

Toxoplasma serology

Distribution: 4381

Laboratory:

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Dispatch Date: 22-Oct-2018

Intended Result		Your Report	Your Score
Specimen 4685	Toxoplasma IgG antibodies negative	Toxoplasma IgG antibodies negative	2
	Toxoplasma IgG avidity: borderline	Toxoplasma IgG avidity: borderline	Not scored
	Toxoplasma IgM antibodies negative	Toxoplasma IgM antibodies negative	2
Specimen 4686	Toxoplasma IgG antibodies positive	Toxoplasma IgG antibodies positive	2
	Toxoplasma IgG avidity: undetermined	Toxoplasma IgG avidity: undetermined	Not scored
	Toxoplasma IgM antibodies positive	Toxoplasma IgM antibodies positive	2
Specimen 4687	Toxoplasma IgG antibodies positive	Toxoplasma IgG antibodies positive	2
	Toxoplasma IgG avidity: low	Toxoplasma IgG avidity: low	Not scored
	Toxoplasma IgM antibodies positive	Toxoplasma IgM antibodies positive	2

Cumulative score information

Total number of specimens sent to you for **UK NEQAS for Toxoplasma serology** over the last 4 distributions is 12 For these distributions specimen numbers 4163 4164 4165 4378 4379 4380 4510 4511 4512 4685 4686 4687 have been sent.

Number of specimens reported too late for analysis (not scored) 0

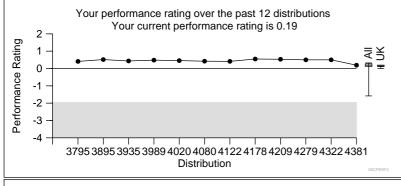
Your cumulative score for these specimens was 48 out of a possible total of 48
The mean score calculated from the reports returned by **UK** laboratories testing the specimen/test combinations you examined was 47.74 with a standard error of

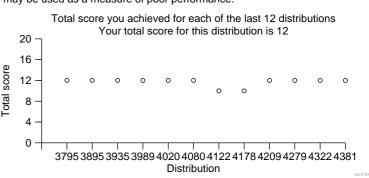
Performance rating

Your performance rating for **UK NEQAS for Toxoplasma serology** (i.e. the number of standard errors by which your cumulative score lies above or below the mean for **UK** laboratories) is 0.19.

A performance rating of more than 1.96 standard errors below the mean indicates possible poor performance. Your performance rating may change if other participants' results are amended.

No score penalty is incurred for non return of reports. However non return of results may be used as a measure of poor performance.





Turn around time: The time taken to report your results was 0 day(s). This information is provided for your own use and does not form part of your performance assessment.

Comments

Participants' results for IgG and IgM analyses were in excellent agreement with the intended results obtained by the Toxoplasma Reference Laboratories.

The figures in the histograms and those in the overall results tables may differ (1) due to exclusion of kits displayed in the histograms resulting in apparently lower numbers of data sets in the histograms or (2) due to participants using more than one kit resulting in higher numbers of data sets in the histograms.

It was noted that many participants did not specify their kit details. Participants are advised to indicate which kit they use so that reports display accurate information. If the kit you use is not specified in the kit list displayed on the web reply form, please email organiser@ukneqasmicro.org.uk to ensure that your kit details are included

At present, the IgG Avidity results are not scored whilst we collect data from different kits. Please note that for sample 4686 there was discrepancy in avidity data results between the different assays during pre-distribution testing (see Teaching notes for further discussions).

Pre-distribution test results are available should you experience a technical failure and wish to discuss the results.

Repeat specimens

For repeat specimens please order using the web form or e-mail organiser@ukneqasmicro.org.uk stating your laboratory identification number, the distribution name and number, and specimen numbers. Participants are asked to request these as soon as the intended results are displayed on the web or on receipt of their

Enquiries: Telephone and written enquiries can be made to Dr. Java Shrivastava: +44(0)20 39081371 E-mail address: organiser@uknegasmicro.org.uk Participants are reminded to quote their Laboratory ID number in all correspondences.

This report was authorised by Professor P L Chiodini, Parasitology Scheme Organiser and Dr Jaya Shrivastava, Parasitology Scheme Manager.

Acknowledgements: We thank colleagues from the Toxoplasma Reference Laboratories and the Department of Medical Microbiology, St George's Hospital, London for the provision of serum samples, for pre- and post-distribution testing and for their expert advice: Drs J.Johnson and D. Padley (London), Dr R. Evans (Inverness), Professor E. Guy, K. Stuart, S.Hadfield (Swansea) and J.Alli-Balogun (St. Georges).

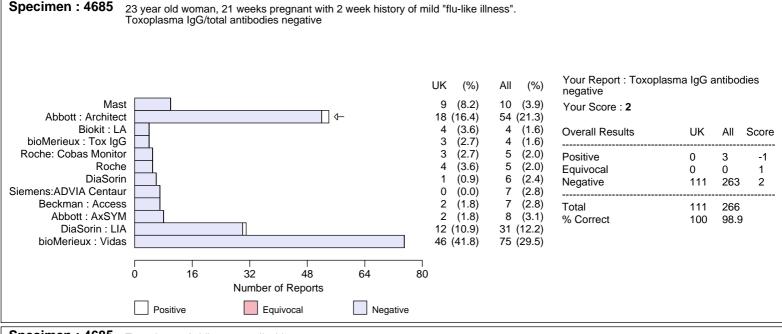


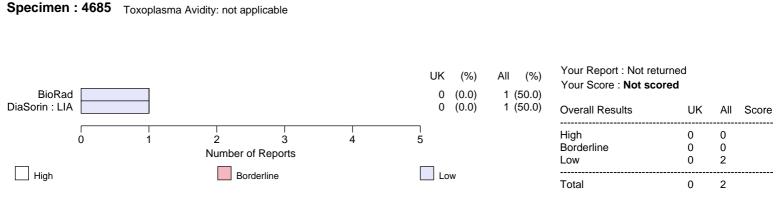


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Three serum specimens were dispatched for the detection of Toxoplasma IgG/total antibodies, IgM antibodies and IgG Avidity. All three specimens were taken from commercially bought single donations.

Specimens returned to the UK NEQAS Parasitology team from 5 randomly selected labs were analysed by a Toxoplasma Reference Lab and gave the intended results



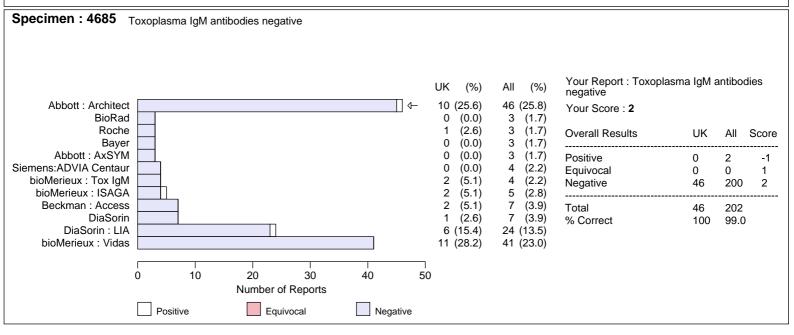




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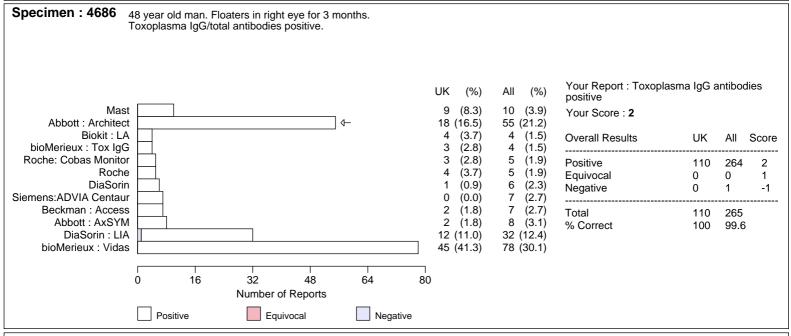


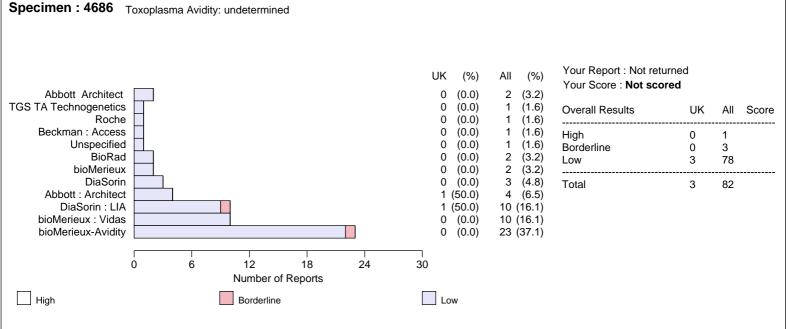


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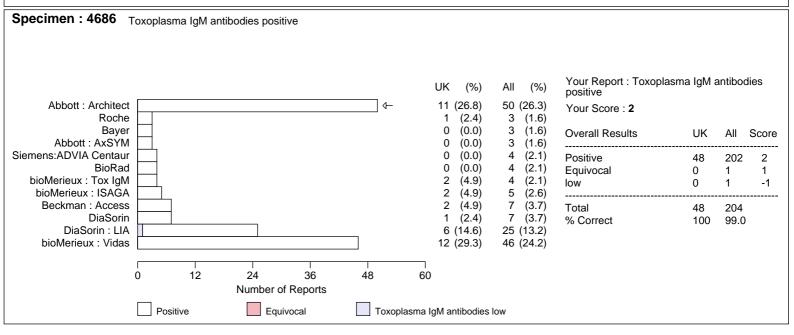




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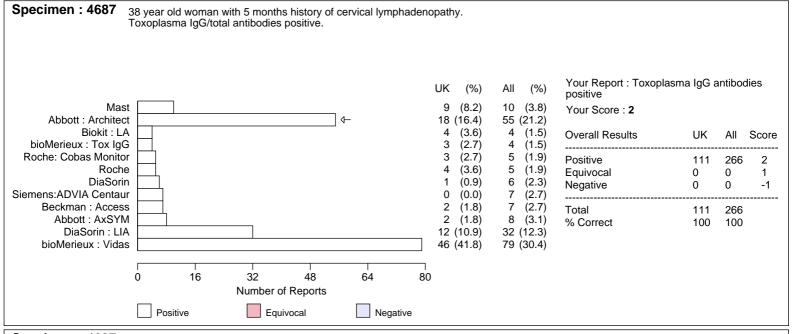
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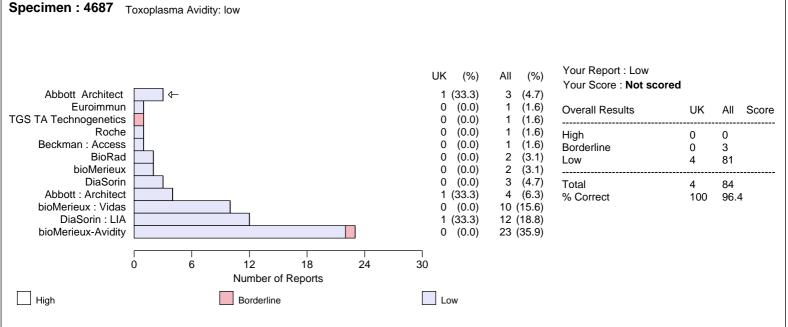
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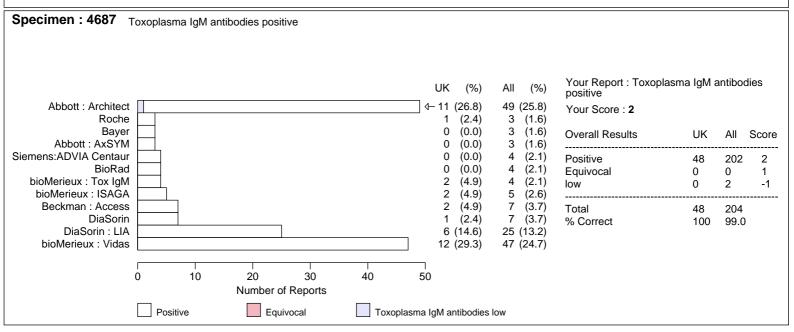




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Specific comments for Distribution 4381

Specimen No 4685

Clinical details 23 year old woman, 21 weeks pregnant with 2- week history of

mild "flu- like illness.

Results Dye test <2 IU/mL

> IgG ELISA (In house) Negative Abbott Architect (IgG) Negative Biokit IgG ELISA Negative

IgG Avidity Not applicable

IgM ELISA (In house) Negative Abbott Architect (IgM) Negative Biomérieux ISAGA Negative

Specific comment No serological evidence of Toxoplasma infection.

Specimen No 4686

Clinical details 48 year old man. Floaters in right eye for 3 months

.Results 500 IU/mL Dye test

> IgG ELISA (In house) Positive Abbott Architect (IgG) Positive Biokit IgG ELISA Positive

IgG Avidity Undetermined

IgM ELISA (In house) Positive Abbott Architect (IgM) Positive Biomérieux ISAGA Positive

Specific comment Serological evidence that Toxoplasma infection has occurred at

some time. Presence of IgM suggests recent infection. These

findings support a diagnosis of ocular toxoplasmosis.

Specimen No 4687

Clinical details 38 year old woman with 5- month history of cervical

lymphadenopathy.





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Results Dye test 250 IU/mL

IgG ELISA (In-House) Positive Abbott Architect (IgG) Positive Positive Biokit IgG ELISA

IgG Avidity Low

IgM ELISA Positive Abbott Architect (IgM) Positive Biomérieux ISAGA Positive

Specific comments Serological evidence that toxoplasmosis infection has occurred at

some time. Presence of IgM suggests recent infection. This is

supported by IgG avidity test result.

USE OF IgG AVIDITY IN ESTIMATING DURATION OF INFECTION

The expected IgG avidity result for Specimen 4686 in the current distribution is classified as 'Undetermined' since the avidity levels obtained in pre-distribution testing were discrepant between the assays used. While one assay consistently returned a 'high' avidity result, the other consistently found the specimen to have "low" avidity. Despite this apparent discrepancy with only one assay finding high avidity, the durations of infection indicated by each assay were not, in fact, mutually exclusive. This is because the high avidity result for the first assay suggests this is not an early infection, while the interpretation for the low avidity result in the second assay was that no significance could be placed on this regarding duration of infection.

One major difficulty in comparing performance of different IgG avidity assays is the lack of standardisation for: Measurement of the IgG avidity index. Definition of low and high avidity levels. and Interpretation of these avidity levels in the context of estimating duration of infection. However, a likely cause for the discrepancy seen in this particular case may be the reportedly poor correlation between low IgG avidity levels and recent infection¹⁻³. Thus, while a high IgG avidity result provides strong evidence to exclude early infection, a low IgG avidity result alone, cannot confirm very recent infection.

Interpretation of all serological results for Specimen 4686 confirms this is a relatively recent infection on the basis of positive IgM and IgG findings, and IgG avidity results are consistent with an infection of greater than 3 months duration. Thus, these findings fit well with the clinical history and support a diagnosis of ocular toxoplasmosis associated with relatively recent infection.

Use of IgG avidity in assessing risk during pregnancy

In most clinical scenarios confirmation of acute or active toxoplasma infection by detection of IgM and IgG, or direct detection of the parasite by PCR, is sufficient to adequately inform clinical In the pregnant mother with acute infection, however, assessing the risk of transmission to the unborn child requires a more precise estimate of when maternal infection was





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acquired in relation to the date of conception. This is because infection acquired before conception presents no significant risk to the pregnancy in the immunocompetent mother while risk of transmission to the unborn child will exist where infection is acquired after conception.

Based on the limitations described above, the use of IgG avidity as a standalone assay is not recommended. However, in investigations during pregnancy, it can be helpful in looking for evidence to exclude recent infection where an expectant mother has been found to be IgG and IgM positive.

References

Jose G. Montoya, Heather B. Huffman, J. S. Remington. Evaluation of the Immunoglobulin G Avidity Test for Diagnosis of Toxoplasmic Lymphadenopathy. Journal of Clinical Microbiology. Oct 2004, 42 (10) 4627-4631.

O. Villard, L. Breit, B. Cimon, J. Franck, H. Fricker-Hidalgo, N. Godineau, S. Houze, L. Paris, H. Pelloux, I. Villena, E. Candolfi the French National Reference Center for Toxoplasmosis Network. Comparison of Four Commercially Available Avidity Tests for Toxoplasma gondii-Specific IgG Antibodies. Clin. Vaccine Immunol. Jan 2013, 20 (2) 197-204.

Findal G, Stray-Pedersen B, Holter EK, Berge T, Jenum PA (2015) Persistent Low Toxoplasma IgG Avidity Is Common in Pregnancy: Experience from Antenatal Testing in Norway. PLoS ONE 10(12): e0145519.



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Suggested algorithm for the use of IgG avidity testing in pregnancy

Pregnant Mother

