

Guideline

Stabilisation and transfer of an infant with exomphalos

1 Scope

For use within the Paediatric and Neonatal Decision Support and Retrieval Service for the East of England.

2 **Purpose**

To provide safe, efficient and practical guidance for stabilisation and transfer of infants with Exomphalos.

3 **Definitions and abbreviations**

PaNDR: Paediatric and Neonatal Decision Support and Retrieval Service

Beckwith-Wiedemann syndrome:

An overgrowth disorder, usually present at birth. Common features include macroglossia (large tongue), macrosomia ('big body'), exomphalos and neonatal hypoglycaemia.

NBM: nil by mouth

NLS: newborn life support

Pentalogy of Cantrell:

A rare syndrome characterized by the following five features: exomphalos, diaphragmatic hernia, sternal cleft, pericardial defect, cardiac defect. Severity varies from mild defects with incomplete expression of the disorder to complete ectopia cordis (heart displaced outside the thoracic cavity so not protected by the chest wall) with large exomphalos.

Trisomies: Genetic abnormalities involving three instances of a particular

chromosome instead of the normal two (e.g. trisomy 18,

trisomy 21).



4 Introduction

- Exomphalos (alternatively: omphalocele) is a congenital abdominal wall defect. Abdominal wall defects occur when a fetus's abdominal wall does not develop fully while in utero, resulting in the intestine developing outside the abdomen. The intestines develop inside the umbilical cord in early pregnancy, and then move inside the abdomen by 12 weeks of pregnancy. An exomphalos occurs if the abdominal contents protrude into the base of the umbilical cord and are covered by a peritoneal membrane. Malrotation of the intestines is commonly seen.
- If the abdominal defect at the base of the exomphalos is less than 5cm it
 is called exomphalos minor and may contain only tissue left over from
 structures in the unborn fetus's digestive tract. If it is larger than 5cm it is
 called exomphalos major. Part of the liver may also be present in the
 sac along with bowel.
- The cause of exomphalos is not always known. Familial occurrence has been described. Up to 80% of babies with exomphalos have other serious abnormalities such as heart defects, gastro-intestinal and renal anomalies and chromosomal abnormalities (e.g. trisomies). It can also occur as part of a syndrome such as Beckwith-Weidemann syndrome or Pentalogy of Cantrell.

Figure 1: Exomphalos minor





5 Management

• At birth

- Ensure careful clamping of the cord, preferably with a plastic cord clamp, at least 10cm away from organs within exomphalos, to avoid damaging any bowel that may be present in it.
- o Resuscitate as required according to NLS guidelines.
- Use only latex free gloves and equipment to reduce risk of latex allergies developing.
- Aim to avoid prolonged mask ventilation, and have a low threshold for intubation. Intubation should be undertaken rather than prolonged mask ventilation or CPAP. This is done to avoid distending the bowel, which may make surgical repair more difficult.

• Care of membranous sac

- Wrap the abdomen securely with cling film and ensure the gut is well supported to avoid compromising its blood supply. It may be necessary to support sides of exomphalos to keep it supported and central.
- Handle the baby carefully to avoid tearing the sac.
 If the sac is torn (at delivery or postnatally) urgent surgical intervention is required.

Nursing position and monitoring

- Nurse supine in the transport incubator so that the gut can be visualised within the sac during transfer.
- Monitor the bowel at least every 15 minutes for dusky or blanching colour changes: if compression, kinking or twisting is suspected remove and rewrap as above, and/or consider positioning right-side up.

NG tube

 Ensure early introduction of a large bore NG tube (size 8 or 10F) on free drainage with at least hourly aspiration to avoid the bowel becoming distended with air.

Access

Do not attempt umbilical lines.

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- Transfer with 2 x peripheral venous linesFluids
 - o NBM
 - Start maintenance IV fluids (10% dextrose) and replace NG losses ml/ml with 0.9% Sodium Chloride with 10mmol/500ml Potassium Chloride.
 - Infants with abdominal wall defects are at risk of dehydration from excessive fluid losses, although the peritoneal membrane covering the intestines in exomphalos offers some protection from this (unlike in gastroschisis where the gut is completely exposed). Observe closely for the need for additional fluid boluses.

Monitor temperature

These babies are at risk of hypothermia

Glucose

 It is important to monitor blood glucose levels regularly and correct any hypoglycaemia with IV dextrose (in view of association with Beckwith-Wiedemann Syndrome)

Sepsis

 Consider antibiotics if risk factors for sepsis present or there are additional clinical concerns.

Associated anomalies

Examine baby for other anomalies.

6 Monitoring compliance with and the effectiveness of this document

The transport team will monitor that the guideline is being adhered to on a case by case basis (very rare condition). This will be fed back to the senior team at a governance meeting.

7 References

London Newborn Transport Service guideline - https://london-nts.nhs.uk/wp-content/uploads/2015/01/Exomphalos-NTS-Guideline.pdf (unavailable)

Southern West Midlands Newborn Network guideline -

https://www.networks.nhs.uk/nhs-networks/southern-west-midlands-newborn-network/documents/Attachment%206%20Exomphalos%20Final%20jan%20201 2.pdf (unavailable)

Starship (NZ) guideline - https://starship.org.nz/guidelines/surgery-management-of-abdominal-wall-defects-in-the-neonate

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UpToDate - https://www.uptodate.com/contents/omphalocele-prenatal-diagnosis-and-pregnancy-management

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