UK NEQAS Update session November 2015

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Update

- Accreditation
- UK NEQAS logo
- Learning points
- Website and web result page improvements
- Pilot scheme updates
- TACT update



ISO 17043

- Assessment June 2015
- Clearances submitted September 2015
- Clearance report received last week
- One outstanding issue
 - Legal entity and logo



UK NEQAS International Quality Expertise

UK NEQAS Feto-Maternal Haemorrhage



Schemes based at Watford GH (BTLP, H, FMH) applied for accreditation as a single unit – same quality management system

UK NEQAS Haematology and Transfusion

West Hertfordshire Hospitals NHS Trust Operating UK NEQAS Haematology and Transfusion



Learning points

- Emergency exercise 15R1
- Mixed field 15R4



15R1 Emergency Exercise

- One extra whole blood sample
- Provide 4 units of red cells in 10 minutes
- Report extent of testing and results of ABO/D if undertaken
- Supplementary questions re component issue and further steps:
 - 4 different patient demographics
 - Women (aged 23 and 75), man (aged 45), child (male 8)



Results

- Return rate 348/396 (87.9%)
- Patient was B D negative



233 (72%) did initial group





Group of FFP issued

ABO group of FFP	% Issuing each ABO group of FFP (n=220) of those undertaking a group			
	Male 45 yrs	Male 8 yrs	Female 23 yrs	Female 75 yrs
Group AB	9%	5%	11%	10%
Group B	88%	82%	88%	89%
Group A	2%	1%	1%	<1%
Group O	1%	0%	0%	1%

Also, 2/92 who did not perform an ABO group selected group O for all 4 patients

BCSH guidelines for use of FFP: Group O FFP should only be given to group O patients



Platelets issued (choice of O D positive and A D positive)

Platelet group	% issuing O or A for each patient type (n=315)			
issueu	Male 45 yrs	Male 8 yrs	Female 23 yrs	Female 75 yrs
A D positive	86%	87%	87%	86%
O D positive	14%	13%	13%	14%

BCSH and SHOT recommend group O as last choice and only if HT neg



15R4 – Mixed Field ABO/D



15R4 – MF ABO/D

Patient 1

• 25:75 A:O (reverse group A)

Patient 2

• 25:75 D pos/neg

Instructions

 Assume all patients have been recently transfused. No patient demographics provided.

AIMS:

- Detection of the MF
- Interpretation of the group if MF is detected



ABO/D dual populations

- Transfusion of ABO/D compatible but non identical blood
- Post HSCT / BMT
- Rarely permanent chimerism or ABO subgroup
- Could be first indication of ABO incompatible transfusion



Sample	Reaction strength recorded (n=383)			
	MF	Strong +	Weak +	Neg
P1: Reaction vs. anti-A	91%	5%	3%	2%
P2:Reaction vs anti-D	57%	5%	10%	28%

Technology	MF anti- A	MIF anti D
DiaMed (n=173 & 177)	91%	72%
BioVue (n=79 & 87)	97%	39%
Capture/LPM (n=8 & 30)	50%	3%
Grifols (n=15 & 15)	100%	67%
Tube (n=14 & 13)	71%	54%

DiaMed Auto/man	MF anti-A	MF anti-D
Auto (n=135)	96%	74%
Manual (n=38)	76%	66%



15R4 P2 MF (D pos/neg 25:75)

Reactions other than MF recorded for Patient 2 vs. anti-D, by technology

Technology	Total	Negative	Weak positive	Strong positive
BioVue	53	33 (62%)	18 (34%)	2 (4%)
DiaMed	49	34 (69%)	3 (6%)	12 (24%)
Grifols	5	5 (100%)	0 (0%)	0 (0%)
LPMP ¹	29	28 (97%)	0 (0%)	1 (3%)²
Tube	6	1 (17%)	3 (50%)	2 (33%)

¹ LPMP = liquid phase microplate, and includes those stating Capture or solid phase ² manual testing



Questions raised

- Why is there a difference in detection rates of MF reactions between ABO and D typing?
- Why is this not consistent within and between technologies?

Antibody affinity Potentiators

Shear forces

Excessive shaking in liquid phase

Centrifugation speeds and time

A combination of these things



Reaction grades vs. interpretation

Reaction grade anti-A	Total	UI	A	Ο
MF	348	70%	30%	0%
Weak +	11	9%	91%	0%
Strong +	17	0%	100%	0%
Neg	9	22%	0%	78%

Reaction grade anti-D	Total	UI	D pos	D variant	D negative
MF	220	85%	11%	3%	<1%
Weak +	39	28%	23%	49%	0%
Strong +	18	0%	100%	0%	0%
Neg	106	0%	0%	1%	99%

BCSH: anomalous reactions should be investigated



Dual populations in clinical practice

- Obvious limitations of EQA... but DP
 - Might not occur in test system used
 - Might not be recognised
 - Might not result in appropriate ABO/D interpretation



Website and data entry developments



Survey data entry

UK NEQAS

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Haematology and Transfusion

P - C Ø UK NEQAS Watford

Who we are

The UK National External Quality Assessment Service (UK NEQAS) is a registered charity offering external quality assessment (EQA) services across all pathology disciplines.

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What we do

The primary aim of UK NEQAS is to maintain and improve performance of diagnostic testing at a high level of proficiency, wherever testing is performed. Participation in EQA is an established part of Quality Assurance and is actively encouraged by professional bodies.

General Haematology Blood Transfusion Laboratory Practice		Feto-Materna	l Haemorrhage
1506RE,1511FBThere are noclose in 1 dayopen distributions		1506F closes in 1 day	
(click for data entry and reports)	(click for data entry and reports)	(click for data	a entry and reports)
More Distributions	General Schedule		Schedule
Exercise Type		Exercise Code	Date of Despatch
ABO/RhD grouping, antibody screen/ider	15R4	20 April 2015	
Antibody screen/identification	15E5	18 May 2015	
Antibody screen/identification	15E6	22 June 2015	
ABO/RhD grouping, antibody screen/ider	ntification, crossmatch and red cell phenotyping	15R7	13 July 2015
Antibody screen/identification		15E8	14 September 2015
ABO/RhD grouping, antibody screen/ider	15R9	12 October 2015	
Antibody screen/identification	15E10	23 November 2015	
ABO/RhD grouping, antibody screen/ider	16R1	18 January 2016	
Antibody screen/identification	16E2	15 February 2016	
Antibody screen/identification	16E3	14 March 2016	

#btsched

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Documents & Downloads

Reports	*
Participant Manuals	*
Presentations	^
UK NEQAS (BT) Symposium 2014 Anti-D Failures - Jane Keidan (PDF)	*
UK NEQAS (BT) Symposium 2014 Do Rules and Regulation Improve Quality? – Jonathan Wallis (PDF)	±
UK NEQAS (BT) Symposium 2014 Immune or Prophylactic Anti-D –Jenny White (PDF)	*
UK NEQAS (BT) Symposium 2014 Implications of the PQA Review for Clinical Laboratories - Ian Barnes ((PDF)
UK NEQAS (BT) Symposium 2014 Interactive Session - Megan Rowley & Clare Milkins (PDF)	*
UK NEQAS (BT) Symposium 2014 Measurement of Anti-D in Pregnancy - D Bruce & F Green (PDF)	*
UK NEQAS (BT) Symposium 2014 NCA Anti-D lg - Megan Rowley (PDF)	*
UK NEQAS (BT) Symposium 2014 NEQAS Update 2014 (PDF)	±
UK NEQAS (BT) Symposium 2014 Results from Anonymous Questionnaire - Clare Milkins (PDF)	*
UK NEQAS (BT) Symposium 2014 Rules and Regulations - James Taylor (PDF)	*

The Organisation Schemes ▼ Schedule F.A.Q. Contact Us Documents & Downloads

Everything else

Surveys

Instructions from manufacturers

General Haematology	Blood Transfusion
🖂 technical queries email	🖂 technical queries er
🖂 other queries email	🖂 other queries em
L (+44) (0) 1923 217878	(+44) (0) 1923 2179
🔓 (+44) (0) 1923 217879	🗎 (+44) (0) 1923 2179

Feto-Maternal Haemorrhage
🖂 technical queries ema
🖂 other queries email
\$ (+44) (0) 1923 217933

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🗎 (+44) (0) 1923 217879

						Q Search
UK NEQAS Haematology Scheme	S es	UK NEQA	AS National Exte	ernal Quality As	sessment Site	version:
Sample Entry Deta	ails Ba	ack to List Summary		<u>Blank data entry form</u> <u>Email scheme</u>	Antibody ID UI Rules a Data Entry Instruction	and Faxheader 15
	В	LOOD TRANSFU	ISION LABOR	ATORY PRACT	FICE	
Distribution Number: 15R4	Participant: 26000	Issued: 20/04/2015 Closi	ing: 22/04/2015 Receiv	ved Date (dd/mm/yyyy):	Analysed Date	e (dd/mm/yyyy):
Sample Quality P	Patient 1 Pa	tient 2 Patient 3	Techniques	Phenotyping 20.	/04/2015	
		Patie	ent 1 - ABO/D Ty	yping		
anti A	anti B	anti D1	Reaction grade vs.	Ctrl	anti iw	A Colle
Not Stated • N	lot Stated 🔹	Not Stated -	Not Stated 🔹	Not Stated 🔹	Not Stated -	Not Stated
Not Stated	Str	ong = strong positive (3-4+), We	ak = weak positive (w-2+),	MF = mixed field (dual popul	ation)	
Negative		А	BO / D Interpretation			
Strong Positive		•		Pos 👻		
Weak Positive			Notes			
2++	tial D					

³ Unable to test is only to be used in situations where the sample is unsuitable for testing and a repeat cannot be obtained before the closing date

NB - Where more than one technique is used to arrive at an interpretation, mark the most appropriate reaction grades for the final interpretation, and if reaction grades differ, please email the scheme via the link above

Haematology	Schemes		UK NEQA	S National Ex	cternal Quali	ty Assessing	ent Site	versior
Sample Ent	try Details	Back to Lis	st Summary	R	eports Blank data	entry form Email Australia Email	ntibody ID UI Rules ata Entry Instruction	and Faxheader 15
		BLOO	D TRANSFL	JSION LABO	DRATORY P	RACTICE		
istribution Numbe	: 15R9 Participa	nt: 26000 Issued:	12/10/2015 Closin	ng: 26/10/2015 Re	eceived Date (dd/mm/y	уууу):	Analysed Date (d	d/mm/yyyy):
ample Quality	Patient 1	Patie	ent 2	Patient 3	Technique	s Phe	notyping	13/10/2015
			Palle	ent I - ABO/D	ryping			
			Reaction grade v	S.			ABO/D	Interpretation
anti-A	anti-B	anti-D1	Reaction grade v anti-D2	s. Ctrl	A Cells	B Cells	ABO / D	Interpretation



Antibody ID data entry

Patient 2 -	Antibody I	dentification							
Antibody specificities positively identified (currently a maximum of 2 in any sample)		Specificities that cannot be excluded	Cannot exclude						
\checkmark D Le^a C Le^b $c+/-E$ \checkmark E Fy^b $e+/-C$ Jk^a M Jk^b N Ul^{-1} S C^w s Kp^a P_1 Enz non-specific Lu^a $VVra$ K vra	D Fyra	□ D S □ C ▼ K □ c+/-E □ Fy ^a □ E □ Fy ^b □ e+/-C □ Jk ^a □ M □ Jk ^b □ S	к						
1	UI = Unable to inte	rpret							
You may indicate commonly encountered antibodies of potential clinical significance that cannot be positively identified but might be present (cannot be excluded) based on your testing and the phenotype provided.									
If you wish these to be taken into account for scoring and performance monitoring you must the instruct	st also make a UI s tions on the down	ubmission by clicking on the " Antibody ID UI rules and faxheader " link above a loadable form.	nd following						



Pilot scheme developments



ABO titration pilot (1)

- 2010 2015 pilot aim to support ABOi renal transplant
- 'Standard' technique initially to facilitate EQA
- Still variation by technique... looking for trends 2014/15 data





ABO titration pilot (2)

- Work with renal specialists & NHSBT living donor strategy group to find a way to achieve standardisation across centres
 - equitable access to ABOi renal transplant programmes
 - safe cut-off limits on day of transplant
- NIBSC reference preparations HT anti-A and HT anti-B
 - Now accepted as WHO standard reference preparations
- Shadow scoring for standard technique in 2014/15 (and any other group large enough)
 - Based on distance from median result (one dilution either side OK)
 - Start in 2016... will allow us to move ABOT to a full UK NEQAS scheme



Red cell genotyping (pre-pilot UK NEQAS / ISBT)

- 14G1, 3 samples (unselected)
 - 55 labs 30 countries, 52 returns
 - D, Cc, Ee, MN, Ss, Kk, Fy^a Fy^b Fy, Jk^a Jk^b, Do^a Do^b
 - Genotype / predicted phenotype / Qs on practice
 - 6 labs with errors testing, interpretation, procedure
 - Terminology!!!
- 15G1, 1 sample (selected Fy(a-b-))
 - Same set of labs + others, same tests, 52 returns
 - New format for results, following terminology nightmare last time!
 - 8 labs with errors (4 to do with Fy)
- Rh variant found in one sample in each exercise!
- EQA Scheme needed for routine testing
- Pilot planned for 2016 UK and non-UK



nternational Society of Blood Transfusion



DAT pre-pilot

Summary of findings from 15R7 (distributed 13/07/2015) 15R9 (distributed 12/10/2015)



Introduction

- Two sets of pre-pilot samples for DAT sent to laboratories in the UK and ROI and some overseas with 15R7 and 15R9
- Initially to assess sample stability over a period of 2 weeks
- Participants were asked to test samples on receipt and again a week later
- 15R7
 - DAT 1: Negative
 - DAT 2: Positive (4+) coated with monoclonal anti-D
- 15R9
 - DAT 1: Positive (2+) coated with a weak monoclonal anti-D
 - DAT 2: Positive (2+) coated with polyclonal anti-K



15R7 DAT 2

DAT 2 vs Poly	/specific AHG
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Re		Re	sult wee	ek 2		
Result	No. labs	4+	3+	2+	1+	Neg
4+	77 ¹	65	9	1	0	1 ²
3+	11	4	6	1	0	0
2+	1	0	0	1	0	0

¹One laboratory did not report a result vs. polyspecific AHG in week 2 ²One laboratory reported a negative reaction for week 2, due to a transcription error.

87.4% of laboratories reported either the same reaction strength or stronger in week 2 as in week 1



Conclusions

- Results from both pre-pilot surveys have shown good sample stability for DAT
- Reported data for sample quality shows even where some haemolysis of the samples had occurred, the laboratories reporting unsatisfactory sample quality obtained the expected results
- A full report with data analysis will be issued soon
- UK NEQAS BTLP will continue piloting next year to include some complement coated cells
- Proceeding to a full DAT pilot in the near future





Claire Whitham MSc MIBMS, Snr EQA Scientist, UK NEQAS BTLP On behalf of the TACT team, UK NEQAS BTLP.

TACT Membership



- As of 26/10/2015, TACT has more than 1400 members
- Funding is available for developments into the next financial year
- Cost of memberships can be incorporated into BTLP re-registration for 2015/2016
- Half price membership offered between 01/10/2015 to 31/03/2016



Recent developments



- Performance Dashboard enhancement
- Manager's drill down review
- Enzyme panel functionality
- Final interpretation stage for antibody ID



Individual D	ashboard BTLP-T 🛪	+											(or		^
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Scenarios	Individual Dashboard	d Lab Overview	Resources											1	
	Name	Test Admin accou	int	Assessme	ent window		1 year(s)		Scenarios	incomplete	51				
	Staff ID	testadmin		Assessme	nt window start		🟥 10 Aug 2014								
	Site	WHT		Scenarios	attempted		127								
						_									
			(Engagement & Outcom	e Participations										
				Section	rarget	Correct	Incorrect	Engagement	Outcome						
				Request Acceptance	40	1	5 6	•							
				Grouping (ABO)	40	5	5 21	•	۲						
				Rh/D		5	5 20	•							
				Antibody Screen		5		•							
				Antibody Identification	10	1	18	•	۲						
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I Individual Dashboard | BTLP-T... × +

Sc Scenarios Individual Dashboard Lab Overview Resources

You are viewing the results for the selected participation.

Return to member overview

		Participant te	started 07/08/2015 12:21:41 stadmin Completed 07/08/2015 12:29:23		
Location	Action	Request	Submitted data	Elapsed time	Visual data
Laboratory	Scenario continued			00:00:01	
Laboratory	Move to in-tray			00:06:11	~
In-tray	Accept Request	80086	Patient request 80086 accepted.	00:06:17	/ • \
In-tray	Move to LIMS			00:06:19	
LIMS	Commissioned grouping test	80086		00:06:26	
LIMS	Commissioned antibody screening test	80086		00:06:26	
LIMS	Selected request for issue	80086		00:06:29	
LIMS	Move to Analyser			00:06:32	
Analyser	Grouping	80086	ABO: B, RhD: POS	00:07:08	1
Analyser	Antibody Screening	80086	Screening status: NEG	00:07:15	• /
Analyser	Move to Fridge			00:07:19	
Stock fridge		80086	ABO: B, RhD: Neg, Phenotype: D-C-E-c+e+, Negative antigens: K, Fy ⁸ , CMV	00:07:27	P
Stock fridge	Stock Issue	80086	ABO: O, RhD: Neg. Phenotype: D-C-E-c+e+, Negative antigens: HT, K, Fy ^a , Jk ^a , M, CMV	00:07:35	•/
Fridge	Move to Laboratory			00:07:40	Y
Laboratory	Exit scenario			00:07:41	





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15/09/2015

Scenario | BTLP-TACT - Mozilla Firefox Tact.dev.certus-tech.com/tact/participate/participate?participationId=73165 Laboratory In-tray Stock Fridge Analyse testadmin Requests Scenario Time 11:23:22am × 1 2 3 4 5 6 7 8 9 10 Laboratory In-tray LIMS Analyser Stock Telephone testadmin Fridge * III. Enzyme antibody identification result for request 84415: Determine which antibodies are present from the Gel cards a An antibody ID panel profile is available (button below) to as Analyser Result list -If you cannot confirm any antibody specificitiy(ies) even thou below: O Antibody identification result was submitted. Click a request number to view the test results. Hospital Number Hospital **Request Number** Request Final Ab ID NEQ02468 **BTLP TACT** 96174 NEQ02468 **BTLP TACT** HUUJU Ab screen NEQ02468 **BTLP TACT** W- P-11:23 e 63

Future developments

- More ABO/D grouping anomalies
- Manager's ability to alter the outcome indicator where a red mark was awarded
- Increased variety and new combinations of antibody specificities
- Group check sample requesting
- Manual testing
- New scenario type MH/emergency situation

