

Working To drive excellence in care for **together** our patients and communities

Jennifer Mills 17/03/2024



Session Outline



How do we get from a sample, to a bag of blood?

- Sample Requirements.
- Sample Testing:
 - ABO, RhD, Antibody ID etc.
- Crossmatching Red Cells.
- Issuing Blood Components and Products.
 - Platelets, Plasma, Drugs.



Guideline



Official Journal of the British Blood Transfusion Society





Transfusion Medicine

GUIDELINES

Guidelines for pre-transfusion compatibility procedures in blood transfusion laboratories*

British Committee for Standards in Haematology

C. Milkins, ¹ J. Berryman, ² C. Cantwell, ³ C. Elliott, ⁴ R. Haggas, ⁵ J. Jones, ⁶ M. Rowley, ^{3,7} M. Williams ⁸ & N. Win ⁹

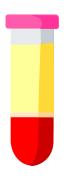
¹UK NEQAS (BTLP), West Herts Hospitals NHS Trust, Watford, UK, ²Department of Blood Transfusion, University College London Hospitals, NHS Foundation Trust, London, UK, ³Department of Blood Transfusion, Imperial College Healthcare NHS Trust, London, UK, ⁴Department of Blood Transfusion, South Tees Healthcare Trust, Middlesborough, UK, ⁵Department of Blood Transfusion, Leeds teaching Hospital NHS Trust, Leeds, UK, ⁶Welsh Blood Service, Cardiff, UK, ⁷Colindale Centre, NHSBT, London, UK, ⁸Leeds Centre, NHSBT, Leeds, UK, and ⁹Tooting Centre, NHSBT, Tooting, UK

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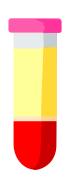
Part 1: Sample Requirements



- EDTA Anti-coagulated Blood Samples (Pink Top)
- Following ESSENTIAL Information:
 - First Name and Surname (No Middle Name Required)
 - Date of Birth
 - NHS Number
 - Date of Collection
 - Time of Collection
 - Signature
- There can be NO AMMENDMENTS to the essential Information.



Part 1: Why is this important?



Essential Information	Reason
First Name, Surname and DOB.	Identifies the patient.
NHS Number	NHS numbers are unique. Other numbers get
Incomplete/Amende	d Information — Increased rick of WRIT



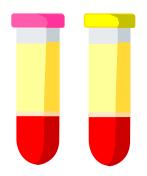
Increased risk of WBIT = Increased risk of DEATH!



Time of Collection	Ensures the lab can provide blood for the maximum amount of time.
Signature	Audit trail for the person providing the sample, in case of any errors discovered



Part 1: Why 2 Samples?



1 in 2000
samples are
WBITs
Even when all
the information
is perfect!

2 samples protects patients!

- Confirms that the 1st sample wasn't a WBIT.
- ABO matched blood is provided only after 2 samples.
 - ABO and Rh groups must match.

0-1 Samples

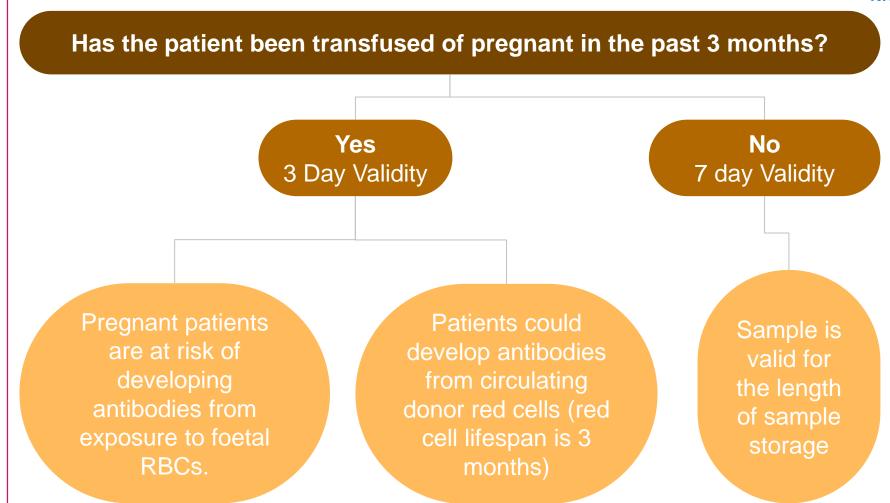
- Patients ONLY get group O.
- Patients who can be pregnant get RhD-.
- Other patients can receive RhD+ units.

Yellow Sample Rule

- Yellow top samples (provided by the lab) for patients on the same ward within 12 hours.
- This ensures 2 samples taken separately!



Part 1: Sample Validity.





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NHS Trust

Part 2: Types of Testing.



These are the most important tests for blood selection.

3 cells to ID significant alloantibodies.

Antibody Identification 10 cells+ to determining antibody specificity.

Extended Phenotype ID red cell phenotype for other antigens e.g., Duffy, MNS

ABO and RhD group and —

Direct Antiglobulin Test (DAT) ID autoantibodies.

Rh/K Phenotype Gives C/c, E/e and K/k phenotype.

Referrals

Samples sent to reference labs for complex and specialist testing.



Part 2: Methods of Testing



- Fully automated ABO/D group & antibody screen
 - Safer
 - Reduced Pipette errors
 - Automated Transcription of results
 - Difficult to mix up patient samples.
- Centrifugation ~ 6 mins
- ABO/D group ~ 8 mins
- Antibody screen ~ 20 mins

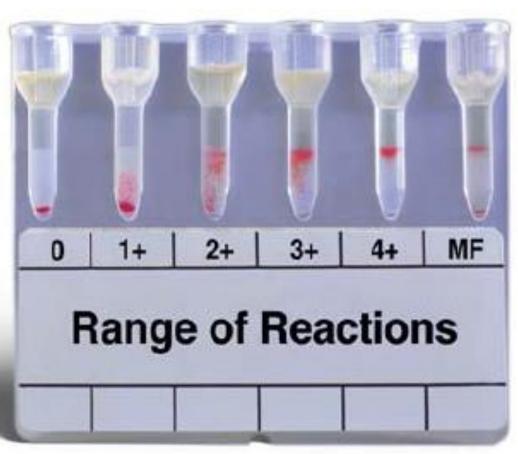




Part 2: Methods of Testing



- Sample needed
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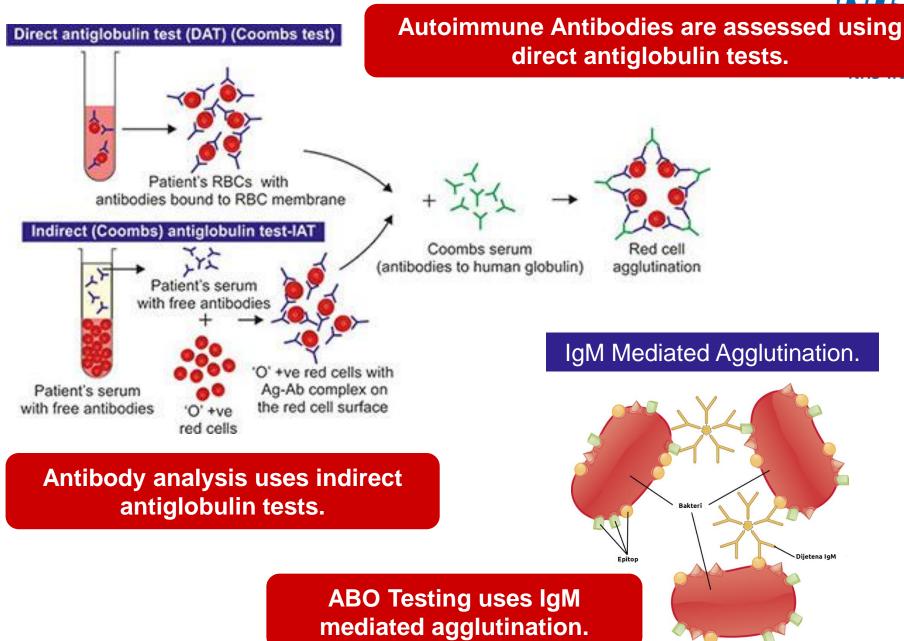
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Part 2: Methods of Testing







Part 2: ABO/RhD



- Identifies the patients ABO and RhD antigens.
- Identifies the presence of innate Anti-A or Anti-B using cells.
 - Should correlate with the patient's Antigens.

PATIENT RED CELLS

PATIENT PLASMA





Part 2: ABO/RhD





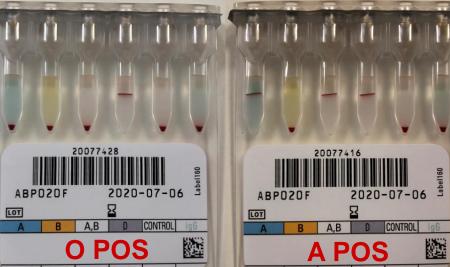
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Part 2: ABO/RhD



Abnormal Results:

- Transfusion with different ABO groups (Emergency units etc)
- Transplant patients
- Elderly/Very young (>4 months)
- Strong autoantibodies
- RhD and ABO Variants
- Abnormal results should be resolved before blood can be crossmatched.
 - If it can't be resolved, Group O will be given.
 - Most abnormal results excludes a patient from electronic issue.



Part 2: Antibody Screen



- 3 Cell Screen.
 - Uses to identify the presence of antibodies
- Uses IAT technique
- Reagent Cells are selected to ensure:
 - All clinically significant antigens are present
 - Homozygous expression of antigens for: Duffy, Kidd, S/s and k
 - All Group O (to avoid Anti-A and B interference)

						Rh	ı-hr				•			KELL			DI	JFFY		KIDD	Sex Links	Ĺ	EWIS		М	NS		P	LUT	HERAN	Special AntigenTyping		Te
Cell#	Rh-hr	Donor Number	D	С	Ε	С	е	f	Cv	v	К	k	K	(ра Кр	b Js	a Js	Fy	a Fyl	Jk	Jkb	Xg ⁸	Le	a Leb	s	s	М	N	P ₁	Lu	Lu ^b		Cell#	
1	R1wR1	324414	+	+	0	0	+	0	+	0	+	+	-	0 +	1	+	+	0	+	+	+	0	+	+	0	+	0	0	0	+		1	
2	R2R2	313967	+	0	+	+	0	0	0	0	+	+	.	0 +	1	+	+	+	0	+	+	+	0	+	+	+	+	+	0	+		2	
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Reagent Red Blood Cells 0.8% Surgiscreen® ©Ortho Clinical Diagnostics 2010

8SS509 EXP. DATE 2020-03-17 CCYY-MM-DD

ANTIGRAM® Antigen Profile 635201161



Part 2: Antibody Identification



- Uses 10+ cells with known phenotypes to identify the antibody/antibodies.
 - Also includes an "auto" cell.
- IAT Technique
- Reagent red cells have the same requirements as screening cells.
- Antibody specificity can be assigned IF:
 - The plasma is reactive with at least two examples of reagent red cells expressing the antigen
 - The plasma is non- reactive with at least two examples of reagent red cells lacking the antigen.



Part 2: Antibody Identification



- Antibodies can be hard to identify, especially if there's more than one present.
- In addition to the basic IAT, we also use enzyme treated cells.
 - Papain, Bromelain, Ficin digest extracellular proteins.

Enhanced	Decreased	Unaffected
ABO-related -ABO/H Systems -Lewis System -I System -P1PK/GLOB Rh System Kidd System	MNS System Duffy System	Kell System



Part 2: **Antibody** Identification



Patient's Name	Patient 1	Ref. No.	Sample No.	Conclusion		
D.O.B.				Tested by	Date	

	Rh	С	D	E	С	e	Cw	м	N	s	s	P1	Lua	к	k	Kpa	Lea	Leb	Fya	Fyb	Jka	Jkb	Other	IAT			
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1	R ₁ WR ₁	+	+	0	0	+	+	0	+	0	+	0	0	0	+	0	0	+	+	0	0	+		4	4		
2	R ₁ R ₁	+	+	0	0	+	0	+	0	+	0	2	0	+	+	0	0	+	0	+	+	0		4	4		
3	R ₂ R ₂	0	+	+	+	0	0	0	+	0	+	2	0	0	+	0	0	+	0	+	0	+		4	4		
4	r'r	+	0	0	+	+	0	+	0	+	0	1	0	0	+	0	0	+	0	+	0	+		0	0	П	
5	r"r	0	0	+	+	+	0	+	0	+	0	4	0	0	+	0	+	0	+	0	+	0	Cob+ HLA+	0	0	П	
6	rr	0	0	0	+	+	0	+	+	0	+	2	0	+	0	0	0	0	+	0	0	+		0	0		
7	rr	0	0	0	+	+	0	0	+	0	+	0	0	+	+	0	0	+	0	+	+	0		0	0		
8	rr	0	0	0	+	+	0	0	+	0	+	3	0	0	+	+	+	0	+	0	+	0		0	0		
9	rr	0	0	0	+	+	0	0	+	+	0	2	0	0	+	0	0	0	0	+	0	+	Cob+	0	0		
10	rr	0	0	0	+	+	0	+	0	0	+	2	+	0	+	0	+	0	+	0	0	+		0	0		
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Anti-D Detected in IAT and Enzyme.



Part 2: Antibody Identification



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.O.B.					-	_													Teste	d by			Date			
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	Rh	С	D	Е	С	е	Cw	М	N	s	s	P1	Lu³	к	k	Kpa	Lea	Leb	Fya	Fyb	Jka	Jkb	Other			eve:
1	R ₁ WR ₁	+	+	0	0	+	+	0	+	0	+	0	0	0	+	0	0	+	+	0	0	+		1	3	0
2	R ₁ R ₁	+	+	0	0	+	0	+	0	+	0	2	0	+	+	0	0	+	0	+	+	0			5	0
3	R ₂ R ₂	0	+	+	+	0	0	0	+	0	+	2	0	0	+	0	0	+	0	+	0	+			0	0
4	r'r	+	0	0	+	+	0	+	0	+	0	1	0	0	+	0	0	+	0	+	0	+		_	5	0
5	r''r	0	0	+	+	+	0	+	0	+	0	4	0	0	+	0	+	0	+	0	+	0	Cob+ HLA+	3	3	0
6	rr	0	0	0	+	+	0	+	+	0	+	2	0	+	0	0	0	0	+	0	0	+		1	3	0
7	rr	0	0	0	+	+	0	0	+	0	+	0	0	+	+	0	0	+	0	+	+	0		-	5	0
8	rr	0	0	0	+	+	0	0	+	0	+	3	0	0	+	+	+	0	+	0	+	0		- 4	3	0
9	rr	0	0	0	+	+	0	0	+	+	0	2	0	0	+	0	0	0	0	+	0	+	Cob+	(5	Ö
10	rr	0	0	0	+	+	0	+	0	0	+	2	+	0	+	0	+	0	+	0	0	+		1	3	0
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Anti-Fya detected in IAT. No antibodies detected in Enzyme. This confirms the Anti-Fya, and excludes many other possible specificities e.g. Anti-Cw.



Part 2: Antibody Identification

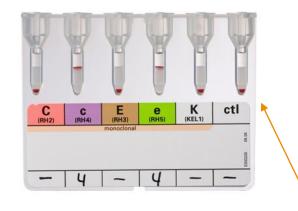


Patient	's Name	П			•	2				Ref.	No.				San	ple No.			Conc	lusion					
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1	R ₁ WR ₁	+	+	0	0	+	+	0	+	. 0	+	0	0	0	+	0	0	+	+	0	. 0	+		4	4
2	R ₁ R ₁	+	+	0	0	+	0	+	0	+	0	2	0	+	+	0	0	+	0	+	+	0		4	4
3	R ₂ R ₂	0	+	+	+	0	0	0	+	0	+	2	0	0	+	0	0	+	0	+	0	+		4	4
4	r'r	+	0	0	+	+	0	+	0	+	0	1	0	0	+	0	0	+	0	+	0	+		2	0
5	r"r	0	0	+	+	+	0	+	0	+	0	4	0	0	+	0	+	0	+	0	+	0	Cob+ HLA+	2	0
6	rr	0	0	0	+	+	0	+	+	0	+	2	0	+	0	0	0	0	+	0	0	+		2	0
7	rr	0	0	0	+	+	0	0	+	0	+	0	0	+	+	0	0	+	0	+	+	0		0	0
8	rr	0	0	0	+	+	0	0	+	0	+	3	0	0	+	+	+	0	+	0	+	0	S.	0	0
9	rr	0	0	0	+	+	0	0	+	+	0	2	0	0	+	0	0	0	0	+	0	+	Cob+	0	0
10	rr	0	0	0	+	+	0	+	0	0	+	2	+	0	+	0	+	0	+	0	0	+		2	0
																							AUTO	0	/

Anti-D detected by IAT and Enzyme. Anti-M detected in IAT only.

Part 2: Other Testing







	Rh/K Phenotype	Extended Phenotype	DAT
Principle	IAT	IAT	DAT
Value of Test	Provides phenotype for RhCE and K.	Phenotype for MNS, Kidd, Duffy etc.	Confirms autoimmune haemolytic anaemia and gives specificity.
Cause for atypical results.	Recent transfusionStem cell transplanVariants		IgM autoantibodies can cause pan-reactive agglutination





Part 3: Crossmatching



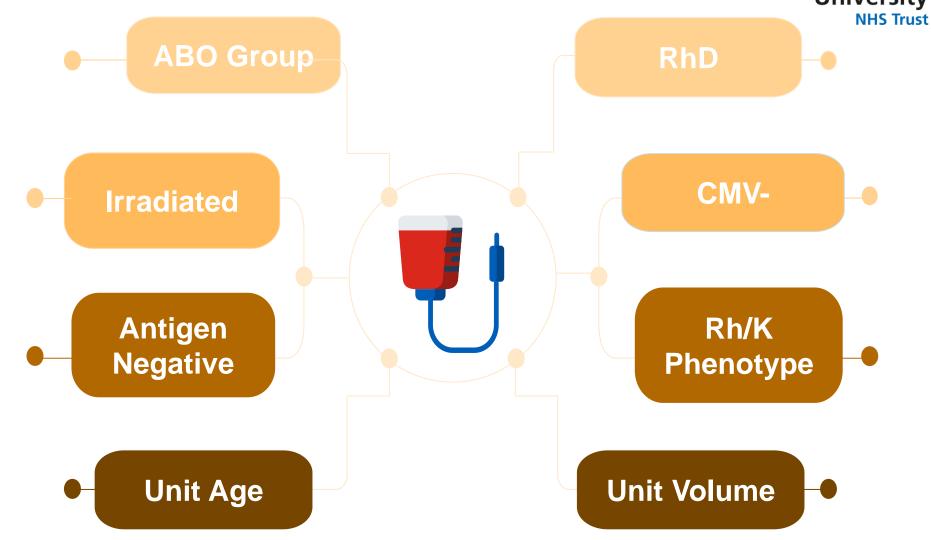
- Crossmatching refers to Red Cells ONLY.
 - All other components are "issued"
- Checking that donor red cells will not react with recipient plasma.
- Requests can be made via:
 - Initial Request (ICE)
 - Minestrone Add on System
 - Phone calls to the Lab
- Crossmatches can take 10mins to 24hours+
 - Emergency units are always available!



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Part 3: Selecting Red Cells.







Part 3: Electronic Issue



- "Theoretical Crossmatch"
 - Assumption that units will be safe based patient info.
 - The lab still selects the units and perform ID checks etc.
- Patients and Units must meet very strict requirements:
 - Units booked into the lab using barcodes
 - Unaltered, automated tests results ONLY
 - Historical blood group required
 - NO ANTIBODIES!
- Units can be issued in 10 minutes.
 - But is it safe?

HTRs with EI are estimated at 1 in 500,000 to 1 in 1,000,000 transfusions (ISBT 2017)



Part 3: Serological Crossmatch



- IAT Test.
 - Physically mix patient plasma with aliquot of donor red cells.
- Only performed if patient is unsuitable for EI.

When?

Historical or current antibody

ABO incompatible solid organ transplant for 3 months.

HSCT transplant

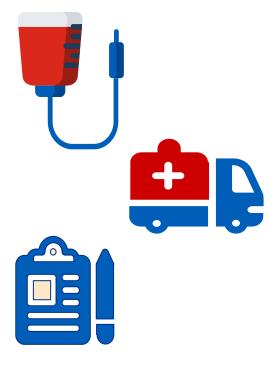
Neonates with maternal antibody present

ABO/D grouping anomalies

- Requires a negative and weak control to ensure accuracy.
- Takes 45 mins to 1 hour.



Part 3: Difficult Crossmatches



- Sometimes we can't crossmatch units in the lab.
 - Pan-reactive Antibodies.
 - Autoantibodies
 - Complex Antibody Profiles.
- So, what can we do?

Is it Urgent?

No Send to RCI

Red cell Immunohematology in Bristol can complete further testing to identify all relevant antibodies and crossmatch "safe" units. Concessionary Release

Yes

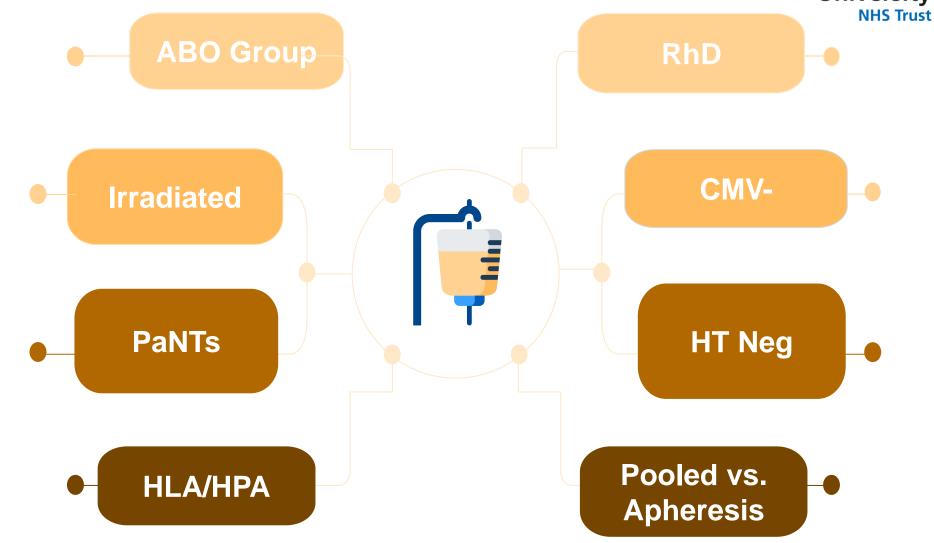
No more incompatible than own or best matched units with haematology approval.



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Part 3: Selecting Other Components.

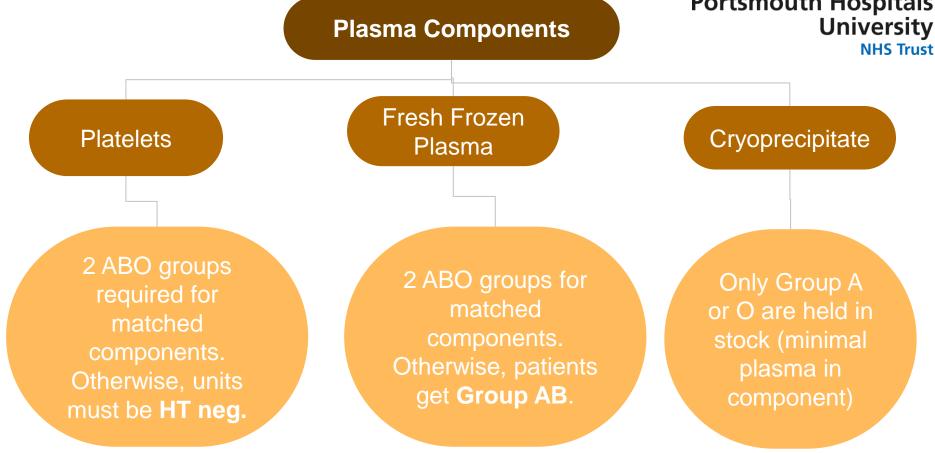






Part 4: Issuing **Other** Components



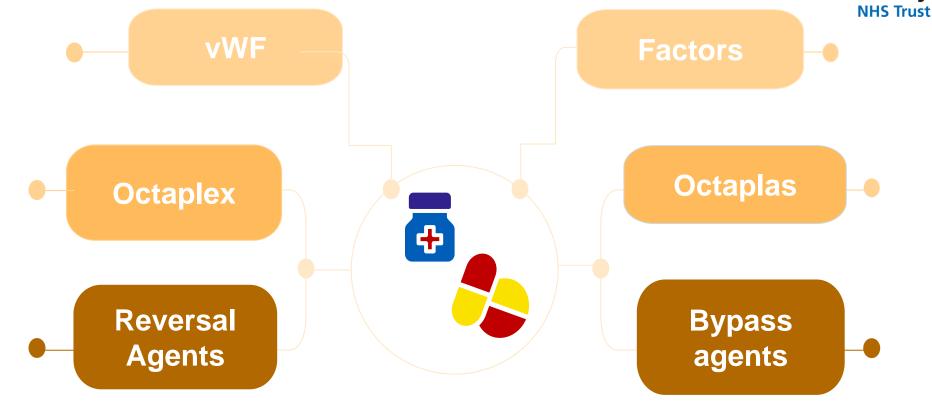


- Units are not crossmatched.
- Issuing takes 10-15 mins.
- Authorisation from Haematology required (with exceptions!)



Part 5: Issuing Blood Products.





- Blood derived products.
- Agreed by haematologists or haemostasis CNS'.
- Octaplex is also stored in Resus.



Key Points to Remember



- The lab needs two samples to provide ABO matched units.
 - But Emergency group O is always available.
- Laboratory testing determines:
 - ABO and RhD
 - Antibody presence and specificity
 - Other red cell phenotypes needed for safe transfusion
- Biomedical Scientists are responsible for meeting patient requirements.
 - So, communicate if you think something is wrong!
- Red cell crossmatching can take up to 24 hours.
 - So, if your patient is complex- give us plenty of warning.

No one should die from a lack of blood!
We will always provide something for your patient!



The End!

Thank you for listening.

Any Questions?



Working To drive excellence in care for together our patients and communities