



Department of Infectious Parasitic and Immunomediated Diseases
Unit of Gastroenteric and Tissue Parasitic Diseases
European Union Reference Laboratory for Parasites



Final report PT-Tx 1/2015

PT report on "<u>Detection of anti-Toxoplasma IgG in ovine serum</u> <u>samples</u>"

Design

Purpose	Evaluation of laboratories in charge for official control on food		
Scheme type	Single		
Participants	NRL		
N. of participants	Depending on request		
Method	Not regulated		
Test method	Chosen by the participant		
PT items	Matrix	Serum	
	Item	anti-Toxoplasma IgG	
	N. of samples	10 for each participant	
	Distribution	Immediate shipment after preparation	
Subcontracted activities	PT item transport and delivery		
Results evaluation	Qualitative		

Implementation

N. of participants	16	PT items	Serum samples	10
Public laboratories	16			
Private laboratories	0		PT panel composition	6 samples with anti- <i>T.</i> gondi IgG 4 samples without anti- <i>T.</i> gondi IgG
NRL	16		Subcontractor	TNT Express
Shipping dates	16/03/201	5		

PT Provider European Union Reference Laboratory for Parasites Istituto Superiore di Sanità In charge of the PT
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Qualitative results

The PT final evaluation was qualitative only. The PT was considered passed if all positive and all negative samples were correctly identified by the participant,

Laboratory code	N° False positives	N° False negatives	Final evaluation
В	0	0	Passed
С	0	0	Passed
D	0	0	Passed
Е	0	0	Passed
F	0	0	Passed
G	0	0	Passed
Н	0	0	Passed
I	3	0	Failed
J	0	0	Passed
K	0	0	Passed
L	0	0	Passed
M	0	0	Passed
N	0	0	Passed
0	0	0	Passed
Р	0	0	Passed
Q	0	0	Passed

Legend: Laboratories that failed the PT are marked in bold.

Summary of qualitative results:

Total number of PT panels	16
Number of participant laboratories	16
Number of participants that passed the PT	15
Number of participants that failed the PT	1

Comments:

The cause of laboratory failure has been analyzed and can be attributed to the lack of the method (in house developed agglutination test) set up for ovine serum samples.

The Director Dr. E. Pozio

Lance

Istituto Superiore di Sanità



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Date	11/05/2015

Notes:

- 1. To guarantee confidentiality, participant laboratories are identified by alphanumeric codes. PT participant identity is kept confidential and bound by professional secrecy. If PT results have to be provided directly to a competent authority, the organizer shall send a written notice to inform the involved participants.
- 2. The organizer subcontracts PT item transport and delivery to a qualified transportation company.
- 3. Each participating laboratory receive a PT panel according to the PT scheme. Each PT item consists of a fish fillet sandwich spiked or not with live Anisakidae larvae. The homogeneity of PT items is ensured by an accurate control of the number of larvae spiked into each sample (item), made by two operators using a stereo microscope. PT items are stable for 7 days from the date of preparation (corresponding to the shipping date), provided that they are maintained in suitable conditions.
- 4. At the beginning of each year, the organizer draws up a PT program and makes it known by sending an email to the NRLs
- 5. The final report issue of each PT round shows the PT program implementation.

End of the report

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