Late anaemia in Rh haemolytic disease

As it is clearly stated in the review by Gottstein and Cooke, we consider it unethical to withhold or delay high dose intravenous immunoglobulin (rHEPO) treatment in infants with haemolytic disease of the newborn. Since the study we did in 1995, we have treated 129 patients with Coomb’s positive haemolytic disease of the newborn, with the same method and had to resort to exchange transfusions only in three cases. On the other hand, late anaemia is a frequent problem in these cases, necessitating multiple blood transfusions, with well known complications.

The authors suggest that multiple doses of IVIG may reduce late anaemia. However, our observation in a limited number of cases is that, even multiple doses of IVIG are inefficient in preventing late anaemia. In an earlier unpublished study, we had shown that the erythropoietin levels were low in these infants. Therefore, we had conducted a double blind, randomised pilot study to investigate the effect of recombinant erythropoietin (rHEPO) in these patients.

In this study, rHEPO was administered at a dose of 200 units/kg, subcutaneously, three times a week, starting at the 14th day of life and lasting for six weeks. This protocol reduced the number of erythrocyte transfusions significantly. With the impetus of this pilot study, we have used the same protocol for the subsequent 103 patients and the mean number of transfusions in this group was 1.5, with the majority of patients (55%) not needing any transfusions at all. There were no complications, including changes in neutrophil or platelet counts, and haemorrhagic or infectious complications. The administration of rHEPO to patients with haemolytic disease of the newborn, who had received IVIG early in life, not only decreases the infants’ exposure to multiple blood donors, but also diminishes the need for hospitalisation and hence the cost that is involved. Therefore, rHEPO treatment is a suitable alternative to erythrocyte transfusions in these infants.

Systematic review of intravenous immunoglobulin in haemolytic disease of the newborn

We read with interest the recent review of Gottstein and Cooke. Their systematic review of trials reporting treatment of infants with proven Rh and/or ABO haemolytic disease of the newborn (HDN) treated with high dose intravenous immunoglobulin (HDIVIG) and phototherapy, with phototherapy alone demonstrated that significantly fewer infants required exchange transfusion in the HDIVIG group. The authors point out that anti-D is the commonest red cell antibody responsible for HDN. We have recently treated two children both of whom developed evidence of immune haemolysis due to anti-D antibodies acquired from IVIG.

The first patient, a 4 month old infant with Kawasaki’s disease, was treated with intravenous immunoglobulin (2g/kg) with immediate control of fever and irritability. Ten days later her disease became clinically active again and she was therefore given a second dose of IVIG which showed her haemoglobin had fallen suddenly by 2g/dl to 6.4g/dl, the blood film showed spherocytes and the direct antiglobulin test was positive, evidence of immune haemolysis. Samples that were collected prior to the second dose of IVIG confirmed her blood group to be AB Rh D positive with a negative direct antiglobulin test. Anti-D antibodies were now detected in the patient’s serum; these were not present in her mother whose antibody screen was negative and whose blood group was A Rhesus D positive. The manufacturer of the IVIG investigated the batches used and reported that the IVIG used for the second dose contained anti-D. The second patient, a 1 year old boy with systemic juvenile idiopathic arthritis received a fifth dose of IVIG from the same batch. He was screened for evidence of haemolysis and his antiglobulin test was positive 14 days after treatment. He remained asymptomatic with no fall in haemoglobin.

IVIG is not a blood product not a drug; each batch is made from a pool of plasma collected from several thousand donors. Passive transfer of potentially significant red cell antibodies is a recognised hazard, reported in the company literature but only as a serological phenomenon, not as a clinical warning.

High dose intravenous immunoglobulin in haemolytic disease of neonates

It was encouraging to read article of Gottstein et al on the use of high dose intravenous immunoglobulin (HDIVIG) in cases of haemolytic disease of newborns (HDN) with their conclusion showing the effectiveness of HDIVIG. I have the following observations to make with respect to implications on practice and future research.

Firstly, all the references were mentioned to be between three and ten years old. These trials did not take into consideration the irradiance of the phototherapy used, although they did observe the number of exchange transfusions performed. Presently, a combination of blue and white fluorescent light double surface phototherapy, with effective higher irradiances of 20-40 uW/cm²/mm, can practically eliminate the need for exchange transfusion, even in severe cases of HDN. Irradiance of phototherapy can be increased further by increasing the distance between the phototherapy unit and the patient, especially with an undersurface phototherapy unit, keeping thermal and nursing issues under consideration.

Secondly, the authors did not address enterohepatic recirculation of bilirubin from the gut. Inexpensive measures can decrease the back entry of bilirubin from gut, like early enteral feeds, oral administration of agar agar, isabogol husk and so forth, and further reduce serum bilirubin levels. Further randomised controlled trials are required before administration of HDIVIG becomes routine in HDN. These trials should compare use of current effective phototherapy combinations with the

References


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5. References


Systematic review of intravenous immunoglobulin in haemolytic disease of the newborn

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The first case is a reminder that such complications may have serious clinical consequences. We would agree with the comment of Gottstein and Cooke that the use of IVIG is not without potential risks, including haemolysis. IVIG is not universally effective at treating haemolytic disease in older children and adults where steroids are the first choice.

indications for the use of IVIG must be clear and evidence based, and as with all pooled blood products, including albumin solutions, the individual batch numbers must be recorded in the case notes, so that adverse events can be appropriately and fully investigated.

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highest possible irradiance, agents that de-
crease enterohepatic recirculation of bilirubin 
with or without HDIVIG, and the need for 
exchange transfusion in HDN. They should 
also address cost effectiveness and safety, 
considering the cost of HDIVIG in the developing 
world.

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Authors’ reply
We are grateful to our colleagues for their interest and responses to our paper.1 In response to Dr Ovaly’s comments we agree that late anaemia can be a problem in babies who receive intravenous immunoglobulin (IVIG), as is also demonstrated in our systematic review. Even when infants have received exchange transfusions (XTs) top up red cell 
transfusions may be required. In a recent local audit of XTs, 35% of babies received top up red 
cell transfusions after one or more exchange 
than 20 XTs. Gestation ranged from 22 to 40 completed weeks. Of 26 infants for whom follow up data was available, 
nine (35%) had received top up red cell transfusions.

We read with interest Dr Ovaly and colleagues paper describing a double blind randomised controlled trial of subcutaneous recombinant human erythropoietin (rHbEPO) and its use in this situation.1 We await with interest the outcome of a Cochrane meta-analysis of this therapy in newborn infants (currently at the protocol stage).

We reviewed our database for a three years between December 1999 to December 2002 to postulate what impact IVIG might have on our population of babies with haemolytic disease of the newborn. Two hundred and five babies had a positive direct Coombs test (DCT) result. Of these infants, 12 received XTs. There is a degree of under ascertainment with this database as there were four additional babies who required an XT during this time period. However, we make the assumption that the proportion of those 
missing requiring XTs is similar to the propor-
tions of DCT positive babies who were missed 
from the database. Eighty five babies had moderate or strongly positive DCT. Of these 11 
received an XT, giving an XT rate in this group 
of 13%. After IVIG the relative risk of requiring an XT is 0.28, thus with IVIG the XT rate 
would be reduced to 3.6%, decreasing the number of XTs to three and therefore prevent-
ing eight. If IVIG were administered to all babies with moderate or strongly positive DCT, in our population the number needed to treat would be 10.6 to prevent one XT. The degree of positivity of the DCT is an objective validated assessment of the strength of antigen/antibody reaction, determined by the degree of agglutination in the laboratory.1 During the three year period of this database there was only one infant who had only a 
weakly positive DCT and required an XT. 

We therefore agree that all the usual pro-
cedures regarding documentation of batch number and so on are followed as for any other 
blood product. IVIG has been used pre-
viously even in preterm and low birthweight 
infants2 and is currently being used in the 
INTS Trial.3 As with any drug or blood product 
we will need to remain vigilant for the occur-
rance of any adverse events.

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Discharging twins separately from neonatal units
We recently had a debate in our unit about whether or not it was better for a well twin to 
remain with its sibling in hospital until the latter was fit to be discharged. Our current 
practice is to keep the well twin in the special 
care baby unit until its twin is fit for discharge. The group who favoured separate discharge cited reduced risk of nosocomial infection, decreased costs, cot availability, and the possibility of settling into a routine with one twin at home as supportive factors for their argument. Those against separate dis-
card impaired bonding, breast feeding 
difficulties, and transport issues as their 
reasons.

We took the discussion to the RCPCH 
and NICU-net email discussion groups and found 
no clear consensus. Our American colleagues routinely send multiples home separately and 
cite health insurance companies as a major 
factor in this decision. They find little problem 
with this arrangement. The opinions were split between the two camps. British doctors 
saw in favour of asking the parents’ 
opinion, so we identified 10 sets of twins from the 
last three years who could have been sent 
home separately. We then sent their parents a 
questionnaire exploring their opinions; five 
(50%) were returned.

Most parents agreed that their twins were ready for discharge at different times and said 
that they would have preferred separate discharge. However, they believed that they 
had been given this option and had not taken it. They realise that this would have caused 
problems with visiting, feeding, and bonding with the remaining twin even although they 
all had their own transport. They did not think that having one twin home first would 
have helped them to adjust and settle into a 
routine. Their preferred option would have 
to be to have roomed in with the well twin 
and the other twin stayed on the special care 
baby unit.

Our current practice is that we have an 
informed discussion with the parents when this 
situation arises. As one email respondent 
(a doctor and father of twins) wrote “Bringing 
up twins is full of decisions about when to 
pair them and when to split them up.”

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Role of serum peak levels of vancomycin in neonatal intensive care units
We would like to comment on the article by Tan et al.1 The purpose of measuring serum 
levels of a drug is either to monitor the toxicity of the drug or the therapeutic concentra-
tion for a particular condition. Emergence of infections with β-lactam-resistant Staphylococ-
cus epidermidis, Staphylococcus aureus, and Ente-
roccus sp., has led to the frequent use of van-
comycin in neonates. Vancomycin has 
historically had a reputation for toxicity. Many of its original adverse reactions, includ-
ing ototoxicity and nephrotoxicity, were prob-
ably due impurities in the formulation. Now that a more purified form is available, these 
adverse reactions are uncommon. However, 
concomitant administration with aminogly-
cosides or other nephrotoxins may increase 
the risk of toxicity.1 Effective drug therapy is measured by response, not by achievement 
of a particular circulating drug concentration. 
Because the association between vancomycin 
peak concentrations and toxicity is poor, we 
have recommended measuring trough con-
centration only1 as this study is clearly 
documenting, but others have suggested not 
measuring any concentrations in the majority 
of children with normal renal function.1 
However, in critically ill premature neonates 
with poor glomerular filtration rate, prematu-

rity, and compromised cardiovascular func-
tion, it remains prudent to measure both peak 
and trough concentrations in those with poor 
or changing renal function. Caution must be 
exercised when other nephrotoxic or oto-
toxic drugs such as aminoglycosides are admin-
istered concurrently.1 In this study, the authors 
did not mention the concomitant use of 
aminoglycoside.

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Methaemoglobinemia with concurrent blood isolation of Saccharomyces and Candida

Saccharomyces boulardii is closely related to Saccharomyces cerevisiae and is used as a biotherapeutic agent, although some reports suggest pathogenicity. We present a case of neonatal sepsis with concurrent methaemoglobinemia, occurring after a brief period of treatment with S boulardii. A male infant was born at 30 weeks of gestational age by caesarean section because of intraterine growth restriction and maternal hypertension. The baby was well apart from persistent gastrointestinal symptoms that hampered feeding and forced parenteral support. During the third week of life, administration of S boulardii (Codex DNB; half a capsule a day, equivalent to 2.5 x 10^6 organisms) was started in an attempt to prevent bacterial overgrowth. After four days of treatment, the baby developed symptoms suggesting sepsis and an unexplained methaemoglobinemia (methaemoglobin concentration = 16%). Codex was stopped and empirical antibiotic coverage, including liposomal amphotericin B, was started. Blood cultures showed growth of S cerevisiae and Candida albicans, but the central venous catheter tip was negative. Methaemoglobin levels halved in two days (7%), but remained constant during the following two weeks of antifungal treatment. Blood cultures at that point showed growth of S cerevisiae, which is susceptible to amphotericin B, in the absence of any clinical finding; again, the catheter tip was sterile. Methaemoglobin concentration was still abnormal (6.2%). Liposomal amphotericin B treatment was prolonged for a further six days and then discontinued. At this time, methaemoglobin levels were near normal (3%), and blood cultures were negative. The gastrointestinal symptoms resolved with age and full gastrointestinal function was achieved.

Recovery of Saccharomyces two weeks after administration had been stopped suggests persistence in the gut. It is tempting to think that the methaemoglobinemia was caused by the continued presence of the yeasts, perhaps through excessive host production of nitric oxide. Several studies have shown that nitric oxide plays a pivotal role in the interaction between yeasts and the phagocytic system, and it is well known that this radical readily oxidises haemoglobin. It is also reasonable to link late bloodstream invasion by Saccharomyces to previous enteric mucosal damage caused by a Candida infection, which was itself probably gut related. The recovery of S cerevisiae in place of S boulardii has been reported by others, and can be explained by the similarities between the two. It is ironic that the intervention used to prevent sepsis from enteric overgrowth not only did not succeed but was itself a cause of the problem that it was intended to prevent.

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References

Wafting does work

We were interested to see the article “Oxygen administration in infants”, and the letter responses. The original article and letters were unsure of the efficacy of “non-contact” oxygen delivery or “wafting” as it is more commonly known. Our study “The efficacy of noncontact oxygen delivery methods”, demonstrated how effective wafting oxygen can be. We found that an area of 34cm by 37cm obtained a concentration of >30% when oxygen is delivered by face mask at 10 l/min. Although this is not a substitution for the more reliable methods of administration as detailed by Drs Frey and Shann, in the short term it can be used with confidence.

We caution against holding a self inflating resuscitation bag over an infant’s airway (without manipulation of the bag itself), as it delivers a negligible amount of oxygen. It is much more efficient to use the oxygen tubing without any attachments.

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References

We wish to apologise for an error that occurred in a letter by Daniels et al (Arch Dis Child Fetal Neonatal Ed 2003;88:F257). The first line of the third paragraph should have read: The salient results were that two thirds of granulomas resolved over a three week period without cauterisation.

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