversity of the Rift Valley Fever virus, from the field isolates to a genetic determinant

Association of Institutions for Tropical Veterinary Medicine (AITVM), the Society for Tropical Veterinary Medicine (STVM) virtual event, Tuesday 17 May 2022. Investigation of the inflammatory response induced by a Rift Valley fever infection (RVF) with a specific focus on the central nervous system (CNS) and Impact of genetic diversity on the replication capacity of the Rift Valley Fever virus

c) National conferences:

Journées Françaises de Virologie, Strasbourg, France. 11-12 April 2022. Etude des mécanismes inflammatoires associés à l'infection du système nerveux central humain par le virus de la fièvre de la vallée du Rift (vFVR),

d) Other (Provide website address or link to appropriate information):

<https://umr-astre.cirad.fr/>

## TOR7: SCIENTIFIC AND TECHNICAL TRAINING

17. Did your laboratory provide scientific and technical training to laboratory personnel from other WOAHA Members?

Yes

a) Technical visit :

b) Seminars : 2+38+40

c) Hands-on training courses: 2

d) Internships (> 1 month)

Type of technical training provided (a, b, c or d)	Country of origin of the expert(s) provided with training	No. participants from the corresponding country
c	Senegal	2
c	Burundi	12

## TOR8: QUALITY ASSURANCE

18. Does your laboratory have a Quality Management System?

Yes

Quality management system adopted	Certificate scan (PDF, JPG, PNG format)	
ISO 17025v2017		COFRAC Accreditation Certificate N° 1-2207 rev.15.pdf

19. Is your quality management system accredited?

Yes

Test for which your laboratory is accredited	Accreditation body
ELISA, PCR	COFRAC

20. Does your laboratory maintain a "biorisk management system" for the pathogen and the disease concerned?

Yes

All efforts are being made to work under biosafety level 3 containment facilities in Montpellier and under biosafety level 2 containment under dedicated safe hood cabinet wherever it is available. Personal equipment (dedicated laboratory coat, gloves, masks, glasses) are being used. Senegalese and French rules are followed up. Transport of biological materials considered as infectious substances by air are done according to the international regulation's guidelines developed by the national regulations, ICAO/IATA/CITES\* regulations, through an air carrier company from the country where the samples are collected to CIRAD (Montpellier, France) and vice versa. The reference laboratory is used to receive and send infectious animal substances by air and has persons dedicated to the management of these shipments that are fully aware of the relevant regulations and of the proper process (identification, categorization, packaging, marking, labelling, documenting and refrigerating). When the candidate laboratory will intend to send infectious animal samples, contact will be made with the person in charge to make the shipment and written procedures and assistance will be given. Briefly, the IATA dangerous goods regulation indicate for the packaging instruction 602 for the shipment to arrive in good condition and to present

no hazard to persons or to animals is the following: the package must include • A inner packaging comprising, watertight primary receptacle, a watertight secondary packaging • A list of the content placed between the secondary and the outer packaging • A rigid outer packaging of adequate strength for its capacity, weight and intended use. A special packaging Division 6.2 Infectious Substances must be used and assigned to UN2814 or UN2900 and the words of "Suspected Category A Infectious substances" must be shown.

## **TOR9: SCIENTIFIC MEETINGS**

21. Did your laboratory organise scientific meetings related to the pathogen in question on behalf of WOA?

No

22. Did your laboratory participate in scientific meetings related to the pathogen in question on behalf of WOA?

No

## **TOR10: NETWORK WITH WOA REFERENCE LABORATORIES**

23. Did your laboratory exchange information with other WOA Reference Laboratories designated for the same pathogen or disease?

Yes

24. Are you a member of a network of WOA Reference Laboratories designated for the same pathogen?

Not applicable (Only WOA Reference Laboratory designated for the disease)

25. Did you organise or participate in inter-laboratory proficiency tests with WOA Reference Laboratories designated for the same pathogen?

Yes

PURPOSE OF THE PROFICIENCY TESTS: 1	ROLE OF YOUR REFERENCE LABORATORY (ORGANISER/ PARTICIPANT)	NO. PARTICIPANTS	PARTICIPATING WOA REF. LABS/ ORGANISING WOA REF. LAB.
Serology IgG and IgM	Participant	4	Participating WOA RL

26. Did your laboratory collaborate with other WOA Reference Laboratories for the same disease on scientific research projects for the diagnosis or control of the pathogen of interest?

Yes

TITLE OF THE PROJECT OR CONTRACT	SCOPE	NAME(S) OF RELEVANT WOA REFERENCE LABORATORIES
Update of the Code and Terrestrial Manual	Review and update of the WOA Code and Terrestrial Manual	ARC-OVI, Onderstepoort, South Africa
ELISA IgM serological method WAHO lab for RVFV	Exchange on the sensitivity/specificity of ELISA IgM RVFV method	ARC-OVI, Onderstepoort, South Africa

## **TOR11: OTHER INTERLABORATORY PROFICIENCY TESTING**

27. Did your laboratory organise or participate in inter-laboratory proficiency tests with laboratories other than WOA Reference Laboratories for the same pathogen?

No

## **TOR12: EXPERT CONSULTANTS**

28. Did your laboratory place expert consultants at the disposal of WOA?

Yes

KIND OF CONSULTANCY	Location	SUBJECT (FACULTATIVE)
Adhoc meetings online	Adhoc meetings online	Adhoc meetings online

29. Additional comments regarding your report: