

SUMMARY OF EXERCISE MATERIAL

Patient 1 - O D positive
 Patient 2 - A D negative
 Patient 3 - B D positive

Your results in bold
 Expected results are shaded

Patient 1

Your Result: O D Positive Your Score ≈ 0
 Overall Results: O D Positive 100.00% n=(17)

Patient 2

Your Result: A D Negative Your Score = 0
 Overall Results: A D Negative 100.00% n=(17)

Patient 3

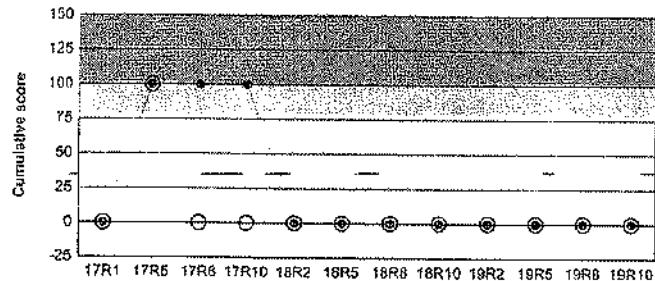
Your Result: B D Positive Your Score = 0
 Overall Results: B D Positive 100.00% n=(17)

Your overall score for this exercise: ABO 0
 RhD 0

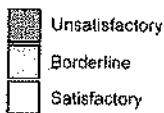
Your last 3 returns contribute to the cumulative scores

Cumulative Score Distribution Penalty Non Return

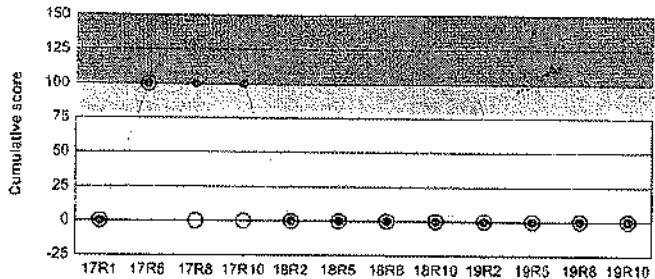
ABO Derived penalty score



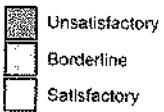
Current Performance: Satisfactory
 Cumulative Score: 0



RhD Derived penalty score



Current Performance: Satisfactory
 Cumulative Score: 0



Antibody Screening and Identification

SUMMARY OF EXERCISE MATERIAL

Patient 1 - Inert
 Patient 2 - Anti-E
 Patient 3 - Inert

Your results in bold
 Expected results are shaded

Antibody Screen

Antibody Identification

Patient 1

Your Result : **No specific antibody detected**
 Your Score = 0

Overall Results : No specific antibody detected 100.00% n=(17)

Patient 2

Your Result : **Antibody present**
 Your Score = 0

No Response
 Your Score = Not Scored

Overall Results : Antibody present 100.00% n=(17) E 100.00% n=(16)

Patient 3

Your Result : **No specific antibody detected**
 Your Score = 0

Overall Results : No specific antibody detected 100.00% n=(17)

Your overall score for this exercise :

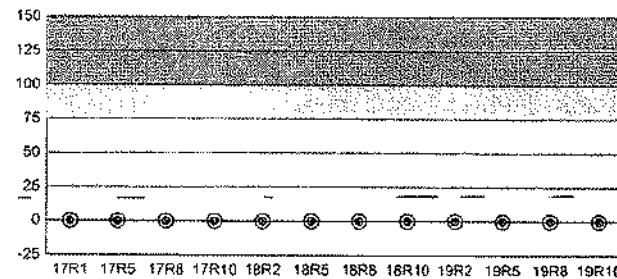
Antibody Screening

0

Your last 3 returns contribute to the cumulative scores

 Cumulative Score Distribution Penalty Non Return

Antibody screen derived penalty score



Current Performance : Satisfactory
 Cumulative Score : 0

- Unatisfactory
- Borderline
- Satisfactory

Printed at 13:38 on Thursday, 12 December, 2019 (Final Report)

For information on data analysis and performance assessment see the UK NEQAS (BTLP) Participants' Manual (www.ukneqasbtlp.org)

Scheme Director: Ms J White

Authorised by: Mr R Haggas

NEQAS (BTLP), PO Box 133, WATFORD WD18 0WP, UK

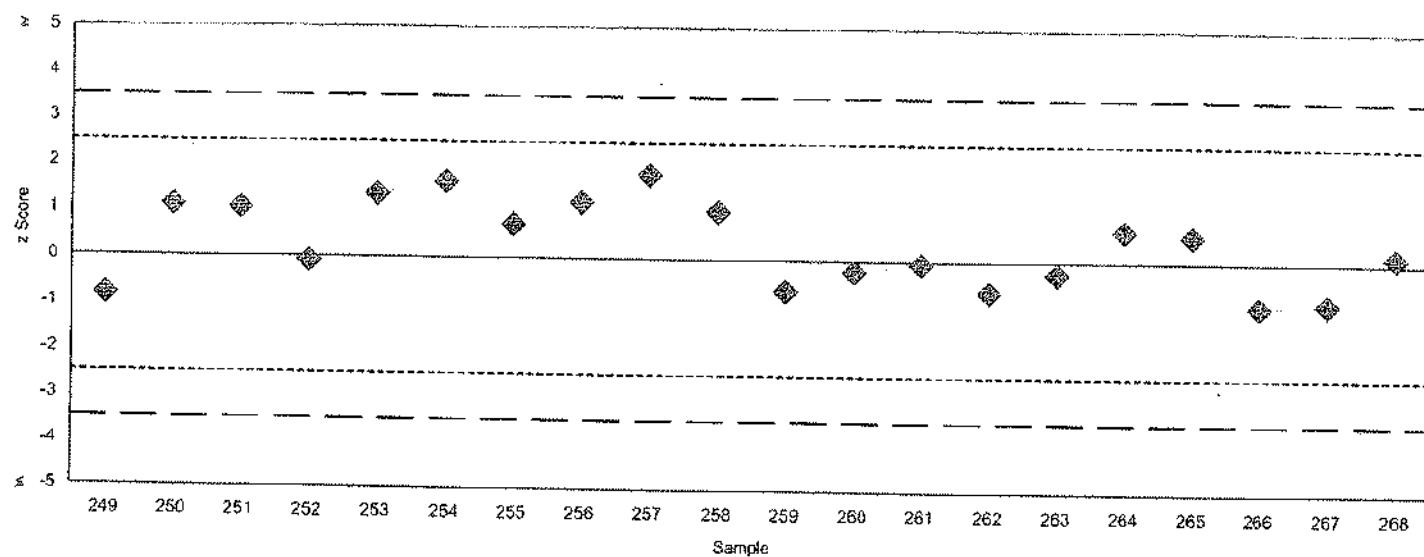
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be published without the Organiser's permission

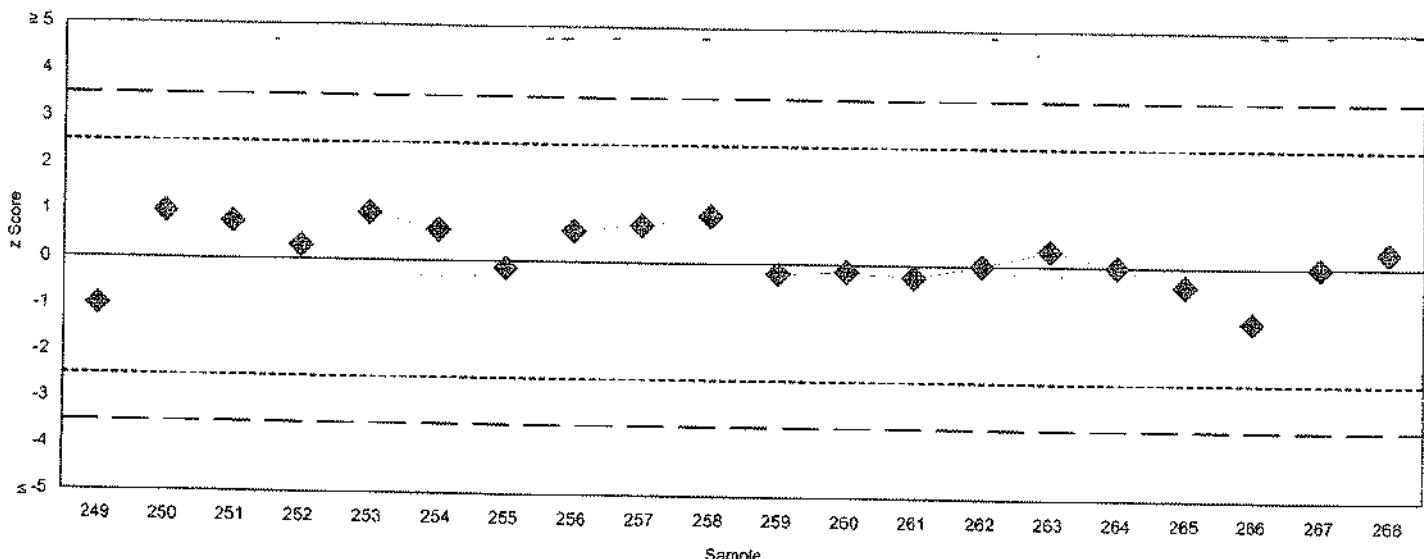
CD34+ Stem Cell Enumeration Programme

Shewhart Control Charts

(Please note each data point represents a single sample)

Absolute Values (cells/ μ L)

Percentage Values (%)





6227

ECAT FOUNDATION

External quality Control for Assays and Tests
With a focus on Thrombosis and Haemostasis

This is to certify that

MDA Blood Services Center

Quality Control Manager

RAMAT GAN

ISRAEL

Under the supervision of

Dr. Katya Shliachterman

Was registered in 2019 in the EQA programme of the ECAT
Foundation for the following modules:

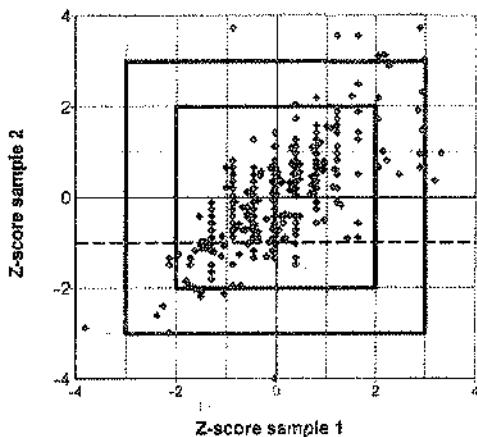
Screen - I; Coagulation Factor - I

Dr. P Meijer
Director ECAT Foundation

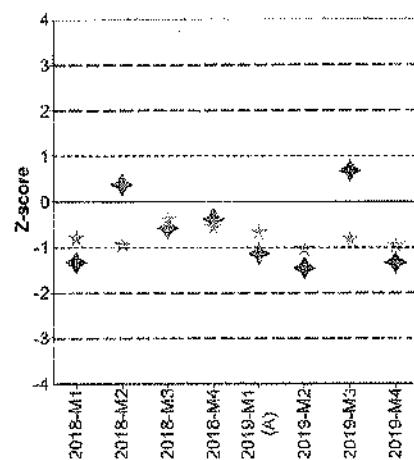
Coagulation Factor
Factor VIII (clotting activity)

Instrument ID	16090357
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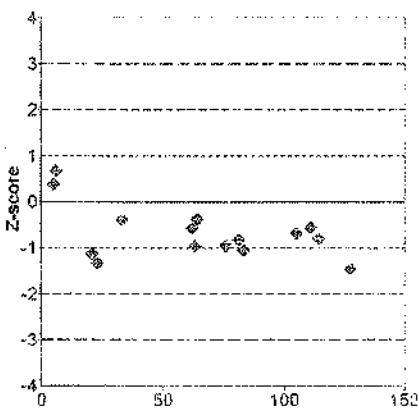
Z-score plot



Z-score history (by survey)

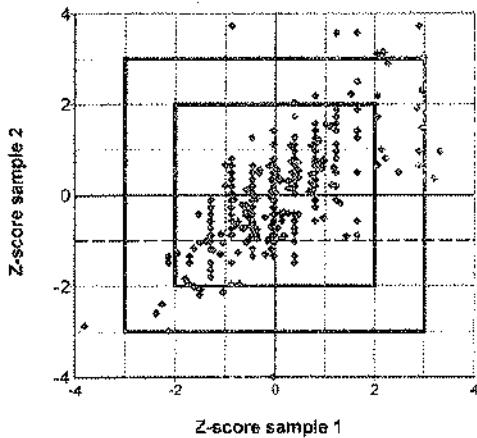


Z-score history (by consensus value)

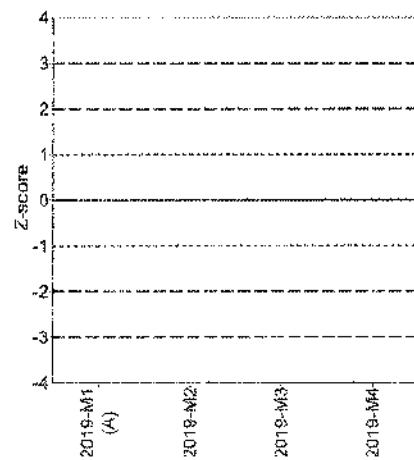


Instrument ID	
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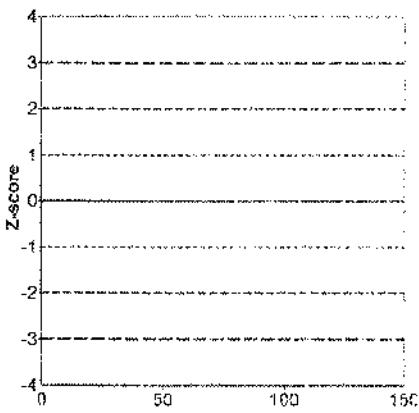
Z-score plot



Z-score history (by survey)

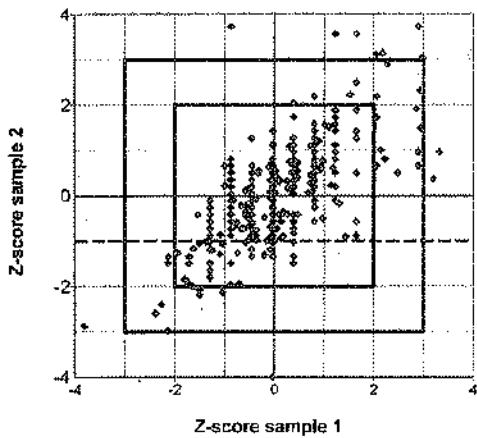


Z-score history (by consensus value)

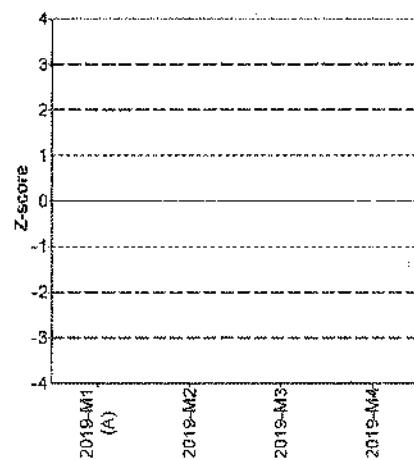


Instrument ID	
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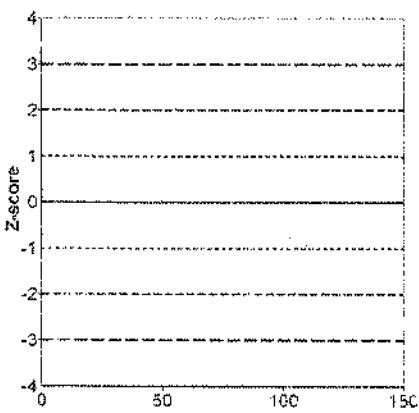
Z-score plot



Z-score history (by survey)



Z-score history (by consensus value)





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This is to certify that

MDA Blood Services Center
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RAMAT GAN
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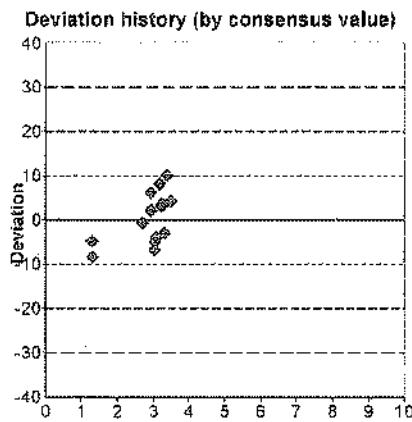
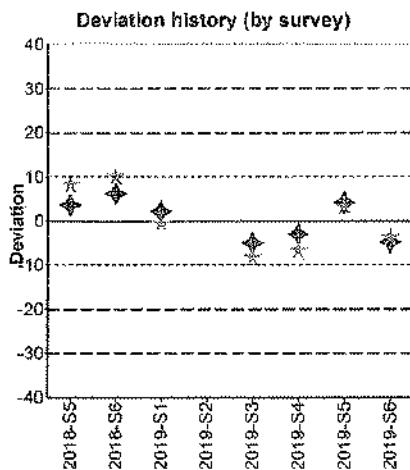
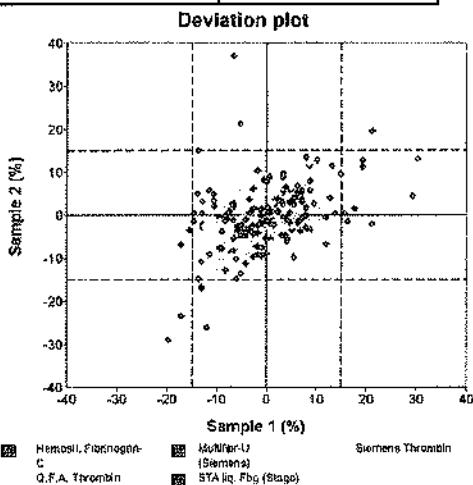
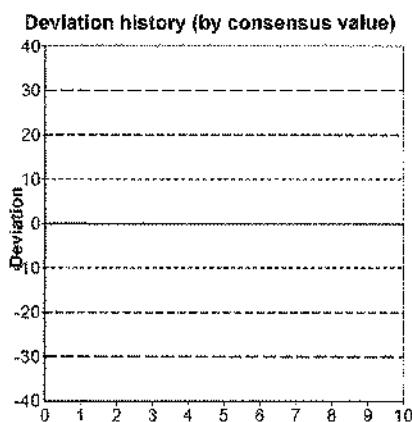
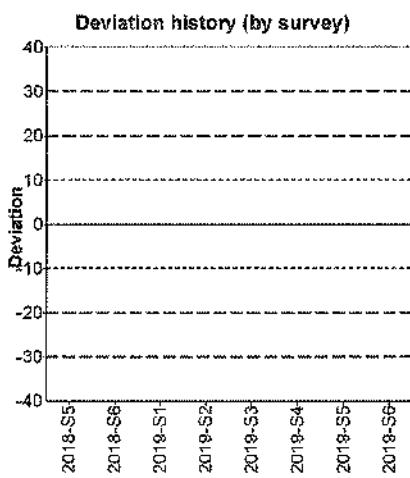
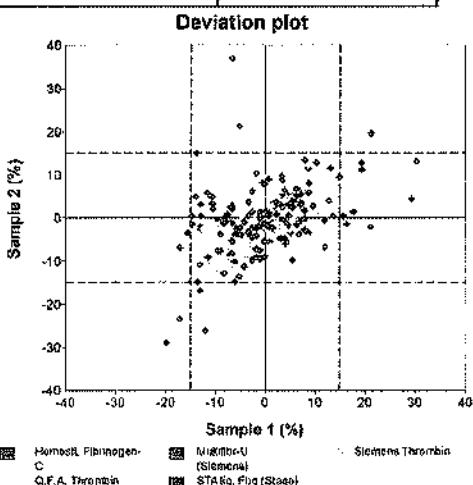
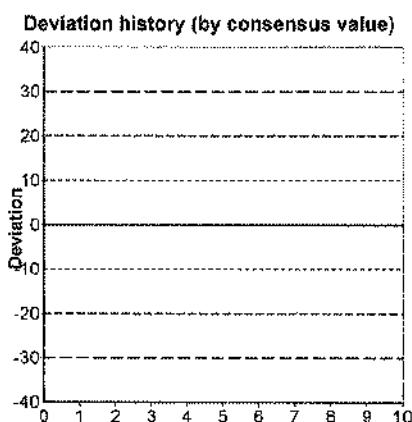
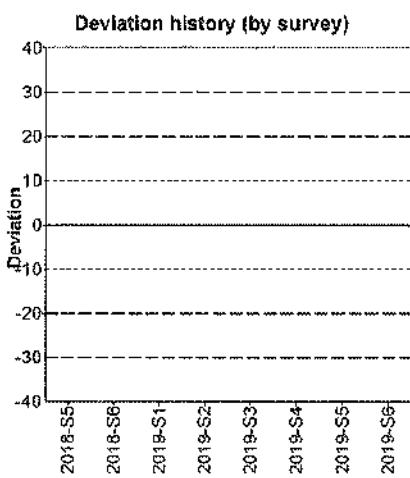
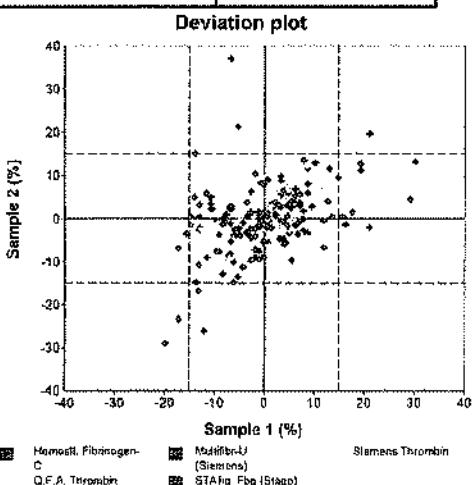
Under the supervision of

Dr. Katya Shliachterman

Was registered in 2019 in the EQA programme of the ECAT
Foundation for the following modules:

Screen - I; Coagulation Factor - I

Dr. P Meijer
Director ECAT Foundation

Screen 1
Fibrinogen (Clotting)
Instrument ID 16090357

Instrument ID

Instrument ID


Survey Contents:

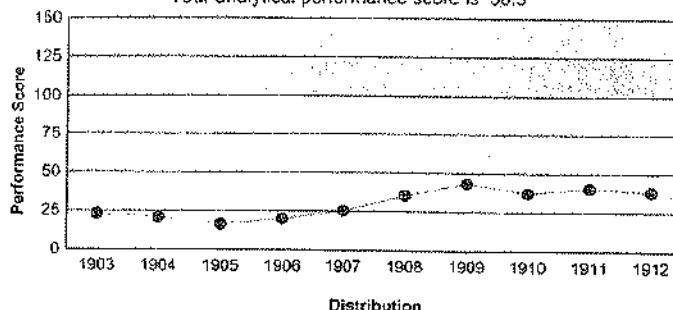
Specimen 1: 1912FB1 Partially fixed human whole blood

Specimen 2: 1912FB2 Partially fixed human whole blood

Non Participation Penalty: 0

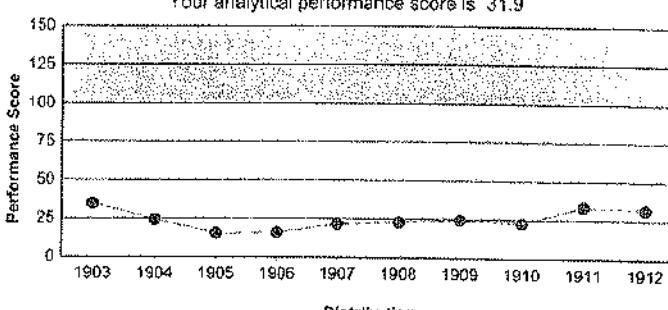
White Blood Count

Your analytical performance score is 38.3



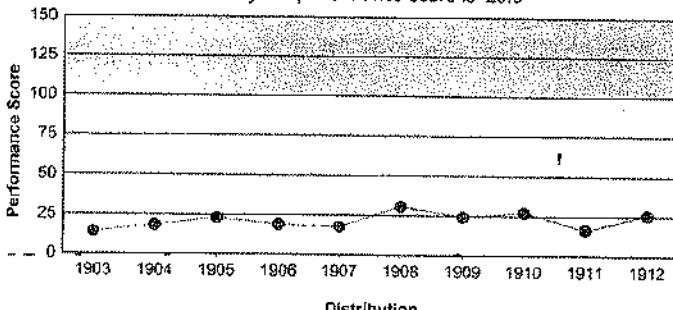
Red Blood Count

Your analytical performance score is 31.9



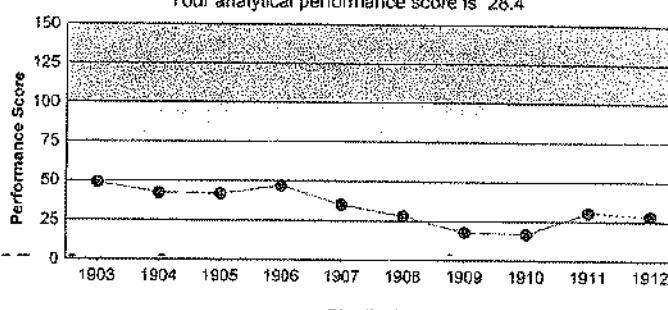
Haemoglobin

Your analytical performance score is 25.8



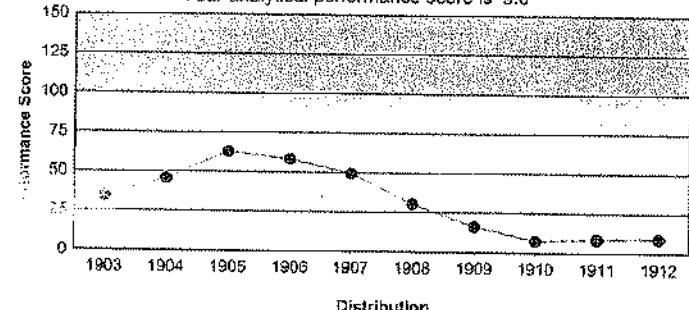
Packed Cell Volume

Your analytical performance score is 28.4



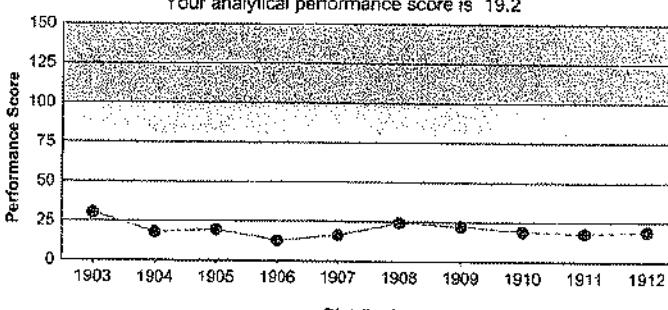
Mean Cell Volume

Your analytical performance score is 9.0



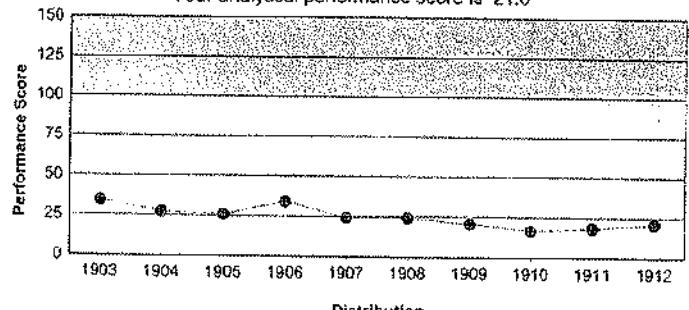
Mean Cell Haemoglobin

Your analytical performance score is 19.2



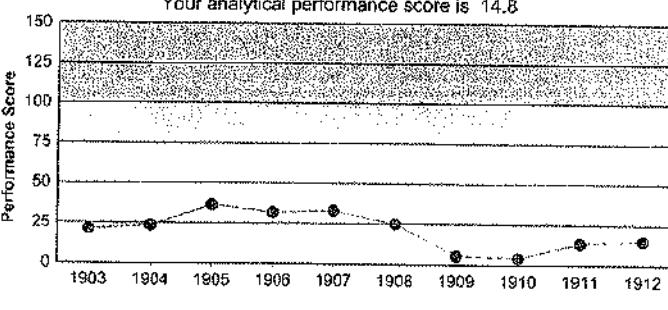
Mean Cell Haem Conc

Your analytical performance score is 21.0



Platelet Count

Your analytical performance score is 14.8

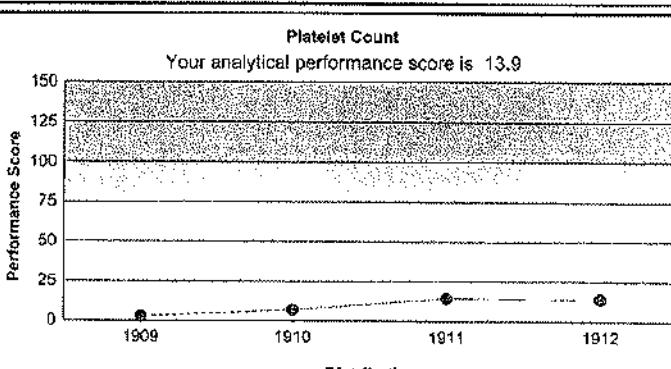
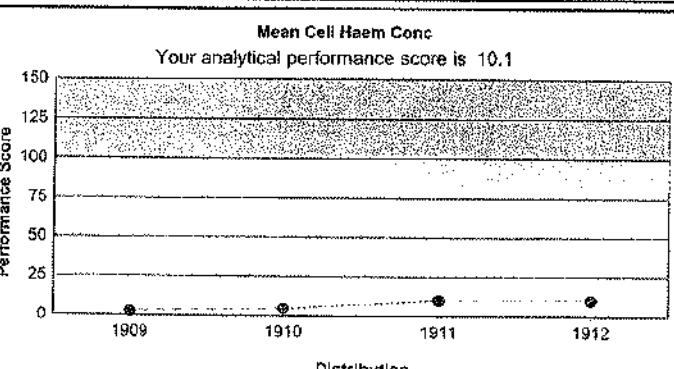
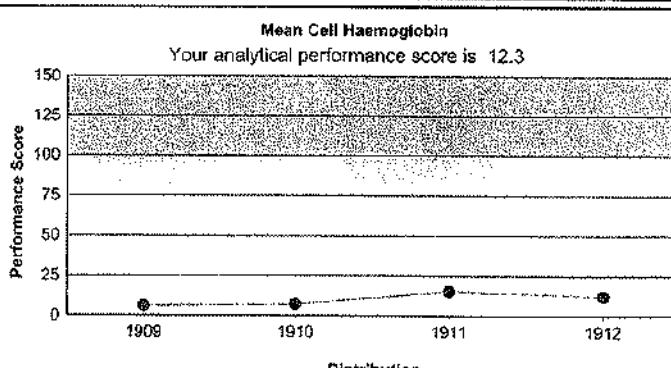
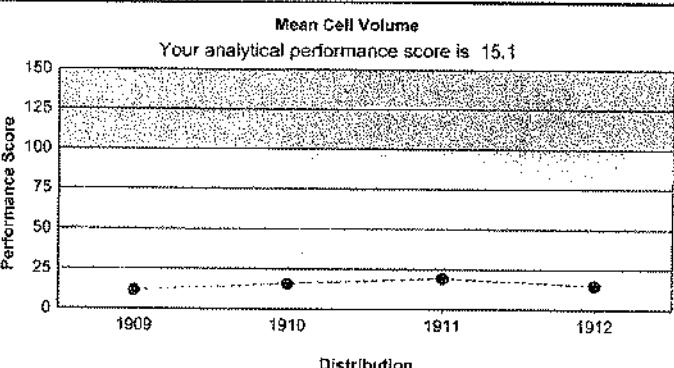
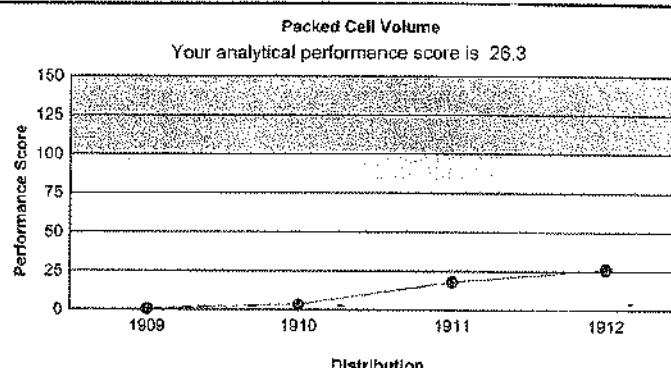
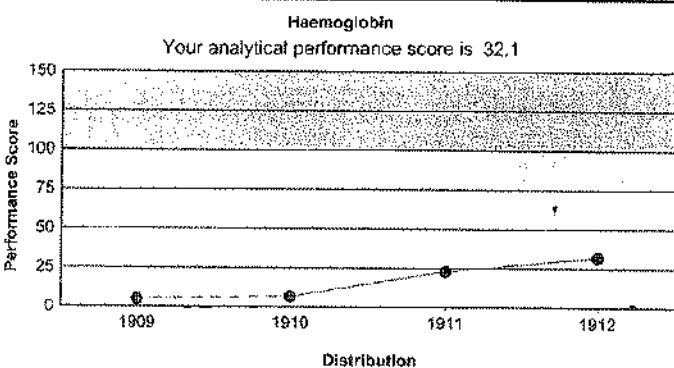
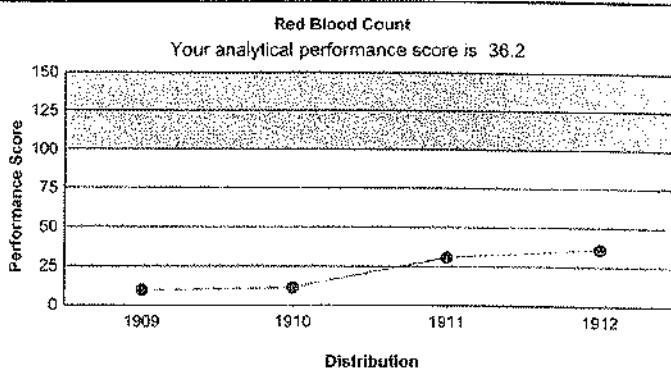
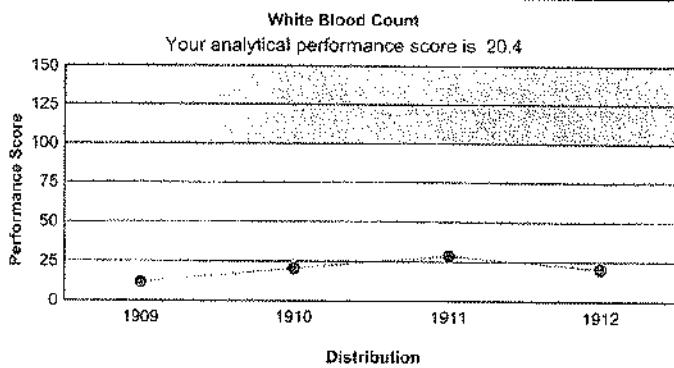


Survey Contents:

Specimen 1: 1912FB1 Partially fixed human whole blood

Specimen 2: 1912FB2 Partially fixed human whole blood

Non Participation Penalty: 0



Printed at 16:12 on Thursday, 12 December, 2019 (Final Report)

For information on data analysis and performance assessment see the UK NEQAS Haematology Participants' Manual (www.ukneqash.org)

Scheme Director: Barbara De la Salle

Authorised by: Yvonne John (Service Manager)

JK NEQAS Haematology, PO Box 14, WATFORD WD18 0FJ, UK

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Magen | David Adom-Blood Services

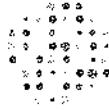
Immunology
7238203-01

As a laboratory demonstrating continuous improvement in quality through participation in the 2019 CAP Surveys, and/or Anatomic Pathology Education Programs.

Patrick Godbey, MD, FCAP

Patrick Godbey, MD, FCAP
CAP President

COLLEGE of AMERICAN
PATHOLOGISTS



The College of American Pathologists recognizes

Magen | David Adom-Blood Services

Immunology
7238203-01

As a laboratory demonstrating continuous improvement in quality through
participation in the 2019 CAP Surveys, and/or Anatomic Pathology
Education Programs.

A handwritten signature in black ink that reads "Patrick Godbey, MD, FCAP".

Patrick Godbey, MD, FCAP
CAP President

**COLLEGE of AMERICAN
PATHOLOGISTS**
325 Waukegan Road, Northfield, Illinois 60093-2750
800-223-4240 • cap.org

CAP Number: 7238203-01 Kit# 1
Institution: Magen David Adom-Blood Services
Attention: Liora Muncher
City / State: Tel Hashomer IS

Kit ID: 32323439
Kit Mailed: 9/9/2019
Original Evaluation: 10/24/2019

**EVALUATION
ORIGINAL**

J-C 2019 Transfusion Medicine (Comp)

CAP

CAP #: 7238203

Subspecialty : Unexpected Antibody Det

Regulated Analyte	Proficiency Event		Proficiency Event		Proficiency Event		Current Event		Cumulative CLIA '88 Performance Interpretation	
	2019	1	2019	2	2019	3	Performance	Performance	'88 Performance	Interpretation
Antibody Detection, Unexpected	Test Event	Score %	Test Event	Score %	Test Event	Score %	Test	Interpretation	Satisfactory	Successful
Unexpected Antibody Det	JA	5/5	JB	5/5	JC	5/5	100	FC	100	Successful



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800-323-6060 • cap.org

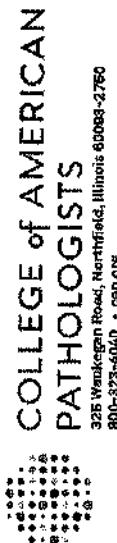
EVALUATION
ORIGINAL

CAP Number: 7238203-01 Kit# 1
Institution: Magen David Adom-Blood Services
Attention: Liora Muncher
City / State: Tel Hashomer IS
Kit ID: 32333439
Kit Mailed: 9/9/2019
Original Evaluation: 10/24/2019

J-C 2019 Transfusion Medicine (Comp)

CAP

CAP #:		Subspecialty : ABO Group/Rh Type					
Regulated Analyte	Test Event	Proficiency Event		Proficiency Event		Current Event	'88 Performance Interpretation
		2019	1	2019	2		
ABO Group	J-A	5/5	100	J-B	5/5	100	J-C
Rh (D) Type	J-A	5/5	100	J-B	5/5	100	J-C
ABO Group/Rh Type		10/10	100		10/10	100	
						10/10	100
						Satisfactory	Successful
							Successful



**COLLEGE of AMERICAN
PATHOLOGISTS**
325 Waukegan Road, Northfield, Illinois 60061-2760
800-323-4040 • cap.org

EVALUATION
ORIGINAL

CAP Number: 7238203-01 Kit# 1
Institution: Magen David Adom-Blood Services
Attention: Iora Muncher
City / State: Tel Hashomer IS

Kit ID: 32323439
Kit Mailed: 9/9/2019
Original Evaluation: 10/24/2019

J-C 2019 Transfusion Medicine (Comp)

CAP

CAP #:		Subspecialty : Antibody Identification				Cumulative CLIA '88 Performance Interpretation	
Regulated Analyte	Proficiency Event 2019	Proficiency Event		Proficiency Event		Current Event Performance	Cumulative CLIA '88 Performance Interpretation
		Test Event	Score %	Test Event	Score %		
Antibody Identification	J-A	5/5	100	J-B	5/5	100	J-C 100 Satisfactory Successful
Antibody Identification		5/5	100	5/5	100	5/5	100 Satisfactory Successful



CERTIFICATE

Survey of 23 October 2019

You have fulfilled the requirements of the External Quality Assessment with the following analysis

Immunohaematology 06 - Molecular Diagnostic (235):

Validity 12 months:

ABO
MNS
RHD Zygote
RHD
RHCE
KEL (k/K)
FY
JK

(R) analysis is subject to the RiliBAK

Participant:

33144
Dr. Liora Müncher Immunohematology MDA Blood
Service
Tel Hashomer
5262000 Ramat Gan



DAkkS
DKEK-Akkreditierung
DKEK-Akkreditierung



Düsseldorf, 20 November 2019

Prof. Dr. med. Michael Spannagl
(Head of Reference Institution)

Prof. Dr. med. Gregor Bein
(Adviser)

Prof. Dr. med. Willy A. Flegel
(Adviser)

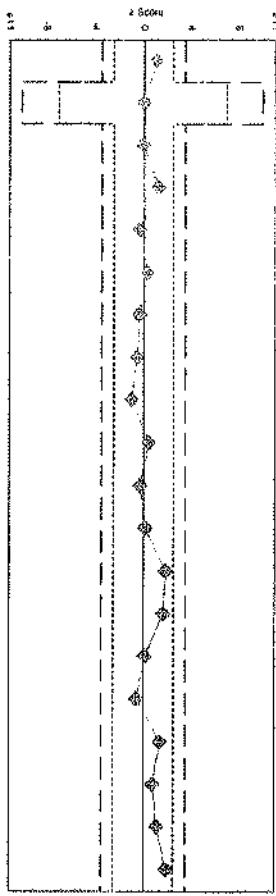
Low Level Leucocyte Enumeration Programme

Low Level Leucocyte Enumeration Programme

Shewhart Control Charts

(Please note each data point represents 15.5 Sample Spots)

Absolute Values (cells/µL) - Machine 2



Flow Cytometer Specific Statistics

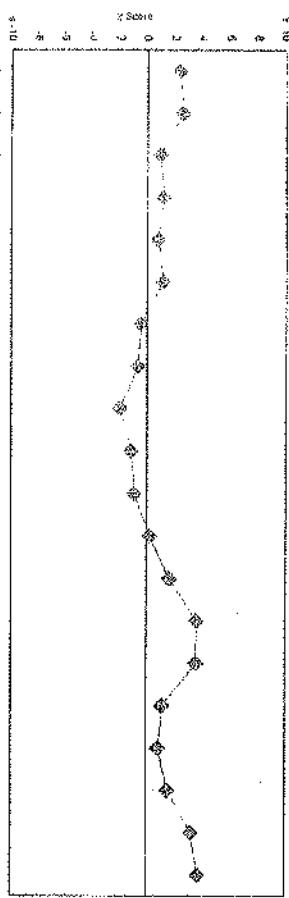
(Please note only groups of >20 counts are displayed)

Method	Absolute Count Values (cells/µL)		
	Returns	Robust	Robust
BD Leucocount	74	11.74	1.19
FDS-Canto II	32	11.93	0.93
FlowCam	23	11.35	1.04
Navios	20	12.07	1.58

Cusum Control Charts

(Please note each data point represents the sum of the current sample and the two previous samples)

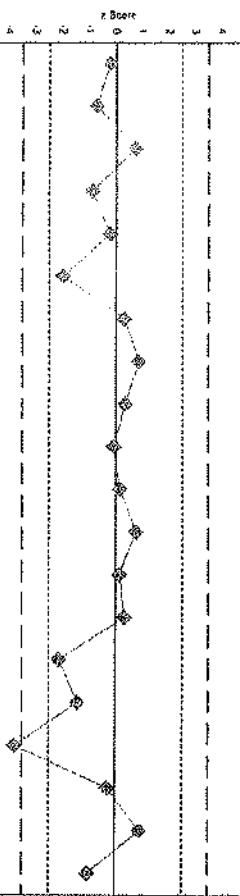
Absolute Values (cells/µL) - Machine 2



Low Level Leucocyte Enumeration Programme

Shewhart Control Charts
(Please note each data point represents a single sample)

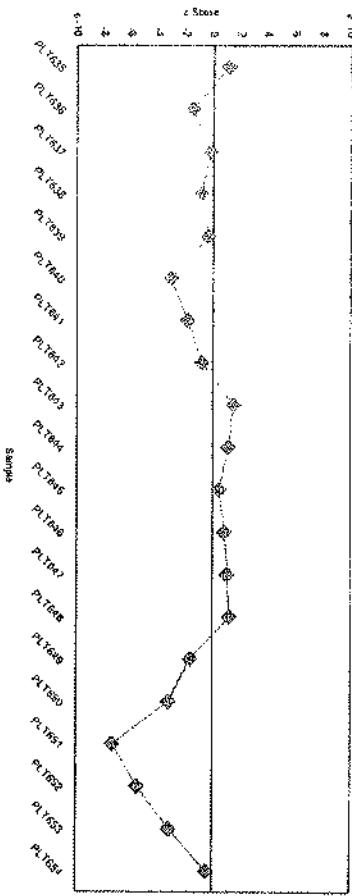
Absolute Values (cells/ μ l) • Machine 2



Custom Control Charts

(Please note each data point represents the sum of the z-scores of the current sample and the two previous samples)

Absolute Values (cells/ μ l) • Machine 2



Low Level Leucocyte Enumeration Programme

Staining Method Specific Statistics
(Please note only groups of >20 returns are displayed)

Method	Absolute Count Values (cells/ μ l)		
	Returns	Robust Mean	Robust SD
BD Luminex	74	2.42	0.37

Flow Cytometer Specific Statistics

(Please note only groups of >20 returns are displayed)

Method	Absolute Count Values (cells/ μ l)		
	Returns	Robust Mean	Robust SD
Facs Canto II	32	2.38	0.38
Facscalibur	23	2.38	0.36
Mavics	20	2.40	0.38

Specimen ID	Specimen Details	Your Report	Your Score
imen 5163	Reagins negative TP Agglutination positive FTA-Abs positive Immunoassay IgG/total Ab positive Immunoassay IgM negative Immunoblot IgG positive Immunoblot IgM negative	Not examined TP Agglutination positive Not examined Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored Not scored
imen 5164	Reagins positive TP Agglutination negative FTA-Abs negative Immunoassay IgG/total Ab negative Immunoassay IgM negative Immunoblot IgG negative Immunoblot IgM negative	Not examined TP Agglutination negative Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored
imen 5165	Reagins negative TP Agglutination positive FTA-Abs positive Immunoassay IgG/total Ab positive Immunoassay IgM negative Immunoblot IgG positive Immunoblot IgM negative	Not examined TP Agglutination negative Not examined Not examined Not examined Not examined	Not scored Not scored Not scored Not scored Not scored Not scored
imen 5166	Reagins positive TP Agglutination positive FTA-Abs positive Immunoassay IgG/total Ab positive Immunoassay IgM positive Immunoblot IgG positive Immunoblot IgM positive	Not examined TP Agglutination positive Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored
imen 5167	Reagins negative TP Agglutination positive FTA-Abs positive Immunoassay IgG/total Ab positive Immunoassay IgM negative Immunoblot IgG positive Immunoblot IgM negative	Not examined TP Agglutination positive Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored
imen 5168	Reagins negative TP Agglutination negative FTA-Abs negative Immunoassay IgG/total Ab negative Immunoassay IgM negative Immunoblot IgG negative Immunoblot IgM negative	Not examined TP Agglutination negative Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored

Comments: Performance information relating to this distribution is available in the commentary section page 21.

Results for each marker by methods used by participants are presented on subsequent pages.

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.

Acknowledgements: We thank colleagues at PHE- Public Health Laboratories in Birmingham, Bristol, Manchester and Newcastle; and the Department of Microbiology, Sheffield for assistance with pre-distribution tests.

Queries: Any queries about this distribution please contact Habib Seyedzadeh at the e-mail address below. Pre-distribution test results are available should you experience a technical failure and wish to discuss the results.

Repeat specimens please order using the web form or e-mail organiser@ukneqasmicro.org.uk stating your laboratory identification number, the distribution number and specimen numbers.

Approved by: Dr. Sanjiv Rughooputh, Director.

Distribution : 4422

Page 1 of 23

Dispatch Date : 04-Feb-2019

Entered Result		Your Report	Your Score
Specimen 4815	Reagins negative TP Agglutination negative FTA-Abs negative Immunoassay IgG/total Ab negative Immunoblot IgG negative Immunoassay IgM negative Immunoblot IgM negative	Not examined TP Agglutination negative Not examined Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored Not scored Not scored
Specimen 4816	Reagins negative TP Agglutination positive FTA-Abs positive Immunoassay IgG/total Ab positive Immunoblot IgG positive Immunoassay IgM negative Immunoblot IgM negative	Not examined TP Agglutination positive Not examined Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored Not scored
Specimen 4817	Reagins negative TP Agglutination positive FTA-Abs positive Immunoassay IgG/total Ab positive Immunoblot IgG positive Immunoassay IgM negative Immunoblot IgM negative	Not examined TP Agglutination positive Not examined Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored Not scored
Specimen 4818	Reagins positive TP Agglutination positive FTA-Abs positive Immunoassay IgG/total Ab positive Immunoblot IgG positive Immunoassay IgM positive Immunoblot IgM positive	Not examined TP Agglutination positive Not examined Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored Not scored
Specimen 4819	Reagins negative TP Agglutination positive FTA-Abs positive Immunoassay IgG/total Ab positive Immunoblot IgG positive Immunoassay IgM negative Immunoblot IgM negative	Not examined TP Agglutination positive Not examined Not examined Not examined Not examined Not examined	Not scored Not scored Not scored Not scored Not scored Not scored Not scored
Specimen 4820	Reagins negative TP Agglutination negative FTA-Abs negative Immunoassay IgG/total Ab negative Immunoblot IgG negative Immunoassay IgM negative Immunoblot IgM negative	Not examined TP Agglutination negative Not examined Not examined Not examined Not examined Not examined	Not scored 2 Not scored Not scored Not scored Not scored Not scored

Comments: Please check last pages (Pages 21-23) for additional comments.

Results for each marker by methods used by participants are presented on subsequent pages.

Results of the Fujirebio INNO-LIA IgG line assay of the positive specimens can be viewed on: www.ukneqasmicro.org.uk, by logging onto the Results and Reports section and selecting the 'Dist' button for the scheme.

Turnaround 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.

Acknowledgements: We thank colleagues at PHE- Public Health Laboratories in Birmingham, Bristol, Manchester and Newcastle; and the Department of Microbiology, Sheffield for assistance with pre-distribution tests.

Queries: Any queries about this distribution please contact Dr Beatrix Kele at the e-mail address below. Pre-distribution test results are available should you experience a technical failure and wish to discuss the results.

To repeat specimens please order using the web form or e-mail organiser@ukneqasmicro.org.uk stating your laboratory identification number, the distribution number and specimen numbers.

This report authorised by: Dr. Sanjiv Rughooputh, Scheme Organiser.



Certificate of Acceptable Performance



THIS IS TO CERTIFY THAT:

**BLOOD-TRANSFUSED INFECTIOUS DISEASES LAB
MADA
Lab 124442**

HAS SUCCESSFULLY COMPLETED THE EXTERNAL QUALITY ASSESSMENT FOR:

**EQAS HIV/Hepatitis Program (Quantitative)
Cycle 4: March 2019 – March 2020**

FOR THE FOLLOWING TESTS:

HBC Ab, Total
HBs Ab, Total-Unit
HIV-1/2 Ab
HTLV-I Ab, Total

Acceptable performance criteria: >= 75% response rate with results within + - 3 Z-score



Bio-Rad Laboratories is fully accredited by the American Association for Laboratory Accreditation (A2LA) to the ISO/IEC 17043:2010 standard.

Azita Mokhtari

Azita Mokhtari, EQAS Program Manager

**Bio-Rad Laboratories
EQAS Program Enrollment**

9500 Jeronimo Road
Irvine, CA 92618-2017, USA
email: eqas@bio-rad.com

Intended Result	Your Report	Your Score
Specimen 5419 HBsAg positive Anti-HBc positive HCV Ag/Ab negative HIV 1/2 Ag/Ab negative Anti-HTLV I/II negative Tp Immunoassay IgG/total Ab negative	HBsAg positive Anti-HBc positive HCV Ag/Ab negative HIV 1/2 Ag/Ab negative Anti-HTLV I/II negative Not examined	2 2 2 2 2 Not scored
Specimen 5420 HBsAg negative Anti-HBc negative HCV Ag/Ab positive HIV 1/2 Ag/Ab negative Anti-HTLV I/II negative Tp Immunoassay IgG/total Ab negative	HBsAg negative Anti-HBc negative HCV Ag/Ab positive HIV 1/2 Ag/Ab negative Anti-HTLV I/II negative Not examined	2 2 2 2 2 Not scored
Specimen 5421 HBsAg negative Anti-HBc negative HCV Ag/Ab negative HIV 1/2 Ag/Ab presumptive positive Anti-HTLV I/II negative Tp Immunoassay IgG/total Ab negative	HBsAg negative Anti-HBc negative HCV Ag/Ab negative HIV 1/2 Ag/Ab presumptive positive Anti-HTLV I/II negative Not examined	2 2 2 2 2 Not scored

Cumulative score information

Total number of specimens sent to you for UK NEQAS for Blood donor screen over the last 6 distributions is 18

For these distributions specimen numbers 4971 4972 4973 5053 5054 5055 5148 5149 5150 5230 5231 5232 5321 5323 5419 5420 5421 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 148 out of a possible total of 148

The mean score calculated from the reports returned by ALL laboratories testing the specimen/test combinations you examined was 147.40 (with a standard error of 1.78).

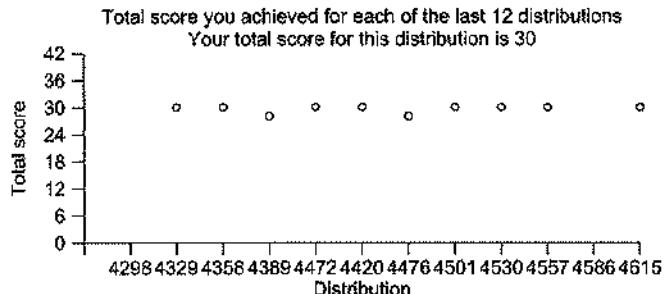
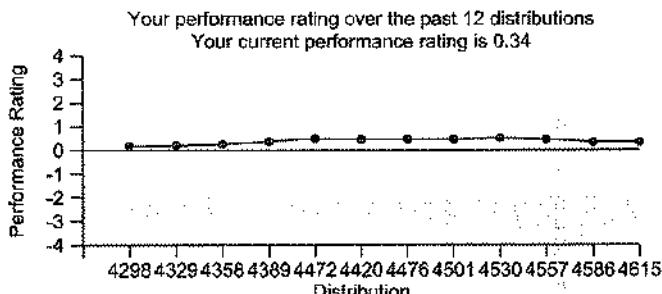
Performance rating

Your performance rating for UK NEQAS for Blood donor screen (i.e. the number of standard errors by which your cumulative score lies above or below the mean for ALL laboratories) is 0.34.

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.



Comments

Overall performance for this distribution was excellent with 100% of participants reporting correct results for HBsAg, Anti-HBc, Anti-HTLV I/II and Treponemal antibodies. 99.7% of participants reported correct results for HIV-1/2 Ag/Ab and HCV Ag/Ab.

This scheme is intended for laboratories performing screening assays only. Results of confirmatory assays (e.g. Immunoblot assays) are not included in the analysis. In the histograms on page 2 and subsequent pages a maximum of 10 kit results are displayed: this includes 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.

The time taken to report your results was 9-days. This information is provided for your own use and does not form part of your performance assessment.

Enquiries: 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, specimen number(s), and reason for request. 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 Rugoophut at the above email address.

Acknowledgments: We thank colleagues at PHE Manchester, PHE Virus Reference Division in Colindale, Gloucestershire Royal Hospital, and NHS Blood and Transplant for their assistance with pre-distribution testing.

Report authorised by: Dr Sanjiv Rugoophut, Director.



Analyte Scorecard

CAP #	CLIA #	Name / Address	CAP Accreditation Participant	Demographic Group
6385501		Magen David Adom-Blood Services Tel Hashomer, 52621 Israel		

Bolded analytes are regulated analytes as defined in CLIA 88, Subpart I; Proficiency Testing Programs for Nonwaived Testing; **Red** highlight indicates unsatisfactory performance; Yellow highlight indicates satisfactory performance <100%.

Analyte	CAP #	Program Year	Event Evaluation Date	Kit #	Mailing	Score	% Performance
Anti-CMV	638550101	2019	1 3/1/2019	32320627-1	VNA-A 2019	3/3	100 Satisfactory
Anti-HTLV VI	638550101	2019	1 3/1/2019	32320627-1	VNA-A 2019	3/3	100 Satisfactory
Anti-HTLV VII	638550101	2019	2 6/14/2019	32320628-1	VNA-B 2019	2/3	67 Satisfactory
HBV by NAT	638550101	2019	2 6/14/2019	32320628-1	VNB-B 2019	3/3	100 Satisfactory
HCV by NAT	638550101	2019	1 4/17/2019	32330643-1	NATA 2019	3/5	100 Satisfactory
HIV by NAT	638550101	2019	1 4/17/2019	32330643-1	NATA 2019	5/5	100 Satisfactory
Multiplex by NAT	638550101	2019	1 4/17/2019	32330643-1	NATA 2019	5/5	100 Satisfactory
HBV by NAT	638550101	2019	2 8/13/2019	32330646-1	NAT-B 2019	4/4	100 Satisfactory
HCV by NAT	638550101	2019	2 8/13/2019	32330646-1	NAT-B 2019	4/4	100 Satisfactory
HIV by NAT	638550101	2019	2 8/13/2019	32330646-1	NAT-B 2019	4/4	100 Satisfactory
Multiplex by NAT	638550101	2019	2 8/13/2019	32330646-1	NAT-B 2019	5/5	100 Satisfactory
West Nile Virus by NAT	638550101	2019	2 8/13/2019	32330646-1	NAT-B 2019	5/5	100 Satisfactory
HBV by NAT	638550101	2019	3 11/12/2019	32330649-1	NAT-C 2019	5/5	100 Satisfactory
HCV by NAT	638550101	2019	3 11/12/2019	32330649-1	NAT-C 2019	5/5	100 Satisfactory
HIV by NAT	638550101	2019	3 11/12/2019	32330649-1	NAT-C 2019	5/5	100 Satisfactory
Multiplex by NAT	638550101	2019	3 11/12/2019	32330649-1	NAT-C 2019	5/5	100 Satisfactory
West Nile Virus by NAT	638550101	2019	3 11/12/2019	32330649-1	NAT-C 2019	5/5	100 Satisfactory