

OIE Reference Laboratory Reports Activities

Activities in 2021

This report has been submitted : 2022-01-12 10:53:43

| | |
|--|---|
| Name of disease (or topic) for which you are a designated OIE Reference Laboratory: | Bluetongue |
| Address of laboratory: | Ash Road, Pirbright Woking, Surrey, GU24 0NF UNITED KINGDOM |
| Tel.: | +44-1483 23 24 41 |
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| E-mail address: | carrie.batten@pirbright.ac.uk |
| Website: | https://www.pirbright.ac.uk/our-science/vector-borne-viral-diseases/non-vesicular-disease-reference-laboratory |
| Name (including Title) of Head of Laboratory (Responsible Official): | Prof Bryan Charleston, Institute Director |
| Name (including Title and Position) of OIE Reference Expert: | Dr Carrie Batten, Head of non vesicular reference laboratories |
| Which of the following defines your laboratory? Check all that apply: | Other: Research institute |

ToR 1: To use, promote and disseminate diagnostic methods validated according to OIE Standards

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 OIE Manual (Yes/No) | Total number of test performed last year | |
|---------------------------|----------------------------------|--|-----------------|
| | | Nationally | Internationally |
| Indirect diagnostic tests | | Nationally | Internationally |
| C-ELISA | Yes | 3271 | 9 |
| Direct diagnostic tests | | Nationally | Internationally |
| Real-Time RT-PCR | Yes | 11562 | 9 |
| Virus Isolation | Yes | 1 | 3 |

**ToR 2: To develop reference material in accordance with OIE requirements, and implement and promote the application of OIE Standards.
To store and distribute to national laboratories biological reference products and any other reagents used in the diagnosis and control of the designated pathogens or disease.**

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

No

3. Did your laboratory supply standard reference reagents (non OIE-approved) and/or other diagnostic reagents to OIE Member Countries?

Yes

| Type of reagent available | Related diagnostic test | Produced/ provide | Amount supplied nationally (ml, mg) | Amount supplied internationally (ml, mg) | No. of recipient OIE Member Countries | Region of recipients |
|--|-------------------------|-------------------|-------------------------------------|--|---------------------------------------|--|
| EHDV (BTV) ILC panel 8 x 300ul sera and 8 x 1ml EDTA | ELISA/PCR | Provide | 10.4ml | 31.2ml | 4 | <input type="checkbox"/> Africa <input checked="" type="checkbox"/> Americas <input type="checkbox"/> Asia and Pacific <input checked="" type="checkbox"/> Europe <input type="checkbox"/> Middle East |
| 7 various BTV strains | Cell culture | Provide | 0 | 14 ml total | 1 | <input type="checkbox"/> Africa <input type="checkbox"/> Americas <input type="checkbox"/> Asia and Pacific <input checked="" type="checkbox"/> Europe <input type="checkbox"/> Middle East |

4. Did your laboratory produce vaccines?

No

5. Did your laboratory supply vaccines to OIE Member Countries?

No

ToR 3: To develop, standardise and validate, according to OIE Standards, new procedures for diagnosis and control of the designated pathogens or diseases

6. Did your laboratory develop new diagnostic methods validated according to OIE Standards for the designated pathogen or disease?

No

7. Did your laboratory develop new vaccines according to OIE Standards for the designated pathogen or disease?

No

ToR 4: To provide diagnostic testing facilities, and, where appropriate, scientific and technical advice on disease control measures to OIE Member Countries

8. Did your laboratory carry out diagnostic testing for other OIE Member Countries?

No

9. Did your laboratory provide expert advice in technical consultancies on the request of an OIE Member Country?

Yes

| Name of the OIE Member Country receiving a technical consultancy | Purpose | How the advice was provided |
|--|-------------------------------------|-----------------------------|
| CANADA | Virus isolation of BTV/EHDV methods | Email and virtual meeting |
| MEXICO | advice on BTV control material | Email |
| FRANCE | Discussions regarding ILC | Email |

ToR 5: To carry out and/or coordinate scientific and technical studies in collaboration with other laboratories, centres or organisations

10. Did your laboratory participate in international scientific studies in collaboration with OIE Member Countries other than the own?

Yes

Bluetongue - Carrie Batten - united kingdom

| Title of the study | Duration | Purpose of the study | Partners (Institutions) | OIE Member Countries involved other than your country |
|---|-----------|---|--|---|
| PALE-Blu: Understanding pathogen, livestock, environment interactions involving bluetongue virus | 5 Years | Full-genome sequence analyses will increase the accuracy of BTV strain distribution maps, to identify pathways and mechanisms for spread into and within Europe, as well as appropriate prevention strategies. PALE-Blu will analyse the genetic connectivity of Culicoides vector populations in different regions, as well as the movements of individual BTV lineages and genes. Together with reverse genetics technologies and infection/replication studies in new Culicoides cell lines, or adults from different Culicoides species, this will elucidate the genetic basis for geographic localisation/movement of BTV strains and serotypes. We will analyse differences in saliva proteins from Culicoides species, their ability to modify the BTV surface proteins (proteases) and effects on efficiency of transmission (in both directions) between vertebrate hosts and insect-vectors. These studies will provide a better understanding of incursion risks for different BTV strains, supporting effective control strategies. PALEBLU will explore more effective and crossserotype subunitvaccines that are DIVA assay compatible and generate a stronger immune response from a single inoculation. We will also explore the potential for use of antiviral agents to induce immediate protection post vaccination. More effective diagnostic systems to better detect mixed infections will also be developed by multiplexing existing or novel diagnostic assay systems | University of Nottingham - UK; AGENCE NATIONALE DE SECURITE SANITAIRE DE L'ALIMENTATION, DE L'ENVIRONNEMENT ET DU TRAVAIL - France; CENTRE DE COOPERATION INTERNATIONALE EN RECHERCHE AGRONOMIQUE POUR LE DEVELOPPEMENT - France; ISTITUTO ZOOPROFILATTICO SPERIMENTALE DELL'ABRUZZO E DEL MOLISE "G. CAPORALE" DI TERAMO - Italy; FRIEDRICH LOEFFLER INSTITUT - BUNDESFORSCHUNGSINSTITUT FUER TIERGESUNDHEIT - Germany; ENVIRONMENTAL RESEARCH GROUP OXFORD LIMITED - UK; UNIVERSITE LIBRE DE BRUXELLES - Belgium; INSTITUTO NACIONAL DE INVESTIGACION Y TECNOLOGIA AGRARIA Y ALIMENTARIA - Spain; STICHTING DIENST LANDBOUWKUNDIG ONDERZOEK - Netherlands; UNIVERSITY OF GLASGOW - UK; KIMRON VETERINARY INSTITUTE - Israel; UNIVERSIDAD COMPLUTENSE DE MADRID - Spain; STATENS VETERINAERMEDICINSKA ANSTALT - Sweden; KAFKAS UNIVERSITESI - Turkey; INSTITUT AGRONOMIQUE ET VETERINAIRE HASSAN II - Morocco; THE PIRBRIGHT INSTITUTE LBG - UK; International Livestock Research Institute - Kenya; INSTITUT SENEGALAIS DE RECHERCHES AGRICOLES - Kenya; INSTITUT PASTEUR DE TUNIS - Tunisia | BELGIUM FRANCE GERMANY ISRAEL ITALY KENYA MOROCCO SENEGAL SPAIN SWEDEN THAILAND THE NETHERLANDS TUNISIA TURKEY UNITED KINGDOM |
| GNAT work | 3 years | Understanding the biology of vectors | London School Hygiene and Tropical Medicine, Universidade Federal do Minas Gerais, ICCDR, B | BANGLADESH BRAZIL UNITED KINGDOM |
| INFRAVEC 2 | 4 years | Harmonisation of vector competence studies | Institut Pasteur, Centro Agricultura E Ambiente Giorgio Nicoli SRL, Cirad, Ministere de la Sante, EMBL, FORTH, Imperial College, IRD, LSTM, Max Planck, POLO GGB, Radboud University, Tropiq Health Sciences, USTTB, University of Glasgow, University of Karlova, University of Zurich, Wageningen University | BURKINA FASO CZECH REPUBLIC FRANCE GERMANY GREECE ITALY MALI NEW CALEDONIA SENEGAL SWITZERLAND THE NETHERLANDS UNITED KINGDOM |
| Sharing of viral communities between individuals and groups of macaques at the wildlife-livestock-human interface | 6 years | Investigation into virus diversity in Macques | University of Surrey, UK University of Columbia/UC Davis, USA plus contacts in Bangladesh | BANGLADESH UNITED KINGDOM UNITED STATES OF AMERICA |
| Influence of virus and host factors on the vector competence of Culicoides biting midges to Bluetongue virus | 3.5 years | Investigate the influence of virus and host factors on the vector competence of Culicoides biting midges to Bluetongue virus | FLI, Germany | GERMANY |

ToR 6: To collect, process, analyse, publish and disseminate epizootiological data relevant to the designated pathogens or diseases

11. Did your Laboratory collect epizootiological data relevant to international disease control?

Yes

If the answer is yes, please provide details of the data collected:

We characterized the BTV-4 strain from North Macedonia and published the genome sequence. As part of an ongoing project we are sequencing historical strains of BTV and related orbiviruses for future publication.

12. Did your laboratory disseminate epizootiological data that had been processed and analysed?

Yes

If the answer is yes, please provide details of the data collected:

John Flannery, Simon King, Paulina Rajko-Nenow, Zagorka Popova, Kiril Krstevski, Igor Djadjovski and Carrie Batten (2021). Re-emergence of BTV serotype 4 in North Macedonia, July 2020. *Transboundary and emerging diseases*, 68: 220-223. DOI: 10.1111/tbed.13900

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

a) Articles published in peer-reviewed journals: 4

Development of real-time RT-qPCR assays for the typing of two novel bluetongue virus genotypes derived from sheeppox vaccine (2021). Simon King, John Flannery; Carrie Batten; Paulina Rajko-Nenow. *Journal of virological methods*, 298. <https://doi.org/10.1016/j.jviromet.2021.114288>

Serological cross-reactions between expressed VP2 proteins from different bluetongue virus serotypes (2021). Petra C. Fay, Fauziah Mohd Jaafar, Carrie Batten, Houssam Attoui, Keith Saunders, George Lomonosoff, Elizabeth Reid, Daniel Horton, Sushila Maan, David Haig, Janet Daly, Peter P. C. Mertens. *Viruses*, 13(8), 1455; <https://doi.org/10.3390/v13081455>.

Identification of a BTV-Strain-Specific Single Gene That Increases *Culicoides* Vector Infection Rate (2021). Honorata Ropiak, Simon King, Marc Guimera, Kerry Newbrook, Gillian Pullinger, Hannah Brown, John Flannery, Simon Gubbins, Carrie Batten, Paulina Rajko-Nenow and Karin Darpel. *Viruses* 2021, 13(9), 1781; <https://doi.org/10.3390/v13091781>

Development of a novel loop mediated isothermal amplification assay (LAMP) for the rapid detection of epizootic haemorrhagic disease virus (2021). Paulina Rajko-Nenow, Emma L A Howson, Duncan Clark, Natasha Hilton, Aruna Ambagala, Nicholas Svitek, John Flannery and Carrie Batten. *Viruses* 2021, 13, 2187. <https://doi.org/10.3390/v13112187>

b) International conferences: 0

c) National conferences: 0

d) Other:

(Provide website address or link to appropriate information) 0

ToR 7: To provide scientific and technical training for personnel from OIE Member Countries**To recommend the prescribed and alternative tests or vaccines as OIE Standards**

14. Did your laboratory provide scientific and technical training to laboratory personnel from other OIE Member Countries?

No

ToR 8: To maintain a system of quality assurance, biosafety and biosecurity relevant for the pathogen and the disease concerned

15. Does your laboratory have a Quality Management System?

Yes

| Quality management system adopted | Certificate scan (PDF, JPG, PNG format) |
|-----------------------------------|---|
| ISO/IEC17025 | UKAS Cer 2021.pdf |

16. Is your quality management system accredited?

Yes

| Test for which your laboratory is accredited | Accreditation body |
|--|--------------------|
| Real-time RT-PCR | UKAS |
| C-ELISA | UKAS |
| Virus isolation | UKAS |

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

Yes

(See *Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, Chapter 1.1.4*)

ToR 9: To organise and participate in scientific meetings on behalf of the OIE

18. Did your laboratory organise scientific meetings on behalf of the OIE?

No

19. Did your laboratory participate in scientific meetings on behalf of the OIE?

No

ToR 10: To establish and maintain a network with other OIE Reference Laboratories designated for the same pathogen or disease and organise regular inter-laboratory proficiency testing to ensure comparability of results

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

Yes

21. Was your laboratory involved in maintaining a network with OIE Reference Laboratories designated for the same pathogen or disease by organising or participating in proficiency tests?

Yes

| Purpose of the proficiency tests: ¹ | Role of your Reference Laboratory (organiser/participant) | No. participants | Participating OIE Ref. Labs/ organising OIE Ref. Lab. |
|--|---|------------------|---|
| Harmonization of diagnostic tests | Participant | 18 | Organiser: IZS, Italy |
| Harmonisation of ELISA and PCR tests for BTV (This years also includes a VI panel) | Participant | 49 | Participants: IZS, Italy and OVI, South Africa |

¹ validation of a diagnostic protocol: specify the test; quality control of vaccines: specify the vaccine type, etc.

22. Did your laboratory collaborate with other OIE 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 OIE Reference Laboratories |
|--|---|--|
| PALE-Blu: Understanding pathogen, livestock, environment interactions involving bluetongue virus | <p>Full-genome sequence analyses will increase the accuracy of BTV strain distribution maps, to identify pathways and mechanisms for spread into and within Europe, as well as appropriate prevention strategies. PALE-Blu will analyse the genetic connectivity of Culicoides vector populations in different regions, as well as the movements of individual BTV lineages and genes. Together with reverse genetics technologies and infection/replication studies in new Culicoides cell lines, or adults from different Culicoides species, this will elucidate the genetic basis for geographic localisation/movement of BTV strains and serotypes. We will analyse differences in saliva proteins from Culicoides species, their ability to modify the BTV surface proteins (proteases) and effects on efficiency of transmission (in both directions) between vertebrate hosts and insect-vectors. These studies will provide a better understanding of incursion risks for different BTV strains, supporting effective control strategies. PALEBLU will explore more effective and cross-serotype subunit vaccines that are DIVA assay compatible and generate a stronger immune response from a single inoculation. We will also explore the potential for use of antiviral agents to induce immediate protection post vaccination. More effective diagnostic systems to better detect mixed infections will also be developed by multiplexing existing or</p> | IZS, Italy |

ToR 11: To organise inter-laboratory proficiency testing with laboratories other than OIE Reference Laboratories for the same pathogens and diseases to ensure equivalence of results

23. Did your laboratory organise or participate in inter-laboratory proficiency tests with laboratories other than OIE Reference Laboratories for the same disease?

Yes

Note: See Interlaboratory test comparisons in: Laboratory Proficiency Testing at: <http://www.oie.int/en/our-scientific-expertise/reference-laboratories/proficiency-testing> see point 1.3

| Purpose for inter-laboratory test comparisons ¹ | No. participating laboratories | Region(s) of participating OIE Member Countries |
|--|--------------------------------|---|
| Harmonisation of ELISA and PCR tests for BTV (This years also includes a VI panel) | 49 | <input checked="" type="checkbox"/> Africa <input checked="" type="checkbox"/> Americas <input checked="" type="checkbox"/> Asia and Pacific <input checked="" type="checkbox"/> Europe <input checked="" type="checkbox"/> Middle East |

ToR 12: To place expert consultants at the disposal of the OIE

24. Did your laboratory place expert consultants at the disposal of the OIE?

No

25. Additional comments regarding your report: