

# OIE Reference Laboratory Reports Activities

## *Activities in 2019*

**This report has been submitted : 2020-01-15 19:47:57**

|  |  |
|--|--|
| <b>Name of disease (or topic) for which you are a designated OIE Reference Laboratory:</b> | Highly and low pathogenic avian influenza              |
| <b>Address of laboratory:</b>  | Südufer 10 D-17493 Greifswald Insel Riems<br>GERMANY   |
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| <b>Name (including Title) of Head of Laboratory (Responsible Official):</b>                | Prof. Dr. Martin Beer, director                        |
| <b>Name (including Title and Position) of OIE Reference Expert:</b>                        | Prof. Dr. Timm C. Harder, head AI reference laboratory |
| <b>Which of the following defines your laboratory? Check all that apply:</b>               | Governmental<br>Research                               |

**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 |
| ELISA                     | Yes                              | 670                                      | 0               |
| HI                        | Yes                              | 70                                       | 0               |
| Direct diagnostic tests   |                                  | Nationally                               | Internationally |
| Virus isolation           | Yes                              | 54                                       | 42              |
| RT-qPCR                   | Yes                              | 2300                                     | 3800            |
| RT-PCR                    | Yes                              | 120                                      | 550             |
| Antigen-ELISA             | Yes                              | 25                                       | 0               |
| Sanger sequencing         | Yes                              | 80                                       | 590             |
| IVPI                      | Yes                              | 2  | 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  |
|---------------------------|-------------------------|-------------------|-------------------------------------|--|---------------------------------------|---|
| Viral RNA                 | RT-qPCR                 | on demand         | 12                                  | 3  | 2                                     | <input checked="" type="checkbox"/> Africa<br><input type="checkbox"/> Americas<br><input type="checkbox"/> Asia and Pacific<br><input type="checkbox"/> Europe<br><input checked="" type="checkbox"/> Middle East            |
| Immune serum              | ELISA; HI               | archived          | 15                                  | 1  | 1                                     | <input type="checkbox"/> Africa<br><input type="checkbox"/> Americas<br><input type="checkbox"/> Asia and Pacific<br><input type="checkbox"/> Europe<br><input checked="" type="checkbox"/> Middle East                       |
| Primers, probes           | RT-qPCR                 | archived          | 0                                   | 3  |                                       | <input checked="" type="checkbox"/> Africa<br><input type="checkbox"/> Americas<br><input type="checkbox"/> Asia and Pacific<br><input checked="" type="checkbox"/> Europe<br><input checked="" 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?

Yes

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

No

| Name of the new test or diagnostic method or vaccine developed | Description and References (Publication, website, etc.)  |
|--|--|
| Generic Influenza multiplex RT-qPCR                            | Simultaneously detects RNA of Influenza Virus types A, B, C and D; Henritzi et al., 2019 (PMID: 30264926). |

***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 |
|---|-----------------|--|--|
| EGYPT   | February, April | 25   | 25   |
| BANGLADESH                                    | May             | 500  |  |
| UKRAINE                                       | March           | 20   |  |

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

No

***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                       |
|---|----------|--|--|---|
| Use of stable isotopes for investigating avian Influenza epidemiology | 4 years  | Relate the origin of wild birds by stable isotope Patterns in feathers and overlay with Influenza surveillance data. | Several national avian influenza reference laboratories. | CANADA<br>KOREA (REP. OF)<br>NIGERIA<br>ROMANIA<br>RUSSIA<br>UNITED KINGDOM |

***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: |
|---|
| Active surveillance data on avian viral pathogens in Bangladesh.    |

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: |
|---|
| Scientific publications in peer-reviewed journals                   |

**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: 8

Henritzi D, Hoffmann B, Wacheck S, Pesch S, Herrler G, Beer M, Harder T. A newly developed tetraplex real-time RT-PCR for simultaneous screening of influenza virus types A, B, C and D. *Influenza Other Respir Viruses*. 2019; 13: 71-82. PMID: 30264926.

Scheibner D, Ulrich R, Fatola OI, Graaf A, Gischke M, Salaheldin AH, Harder TC, Veits J, Mettenleiter TC, Abdelwhab EM. Variable impact of the hemagglutinin polybasic cleavage site on virulence and pathogenesis of avian influenza H7N7 virus in chickens, turkeys and ducks. *Sci Rep*. 2019; 9: 11556. PMID: 31399610.

Hassan KE, El-Kady MF, El-Sawah AAA, Luttermann C, Parvin R, Shany S, Beer M, Harder T. Respiratory disease due to mixed viral infections in poultry flocks in Egypt between 2017 and 2018: Upsurge of highly pathogenic avian influenza virus subtype H5N8 since 2018. *Transbound Emerg Dis*. 2019 [Epub ahead of print] PubMed PMID: 31297991.

Parvin R, Begum JA, Chowdhury EH, Islam MR, Beer M, Harder T. Co- subsistence of avian influenza virus subtypes of low and high pathogenicity in Bangladesh: Challenges for diagnosis, risk assessment and control. Sci Rep. 2019; 9: 8306. PMID: 31165743

Moharam I, Razik AAE, Sultan H, Ghezlan M, Meseko C, Franzke K, Harder T, Beer M, Grund C. Investigation of suspected Newcastle disease (ND) outbreaks in Egypt uncovers a high virus velogenic ND virus burden in small-scale holdings and the presence of multiple pathogens. Avian Pathol. 2019; 25:1-10. PMID: 31090444.

Ayim-Akonor M, May J, Krumkamp R, Harder T, Mertens E. Molecular and serological prevalence of influenza A viruses in poultry and poultry farmers in the Ashanti region, Ghana. Infection Ecology & Epidemiology 2019; 9: 1698904.

Graaf A, Henritzi D, Harder T. Influenza bei Schweinen in Deutschland und Europa: Neue Herausforderungen für Diagnose und Prävention. Tierärztl Umschau. 2019; 74: xx-xx.

Pohlmann A, Hoffmann D, Grund C, Koethe S, Hüsey D, Meier SM, King J, Schinköthe J, Ulrich R, Harder T, Beer M. Genetic Characterization and Zoonotic Potential of Highly Pathogenic Avian Influenza Virus A(H5N6/H5N5), Germany, 2017-2018. Emerg Infect Dis. 2019; 25: 1973-1976. PMID: 31538926

b) International conferences: 4

Congress of the International Association for Biological Standardization, Wiesbaden, April 2019: Invited presentation.

Annual EU-reference laboratory Meeting for AI and ND: Two scientific presentations, Padova, Italy

Continuing education for swine practioners, invited presentation, Gent, Belgium

International Meeting on Veterinary Virus diagnostics, co-authored three presentations, Moderation, Greifswald, Germany

c) National conferences: 2

Annual Meeting of the German Society for Virology, three poster presentation, Düsseldorf, Germany

Continuing education for swine practioners, invited presentation, Muenster, Germany

d) Other:

(Provide website address or link to appropriate information) 12

Direct e-Mail contacts.

**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?

Yes

a) Technical visits: 2

b) Seminars: 0

c) Hands-on training courses: 0

d) Internships (>1 month): 2

| 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 |
|--|---|---|
| a  | Kenia   | 1   |
| a  | Egypt   | 1   |
| d  | Egypt   | 1   |
| d  | Bangladesh  | 1   |

**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                         | Akkreditierungsurkunde_FLI_Riems-Jena-2015.pdf |

16. Is your quality management system accredited?

Yes

| Test for which your laboratory is accredited | Accreditation body |
|--|--------------------|
| Virus isolation                              | DAKKS              |
| Antigen detection                            | DAKKS              |
| Antibody detection                           | DAKKS              |
| RNA detection                                | DAKKS              |
| Nucleic acid sequencing                      | DAKKS              |

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: <sup>1</sup> | Role of your Reference Laboratory (organiser/participant) | No. participants | Participating OIE Ref. Labs/organising OIE Ref. Lab. |
|--|---|------------------|--|
| Antibody detection                             | participant   | >40              | EU-RL, Padova; OIE_RL Weybridge                      |
| Antigen characterization                       | participant   | >40              | EU-RL, Padova; OIE_RL Weybridge                      |
| RNA detection                                  | participant   | >40              | EU-RL, Padova; OIE_RL Weybridge; OIE-RL Geelong;     |
| Pathotyping, molecular                         | participant   | >40              | EU-RL, Padova; OIE_RL Weybridge; OIE-RL Geelong      |

<sup>1</sup> 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 |
|----------------------------------|---|--|
| Delta-Flu                        | Advancement of knowledge re avian influenza | Various European RLs.                          |

**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 <sup>1</sup> | No. participating laboratories | Region(s) of participating OIE Member Countries   |
|--|--------------------------------|---|
| Antibody detection   | >30                            | <input type="checkbox"/> Africa<br><input type="checkbox"/> Americas<br><input type="checkbox"/> Asia and Pacific<br><input checked="" type="checkbox"/> Europe<br><input type="checkbox"/> Middle East |
| RNA detection  | >30                            | <input type="checkbox"/> Africa<br><input type="checkbox"/> Americas<br><input type="checkbox"/> Asia and Pacific<br><input checked="" type="checkbox"/> Europe<br><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: