SHOT & Anti-D

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Anti-D Ig Prophylaxis

- Post-delivery anti-D Ig prophylaxis for RhD negative women began in the UK in 1969
- The programme has been a huge success
- Deaths due to haemolytic disease:
 - 320 / 100,000 births in the 1940s
 - -46 / 100,000 births pre-1969
 - -18.4 / 100,000 births by 1977
 - -1.6 / 100,000 births by 1990

....but RhD alloimmunisation continues to occur

 Maternal anti-D level 10.5iu/ml

• Unmonitored pregnancy

 Bilirubin reached exchange trigger by end of day 1 Liumbruno et al Review of antenatal prophylaxis Blood Transfusion 2010; 8: 8-16

- Risk of sensitisation if Baby RhD pos and ABOcompatible – 16%
- Risk of sensitisation if baby RhD pos and ABOincompatible – 2% (so ABO antibodies are protective)
- 92% women develop immune anti-D AFTER 28 weeks gestation

Liumbruno et al Review of antenatal prophylaxis Blood Transfusion 2010; 8: 8-16

- Foetal red blood cells (typically <0.1mL) are found in;
 - 3% women in first trimester
 - 12% women in second trimester
 - 45% women in 3rd trimester
 - ->50% women at delivery

Feto-Maternal Haemorrhages (FMH)

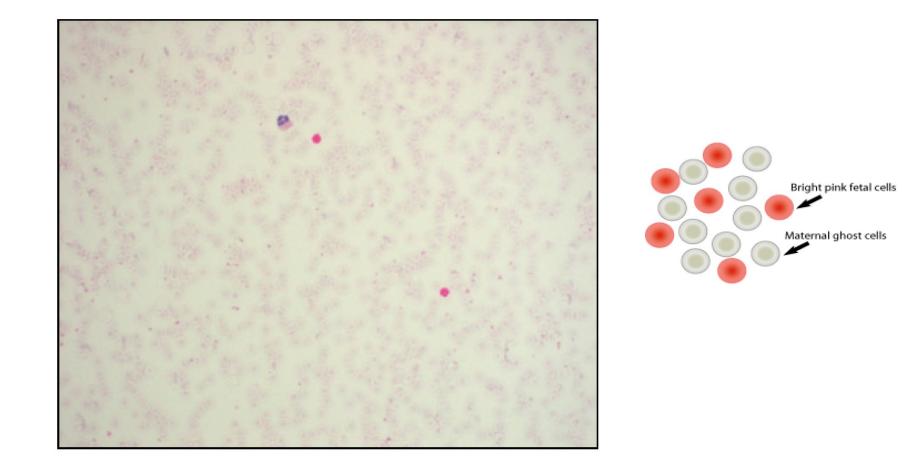
• 1st and 2nd Trimester:

- Miscarriage / Ectopic / TOP / CVS / Amnio
- 0.05mL in 5% 0.5mL in 2%

• After 20 weeks:

- APH / Abruption / Trauma / Amnio / ECV
- <2mL in 98% >30mL in 0.03%
- At Delivery:
 - > 3mL in 1% >10mL in 0.03%
 - Increased by C-section, Cell Salvage or manual removal of placenta

The FMH (Kleihauer / Acid Elution) Test



Anti-D - What Guidelines are there ?

- BCSH Dec 2013
- RCOG
 - Green Top 22, 2011– Use of anti-D
 - Green Top 47, 2008/9 Transfusion in Obstetrics
 - Green Top 38, 2010 (molar pregnancies)
- NICE
 - Antenatal Care / Postnatal Care
 - Routine Antenatal Anti-D Prophylaxis
 - Early miscarriage & Ectopic pregnancy

What SHOULD we be doing.....

- <12/13 weeks gestation, give at least 250iu anti-D for surgical interventions (ectopic, molar, TOP) or persistent painful bleeding, within 72 hrs of the event
- 12/13 20 weeks, give at least 250iu anti-D for PSEs such as bleeding, trauma etc

What SHOULD we be doing.....

- >20 weeks give at least 500iu anti-D for PSEs and perform Kleihauer test in case more is needed
- 28 34 weeks Routine Antenatal Anti-D Prophylaxis RAADP,
 - either at least 500iu at 28 and at 34 weeks
 - OR 1 x 1500iu at 28-30 weeks
- At birth (or stillbirth) If baby is RhD pos (or unknown) give at least 500iu anti-D and perform Kleihauer test in case more is needed

Cell Salvage in Obstetrics

An example of evolving practice, with increasing use of intra-operative cell salvage in the obstetric setting and the potential for relatively large volumes of foetal cells mixing with maternal circulation;

 Remember appropriate dosage of anti-D in RhD negative women – BCSH recommend *at least* 1500iu following cell salvage reinfusion, more if the Kleihauer indicates it

What Presentations of anti-D Ig are there ?

- In the UK:
 - CSL Behring: 'Rhophylac' 1500 IU pre-loaded syringe – IM/SC or IV
 - BPL: 'D-Gam'
 - 250 IU IM/SC only
 - 500 IU IM/SC only
 - 1500 IU IM/SC only
 - 2500 IU IM/SC only
- In Eire;
 - Octapharma: 'Rhesonativ' 625 IU or 1250 IU vials
 - CSL Behring: 'Rhophylac' 1500 IU pre-loaded syringe

Should we give Rhophylac 1500IU IV in women with BMI >30 ?

- Recent change in SmPC in Jan 2014 following concerns raised to the German licensing authority about POTENTIAL poor uptake following IM administration
- No clear evidence, no information about sites of injection etc etc
- Advise from BCSH, endorsed by RCOG members is to risk-assess process – very small risk of poor uptake balanced against training, calling women in to be cannulated, potential for error using BPL anti-D IV etc etc

What Dose should we be using ?

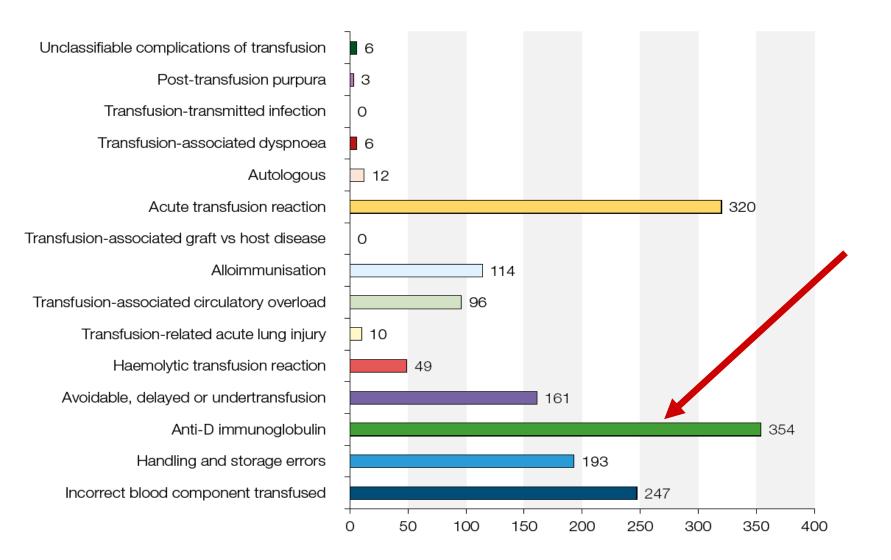
Anti-D Ig given IV

- 100 IU will clear 1 ml of foetal red cells
- is instantly available

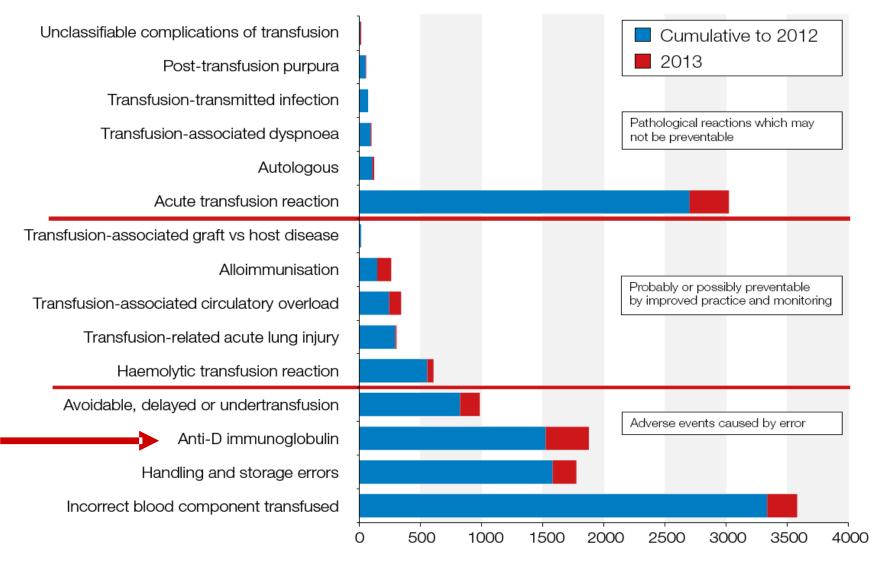
Anti-D Ig given IM

- 125 IU will clear 1 ml of foetal red cells
- will take hours to get into bloodstream via muscle, much longer (if at all) via fat and will lose some on the way

Cases reviewed by SHOT in 2013



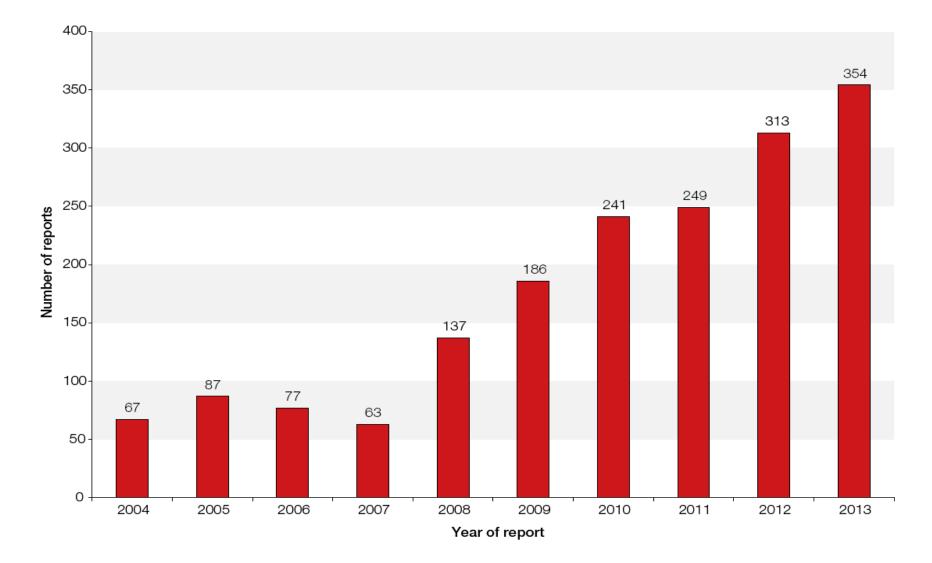
SHOT cumulative data 1996-2013



Anti-D Ig Reporting Categories

- Omission or late administration of anti-D immunoglobulin
- Inappropriate administration of anti-D immunoglobulin to:
 - a RhD positive woman
 - a woman who already has immune anti-D
 - a mother of a RhD negative infant (erroneously)
 - a different woman from the woman it was issued for
- Incorrect dose of anti-D Ig given
- Handling and storage errors
 - administration of expired, or otherwise out of temperature control, anti-D immunoglobulin

Trend in Anti-D Ig reports



Anti-D lg reports in 2013 (n = 354)

- **59** cases where anti-D Ig was inappropriately administered unnecessary exposure to a human medicinal blood product
- 277 cases where anti-D Ig was delayed or omitted, putting patient at risk of sensitisation to the D antigen *potential Major Morbidity*
- 9 cases where the wrong dose of anti-D Ig was administered (usually too little)
- 9 handling and storage errors

System Failures from SHOT cases (1)

Communication

- Lack of comms between hospital midwifery teams and those in the community – failure of RAADP in the community noted in 63 cases
- Assumption / Failing to take responsibility or ownership
 - Lack of robust systems to identify outstanding work in the laboratory
 - Lack of robust systems for identifying women eligible for RAADP
 - Lack of robust systems for women booking late or transferring care
 - Assumptions that someone else is sorting out a particular issue

System Failures from SHOT cases (2)

• Lack of knowledge or training

- Failure of lab staff to consider the need to issue anti-D Ig when giving RhD positive platelets to RhD negative patients of childbearing potential
- A lack of understanding behind the principles of anti-D Ig prophylaxis, compounded by availability of uncontrolled stocks held by clinics
- Increasing trend in poor advice being offered to women by (relatively senior) medical staff
- Decision making without reference to blood grouping results in both lab and clinical area
- Misinterpretation of FMH (Kleihauer) tests in labs leading to dosing errors
- Failure of inventory management in lab and clinical area, especially in the community

System Failures from SHOT cases (3)

• Pressures of work / staffing issues

 Understaffing and availability of senior staff in both the laboratory and clinical area leading to pressurised and poor decision making

• Poor practice / culture

- Manual transcription of blood grouping results onto notes, care plans and discharge sheets in the clinical area – repeatedly highlighted by SHOT but persists as poor practice
- A culture of completing paperwork when the interventions have not actually been performed
- Devolving responsibility to the pregnant/delivered woman to return at a later date for anti-D Ig administration, when they are obviously in a vulnerable and distressed state, instead of managing it at the presentation visit, be that in the ED, day unit or clinic
- Use of the Kleihauer Test to determine whether anti-D Ig should be given in the first place

Always confirm; • the woman's identity • that the woman is RhD Negative using the latest availab • that the woman does not have immune anti-D using the • that a blood sample has been taken to confirm group & administration of anti-D Ig) • that informed consent for administration of anti-D Ig is re • product / dose / expiry and patient ID pre-administration	latest available laboratory report antibody screen, (but do not wait for results before scorded in notes	• `
	vents (PSEs) during pregnancy	
Gestation up to 13 completed weeks	Gestation up to 13 completed weeks	
Surgical procedure to manage an ectopic pregnancy or miscarriage	Administer at least 250 units anti-D lg within 72 hours of event. No need for a Kleihauer	
Regardless of Gestation	Gestation 13 completed weeks to 20 weeks	
Amniocentesis, chorionic villus biopsy /cordocentesis Antepartum haemorrhage / PV bleeding External cephalic version Fall or abdominal trauma (sharp / blunt, open or	Administer at least 250 units anti-D Ig within 72 hours of event. No need for a Kleihauer under 20 weeks	
 Intrauterine death (at diagnosis and delivery) 	Gestation 20 weeks to term	
• <i>In-utero</i> therapeutic interventions (transfusion, surgery, insertion of shunts, laser)	Request a Kleihauer and administer at least 500 units anti-D Ig within 72 hours of event.	• [
Administer anti-D Ig for a PSE irrespec	tive of whether RAADP has already been given	
Does the Kleihauer indicate that further anti-D Ig is required ?	Administer more anti-D Ig following discussion with laboratory	t
Recurrent bleeding should be clinically review	wed by an Obstetrician for further anti-D requirements	
Size of FMH	Dose of anti-D required	k
Less than 2mL	500 units	
2.1 – 12mL	1500 units	b
Greater than 12mL	Specialist advice required	
Routine Antenatal A	nti-D Prophylaxis (RAADP)	S
For Routine Antenatal Anti-D Prophylaxis (Irrespective of whether anti-D Ig already given for PSE)	Take 28 week blood sample to confirm group & check antibodies	_
	Administer 1500 units anti-D lg at 28 – 30 weeks	a
At Delivery (or at diagnosis of Intra Ute	rine Death more than 20 weeks and at delivery)	
Is the baby's group confirmed as RhD positive ? Or are cord samples not available ?	Request a Kleihauer and administer at least 500 units anti-D lg within 72 hours of event	•
Transfusion Laboratory staff will advise if further anti-D Ig is required	Administer more anti-D following discussion with laboratory	· ·
Has Cell Salvage been used ?	Request a Kleihauer after re-infusion of red cells, inform	

- Your Trust policy can be a mixture of NICE and BCSH guidance, but BE CONSISTENT
- DO NOT wait for the result of a Kleihauer test before giving a standard dose of anti-D Ig
- If in doubt GIVE IT

Anti-D Blood and Transplant When and How Much?

This poster gives recommended dosages of anti-D immunoglobulin at different stages during pregnancy for women with an RhD negative blood type who do not already have immune anti-D antibodies.

NHS

05/13

At less than 12 weeks

 Anti-D is NOT usually indicated unless there has been therapeutic termination, molar or ectopic pregnancy, surgical intervention associated with miscarriage, or continued painful vaginal bleeding (request at least 250iu within 72 hours in these cases).

Between 12 and 20 weeks

Administer at least 250lu anti-D ig within 72 hours of a potentially sensitising event.

Between 20 weeks and delivery

- Administer at least 500iu anti-D ig within 72 hours of a potentially sensitising event.
- Send a sample for a Kleihauer Test in case additional anti-D ig is needed.
- Anti-D ig should be given for potentially sensitising events, even if RAADP has been given already.

Routine Antenatal Anti-D Prophylaxis (RAADP) should be administered between 28 and 30 weeks

- Send a sample for blood group and antibody screen and then administer RAADP according to local policy, even if anti-D ig has been recently given for a sensitising event.
- SINGLE DOSE: Administer 1,500iu anti-D ig at 28-30 weeks.
- TWO-DOSE: Administer at least 500iu anti-D ig at 28 and 34 weeks.

After delivery

- Send 'Mother & Cord' samples for testing.
- Where the baby is RhD positive, administer at least 500iu anti-D ig within 72 hours of delivery.
- Administer further anti-D ig on the advice of the laboratory if the Kleihauer shows a large fetomaternal haemorrhage.

For further information please refer to your local policy

British Committee for Standards in Haematology (BCSH) Guidelines for the use of prophytactic anti-D immunoglobulin 2006.

Royal College of Obstetricians and Gynaecologists (RCOG) Green Top Guideline No 22 The Use of Anti-D immunoglobulin for Rhesus D Prophylaxis 2011.

BLC680.1

 If outside 72 hrs still give anti-D, as a dose up to 10 days may provide some protection

 Give RAADP in addition to prophylaxis for sensitising events, and vice versa

INHS Blood and Transplant

Women who are already sensitised (have Anti-D

· Women who have already made Anti-D or other antibodies must be referred to a Consultant Obstetrician

as they may need specialised care. The Neonatal Team should be

informed when any woman with Anti-D or other antibodies is

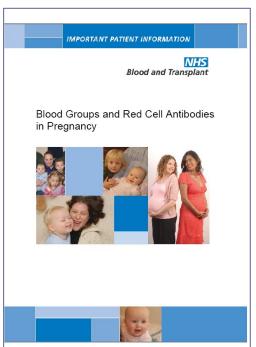
admitted in labour.

antibodies)

ANTI-D QUICK FACTS

RAADP (Routine Antenatal Anti-D)

- Maternal blood sample for antibody check should always be taken at 28 weeks before giving Anti-D. If a women has a PSE close to the date of her RAADP, both RAADP and Anti-D to treat the PSE should still be given.
- Postnatal Care
- Only women who have an RhD positive baby will require Anti-D Do not weit for the Kleihauer Test result before giving the standard dose.
- · Some women may require more than one postnatal Anti-D
- iniarting
- . This depends on the results of the Kleihauer Test done on
- maternal samples taken at delivery.





HDN awareness

Reducing the impact of haemolytic disease of the (fetus and) newborn



Key Issues with Anti-D Ig

- Ignorance
- Failure to adhere to local protocol
- Failure to utilise IT systems properly / fully
- Late bookers
- Transfers of care
- Assumptions
- Failures of communication
- Failure to obtain valid consent for anti-D Ig

Anti-D Summary

- Effective anti-D prophylaxis is a *partnership* between the laboratory and the clinical area – *work together* to produce robust Trust guidelines, and follow them !
- Requests for anti-D should be driven by the clinicians, especially in early pregnancy

..... and finally

Thanks to;

- SHOT Team
- UK NEQAS
- Megan Rowley
- YOU for listening

www.shotuk.org