Fifteen-minute consultation: stabilisation of the high-risk newborn infant beside the mother

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ABSTRACT
Paediatric and adult resuscitation is often performed with family present. Current guidelines recommend deferred umbilical cord clamping as part of immediate neonatal care, requiring neonatal assessment next to the mother. This paper describes strategies for providing care beside the mother using both standard resuscitation equipment and a trolley designed for this purpose.

INTRODUCTION
Family presence at both paediatric and adult resuscitation is common and part of international resuscitation guidelines.1 2 Being present at resuscitation can decrease anxiety and feelings of helplessness and improve grief for the family.3 4 Having the family present does not seem to interfere with or impede clinical care.5

For parents of high-risk infants, the first experience of their baby may be on the neonatal unit, and many parents describe touching their baby for the first time as a particularly powerful experience.5 Delay in this first contact may be due to concerns around maternal or infant health or simple physical separation.6

Current guidance is to defer umbilical cord clamping for uncompromised term and preterm infants.7 8 For high-risk infants, assessment following newborn life support (NLS) algorithms of whether stabilisation is needed must therefore take place while the cord is intact. Having the infant beside their mother facilitates this assessment.

We have developed two strategies for providing stabilisation beside the mother: the first using standard equipment and the second using a mobile trolley (LifeStart).9 10 These were developed as part of a programme of work aimed at improving the quality of care an outcome at very preterm birth. Working with an industry partner, we developed the LifeStart trolley (figure 1). This is a small, mobile, height-adjustable trolley with a horizontal platform and heated mattress; additional resuscitation equipment, including gas cylinders, can be mounted on side rails.11

Both strategies for providing care beside the mother were used for infants born before 32 weeks gestation in the Cord Pilot Trial12 and for other high-risk births.9 The Cord Pilot Trial compared umbilical cord clamping after at least 2 min with clamping within 20 s. Providing care and stabilisation beside the mother ensured that neonatal care was the same regardless of when the cord was clamped.

PLANNING BEFORE THE BIRTH
To facilitate neonatal care beside the mother at birth, significant planning involving the multidisciplinary team (MDT) is needed. Guidance should be drawn up with input from all those involved in the care of the woman and her baby (figure 2). This should include planning for births within the labour suite and the obstetric theatres.

Training materials should be developed and can include presentations, documents, videos and simulation scenarios. In our experience, simulations with all members of the MDT, based on NLS scenarios and protocols, adapted with care beside the mother, were the most effective at embedding practice. Continual training, integrated into changeover inductions for rotating junior doctors, ensures new members of the team are familiar with procedures.

WHAT EQUIPMENT TO USE
Different types of equipment can be used to provide neonatal care beside the mother. We have experience of using both...
standard equipment (the Draeger Resuscitaire and Fisher and Paykel CosyCot) and the LifeStart trolley. The set up required for each of these platforms is slightly different.

**Standard equipment**

For standard equipment, a solid-based mattress should be used, and the side barriers of the platform lowered to create a flat surface. At vaginal births, wheel the platform longitudinally alongside the mother’s bed and lock the wheels. Adjust the height of both the platform and the mother’s bed so they are at the same level, and slide the mattress part-way across the side of the mother’s bed to provide a stable surface for stabilisation of the neonate. Rotate the overhead heater to cover the area directly over the infant (figure 3).

At caesarean section place the platform end on to the operating table and drape it with sterile towels (figure 4). Adjust the height of the platform and the operating table so they are at identical levels.

**The LifeStart trolley**

At vaginal births adjust the height of the platform so that it can rest on the mother’s bed (figure 5). At caesarean section wheel the trolley end on to the operating table and adjust the height of the platform to the mother’s thighs (figure 6). For both types of birth, the trolley is located with the neonatal interface facing the neonatal team.

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**Figure 1** The LifeStart trolley.

**Figure 2** Members of the MDT required for planning neonatal stabilisation beside the mother. MDT, multidisciplinary team.

**Figure 3** Demonstration of how to set up standard equipment for neonatal care beside the mother immediately after birth at vaginal births.

**Figure 4** Demonstration of how to set up standard equipment for neonatal care beside the mother immediately after birth a caesarean section.
**Best practice**

**PROVIDING CARE WITH EITHER EQUIPMENT**

Prior to birth, the equipment should be moved to the agreed position and the wheels should be locked. If deferred umbilical cord clamping is taking place, the position of the umbilical cord needs to be considered. If the woman gives birth in a semirecumbent position, the baby should be placed on the mattress with the cord over mother’s thigh, but if giving birth in the lithotomy position, the baby should be passed underneath the thigh. Care is needed to ensure that no pressure is applied to the cord.

**Caesarean section**

Prior to birth, the optimal positioning of the neonatal resuscitation equipment should be agreed with the theatre team. This is usually between the operating obstetrician and the diathermy machine with the surgical assistant and the scrub nurse on the opposite side of the operating table.

Maintaining the sterile field is essential. Prebirth equipment checks of all non-sterile equipment are needed, and then the platform covered in sterile drapes to ensure continuation of the sterile field. At least two members of the neonatal team should scrub, another member of the team opens packets and handles non-sterile equipment. For example, inserting the end of the suction catheter into the tubing.

Shortly before birth, the platform or trolley is moved into the agreed position and its wheels locked. At birth, the obstetrician either passes the baby to the neonatal team or for a very preterm birth places the baby’s feet first into a sterile plastic bag or wrap. The stethoscope, hat and laryngoscope handle are not placed on the sterile field until after the obstetrician has placed the baby on the platform to minimise the risk of contact with non-sterile equipment.

**CURRENT EVIDENCE AND FUTURE RESEARCH**

With good planning and training we have shown that stabilisation of very preterm infants can be conducted beside the mother using both forms of resuscitation platform. Despite initial anxieties about the equipment and providing care in front of parents, preliminary feedback from parents and clinical staff about care beside the mother appears largely positive. Parents who expressed their opinion commented that they were pleased that the baby was so close to them and appreciated being able to witness management and converse and interact with clinicians. Many mothers spontaneously touched their baby and others did when invited to do so. Some parents found it stressful watching their baby receive treatment, but none regretted the decision to have their baby beside them. Parents raised concerns about staff experiences including whether they felt pressure from resuscitating at the bedside.

Clinicians wondered about the impact on parents of watching stabilisation at birth. They felt, however, that parents being closer to their baby was important,
and the close proximity aided communication between the clinical team and the parents. Access to the baby and the ability to assess the baby were undiminished, although access to equipment has been raised as a potential concern.

Further evaluation of care beside the mother in a wide range of settings is needed and should include use of standard equipment as well as the LifeStart trolley. As well as further evaluation of parents’ and clinicians’ experiences, this should include assessment of outcome for babies and parents including, but not restricted to, temperature on admission to neonatal care can be provided beside the mother. It is With good MDT planning and working delivery room

CONCLUSION With good MDT planning and working delivery room neonatal care can be provided beside the mother. It is important to develop a framework of teaching packages and to provide continual MDT training.

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REFERENCES


