OIE Reference Laboratory Reports ActivitiesActivities in 2016

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Name of disease (or topic) for which you are a designated OIE Reference Laboratory:	Myxomatosis
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Website:	http://www.izsler.it/izs_home_page/servizi/00003931_OIE_Reference_Laboratory_for_Myxomatosis_of_Rabbits.html
Name (including Title) of Head of Laboratory (Responsible Official):	prof. Stefano Cinotti Genaral Director
Name (including Title and Position) of OIE Reference Expert:	Dr. Antonio Lavazza, DVM Deputy head Virology Unit
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 yea	
Indirect diagnostic tests		Nationally	Internationally
c-ELISA	yes	33	219
Direct diagnostic tests		Nationally	Internationally
negative staining EM	yes	29	0
cell culture isolation	yes	1	0
immunoperoxidase	yes	0	0
PCR	yes	26	0
Immunofluorescence	yes	0	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?

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
Positive control for Myxoma virus	PCR	provided	0	1ml	1	□Africa □Americas □Asia and Pacific □Europe □Middle East
Panel of 11 positive field sera at known titre for test validation	serological ELISA	provided	0	11 ml	1	□Africa □Americas □Asia and Pacific □Europe □Middle East
DNA positive control for Myxoma virus	PCR	provided	0	1 ml	1	□Africa □Americas □Asia and Pacific □Europe ⊠Middle East

4	Did v	vour	laboratory	nroduce	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?

Yes

Name of OIE Member Country seeking assistance	Date (month)	No. samples received for provision of diagnostic support	No. samples received for provision of confirmatory diagnoses
BULGARIA	May	219	0

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
VIETNAM	To give indication on the procedures of cell isolation of myxomavirus	By sending multiple email containing the protocols and answers to specific questions
SWEDEN	Consultation on resistence of Myxovirus particles in the environment	By sending known data and literature available by 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?

Title of the study	Duration	Purpose of the study	Partners (Institutions)	OIE Member Countries involved other than your country
study and examination of rabbit and brown hare diseases	3 years	Virological and serological studies on wild and domestic rabbit and brown hare diseases in Bulgaria	University of Forestry, Faculty of Veterinary Medicine	BULGARIA

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

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

Yes

- 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: 0
- b) International conferences: 0
- c) National conferences: 1

BADAGLIACCA PIETRO, CAVADINI PATRIZIA, CAPUCCI LORENZO, VANTINI CASSANDRA, ROMEO GIANFRANCO, SAVINI GIOVANNI, LAVAZZA ANTONIO. Utilizzo di tecniche d'indagine molecolare su un ceppo di myxomavirus evidenziato in un coniglio mummificato in corso di indagine forense. Atti 6° Workshop di Virologia Veterinaria. 13-14 Ottobre 2016 Torino. pag. 14.

d) Other:

(Provide website address or link to appropriate information) 2

- 1. LAVAZZA A., COOKE B.D. " Lagomorphs Diseases Chapter " in: Smith A.T., Johnstione C.H. Alves P., Hackländer
- K. "Lagomorphs of the world" Johns Hopkins University Press, Baltimore, Maryland (USA)
- 2. Web-site link:

http://www.izsler.it/izs_home_page/servizi/00003931_OIE_Reference_Laboratory_for_Myxomatosis_of_Rabbits.html

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?

- a) Technical visits: 1
- b) Seminars: 0
- c) Hands-on training courses: 2
- d) Internships (>1 month): 0

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
С	Germany/USA	1
a, c	United Arabian Emirates (UAE)	2

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 certified according to an International Standard?

Yes

Quality management system adopted	Certificate scan (PDF, JPG, PNG format)
UNI CEI EN ISO/IEC 17025	CERTIFICATO ACCREDITAMENTO 20150928.pdf

16. Is your laboratory accredited by an international accreditation body?

Yes

Test for which your laboratory is accredited	Accreditation body
PCR	ILAC MRA, ACCREDIA
Serological Competitive ELISA	ILAC MRA, ACCREDIA
Immunohistochemistry	ILAC MRA, ACCREDIA
EM methods	ILAC MRA, ACCREDIA

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?

Yes

Title of event	Date (mm/yy)	Location	Role (speaker, presenting poster, short communications)	Title of the work presented
84°OIE GENERAL SESSION	May 2016	Paris (France)	Member of the Italian delegation	-
5° World Lagomorph Conefrence	July 2016	Turlock (CA, USA)		
12° Conference of the European Wildlife Disease Association (EWDA)	August 2016	Berlin (Germany)		

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?

Not applicable (Only OIE Reference Lab. designated for disease)

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?

Not applicable (Only OIE Reference Lab. designated for disease)

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?

Not applicable (Only OIE Reference Lab. designated for disease)

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?

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
Use of provided internal reference sera for the re- evaluation of an ELISA kit for detection of antibodies specific of Mixomatosis.	1	□Africa □Americas □Asia and Pacific ⊠Europe □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:

This OIE Reference Lab for Myxomatosis at IZSLER was recognised five years ago. The second year (2013) was mostly dedicated to the revision of the chapter of the OIE Terrestrial Manual that was then approved by the Standard Commission and adopted on May 2014. During the last three years we increased the number of informal contacts with laboratories of member countries for supplying PCR methods and reference samples, and for performing diagnostic tests.