Anti-D Prophylaxis Reviewed in the Erea of Foetal RHD Genotyping

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Received date: Aug 05, 2015, Accepted date: Aug 31, 2015, Publication date: Sep 04, 2015

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Abstract

A few years ago, the prevention of anti-D immunization was currently based on systematic postnatal prophylaxis associated with targeted antenatal injection in high-risk situations of foeto-maternal haemorrhage. The failures of prevention are mainly due to the non-respect of established guidelines for RhIG prophylaxis, and to spontaneous undetected foetal-maternal haemorrhages without any obvious cause during the third trimester of pregnancy.

In order to reduce the rate of residual post-pregnancy anti-D immunization, several countries decided to associate the classical prophylaxis to a routine antenatal anti-D prophylaxis (RAADP) during the 28th or 29th week of gestation. Since about ten years, the foetal RHD genotyping in maternal plasma enables us to limit the antenatal prophylaxis only to those D- women carrying a D+ foetus.

This paper deals with: the advantages of an antenatal prevention in the light of non invasive foetal RHD genotyping, the rules rendering prevention protocols efficient whatever the algorithm applied, and the recommended immuno-haematology follow-up of women who have received RhIG.

Keywords: Anti-D prophylaxis; Foetal RHD genotype; Haemolytic Disease of the foetus and Newborn

Introduction

The pathogenesis of haemolytic disease of the foetus and newborn (HDFN) was first elucidated by Levine (1941) thanks to the discovery of the Rh (Rhesus) blood group by Landsteiner and Wiener in 1940 [1,2]. He demonstrated the possibility of maternal immunization through placenta against foetal antigen(s) inherited from the father and lacking in the mother; in this particular case, the D antigen.

In 1940, prenatal mortality due to HDFN was around 4 out of 100 births, representing 10 percents of the global childhood mortality. Half of the affected foetuses died from hydrops foetalis or nuclear icterus. The improvement of the management of this pathology by experts reduced the mortality rate to less than 5 percent. The foetal RHD genotyping, the rules of prevention are mainly due to the non-respect of established guidelines for RhIG prophylaxis, and to spontaneous undetected foetal-maternal haemorrhages without any obvious cause during the third trimester of pregnancy.

Despite the application of this immuno-prophylaxis to RhD negative patients, HDFN due to anti-D still remains today the most frequent and severe in Europe [4].

In the absence of anti-D injection, a D negative mother bearing an ABO compatible D positive foetus has a 16% possibility of developing alloimmune anti-D. When the mother and the foetus are ABO incompatible (20% of pregnancies), this rate falls to 2%. So, globally, the risk for a D-negative woman to become immunized by a D-positive foetus stands at around 13.2% [3].

However, when appropriate doses of anti-D immunoglobulin’s are injected within 72 hours after delivery, the rate of immunisation is reduced by 90% and the residual risk is around 1 to 2% [5].

Most failures are due on one hand, to undetected foeto-maternal haemorrhages (FMH) that occurs during the third trimester of pregnancy, and, on the other hand, in more than 30% of cases, to disregard for prophylaxis rules [6].

When D negative patients are systematically injected at 28 weeks of pregnancy, the residual risk is further reduced by more than 60% [7].

Throughout pregnancy, foeto-maternal haemorrhage increases in frequency (from 3% during the first trimester to 45% during the third one), and in volume (<0.1 mL during the first trimester and in variable amounts during the third one) [8]. A dose-dependant correlation is known to exist between the volume of foeto-maternal haemorrhage and the occurrence of allo-immunisation. This is the reason why an adequate and timely administration of anti-D immunoprophylaxis remains essential. These practical aspects, including guidance regarding to routine antenatal anti-D prophylaxis (RAADP), are reviewed in this paper.

Systematic Antenatal Prophylaxis

Cost/efficiency

Prevention of anti-D allo-immunisation can be applied following at least two different approaches: the first one, as a postnatal prophylaxis after delivery of a RhD positive baby, in addition with targeted prophylaxis following potentially sensitizing events during pregnancy,
and the second one, as a systematic antenatal prophylaxis at 28 weeks of gestation.

This matter has been discussed in a review published by Parant, who concluded that in the group without systematic antenatal prophylaxis, the global rate of immunisation turned around 1.5% (1.2 to 1.9%), whereas in the group of patients who benefited from a systematic antenatal prophylaxis, the average rate of allo-immunization was around 0.3% (0 to 0.9%) [9]. These data are in favour of systematic prevention at 28-29 weeks of gestation [10].

From an economical point of view, Canadian and English studies tend to show that systematic prevention involves greater drug and administrative costs than the targeted ones, but the cost/efficiency ratio still remains favourable [11]. The benefits of RAADP to be considered include: avoidance of anti-D sensitisation, reduction of incidence of haemolytic diseases, reduction of foetal/neonatal losses, avoidance of disability of the child, and positive effects of these outcome measures on the quality of life of the mother [12]. Most western countries recommend a systematic antenatal prophylaxis for all Rh D negative patients. However, so far, none of these studies ever organized an antenatal prevention which would be dedicated solely to those women bearing Rh D+ foetuses. This eventuality is now rendered possible owing to the determination of foetal RHD from maternal plasma. It allows us to reduce, by around 40%, the cost of unnecessary RhIG prenatal injections. This genotyping approach may offer additional cost benefits by reducing the immuno-haematological and sonographic follow-up of pregnancy [13]. The cost-effectiveness calculations of this approach will depend on the cost of foetal genotyping in maternal plasma. At present, this test is not refunded by the Health Service in Belgium except in anti-D alloimmunized mothers.

Search for RHD foetal gene from maternal plasma

Since 2002 in our hospital, the determination of foetal RHD from maternal plasma has been included in the biological follow-up of Rh D negative patient as early as the 12th week of gestation. We have reported 100% diagnostic accuracy in our non invasive foetal RHD genotyping assay by targeting multiple exons with real-time PCR [14,15]. The overall diagnostic accuracy was 96.5% (95% CI 95.6-97.2) in the meta-analysis of Geifman-Holtzman, several studies also showing 100% diagnostic accuracy [16].

In order to avoid reporting a false negative result, the Y-chromosome linked SRY gene sequence is amplified to confirm that foetal DNA is present in maternal plasma. A positive PCR signal ensures that the foetal DNA is present. This SRY-based internal control is only applicable for pregnancies carrying a male foetus. So, when the search for foetal RHD is negative and SRY gene absent, it is recommended to confirm the first result by using either a new blood sample a few weeks later or amniocytes when an amniotic liquid puncture has been programmed for another reason. This careful approach is justified by the absence of internal control for the Rh D negative female foetuses. It does not significantly differ from the basic rules in the matter of ABO/D blood groups that require two different determinations before being validated.

The knowledge of foetal RHD status allows us to save up the injection of anti-D immunoglobulins at 28-29 weeks as well as those applied in the situations at risk of FMH among the 40% of Rh D negative women bearing an RHD negative foetus.

On the other hand, genotyping of the foetal D gene also exempts D negative patients, bearing a D- foetus, to be exposed to any risk that might be associated with the administration of a human plasma-derived product. Moreover, there is worldwide shortage of anti-D [17].

As far as the post-partum immunoprophylaxis is concerned, determining the Rh D phenotype of a newborn from a D negative woman must continue to be determined even when the RHD foetal genotype has been searched during gestation [18].

Because samples collected in operating rooms are frequently mislabelled (newborn/mother inversion, cord blood contaminated by maternal blood, requisition mismatches), the French Society of Blood Transfusion (as well as the Society of Perinatal Medicine in France) recommends confirming the D negative phenotype of the baby determined on cord blood, by a second one performed on peripheral blood [19]. This should be strongly recommended when there was no previous RHD genotyping of the foetus during the pregnancy.

Rules of Good Practice

Indications

D negative women: Prophylaxis concerns all non immunized D negative women during and at the end of gestation. It has therefore no effect and is perfectly useless on previously immunized women. In rare cases of very weak anti-D which are solely detected by ultra sensitive techniques (enzymes) but remain undetectable by the anti-human globulin assay (AHG) in gel phase, the relevancy of prophylaxis must be discussed for each case individually. Usually, the follow-up in time allows us to decide the best approach.

Weak D women (previously named D*): The term weak D (Du) is used to design a weakened expression of a “complete” D antigen (Figure 1). The molecular characteristic of the weak D types is single missense mutations which produce amino-acid exchange in the intracellular or transmembrane region of the D antigen [20]. The molecular techniques to predict foetal D status using maternal plasma, don't detect maternal weak D types and the maternal RHD gene appears “intact” as in D+ patients. The lower immunoreactivity of D antigen requires the indirect antiglobulin technique (IAT) for detection. However, at present time, most weak D are now detected by direct agglutination using monoclonal IgM anti-D reagents, or in the gel matrix technique and are considered as D positive [21]. The majority (90%) of caucasian individuals presenting a weak D phenotype are weak types 1, 2, 3. Among European persons, these weak D types are always associated with the DCe or the DcE haplotypes. They cannot so far be anti-D immunized. Therefore these patients have to be treated as D positive, i.e.: transfused with D positive red cells and, following delivery of a D positive baby, not given anti-D immunoglobulin [22,23].

Partial D women: The term partial D is used to refer to an incomplete D antigen in which some extracellular epitopes are missing (Figure 1). Partial D phenotypes are the results of gene rearrangements and/or senseless mutations in regions of RHD encoding portions of D protein external to the red cell membrane [20]. Individuals with a partial D phenotype can produce anti-D when they are exposed to D positive red blood cells (RBC). Therefore, it is advisable that they should be typed as D negative if they are candidates for transfusion or anti-D prophylaxis. Such a strategy is applied in our laboratories to type partial DVI, the most frequent D partial phenotype amongst white people. DVI presents also a weak expression of D and their
carriers may be readily anti-D immunized [24]. In practice, most other partial D are usually classified as D positive in direct agglutination tests with currently available reagents and can often only be identified as having partial D antigen after and because they have produced anti-D.

Among European populations, about one percent of people carry variant RHD alleles producing weak D including some partial D [26,27]. In Black individuals, most D variants with a weak expression are partial D and are often associated with Dce haplotype [28].

Weak D baby: All serologically D negative babies should be tested for the presence of weak D expression with the indirect antiglobulin technique. If the baby is weak D, prophylaxis must be applied to D negative women, whereas it’s not necessary if the baby is proven to be D negative.

Detection of foetal erythrocytes

The principle of the Kleihauer-Betke assay - the most widely used in Belgium - lies on the greater resistance to acid elution of foetal hemoglobin (HbF) compared to adult hemoglobin (HbA). In practice, HbA is eluted from RBCs fixed on blood films, leaving red cell ghosts, so that the red cells containing HbF can be stained.

The volume of foetal RBC is calculated according to the following formula [29]:

\[
\text{Volume of foetal RBC} = 2400 \times \frac{F}{A} \text{ mL},
\]

where F/A is the proportion of foetal (F) RBC among maternal adult (A) RBC.

- 2400 = 1800 x 1.22 x 1.09 mL
- 1800 mL = volume of maternal red blood cells,
- 1.22 = foetal RBC are 22% larger than adult RBC,
- 1.09 = 92% = proportion of foetal RBC stain darkly.

The volume of foetal blood (assuming a haematocrit of 50%) can be deduced by multiplying the volume of foetal RBC by two.

Clinicians should be aware that the results of a Kleihauer-Betke test can be expressed on laboratory protocols either in terms of mL of foetal RBC or in mL of foetal whole blood. This is particularly important when the dose of anti-D immunoglobulin has to be adjusted according to the volume of the FMH. The general principle is that 100 µg of RhIG is capable of suppressing sensitization by 4-5 mL of foetal RhD positive red blood cells or by 8-10 mL of foetal RhD positive whole blood.

The Kleihauer test’ sensitivity allows to detect 7.2 mL of foetal RBC in the maternal circulation i.e. around 0.3% of foetal RBC, with a variation coefficient of around 10%.

Some hemoglobinopathies (thalassemia, drepanocytosis, and hereditary persistence of foetal hemoglobin…) present varying resistance to acid elution and may entail false positive results. When such a case is suspected, some laboratories detect D positive foetal RBC by flow cytometry, by using fluorochrome coupled anti-D reagents. Flow cytometry presents a very high sensitivity (0.01 % i.e. 0.2 mL of foetal RBC), but this method is expensive, time consuming and not suitable for emergency cases.

At delivery or in situations at risk of FMH, prophylaxis must always be applied even when the result of the Kleihauer test is negative.

In practice, the search for foetal RBC should be performed within 2 hours of delivery (or a high-risk situation) but at least 30 minutes after

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**Figure 1:** The genetic background for different RhD types (simplified). D negative Caucasians most often have a complete RHD gene deletion, but Africans and Asians have often an inactive RHD. The partial D red blood cells (RBC) demonstrate a normal number of Rh D antigens but with altered D epitopes. Many partial D results from hybrid genes with portions of RHD replaced by the corresponding portions of RHCE. The weak D red blood cells have a reduced number of complete D antigens. Weak D types primarily result from single point mutations in RHD. Some partial D (the most common in Caucasians: DVI) show both weak D and partial D features.
placental evacuation to be sure that all the foetal erythrocytes have entered in maternal circulation [30].

Kleihauer’s test results should always be transmitted to the clinician for those cases necessitating a supplementary dose, which should be administered within 72 hours post-delivery.

Doses

In Belgium, there is only one preparation of anti-D immunoglobulin (RhIgG) available (Rhogam); it contains 300 µg/mL of IM injectable RhIgG. 300 µg is an amount of IgG capable of neutralizing 12 to 15 ml of foetal RBC (i.e. 24 to 30 mL of foetal blood).

The routine antenatal anti-D prophylaxis consists of injecting 300 µg of anti-D at between 28 and 29 weeks of gestation.

After invasive examinations or accidents occurring before the 20th week, one IM dose is sufficient since the blood mass of a 20 weeks old foetus is lower than the volume neutralized by 300 µg.

After 20 weeks, there is a need to assess the volume of FMH and the dosage should be adapted according to the volume of circulating foetal RBC: 10 µg per 0.5 mL foetal RBC or 1 mL foetal blood.

When the risk of FMH is low, for example after extra-placenta amniocentesis, it is not necessary to perform a Kleihauer test, and injection of one standard dose of anti-D is sufficient.

It is recommended to search for irregular antibodies (RAI) before the first injection of anti-D prophylaxis, even if the injection can be made before the RAI result is obtained. This allows for the detection of any anti-D allo-immunisation (and other than anti-D) which would have been undetectable earlier [31,32].

Delay for the injection of RhIgG

To reach optimal efficiency, the injection of RhIgG should be made as soon as possible within 72 hours following delivery or HFM. If it has been omitted, injection of RhIgG can still be performed until 28 days after delivery, but the later it is done, the less efficient it will be [33].

Control of RhIgG efficiency

The sole relevant control test is the search for foetal RBC in maternal blood 48 hours after injection of RhIG [10], RhIG has been efficient when all foetal RBC have disappeared. This control is only necessary when a supplementary dose has to be injected i.e. when the foetal hemorrhage was greater than 15 mL of foetal RBC.

There are no arguments at all to say that the presence of passive anti-D in maternal circulation can be considered as reflecting the efficiency of RhIG injection; it only reflects the fact that the injection has really been made.

Within three to seven days after IM injection of RhIG, the titre of anti-D in maternal blood can reach ¼ (when measured by using the reference method i.e. IAT in saline). Antibody titration decreases thereafter as a function of both the half-life time of IgG (21 days), and the rate of consumption by D positive foetal RBC. However, a residual activity of RhIG can be demonstrated in some maternal plasma during several months.

If any doubt remains concerning the appropriate dosage of RhIgG administered, the clinician can register the situation in the light of both the results of the Kleihauer test and the presence of anti-D in maternal blood (Table 1). Absence of detectable anti-D has – contrarily to its presence—a predictive value in terms of absence of protection.

| Foetal red blood cells (Kleihauer) | Free circulating anti-D’ in maternal blood | Action |
|-----------------------------------|------------------------------------------|--------|
| 0                                 | No                                       | 2nd Injection |
| 0                                 | Yes                                      | No 2rd injection |
| Yes                               | No                                       | Assess FMH Injectable dose |
| Yes                               | Yes                                      | assess FMH at 48 h |

IAT tube method in phosphate buffered saline

Table 1: After an initial injection of RhIG, when a second one should be injected?

One should also keep in mind that several factors can affect the clearance of fetal RBC, including: a laboratory failure at the time of the initial foetal RBC count, a dizygotic twin pregnancy with one D positive and one D negative foetus, a weak D expression or a splenectomized woman [34].

Finally, so many different situations exist - which can entail error, omission or default - that it is more reasonable to promote a systematic injection of RhIG at delivery even when the patient has already received one before.

Immuno-Haematologic Features of RhIG Injections

In maternal blood

Residual anti-D: Sometimes, a weak anti-D activity may be detected in maternal serum for several weeks by IAT and, by more sensitive techniques, for several months after RhIG injection, and persistent passive anti-D may mask an early active allo-immunisation. On the other hand, the clearance of passive anti-D is highly variable from one patient to another. Therefore, a low titration of anti-D (<4 by IAT) does not allow to discriminate passive and active immunisation [25].

To provide a basis for distinguishing between prophylactic and immune anti-D, all D negative pregnant women should benefit before the RAADP, at 28 weeks gestation, from a screening for red cell alloantibodies. Moreover, it is essential for the biologist to be informed to any previous administration of RhIG in the current pregnancy, including date and dose, at the time of analysis, in order to interpret the anti-D titre. It is generally admitted that anti-D titration higher than ¼ is highly predictive of an active immunisation. When titration results are not conclusive, it is necessary to repeat the analysis a few weeks later to be able to conclude to passive or active immunisation. Prophylactic anti-D levels will fall with time whereas immune anti-D levels will remain stable or rise. Some authors propose to perform micro titration to differentiate an active from a passive anti-D in the follow-up of RhIG prophylaxis [35].

In practice, any laboratory request form for the search for irregular antibodies during pregnancy should indicate if the patient has or not
received RhIG, and if so, when. Besides, injection of RhIG should systematically be noted in the medical history of the patient even though this is not (yet) mandatory in Belgium.

*Calendar for the research of irregular antibodies during pregnancy:*

In the absence of any sensitizing event or clinical indication, there is no need to repeat an antibody screening at the 8th or 9th month when a systematic prophylaxis, preceded by a screening for red cell alloantibodies, has been applied at 28 weeks [36].

However, the introduction of RAADP has resulted in a positive antibody screen with the detection of anti-D in samples taken after 28 weeks gestation. A panel of D negative cells should be used to exclude the presence of unexpected alloantibodies of other specificities. If the mother needs blood transfusion in a context of obstetrical haemorrhage in the delivery, this would lengthen the delay before blood product can be delivered. Therefore to ensure an appropriate blood management of obstetrical haemorrhage, which are often sudden and unexpected, a search for irregular antibodies can be suggested within the 7 days preceding delivery in order to identify alloantibodies other than pass if anti-D possibly present in the serum of the mother [37,38]. In presence of clinically relevant alloantibodies, a compatibility testing must be performed [39].

**Hindrances**

Plasma used to prepare anti-D immunoglobulins is of human origin; they are collected from immunized blood donors and therefore may contain Rhesus antibodies other than anti-D: anti-C or anti-E, for example. That is the reason why it may occur that anti-C or anti-E be detected in the serum of injected women [34].

**In the newborn**

Anti-D prenatal prophylaxis has no effect on the foetus. At most, foetal red blood cells can be sensitized, which will lead to a positive direct Coombs test at birth, but without any clinical consequences [40]. However, if multiple doses are given, it is recommended that bilirubinemia should be followed.

**Conclusion**

The possibility to know the RHD foetal genotype from a maternal blood sample during pregnancy should dramatically modify the recommendations in the matter of prophylaxis for foetal-maternal allo-immunization.

Associated with the routine antenatal anti-D prophylaxis at 28-29th week of gestation, it is possible now to propose a new approach of D negative pregnant women (Figure 2).

RHD foetal genotyping during pregnancy allows us to recommend a prevention policy targeted at the 61% of D- women bearing a D+ foetus, by systematically injecting anti-D immunoglobulin during the third trimester of pregnancy. At the same time, knowing the RHD foetal status allows to avoid injections in 39% of women carrying a D- foetus.

It is highly regrettable to see that prevention failures are mainly due to human or functional negligence: default of injection, too late injection, and so on.

At the time of accreditation, it was probably worth reminding ourselves of the best practices for a high-risk pregnancy follow-up: the predictive value of antibody screening, the significance of circulating specific antibodies, the results of Kleihauer, which allows us to measure the range of foetal-maternal haemorrhage and to adjust the dose of RhIG to inject.

Even if any best practice rule will never replace the medical judgement of clinicians, it should be taken into consideration how they have progressed and benefited from technological advances such as the foetal RHD genotyping obtained from a non invasive examination.

**References**

1. Levine P, Katzin E, Burnham L (1941) Isoimmunisation in pregnancy: its bearing on the aetiology of erythroblastosis fetalis. JAMA 116: 825-827.
2. Landsteiner K, Wiener A (1940) An agglutinate factor in human blood recognized by immune sera for rhesus blood. Proc Soc Exp Biol Med 43: 223.
3. Bowman J (2003) Thirty-five years of Rh prophylaxis. Transfusion 43: 1661-1666.
4. Daniels G (2002) Blood group antibodies in haemolytic disease of the fetus and newborn. Alloimmune disorders of pregnancy. Cambridge: University Press: 21-40.
5. Ratsimbazagy V, Alba J, Cohen J (2002) L’allo-immunisation foetomaternelle anti-D. Gynecol Obstet 450: 20-23.
6. [No authors listed] (1999) ACOG practice bulletin. Prevention of Rh D alloimmunization. Number 4, May 1999 (replaces educational bulletin Number 147, October 1990). Clinical management guidelines for obstetrician-gynecologists. American College of Obstetrics and Gynecology. Int J Gynaecol Obstet 66: 63-70.
7. Chilcott J, Lloyd Jones M, Wight J, Forman K, Wray J, et al. (2003) A review of the clinical effectiveness and cost-effectiveness of routine anti-D prophylaxis for pregnant women who are rhesus-negative. Health Technol Assess 7: iii-62.
8. Bowman JM, Pollock JM, Penston LE (1986) Fetomaternal transplacental hemorrhage during pregnancy and after delivery. Vox Sang 51: 117-121.

9. Parant O (2006) Comparaison de l’efficacité des différentes formes de prévention de l’allo-immunisation anti-D au cours de la grossesse: prévention cible limitée aux situations très graves ou associée à une prévention systématique au 3e trimestre. J Gynecol Obstet Biol Reprod (Paris). 35: 1593-159103.

10. Cortey A, Brossard Y (2006) [Prevention of fetomaternal rhesus-D allo-immunization. Practical aspects.]. J Gynecol Obstet Biol Reprod (Paris) 35: 1S123-121S130.

11. Ravinet J, Carbonne B (2006) [Economic analysis of the prevention of anti-D immunization.]. J Gynecol Obstet Biol Reprod (Paris) 35: 1S104-101S111.

12. https://www.nice.org.uk/.

13. Carbonne B, Cortey A, Rouillac-Le Scellour C, Brossard Y (2008) [Non invasive fetal RhD genotyping using maternal blood: time for use in all RhD negative pregnant women]. Gynecol Obstet Fertil 36: 200-203.

14. Minon JM, Schaaps JP, Retz MC, Dricot JF, Foidart JM, et al. (2005) [Prenatal determination of fetal RhD in maternal plasma: two-years experience of routine clinical use.]. J Gynecol Obstet Biol Reprod (Paris) 34: 448-453.

15. Minon JM, Gerard C, Senterre JM, Schaaps JP, Foidart JM (2008) Routine fetal RhD genotyping with maternal plasma: a four-year experience in Belgium. Transfusion 48: 373-381.

16. Geifman-Holtzman O, Grotegut CA, Gaughan JP (2006) Diagnostic accuracy of noninvasive fetal Rh genotyping from maternal blood—a meta-analysis. Am J Obstet Gynecol 195: 1163-1173.

17. van der Schoot CE, Hahn S, Chitty LS (2008) Non-invasive prenatal diagnosis and determination of fetal Rh status. Semin Fetal Neonatal Med 13: 63-68.

18. Cortey A, Brossard Y, Belliard R, Bourel D (2006) [Prevention of fetomaternal rhesus-D allo-immunization. Perspectives.]. J Gynecol Obstet Biol Reprod (Paris) 35: 1S119-111S122.

19. Mennessier L, Ali-Daram S, Roubinet F, Brossard Y; Groupes de travail Immunohématologie de la Société française de transfusion sanguine et de la Société française de médecine périnatale (2000) [Prevention of fetal hemolytic disease: it is time to take action]. Transfus Clin Biol 7: 527-532.

20. Wagner FF, Flegel WA (2004) Review: the molecular basis of the Rh blood group phenotypes. Immunohematology 20: 23-36.

21. Williams M (2000) Monoclonal reagents for rhesus-D typing of Irish patients and donors: a review. Br J Biomed Sci 57: 142-149.

22. Flegel WA, Wagner FF (2002) Molecular biology of partial D and weak D: implications for blood bank practice. Clin Lab 48: 53-59.

23. Flegel WA (2006) Molecular genetics of RH and its clinical application. Transfus Clin Biol 13: 4-12.

24. Wagner FF, Gassner C, Muller TH, Schonitzer D, Schunter F, et al. (1998) Three molecular structures cause rhesus D category V1 phenotypes with distinct immunohematologic features. Blood 91: 2157-2168.

25. https://www.karger.com/Book/Home/231653.

26. Wagner FF, Gassner C, Muller TH, Schonitzer D, Schunter F, et al. (1999) Molecular basis of weak D phenotypes. Blood 93: 385-393.

27. Wagner FF, Kasulke D, Kerowgan M, Flegel WA (1995) Frequencies of the blood groups ABO, Rh, D category V1, Kell, and of clinically relevant high-frequency antigens in south-western Germany. Infusionsther Transfusionsmed 22: 285-290.

28. Chen Q, Flegel WA (2005) Random survey for RhD alleles among D+ European persons. Transfusion 45: 1183-1191.

29. British Committee for Standards in Haematology (2000) Guidelines for blood grouping and red-cell antibody testing during pregnancy and for performing red-cell alloantibody titrations. Oxford: Blackwell Science Ltd. 201-206.

30. Judd WJ, Scientific Section Coordinating Committee of the AABB (2001) Practice guidelines for prenatal and perinatal immunohematology, revisited. Transfusion 41: 1445-1452.

31. Hartwell EA (1998) Use of Rh immune globulin: ASCP practice parameter. American Society of Clinical Pathologists. Am J Clin Pathol 110: 281-292.

32. Banks AA (2002) Hemolytic Disease of the Newborn Methods In: Banks AAoB, ed. Technical Manual. (14thedn) Bethesda 2002 :730-731.

33. Bowman JM (1985) Controversies in Rh prophylaxis. Who needs Rh immune globulin and when should it be given? Am J Obstet Gynecol 151: 289-294.

34. Brossard Y, Parmet-Mathieu F, Larsen M (2000) Incompatibility foeto-maternelles arthryocytaires. John Libbey Eurotext: 294.

35. Dupont M, Gouvitos J, Dettori I, Chiaroni J, Ferrera V (2007) [Microtitration of anti-RH1 antibodies: interest in the follow-up of pregnant women]. Transfus Clin Biol 14: 381-385.

36. British Committee for Standards in Haematology Blood Transfusion Task Force, Gooch A, Parker J, Wray J, Qureshi H (2007) Guideline for blood grouping and antibody testing in pregnancy. Transfus Med 17: 252-262.

37. Sherman SJ, Greenspoon JS, Nelson JM, Paul RH (1993) Obstetric hemorrhage and blood utilization. J Reprod Med 38: 929-934.

38. Chapman JF, Elliott C, Knowles SM, Milkins CE, Poole GD; Working Party of the British Committee for Standards in Haematology Blood Transfusion Task Force (2004) Guidelines for compatibility procedures in blood transfusion laboratories. Transfus Med 14: 59-73.

39. Mennessier L (2007) [Immunohematologic surveillance of the pregnant woman and the new prevention policy of anti-RH1 allo-immunization]. Transfus Clin Biol 14: 112-119.

40. Maayan-Metzger A, Schwartz T, Sulkes J, Merlob P (2001) Maternal anti-D prophylaxis during pregnancy does not cause neonatal haemolysis. Arch Dis Child Fetal Neonatal Ed 84: 660-62.