

OIE Reference Laboratory Reports Activities

Activities in 2020

This report has been submitted : 2021-01-20 18:52:17

Name of disease (or topic) for which you are a designated OIE Reference Laboratory:	Chronic wasting disease
Address of laboratory:	3851 Fallowfield Road P.O. Box 11300 Station H, Nepean Ontario K2H 8P9 CANADA
Tel.:	+1-343 212 0272
Fax:	+1-343 212 0217
E-mail address:	gordon.mitchell2@canada.ca
Website:	www.inspection.gc.ca
Name (including Title) of Head of Laboratory (Responsible Official):	Dr. Abed Harchaoui, Executive Director, Ontario Laboratories Network, Canadian Food Inspection Agency
Name (including Title and Position) of OIE Reference Expert:	Dr. Gordon Mitchell, Head, National and OIE Reference Laboratory for Scrapie and CWD
Which of the following defines your laboratory? Check all that apply:	Governmental

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			
0	0	0	0
Direct diagnostic tests			
PrP ELISA		3325	0
PrP Immunohistochemistry		2401	0
PrP Western Blot		264	0
PRNP Genotyping		2093	0

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
Tissue Homogenates	PrP ELISA / Western Blot	Provide	Multiple	Multiple	3	<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

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
COLOMBIA	TSE pathology and proficiency testing	Email
BRAZIL	TSE diagnostic testing	Email
UNITED STATES OF AMERICA	Surveillance testing, genetic susceptibility	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

Title of the study	Duration	Purpose of the study	Partners (Institutions)	OIE Member Countries involved other than your country
Investigating cross-species transmission of CWD	Ongoing	Characterize transmission of Korean and Canadian CWD Isolates	Animal and Plant Quarantine Agency	KOREA (REP. OF)
Characterization of CWD Isolates	Ongoing	Investigating variability between CWD isolates in Europe and North America	Norwegian Veterinary Institute, Colorado State University	FRANCE GERMANY ITALY NORWAY UNITED STATES OF AMERICA

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:
All positive cases are characterized based on data combined from confirmatory test results.

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:

Data on positive cases is collated and communicated to regulatory agencies.

**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

Nonno R, Di Bari MA, Pirisinu L, D'Agostino C, Vanni I, Chiappini B, Marcon S, Riccardi G, Tran L, Vikøren T, Våge J, Madslie K, Mitchell G, Telling GC, Benestad SL, Agrimi U. Studies in bank voles reveal strain differences between chronic wasting disease prions from norway and north america. Proc Natl Acad Sci U S A. 2020;117(49):31417-26.

Arifin MI, Staskevicius A, Shim SY, Huang Y-, Fenton H, McLoughlin PD, Mitchell G, Cullingham CI, Gilch S. Large-scale prion protein genotyping in canadian caribou populations and potential impact on chronic wasting disease susceptibility. Mol Ecol. 2020;29(20):3830-40.

Sohn HJ, Mitchell G, Lee YH, Kim HJ, Park KJ, Staskevicius A, Walther I, Soutyrine A, Balachandran A. Experimental oral transmission of chronic wasting disease to sika deer (cervus nippon). Prion. 2020;14(1):271-7.

Haley NJ, Donner R, Henderson DM, Tennant J, Hoover EA, Manca M, Caughey B, Kondru N, Manne S, Kanthasamay A, Hannaoui S, Chang SC, Gilch S, Smiley S, Mitchell G, Lehmkuhl AD, Thomsen BV. Cross-validation of the RT-QulC assay for the antemortem detection of chronic wasting disease in elk. Prion. 2020;14(1):47-55.

b) International conferences: 1

Sohn HJ, Park KJ, Mitchell G, Hwang JY, Park HC, Kang HE. Distribution of PrPCWD in tissues of CWD-infected sika deer analyzed by RT-QulC. Annual meeting of the International Union of Microbiological Societies, Daejeon, Korea, 2020.

c) National conferences: 0

d) Other:

(Provide website address or link to appropriate information) 1

Mitchell G. Chronic Wasting Disease Surveillance in Canada. One Health Workshop Series, University of Calgary, Canada. Virtual presentation and panel discussion. October 7, 2020.

**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 17025	Accreditation Certificate.pdf

16. Is your quality management system accredited?

Yes

Test for which your laboratory is accredited	Accreditation body
PrP Immunohistochemistry	Standards Council of Canada (SCC)
PrP ELISA	SCC
PrP Western blot	SCC
PRNP Genotyping	SCC

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?

No

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
Investigating cross-species transmission of CWD	Characterizing transmission of Korean and Canadian CWD isolates	Animal and Plant Quarantine Agency, Republic of Korea
Collaborative framework concerning CWD reference laboratories	Advancing strategies to detect, control and characterize CWD	Norwegian Veterinary Institute

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
PrP Immunohistochemistry proficiency	3	<input type="checkbox"/> Africa <input checked="" type="checkbox"/> Americas <input type="checkbox"/> Asia and Pacific <input type="checkbox"/> Europe <input type="checkbox"/> Middle East
PrP ELISA proficiency	6	<input type="checkbox"/> Africa <input checked="" type="checkbox"/> Americas <input type="checkbox"/> Asia and Pacific <input type="checkbox"/> Europe <input type="checkbox"/> Middle East
PrP Western blot proficiency	1	<input type="checkbox"/> Africa <input checked="" type="checkbox"/> Americas <input type="checkbox"/> Asia and Pacific <input type="checkbox"/> Europe <input type="checkbox"/> Middle East
PRNP Genotyping proficiency	3	<input type="checkbox"/> Africa <input checked="" type="checkbox"/> Americas <input type="checkbox"/> Asia and Pacific <input type="checkbox"/> Europe <input 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: