European Union Reference Laboratory for Parasites Istituto Superiore di Sanità

Proficiency Testing for the detection of anti-*Toxoplasma* IgG in ovine serum samples







The aims of the proficiency testing (PT) are



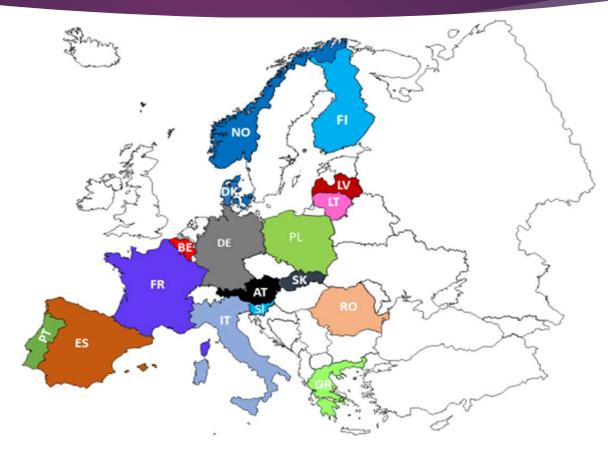
- the laboratories competence in detection of anti-Toxoplasma IgG in ovine serum samples
- the performance of the commercial kits, utilized for the detection of anti-Toxoplasma IgG in ovine







Participating laboratories: Seventeen Labs partecipated to this PT





Eighteenth Annual EURLP Workshop, Rome 16-17 November 2023 Alessandra Ludovisi





Any serological test based on the detection of anti-Toxoplasma IgG could

be used. Each laboratory should choose the test/s routinely used









Materials and Methods 1

The test material forwarded to each laboratory consisted of a panel of 3 serum samples collected from *T. gondii* experimentally infected animals

These serum samples were individually tested for anti-*Toxoplasma* IgG by ID Screen Toxoplasmosis Indirect Multi-species





Test interpretation

Serum sample code	Expected result	S/P
s.1	positive	76%
s.2	positive	>100%
s.3	positive	>100%





Eighteenth Annual EURLP Workshop, Rome 16-17 November 2023

Materials and Methods 2

Serum samples were:

- \succ distributed in 100 µL aliquots,
- > preserved with 1% merthiolate solution,
- Iabelled with a code





Materials and Methods 3

Each laboratory received a link available to NRLs on the EURLP website

to the online survey platform Microsoft Forms, through which PT results

and other details could be submitted







Tests used by the participants

(The more frequently used commercial kits are immunoenzimatic assays)

- IDVETScreen®toxoplasmosis indirect multi-species Ab, 17 labs, coded: A,B,C,D,E,F,G,H,I,J,L,M,N,O,P,Q,R
- Indirect Immunofluoresence antibody test (IFAT), in house methods 1 lab, coded: F (additional test)
- **Toxoreagent Mast Kit**, **2 labs**, coded: G and H (additional test)
- **IDEXX Toxotest 2 labs**, coded: D, J (additional test)
- Pigtype Toxoplasma Ab, Indical Bioscience, 1 lab, coded: A (additional test)
- Immunoblotting full Antigen or P30 Antigen, 1 lab, coded: D (in house, additional test)



Eighteenth Annual EURLP Workshop, Rome 16-17 November 2023



Criteria for the results evaluation:

The participating laboratory had to indicate the positivity or negativity of each sample

 The result of the analysis of each serum sample was reported as

 correctly classified





RESULTS 17/17 NRLs correctly classified as positive all serum samples

Sample code	Expec. results	Lab. code																
Couc	TCSUITS	Α	В	С	D	E	F	G	н	Ι	J	L	M	N	0	Р	Q	R
1	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
2	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
3	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+





Criteria for the final PT evaluation

The result evaluation is qualitative,

IgG titers will be considered as additional information to compare the performance of tests used by participants

Final evaluation is considered as

"positive" if all samples are correctly classified "negative" in all the other cases.





PT evaluation

Laboratory code	N° of samples correctly classified	N° of samples NOT correctly classified	Final evaluation	
Α	3/3	0/3	Positive	
В	3/3	0/3	Positive	
С	3/3	0/3	Positive	
D	3/3	0/3	Positive	
E	3/3	0/3	Positive	
F	3/3	0/3	Positive	
G	3/3	0/3	Positive	
Н	3/3	0/3	Positive	
Ι	3/3	0/3	Positive	
J	3/3	0/3	Positive	
L	3/3	0/3	Positive	
Μ	3/3	0/3	Positive	
Ν	3/3	0/3	Positive	
0	3/3	0/3	Positive	
Р	3/3	0/3	Positive	
Q	3/3	0/3	Positive	
R	3/3	0/3	Positive	

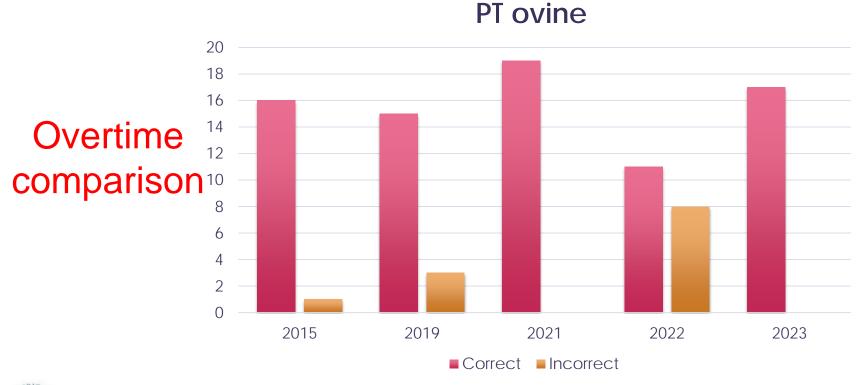






Conclusion

All laboratories (100%) correctly classified all serum samples





Eighteenth Annual EURLP Workshop, Rome 16-17 November 2023



Thanks for your attention!





