UK NEQAS
Leucocyte Immunophenotyping

Sheffield Teaching Hospitals NHS
NHS Foundation Trust

Measurable Residual Disease for CLL by Flow Cytometry (Not Accredited) Programme

All Participant Report

Distribution - 242501 Sample - 067 Participant ID -

Date Issued - 30 April 2024 Closing Date - 21 May 2024 Machine Used - FACSLyric

Trial Comments

This exercise was issued to 89 participants of which 77 (86.5%) returned results at the time of report generation. Of the non returning centres, 4 had requested an extension to the exercise deadline and 1 was a pre notified non return.

Sample Comments

The sample was manufactured by UK NEQAS using a CLL patient sample and a stabilised whole blood unit .

Results and Performance

Percentage MRD Population	Your Results (%)	Robust Mean (%)	Robust SD (%)	Uncertainty of the Assigned Value (Robust Mean)
	0.1525	0.2426	0.0404	± 0.01

Percentage MRD Population	z Score*	Performance Status for this Sample	Performance Status Classification Over 12 Sample Period		
			Satisfactory	Action	Critical
	-2.23	Satisfactory	12	0	0

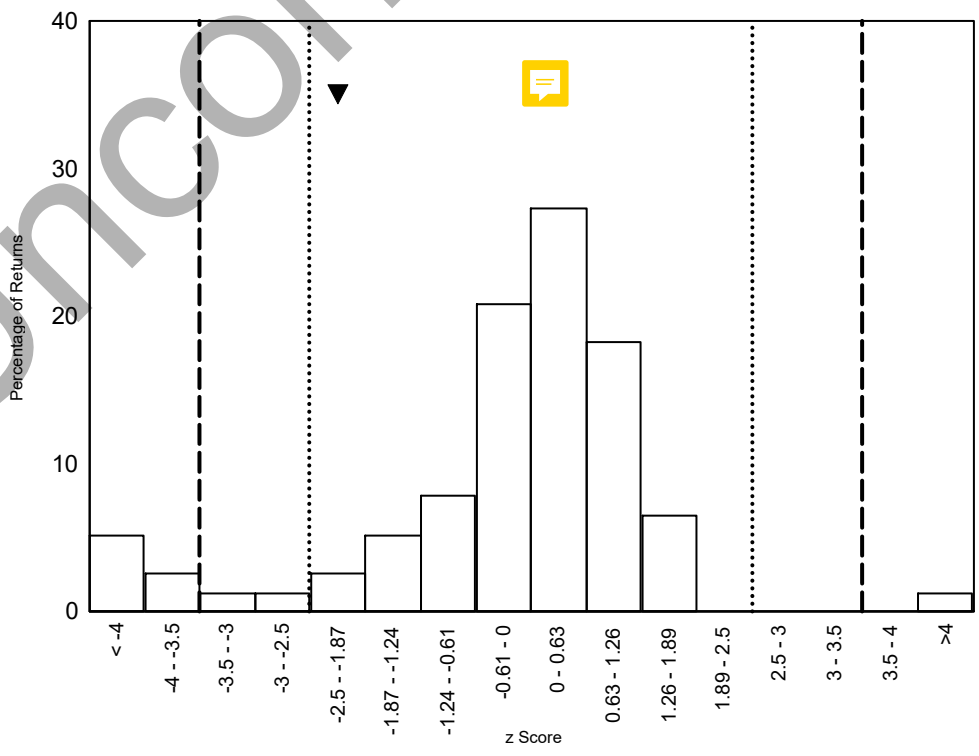
***z Score Limits Definitions**

Please note the scale below is applicable to the tables above and to the z score histograms and Shewhart control charts that follow. It is not applicable to the Cusum control charts.

Histograms of Participant z Scores

Percentage MRD Population -

Please note ▼ denotes your result



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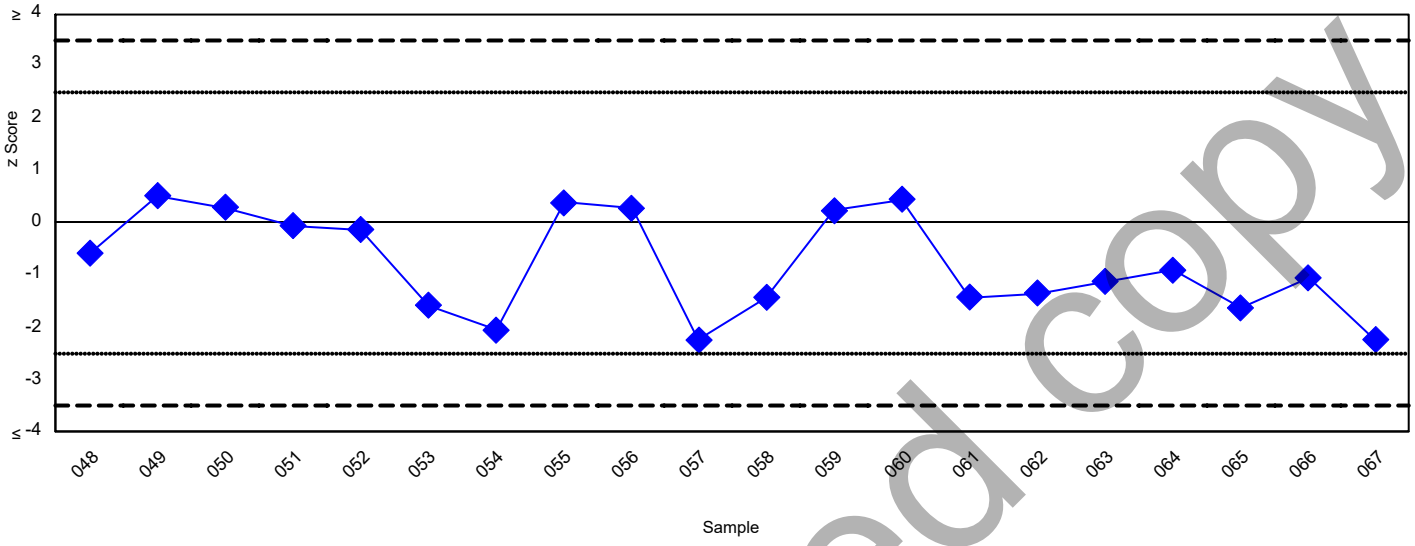
Shewhart Control Charts

(Please note each data point represents a single sample)

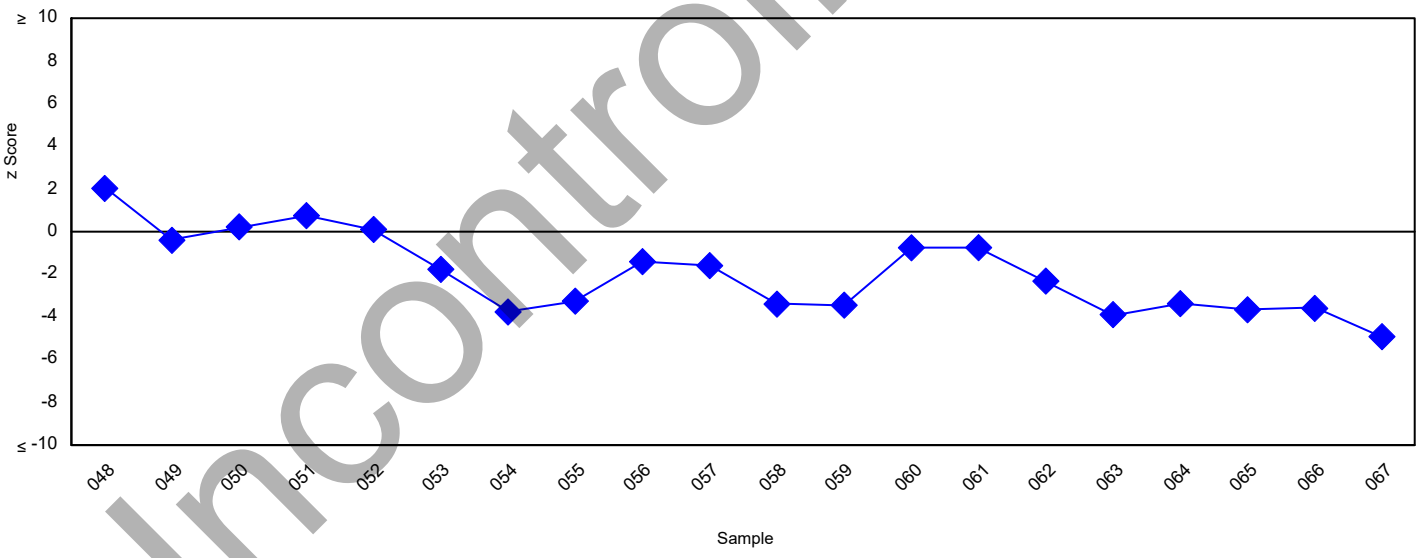
Values (Percentage MRD Population)



Cusum Control Charts

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

Values (Percentage MRD Population)



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Measurable Residual Disease for CLL by Flow Cytometry (Not Accredited) Programme

Flow Cytometer Specific Statistics

(Please note only groups of >0 returns are displayed)

Method	Count Values (%)		
	Returns	Robust Mean	Robust SD
CytoFlex	1	0.2150	0.0000
DxFLEX	7	0.2266	0.0580
FACSCanto	1	0.2500	0.0000
FACSCanto II	22	0.2550	0.0381
FACSLyric	27	0.2481	0.0380
Gallios	1	0.2100	0.0000
LSRFortessa	2	0.2219	0.0612
Navios	16	0.2285	0.0392

MRD Group Specific Statistics

(Please note only groups of >0 returns are displayed)

Method	Count Values (%)		
	Returns	Robust Mean	Robust SD
ERIC	9	0.2667	0.0086
Not-Affiliated	56	0.2361	0.0402
Other	7	0.2547	0.0448

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Measurable Residual Disease for CLL by Flow Cytometry (Not Accredited) Programme

Distribution - 242501 | Sample - 068 | Participant ID -

Date Issued - 30 April 2024 | Closing Date - 21 May 2024 | Machine Used - FACSLyric

Trial Comments

This exercise was issued to 89 participants of which 77 (86.5%) returned results at the time of report generation. Of the non returning centres, 4 had requested an extension to the exercise deadline and 1 was a pre notified non return.

Sample Comments

The sample was manufactured by UK NEQAS using a CLL patient sample and a stabilised whole blood unit .

Results and Performance

Percentage MRD Population	Your Results (%)	Robust Mean (%)	Robust SD (%)	Uncertainty of the Assigned Value (Robust Mean)
	0.0858	0.1218	0.0268	± 0.00

Percentage MRD Population	z Score*	Performance Status for this Sample	Performance Status Classification Over 12 Sample Period		
			Satisfactory	Action	Critical
	-1.34	Satisfactory	12	0	0

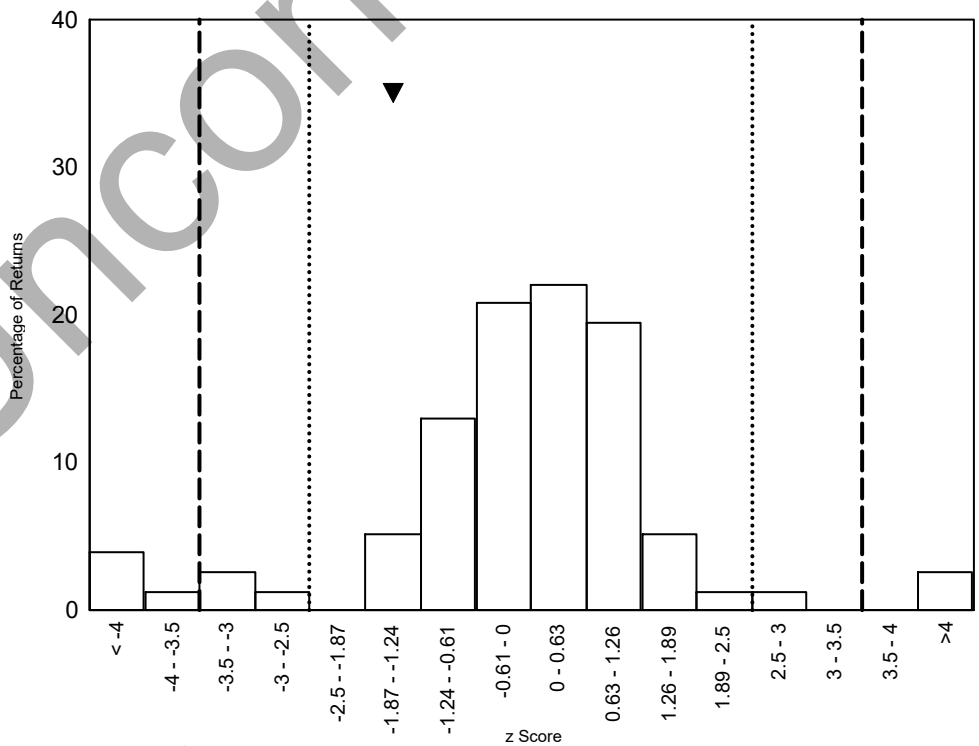
***z Score Limits Definitions**

Please note the scale below is applicable to the tables above and to the z score histograms and Shewhart control charts that follow. It is not applicable to the Cusum control charts.

Histograms of Participant z Scores

Percentage MRD Population -

Please note ▼ denotes your result



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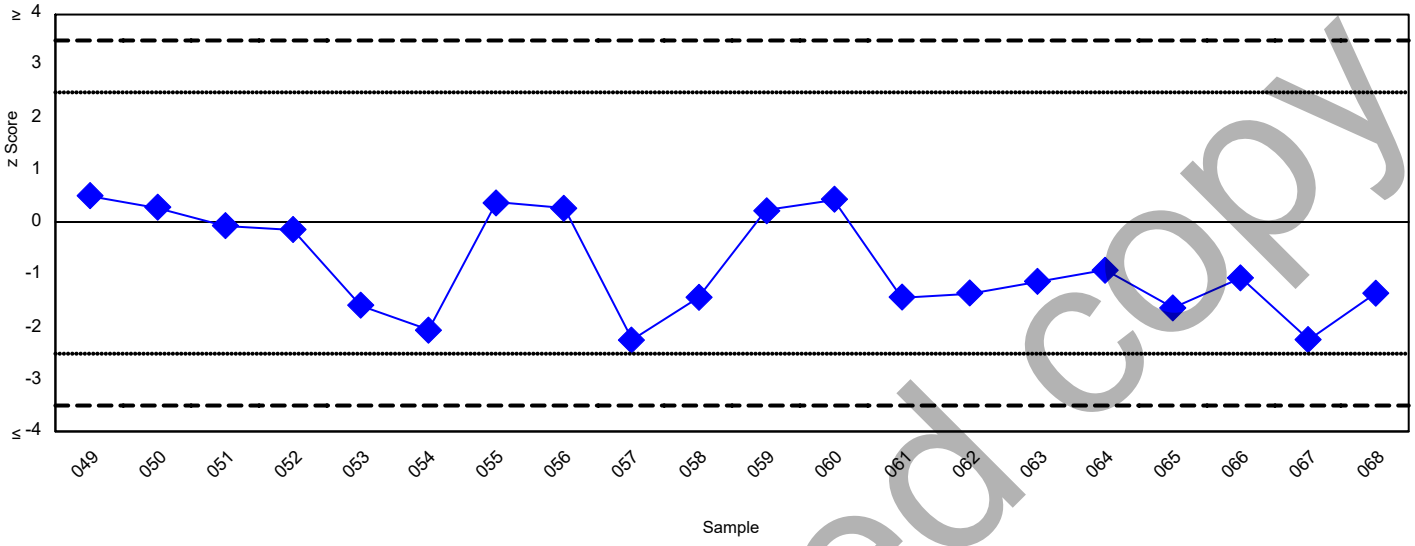
Measurable Residual Disease for CLL by Flow Cytometry (Not Accredited) Programme

Shewhart Control Charts

(Please note each data point represents a single sample)

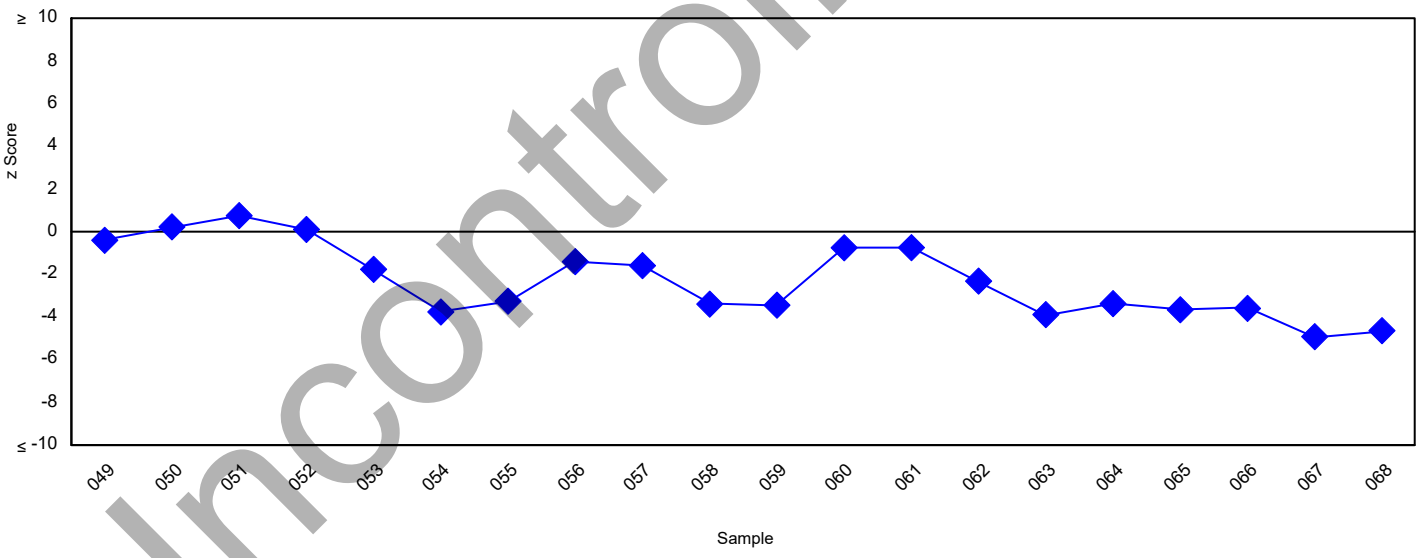
Values (Percentage MRD Population)



Cusum Control Charts

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

Values (Percentage MRD Population)



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Measurable Residual Disease for CLL by Flow Cytometry (Not Accredited) Programme

Flow Cytometer Specific Statistics

(Please note only groups of >0 returns are displayed)

Method	Count Values (%)		
	Returns	Robust Mean	Robust SD
CytoFlex	1	0.1130	0.0000
DxFLEX	7	0.1268	0.0254
FACSCanto II	22	0.1276	0.0207
FACSLyric	28	0.1229	0.0294
Gallios	1	0.1100	0.0000
LSRFortessa	2	0.1241	0.0256
Navios	16	0.1092	0.0335

MRD Group Specific Statistics

(Please note only groups of >0 returns are displayed)

Method	Count Values (%)		
	Returns	Robust Mean	Robust SD
ERIC	9	0.1373	0.0076
Not-Affiliated	55	0.1183	0.0291
Other	8	0.1256	0.0288

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Information with respect to compliance with standards BS EN ISO/IEC 17043:2010

4.8.2 a) The proficiency testing provider for this programme is:
UK NEQAS for Leucocyte Immunophenotyping
Pegasus House, 4th Floor Suite
463A Glossop Road
Sheffield, S10 2QD
United Kingdom
Tel: +44 (0) 114 267 3600
e-mail: amanda.newbould@ukneqasli.co.uk

4.8.2 b) The coordinators of UK NEQAS LI programmes are Mr Liam Whitby (Director) and Mr Stuart Scott (Centre Manager).

4.8.2 c) Person(s) authorizing this report:
Mr Liam Whitby (Director) or Mr Stuart Scott (Centre Manager) of UK NEQAS LI.

4.8.2 d) No activities in relation to this EQA exercise were subcontracted.

4.8.2 g) The UK NEQAS LI Confidentiality Policy can be found in the Quality Manual which is available by contacting the UK NEQAS LI office. Participant details, their results and their performance data remain confidential unless revealed to the relevant NQAAP when a UK participant is identified as having performance issues.

4.8.2 i) All EQA samples are prepared in accordance with strict Standard Operational Procedures by trained personnel proven to ensure homogeneity and stability. Where appropriate/possible EQA samples are tested prior to issue. Where the sample(s) issued is stabilised blood or platelets, pre and post stability testing will have proved sample suitability prior to issue.

4.8.2 l), n), o), r) & s) Please refer to the UK NEQAS LI website at www.ukneqasli.co.uk for detailed information on each programme including the scoring systems applied to assess performance (for BS EN ISO/IEC 17043:2010 accredited programmes only). Where a scoring system refers to the 'consensus result' this means the result reported by most participants for that trial issue. Advice on the interpretation of statistical analyses and the criteria on which performance is measured is also given. Please note that where different methods/procedures are used by different groups of participants these may be displayed within your report, but the same scoring system is applied to all participants irrespective of method/procedure used.

4.8.2 m) We do not assign values against reference materials or calibrants.

4.8.2 q) Details of the programme designs as authorized by The Steering Committee and Specialist Advisory Group can be found on our website at www.ukneqasli.co.uk. The proposed trial issue schedule for each programme is also available.

4.8.2 t) If you would like to discuss the outcomes of this trial issue, please contact UK NEQAS LI using the contact details provided. Alternatively, if you are unhappy with your performance classification for this trial, please find the appeals procedure at www.ukneqasli.co.uk/contact-us/appeals-and-complaints/

4.8.4) The UK NEQAS LI Policy for the Use of Reports by Individuals and Organisations states that all EQA reports are subject to copyright, and, as such, permission must be sought from UK NEQAS LI for the use of any data and/or reports in any media prior to use. See associated policy on the UK NEQAS LI website: http://www.ukneqasli.co.uk/eqa-pt-programmes/new-participant-information/

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