

Final report PT-03: Tm 1/2023

PT “Identification of *Trichinella* larvae at the species level by a molecular method”

Design

Purpose	Evaluation of laboratories competence in molecular identification of <i>Trichinella</i> larvae species	
Scheme type	Single	
Participants	Public and private, European laboratories	
N. of participants	Depending on request	
Method	not regulated	
Test method	chosen by the participant	
PT items	Matrix	not applicable
	Item	<i>Trichinella</i> spp. larvae in 96% ethanol
	N. of samples	4 vials (10 larvae/each)
	Distribution	Preparation and packaging can be performed before shipment
Subcontracted activities	NA	
Results evaluation	Qualitative	

Implementation

N. of participants	25	PT items	PT panel 4 vials	10 larvae for each of the following species: <i>T. spiralis</i> , <i>T. nativa</i> , <i>T. murrelli</i> e <i>T. nelsoni</i>
Public laboratories	-		PT panel 12 vials	-
Private laboratories	-		Shipping	DHL
NRLs	25			
Shipping dates	March 13, 2023			

PT Provider
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Qualitative results

The PT final evaluation was qualitative only. The PT was considered passed if all isolates or, in case of single larvae at least one of them for each isolate, were correctly identified.

Laboratory code	N. right identification	N. wrong identification	N. missed identification	Final evaluation
NRL50	4	0	0	positive
NRL51	4	0	0	positive
NRL52	2	2	0	negative
NRL55	4	0	0	positive
NRL56	4	0	0	positive
NRL57	4	0	0	positive
NRL59	4	0	0	positive
NRL61	4	0	0	positive
NRL62	4	0	0	positive
NRL63	4	0	0	positive
NRL65	4	0	0	positive
NRL66	4	0	0	positive
NRL70	4	0	0	positive
NRL71	4	0	0	positive
NRL72	4	0	0	positive
NRL73	4	0	0	positive
NRL74	4	0	0	positive
NRL76	4	0	0	positive
NRL77	3	1	0	negative
NRL78	4	0	0	positive
NRL79	4	0	0	positive
NRL80	4	0	0	positive
NRL82	4	0	0	positive
NRL83	4	0	0	positive
NRL84	4	0	0	positive

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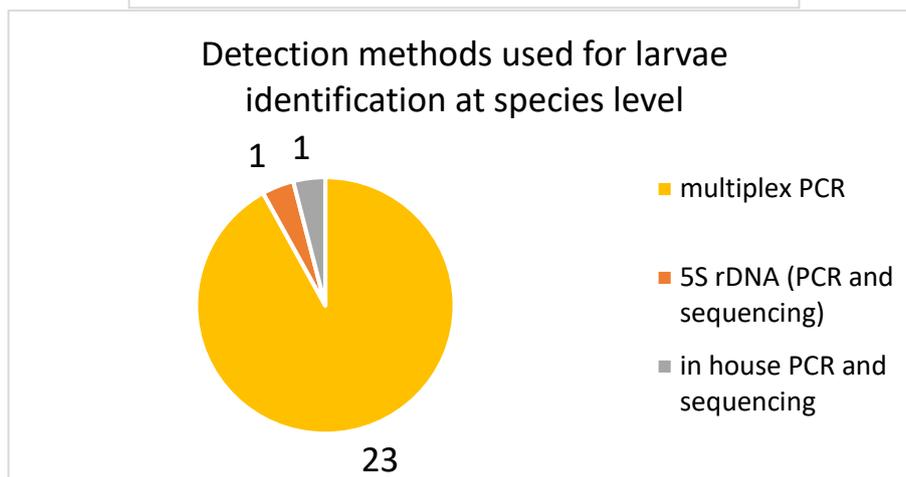
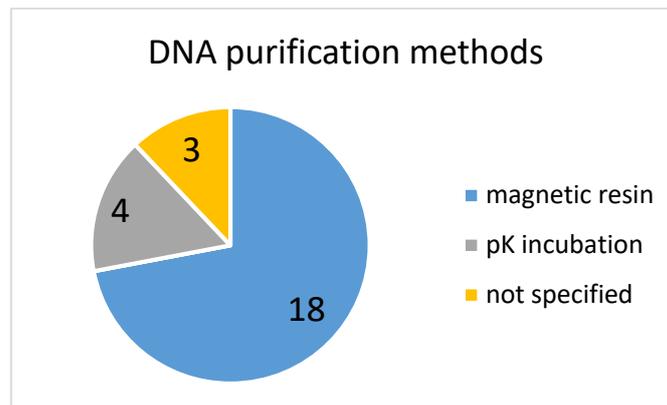
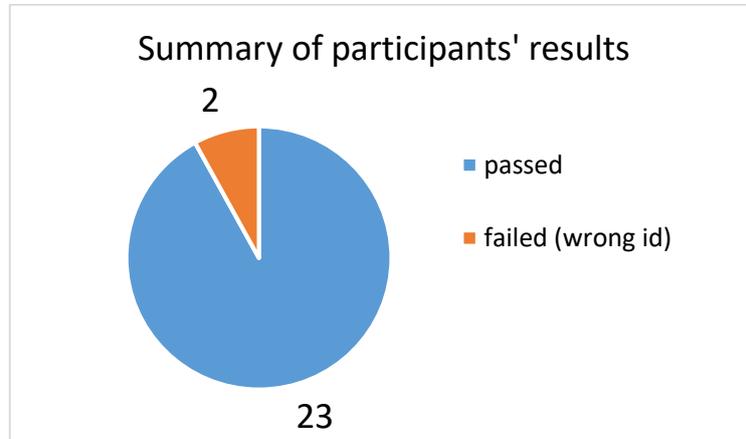


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Graphical summary of PT results



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Overtime comparison of results (last 5 years)

Laboratory code	2019	2020	2021	2022	2023
NRL50	positive	positive	negative	positive	positive
NRL51	positive	positive	positive	positive	positive
NRL52	negative	-	negative	negative	negative
NRL55	positive	positive	positive	positive	positive
NRL56	positive	positive	positive	positive	positive
NRL57	positive	positive	positive	positive	positive
NRL59	positive	positive	positive	positive	positive
NRL61	positive	positive	positive	positive	positive
NRL62	positive	-	positive	positive	positive
NRL63	negative	-	negative	positive	positive
NRL65	positive	positive	negative	positive	positive
NRL66	positive	positive	positive	positive	positive
NRL69	positive	positive	negative	-	positive
NRL70	positive	positive	positive	positive	positive
NRL71	positive	positive	negative	positive	positive
NRL72	positive	positive	positive	positive	positive
NRL73	positive	positive	negative	positive	positive
NRL74	positive	positive	negative	positive	positive
NRL76	positive	-	positive	-	positive
NRL77	positive	positive	positive	positive	negative
NRL78	negative	negative	positive	positive	positive
NRL79	-	-	negative	positive	positive
NRL80	positive	positive	positive	positive	positive
NRL82	positive	positive	positive	positive	positive
NRL83	positive	-	positive	positive	positive
NRL84	-	-	positive	positive	positive

Comments:

The multiplex PCR was the most used method to identify larvae at species level. DNA purification was done mainly by commercial kits based on magnetic beads or silica gel while few participants used a protocol based on incubation in Tris-HCl and pK solutions. Only two participants out of a total of 25 failed because they were not able to correctly identify all the species.

PTP person in charge

Dr. G. Marucci



The Director

Dr. S.M. Cacciò



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Date 28/05/2022

Notes:

1. To guarantee confidentiality, participant laboratories are identified by alphanumeric codes. PT participant identity is kept confidential and bound by professional secrecy. The PTP reserves itself the right to provide the laboratory PT result to the competent authority on request.
2. The organizer designates a qualified company for the transport and delivery of PT items.
3. Each participating laboratory receives a PT panel according to the PT scheme. Each PT item consists of 4 or 12 1.5 ml vials containing four different *Trichinella spp.* The homogeneity of PT items is ensured by an accurate control of the number of larvae spiked into each vial (item), made by two operators using a stereo microscope. PT items are stable for 5 years 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 issued for each PT round shows the PT program implementation.

End of the report

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