

# WOAH Reference Laboratory Reports Activities 2024

This report has been submitted: 30 janvier 2025 12:16

## LABORATORY INFORMATION

|  |   |
|--|---|
| <b>*Name of disease (or topic) for which you are a designated WOAH Reference Laboratory:</b> | Glanders  |
| <b>*Address of laboratory:</b>   | P.O. Box 597 Dubai  |
| <b>*Tel:</b>   | +971-4 337.51.65  |
| <b>*E-mail address:</b>  | cvrl@cvrl.ae  |
| <b>Website:</b>  | www.cvrl.ae   |
| <b>*Name (including Title) of Head of Laboratory (Responsible Official):</b>                 | Priv. Doz. Dr. Dr. habil. Ulrich Wernery                      |
| <b>*Name (including Title and Position) of WOAH Reference Expert:</b>                        | Priv. Doz. Dr. Dr. habil. Ulrich Wernery, Scientific Director |
| <b>*Which of the following defines your laboratory? Check all that apply:</b>                | Semi Governmental   |

## 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 WOAH Manual (Yes/No) | Total number of test performed last year |                 |
|---------------------------|-----------------------------------|--|-----------------|
| Indirect diagnostic tests |                                   | Nationally                               | Internationally |
| CFT                       | Yes                               | 5259                                     | 1754            |
| ELISA                     | Yes                               | 165                                      | 3               |
| Direct diagnostic tests   |                                   | Nationally                               | Internationally |

## TOR2: REFERENCE MATERIAL

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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 |
|---------------------------|-------------------------|-------------------|-------------------------------------|--|---|-----------------------|
| Negative serum control    | CFT/ELISA               | Provided          | 0                                   | 2.5 ml                                   | 1                                       | UNITED KINGDOM,       |
| Positive serum control    | CFT/ELISA               | Provided          | 0                                   | 5.0ml                                    | 1                                       | UNITED KINGDOM,       |

4. Did your laboratory produce vaccines?

No

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

No

## 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?

Yes

| Name of the new test or diagnostic method developed | Description and References (Publication, website, etc.) |
|---|---|
| Indirect ELISA for Glanders from Bioclin, Brazil    | Validation data is available from CVRL                  |

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 |
|---|------------|----------------------------|--|--|
| BAHRAIN   | 2024-01-01 | CFT                        | 564  | 0  |
| BRAZIL  | 2024-01-01 | CFT                        | 0  | 1  |
| EGYPT   | 2024-01-01 | CFT                        | 75   | 0  |
| HONG KONG                                       | 2024-01-01 | CFT                        | 0  | 1  |
|   |            |                            |  |  |

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|              |            |     |     |   |
|--------------|------------|-----|-----|---|
| IRAN         | 2024-01-01 | CFT | 14  | 0 |
| JORDAN       | 2024-01-01 | CFT | 98  | 0 |
| KUWAIT       | 2024-01-01 | CFT | 441 | 0 |
| OMAN         | 2024-01-01 | CFT | 142 | 0 |
| SAUDI ARABIA | 2024-01-01 | CFT | 159 | 0 |
| THAILAND     | 2024-01-01 | CFT | 213 | 0 |
| RUSSIA       | 2024-01-01 | CFT | 12  | 0 |

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

Yes

| Name of the WOAHP Member Country receiving a technical consultancy | Purpose                               | How the advice was provided |
|--|---------------------------------------|-----------------------------|
| BRAZIL   | Validation of Glanders Indirect ELISA | email                       |

## TOR5: COLLABORATIVE SCIENTIFIC AND TECHNICAL STUDIES

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

No

13. In exercising your activities, have you identified any regulatory research needs\* relevant for WOAHP?

No

## 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:

Monitoring disease situation in this region by performing serological investigations on equine sera sent from neighboring countries.

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:

All data produced at CVRL are sent to the authorities at the ministerial level of the UAE and the home countries from where we receive the samples.

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:

0

b) International conferences:

0

c) National conferences:

0

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

0

## TOR7: SCIENTIFIC AND TECHNICAL TRAINING

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

No

## TOR8: QUALITY ASSURANCE

18. Does your laboratory have a Quality Management System?

Yes

| Quality management system adopted | Certificate scan (PDF, JPG, PNG format) |                     |
|-----------------------------------|---|---------------------|
| ISO/IEC 17025:2017                | CVRL-PDF                                | CVRL-IAS.pdf        |
| ISO/IEC 17025:2017                | MBG-Veterinary                          | MBGX-Veterinary.pdf |
| ISO 15189:2012                    | MBG-Medical                             | MBGX-medical.pdf    |

19. Is your quality management system accredited?

Yes

| Test for which your laboratory is accredited | Accreditation body                       |
|--|--|
| Brucella Culture                             | International Accreditation Service, USA |
| CEM- Culture for Taylorella equigenitalis    | International Accreditation Service, USA |
| Brucellosis ELISA, CFT, RBT, SAT             | International Accreditation Service, USA |
| Dourine CFT                                  | International Accreditation Service, USA |
| Glanders CFT                                 | International Accreditation Service, USA |
| West Nile IgM ELISA                          | International Accreditation Service, USA |
| Strangles ELISA                              | International Accreditation Service, USA |

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|   |  |
|---|--|
| AHS ELISA   | International Accreditation Service, USA |
| West Nile ELISA   | International Accreditation Service, USA |
| Theileria equi ELISA  | International Accreditation Service, USA |
| Babesia caballi ELISA   | International Accreditation Service, USA |
| MERS CoV ELISA  | International Accreditation Service, USA |
| Equine infectious Anemia AGID   | International Accreditation Service, USA |
| Virus Neutralisation Test for Equine Viral Arteritis (EVA - VNT)  | International Accreditation Service, USA |
| Influenza A Virus Isolation   | International Accreditation Service, USA |
| Avian paramyxovirus type 1 (APMV-1) Virus Isolation   | International Accreditation Service, USA |
| Indirect Fluorescent Antibody Test (IFAT) for the detection of antibodies to Equine Piroplasmiasis (Theileria equi and Babesia caballi) | International Accreditation Service, USA |
| MERS-Coronavirus (MERS-CoV) Isolation   | International Accreditation Service, USA |
| Fluorescent Antibody Virus Neutralization (FAVN) Test for Rabies  | International Accreditation Service, USA |

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

Yes

The laboratory facility, management practices, and biosecurity procedures are regularly monitored to ensure specific biosafety and laboratory biosecurity at CVRL.

## TOR9: SCIENTIFIC MEETINGS

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

No

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

No

## TOR10: NETWORK WITH WOAHA REFERENCE LABORATORIES

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

No

24. Do you network (collaborate or share information) with other WOAHA Reference Laboratories designated for the same pathogen?

No

25. Did you organise or participate in inter-laboratory proficiency tests with WOAHA Reference Laboratories designated for the same pathogen during the past 2 years?

No

CVRL participated in the interlaboratory proficiency tests conducted by FLI, Germany in 2023.

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

No

## TOR11: OTHER INTERLABORATORY PROFICIENCY TESTING

27. Did your laboratory organise or participate in inter-laboratory proficiency tests with laboratories other than WOAHA Reference Laboratories for the same pathogen during the past 2 years?

Yes

| Purpose for inter-laboratory test comparisons <sup>1</sup> | Role of your reference laboratory (organizer/participant) | No. participating laboratories | Name of the test           | WOAH Member Countries  |
|--|---|--------------------------------|----------------------------|--|
| WOAH international comparison test-<br>Glanders ELISA      | Participate   | 6                              | Glanders Indirect<br>ELISA | AUSTRALIA, BRAZIL, CHINA<br>(PEOPLE'S REP. OF),<br>FRANCE, GERMANY,<br>UNITED ARAB EMIRATES, |

## TOR12: EXPERT CONSULTANTS

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

No

29. Additional comments regarding your report:

No

*In the WOAHA reference laboratory annual report 2025, melioidosis should be added to the disease name. Also, we request to designate CVRL as the reference laboratory for both diseases.*