

Intended Result	Your Report	Your Score
Specimen 8272 HBsAg positive HCV Ag/Ab negative HIV 1/2 Ag/Ab negative	HBsAg positive HCV Ag/Ab negative HIV 1/2 Ag/Ab negative	2 2 2
Specimen 8273 HBsAg negative HCV Ag/Ab negative HIV 1/2 Ag/Ab presumptive positive	HBsAg negative HCV Ag/Ab negative HIV 1/2 Ag/Ab presumptive positive	2 2 2
Specimen 8274 HBsAg negative HCV Ag/Ab positive HIV 1/2 Ag/Ab negative	HBsAg negative HCV Ag/Ab positive HIV 1/2 Ag/Ab negative	2 2 2

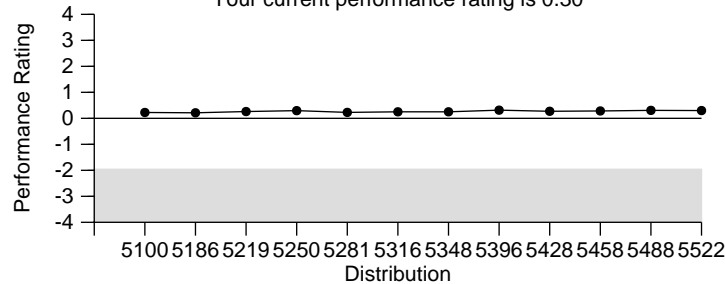
Cumulative score information

Total number of specimens sent to you for **UK NEQAS for Blood borne viruses** over the last 6 distribution is 18
 For these distributions specimen numbers 7731 7732 7733 7880 7881 7882 7979 7980 7981 8072 8073 8074 8164 8165 8166 8272 8273 8274 have been sent.
 Number of specimens reported too late for analysis (not scored) 0
 Your cumulative score for the specimen/test combinations that you reported was 108 out of a possible total of 108
 The mean score calculated from the reports returned by **UK** laboratories testing the specimen/test combinations you examined was 107.51 (with a standard error of 1.63).

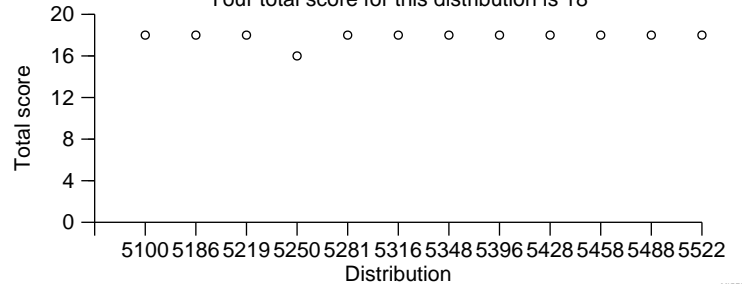
Performance rating

Your performance rating for **UK NEQAS for Blood borne viruses** (i.e. the number of standard errors by which your cumulative score lies above or below the mean for **UK** laboratories) is **0.30**.
 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.

Your performance rating over the past 12 distributions
 Your current performance rating is 0.30



Total score you achieved for each of the last 12 distributions
 Your total score for this distribution is 18



Comments

Overall performance for this distribution was outstanding with 100% of participants returning the intended results for HBsAg and HCV Ag/Ab, and 98.4% of participants returning the intended results for HIV-1/2 Ag/Ab.

This scheme is intended for laboratories performing screening assays. Results of confirmatory assays (e.g. immunoblot assays) are not included in the analysis.

Please see comments page 5 for scheme specific information

In the histograms on page 2 and subsequent pages a maximum of 12 kit/test method results are displayed. These include the most commonly used methods and the method(s) used in your laboratory indicated by an arrow(s). 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.

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

Enquiries: For repeat specimens in case of an EQA failure investigation, please request using the web form at <https://ukneqasmicro.org.uk/participant-info/order-repeat-specimens/>.

Pre-distribution test results are available should you experience a technical failure and wish to discuss the results. Any queries about this distribution can be addressed to Dr Sanjiv Rughooputh at organiser@ukneqasmicro.org.uk.

Acknowledgments

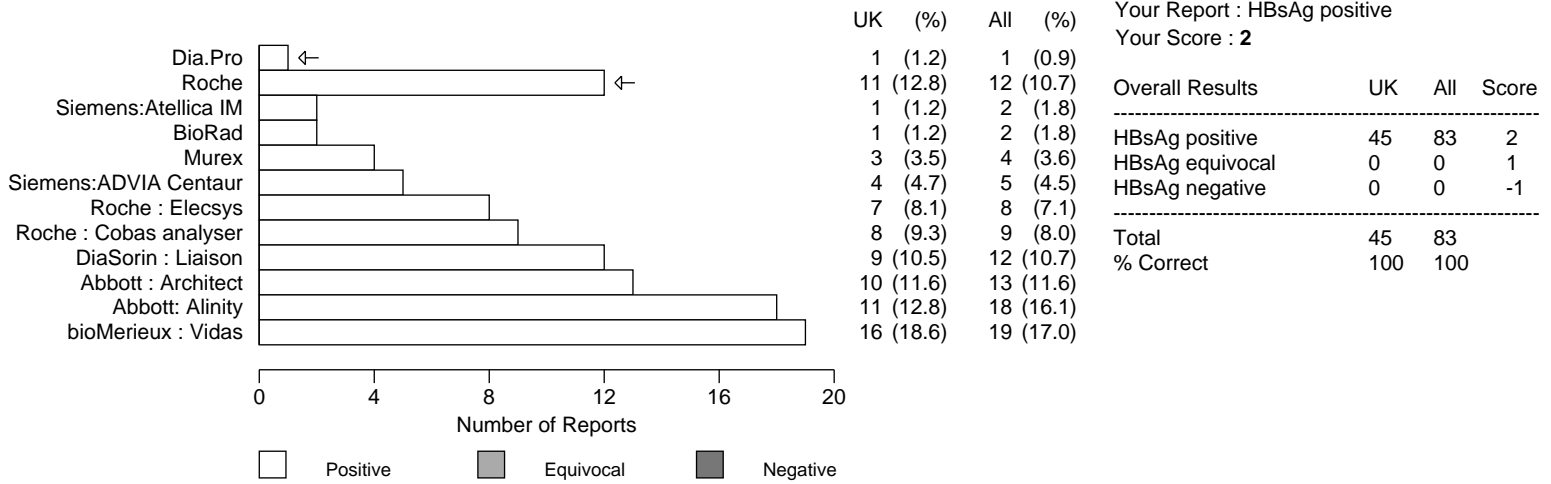
We thank colleagues at UKHSA Manchester, UKHSA Virus Reference Division in Colindale, NHSBT, St George's Hospital, and the Department of Microbiology at Gloucestershire Hospitals for their assistance with pre-distribution testing.

Report authorised by: Dr Sanjiv Rughooputh, Scheme Organiser.

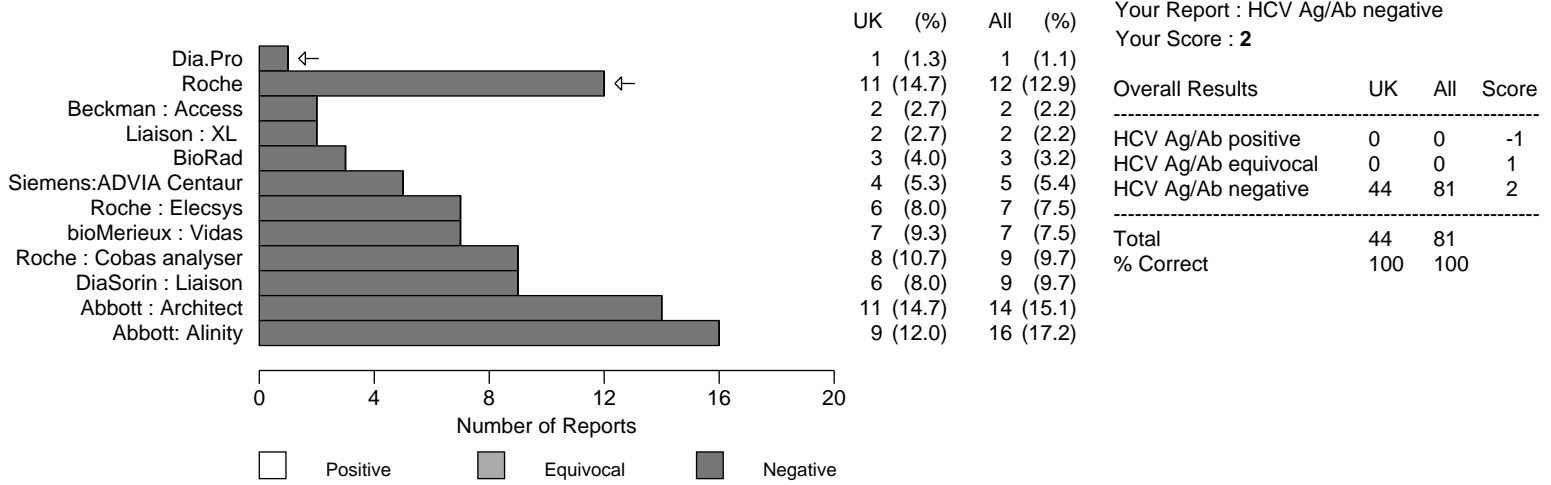


Three specimens derived from thrombinised human plasma (serum) donations were dispatched with a request to report on the presence of hepatitis B surface antigen (HBsAg), hepatitis C antigen/antibody and HIV-1/2 antigen/antibody. Specimens 8272, 8273 and 8274 were diluted 9.6, 17.1 and 4.8, respectively. Positive neat materials used to prepare HBsAg positive specimens were screened negative for HBeAg. Bronidox at a final concentration of 0.05% was added as a preservative.

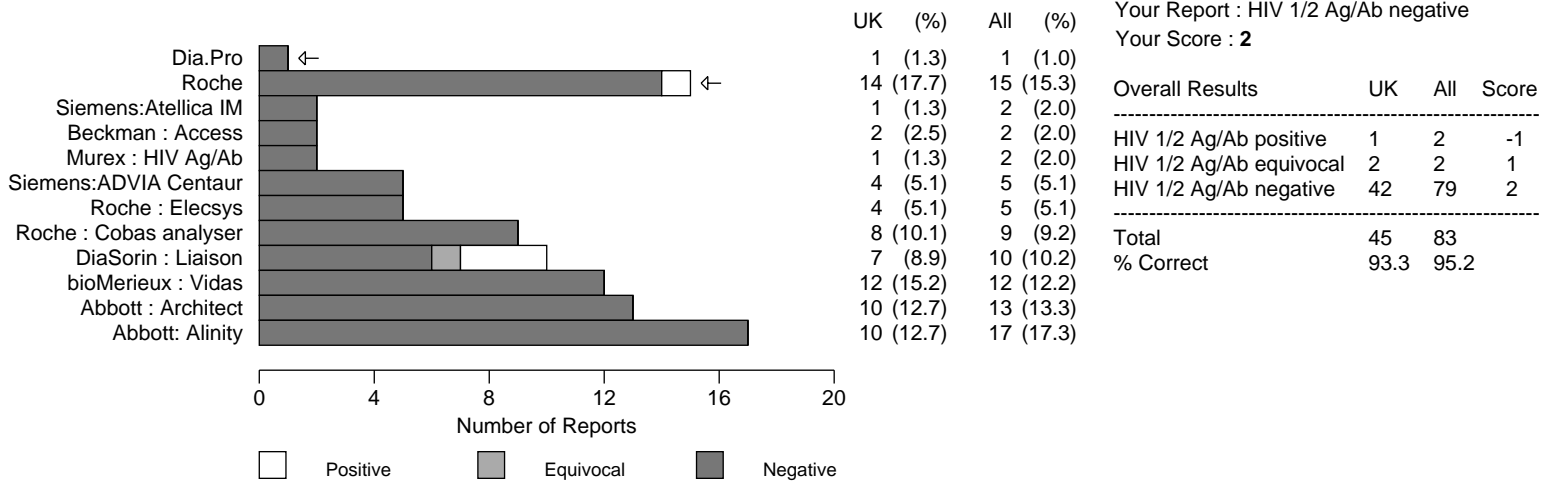
Specimen : 8272 Intended result HBsAg positive



Specimen : 8272 Intended result HCV Ag/Ab negative

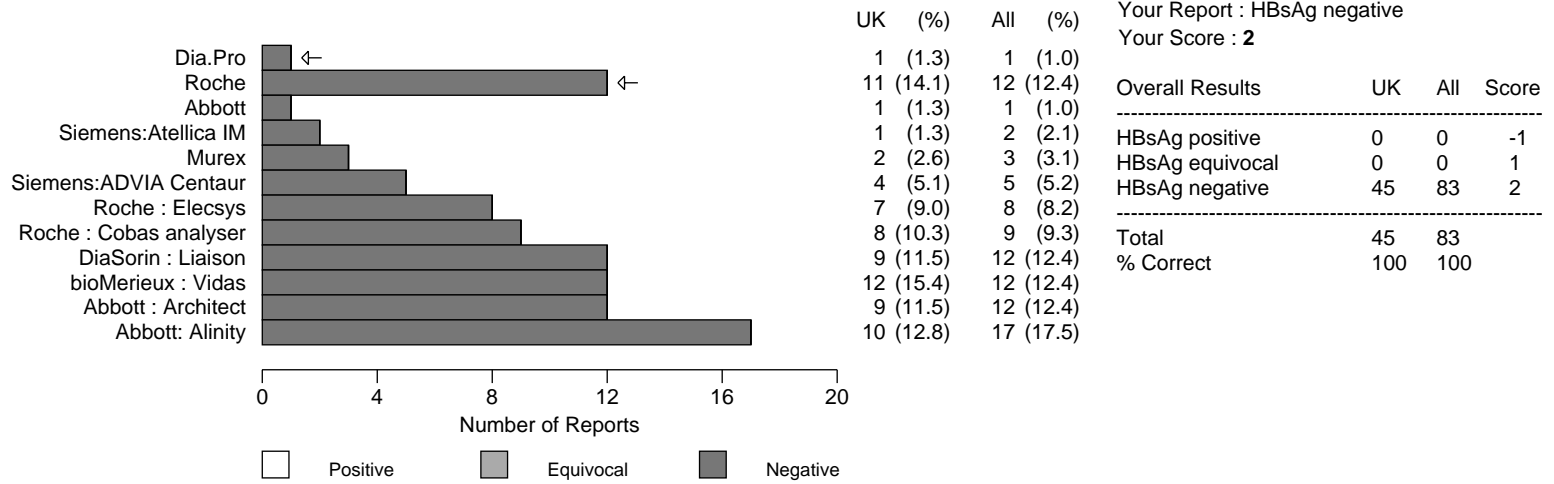


Specimen : 8272 Intended result HIV-1/2 Ag/Ab negative

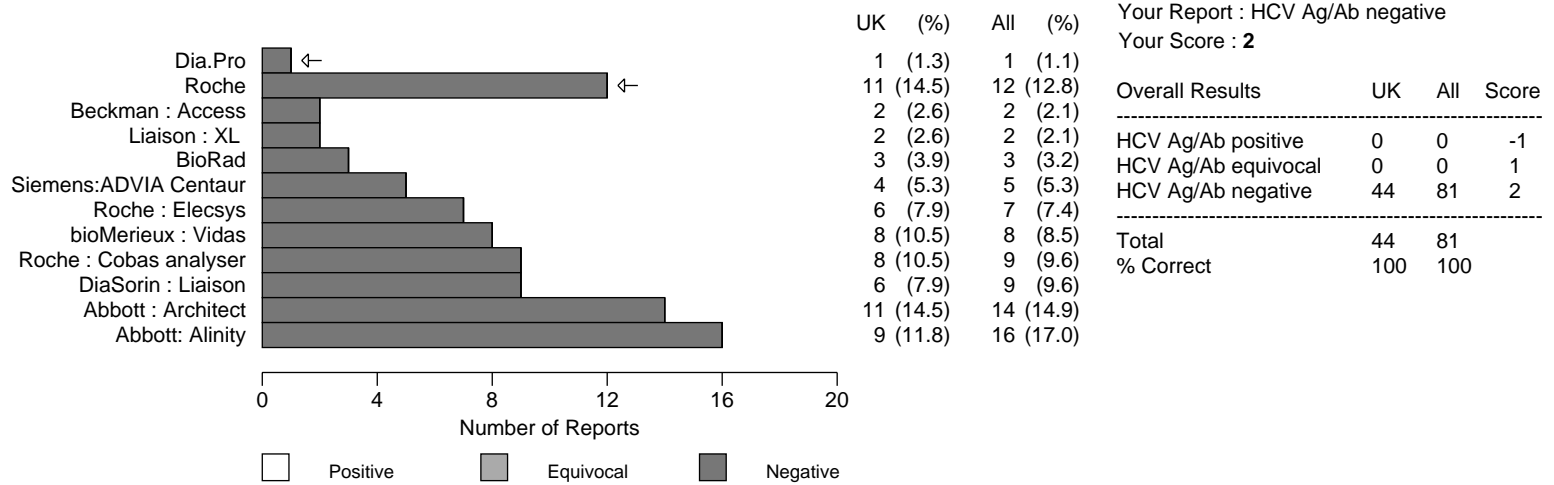


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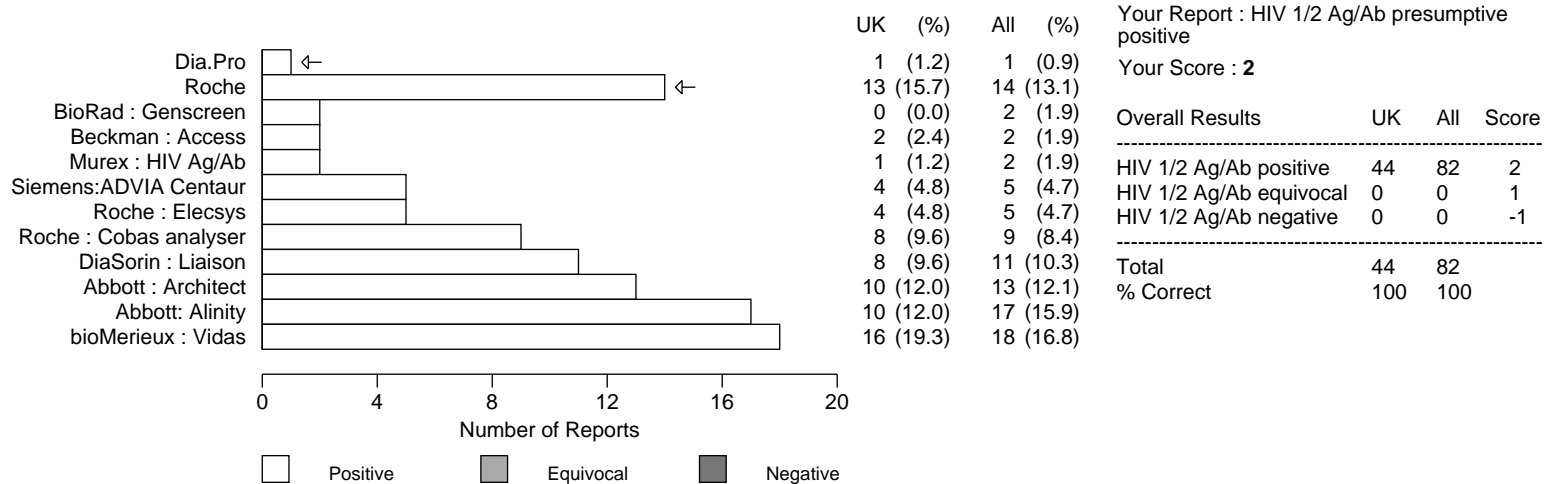
Specimen : 8273 Intended result
HBsAg negative



Specimen : 8273 Intended result
HCV Ag/Ab negative

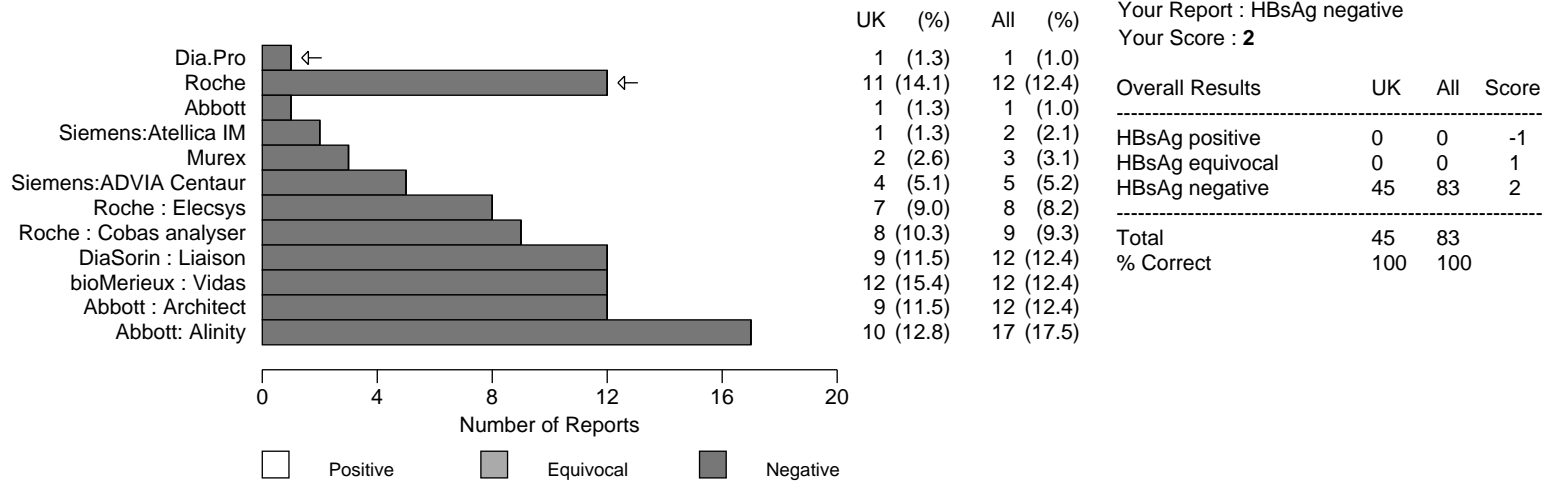


Specimen : 8273 Intended result
HIV-1/2 Ag/Ab presumptive positive

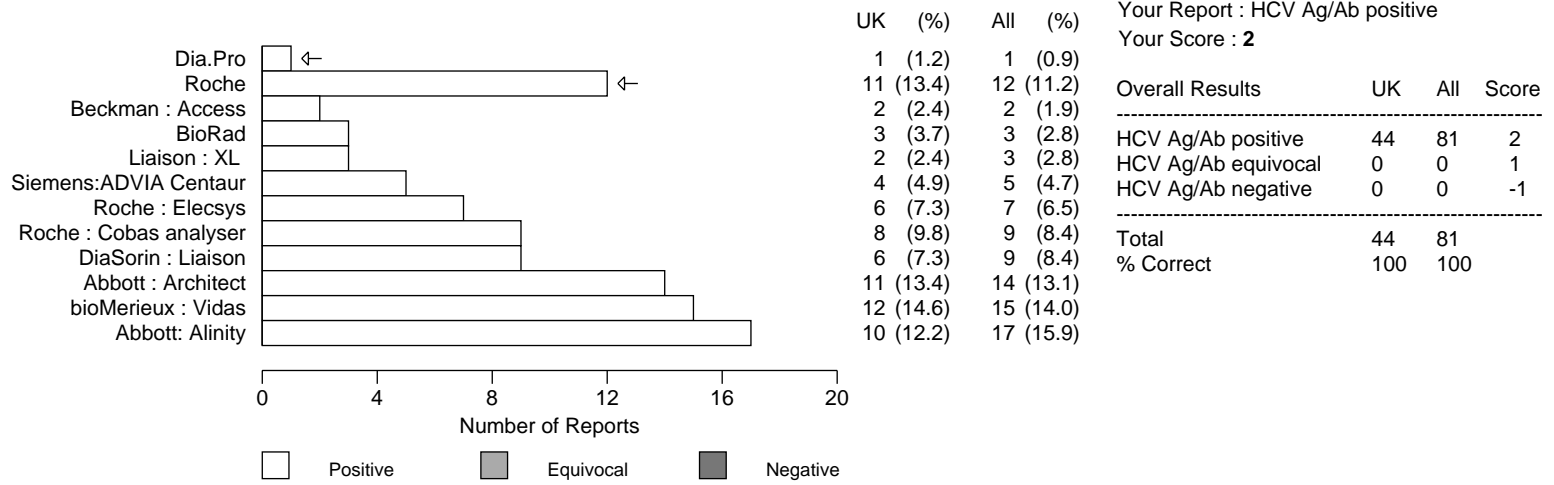


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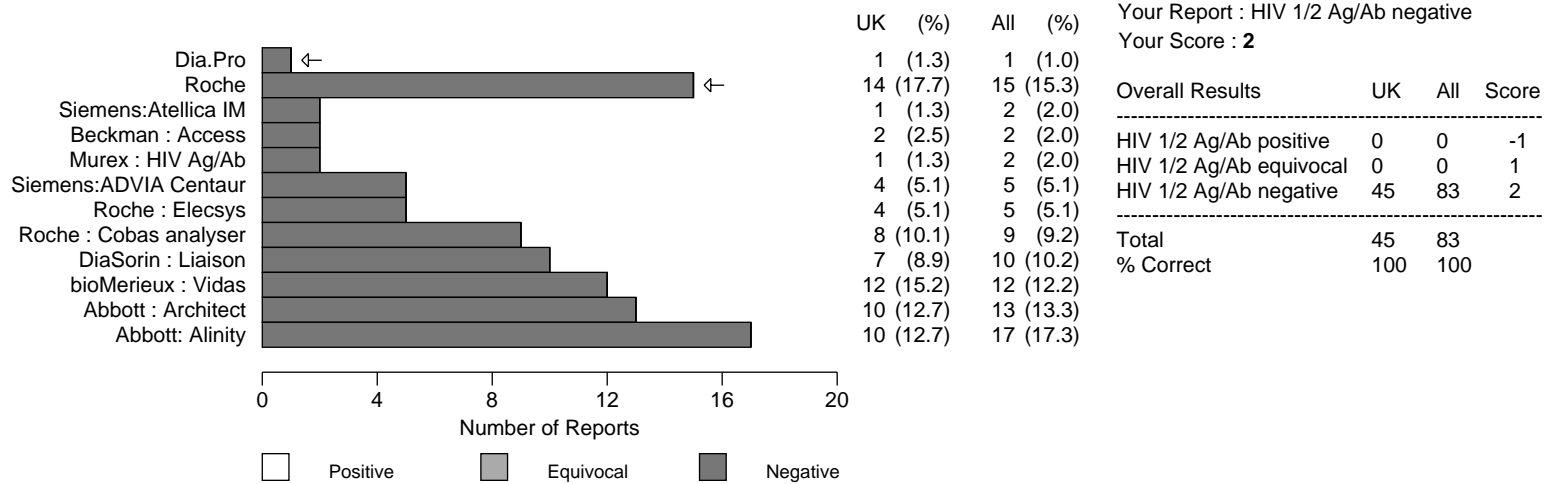
Specimen : 8274 Intended result
HBsAg negative



Specimen : 8274 Intended result
HCV Ag/Ab positive



Specimen : 8274 Intended result
HIV-1/2 Ag/Ab negative



Comments on distribution 5522

Overall performance in this distribution was excellent, with 99.4% of participants reporting the intended results. This is a slight improvement of 0.1% on the previous distribution 5488, where the overall performance was 99.3%.

Eighty-seven sets of specimens were dispatched with 83 participants returning results: representing a return rate of 95.4%. Compared to distribution 5488; where the return rate was 93.1%, this is an improvement of 2.3%.

Participants are requested to provide more details on the methods they use in order to have better platform groupings. This information is very helpful for troubleshooting in case of kit/platform issues.

Participants who did not obtain the intended results are requested to investigate and report the plausible root cause by completing an incident review form (IRF) online within 30 days of this report being published.

Non return of results without a valid reason is considered as poor performance and should also be reported in a similar manner.

IRFs are taken into considerations when poor performance analysis is carried out for the National Quality Assurance Advisory Panel (NQAAP). IRFs can be completed on the following link: <https://ukneqasmicro.org.uk/incident-review-form>

End of report

