

Fluid Balance in RDS

There is a natural contraction of the Extra cellular fluid volume (ECFV) in newborn babies. This is caused by a diuresis that follows a naturesis and lung function improves after this diuresis¹. This natural contraction of ECFV is delayed in babies who have surfactant deficient RDS.² Newborn babies have an impaired ability to excrete sodium thus sodium supplementation should start only after they have had a diuresis which will be borne out by weight loss and increased urine output.³ In neonates who have been given natural surfactants this diuresis may occur sooner⁸. Fluid input for premature infants should be carefully restricted so that physiological needs are met without allowing significant dehydration. This practice could be expected to decrease the risks of patent ductus arteriosus and necrotizing enterocolitis--and perhaps the overall risk of death--without significantly increased risk of adverse consequences (Bell 2001 (Cochrane Review))^{4,5,6,7}.

Daily weighing and regular (twice daily) measurement of serum sodium are important to achieve this particularly in the smallest babies (<1000g).

SUMMARY and level of evidence

1. Give sodium only after a post-natal diuresis A
 2. Restrict fluids to meet physiological requirements without allowing dehydration A
 3. Weigh daily and measure sodium twice daily D
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 2. Modi N, Betremieux P Midgley J,Hartnoll G. 2000 Postnatal weight loss and contraction of extracellular compartment is triggered by atrial natriuretic peptide. Early Hum Dev 59,201-208
 3. Hartnoll G, Betremieux P, Modi N Randomised controlled trail of postnatal sodium supplementation on body composition in 25 to 30 week gestational age infants. Arch Dis Child Fetal Neonatal Ed 2000; 82:F24-F28
 4. Hartnoll G, Betremieux P, Modi N Randomised controlled trail of postnatal sodium supplementation on oxygen dependency and body weight in 25-30 week gestational age infants .Arch Dis Child Fetal Neonatal Ed 2000 ;82:F19-F23
 5. Randomised controlled trial of postnatal sodium supplementation in infants of 25-30 weeks gestational age: effects on cardiopulmonary adaptations Hartnoll G, Betremieux P, Modi N .Arch Dis Child Fetal Neonatal Ed 2001;85 : F29-32
 6. E F Bell etall Effect of fluid administration on development of symptomatic patent ductus arteriosus and congestive heart failure in premature infants . New Engl Journal of medicine 1980;302:598-604
 7. Van Marter LJ, Leviton A et al Hydration during the first days of life and the risk of bronchopulmonary dysplasia in low birth weight infants Journal of Pediatr 1990 :116:942
 8. Murdoch E, Kempley ST, The effects of synthetic and natural surfactants on fluid balance in acute respiratory distress syndrome Eur J Pediatr 2000 ; 159 767-769

Blood transfusions to replace Blood loss

1. Preterm infants with RDS become anaemic due to blood sampling and blood transfusions may be necessary to correct anaemia and or hypotension. However there are significant adverse effects of transfusions (4) that not only include infectious complications but also the risk of fluid overload, skin injury due to tissue IV cannulae, and a possible increased risk of ROP and CLD. Therefore transfusions should be limited to the minimum number required for the infants' optimal well-being. It is useful to note that simple approaches such as reducing the frequency of blood sampling are effective⁵. Replacement therapy should be based on a transfusion guideline which is relatively restrictive to prevent unnecessary transfusion⁶. Measuring the volume of blood lost and replacing after a certain volume is only useful if the infant's status is also taken into consideration when deciding whether to transfuse. Dividing a single donation of red cells into satellite packs reduces donor exposure¹. Routine use of frusemide is not recommended.² Recombinant human erythropoietin is not effective in reducing transfusion requirements of sick neonates who require frequent blood sampling^{3,4}.

Transfusion guidelines for neonate and older children – British Committee for Standards in Haematology Transfusion task force

1. Betremieux P, Hartnoll G, Modi N Eur J pediatric 1997 88-89 Should frusemide be prescribed after packed cell transfusions in newborn ?
2. Franz AR et al Red cell transfusion in very and extremely low birth weight infants under restrictive transfusion guidelines :is exogenous erythropoietin necessary ? Arch Dis Child Fetal Neonatal Ed 84 F96 -100
3. Maier RF et al The effect of epoietin beta on need for transfusions in very-low-birth weight infants European multi-centre Erythropoietin Study Group New Eng J Med 1994 330 1173-1178
4. Maier RF et al High versus low –dose erythropoietin in extremely low birth weight infants European multi-centre Erythropoietin Study Group J Pediatr 1998 132 866-870
5. Bifano EM, Curran TR Minimizing blood donor exposure in the neonatal intensive care unit - current trends and future prospects. Clin Perinatol 1995;22;657-669)
6. Ramasethu J, Luban NLC Red blood cell transfusions in the newborn. Seminars in Neonatology 1999;4; 5-16).

Summary and Level of Evidence

1. Using a restrictive blood transfusion regimen is a useful way to decrease the number of transfusions given B
2. Routine use of frusemide with blood transfusion is not recommended B
3. Recombinant EpO is not effective at reducing blood transfusions in the first two weeks after birth A
4. Dividing a single donation of red cells into satellite packs reduces donor exposure. B