False positive latex tests negative by ELISA for toxoplasma IgG

G M SUTEHALL, T G WREGHITT Clinical Microbiology and Public Health Laboratory, Addenbrooke’s Hospital, Cambridge

SUMMARY An enzyme linked immunosorbent assay (ELISA) kit for detecting IgG class antibody to T gondii was compared with the latex agglutination test to determine the specificity as a screening method in 12 patients who had undergone heart transplantation (recrudescence of T gondii infection n = 3, donor acquired infection n = 3; acute cytomegalovirus infection n = 6). The latex agglutination test detected antibodies to primary T gondii infection much earlier in the infection than the ELISA, but the ELISA method was useful for detecting previous infection.

It is concluded that the ELISA technique is more complex to perform than the latex agglutination test but the use of IgM and IgG assays combined could reduce the number of samples sent to the reference laboratory and thus reduce the time taken to obtain a final result.

It has been reported that a latex agglutination test for detecting antibodies to Toxoplasma gondii (Toxoreagent, Mast Diagnostics), which is widely used as a screening test for toxoplasmosis, can give rise to false positive results with serum samples from heart transplant patients with cytomegalovirus (CMV) infection, which seems to be IgM-mediated.\(^1\) A more recent report shows that false positive results have been found with serum samples from people who are not specifically immunocompromised but who also have CMV infection.\(^2\) We examined an enzyme linked immunosorbent assay (ELISA) kit for detecting IgG class antibody to T gondii to determine whether it was a more specific screening test than the latex agglutination test.

Material and methods

Sequential serum samples from heart transplant patients were examined by means of the CAPTIA Toxo-G kit (Mercia Diagnostics, Guildford). Three patients who had antibodies to T gondii positive at the time of transplantation and who subsequently experienced recrudescence of T gondii after transplantation, three patients with primary donor-acquired T gondii infection, and six cases with acute CMV infection in which the toxoplasma latex agglutination test was falsely reactive were studied. All samples were initially tested by the latex agglutination test and the reactive specimens were further tested in the dye test, haemagglutination test, and \(\mu\)-capture IgM ELISA test at the Public Health Laboratory Service Toxoplasma Reference Laboratory at Leeds.

Results

For the cases with recrudescence of T gondii, we found a good correlation between the latex agglutination test titre and the CAPTIA Toxo-G index (figure).

Samples with false reactivity in the latex agglutination test were not positive in the CAPTIA Toxo-G assay (figure) and neither were samples negative in the latex agglutination test.

In one case of primary toxoplasmosis, however, the ELISA did not detect antibody early in the infection and in a second case the ELISA result was only just positive, at 5% over the cut off value at the start of the antibody response, and was not strongly positive until much later. By contrast, the latex agglutination test detected T gondii antibody early in the infection.

Discussion

The CAPTIA Toxo-G assay is useful for detecting previous toxoplasma infection and would be useful as a screening test in this context. The assay is not subject to the type of non-specific reactions encountered when using the Toxoreagent latex agglutination test. This could be because false positive reactions in the latex agglutination test are associated with an IgM-mediated mechanism\(^1\) to which the ELISA is not
susceptible. For use as a test to monitor transplant patients for donor-acquired toxoplasmosis, however, this ELISA would need to be used in conjunction with a suitable assay for detecting *T. gondii* specific IgM.

The ELISA technique is more complex to perform than that of the latex agglutination test but the use of IgM and IgG assays combined could reduce the number of samples to be sent to the reference laboratory and hence the time taken to obtain a final result. This should be taken into account when selecting the most suitable type of screening assay to use.

References


Requests for reprints to: Mr G M Sutehall, Clinical Microbiology and Public Health Laboratory, Level 6, Addenbrooke’s Hospital, Hills Road, Cambridge CB2 2QW, England.
False positive latex tests negative by ELISA for toxoplasma IgG.

G M Sutehall and T G Wreghitt

*J Clin Pathol* 1989 42: 204-205
doi: 10.1136/jcp.42.2.204

Updated information and services can be found at:
http://jcp.bmj.com/content/42/2/204

**Email alerting service**

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/