What is an implantable port?

An implantable port consists of a titanium or plastic chamber connected to a thin plastic tube (catheter). The entire device is placed under the skin and the catheter is threaded into a vein in the neck until it reaches the right side of the heart. To access the bloodstream, a special type of needle is inserted through the skin into the chamber. An implantable port provides easy and reliable access to your child’s bloodstream.

There are many reasons why your child might benefit from an implantable port, but the most common reasons are for taking regular blood samples for testing or for giving medicines such as chemotherapy or other medicines intravenously on a long term basis.

Implantable ports can be inserted by different teams at GOSH, including the surgical department as well as interventional radiology. In Interventional Radiology, ports are placed by a variety of operators, such as doctors, specialist nurses and radiographers.
What happens before the procedure?
You will already have received information about how to prepare your child for the procedure in your admission letter. You may need to come to GOSH before the procedure so that your child can have a pre-admission assessment to check that they are well enough. The appointment may involve taking blood samples and other tests.

Sometimes an implantable port insertion happens while your child is still in hospital. The doctor will explain the procedure in more detail, discuss any worries you may have and ask you to sign a consent form giving permission for your child to have the procedure. If your child has any medical problems, please tell the doctors.

Many of the procedures we perform involve the use of x-rays. Legally, we are obliged to ask any girls over the age of 12 whether there is any chance they might be pregnant. This is to protect babies in the womb from receiving unnecessary radiation.

What does the procedure involve?
Your child will need to have a general anaesthetic for this procedure. It is important that your child does not eat or drink anything for a few hours before the anaesthetic. This is called ‘fasting’ or ‘nil by mouth’. Fasting reduces the risk of stomach contents entering the lungs during and after the procedure.

You will be informed the night before the procedure of the time that your child should be ‘nil by mouth’ – in other words, have nothing to eat or drink before the anaesthetic. Fasting times are provided in your admissions letter – in broad terms, this is six hours for food (including milk), four hours for breast feeding and two hours for clear fluids before the procedure.

It is equally important to keep giving your child food and drink until those times to ensure they remain well-hydrated and get adequate nutrition. This may involve waking your child in the night to give them a drink which we recommend.

Once your child is under general anaesthetic, the operator will inject some local anaesthetic into the side of the chest and make an incision to create a ‘pocket’ under the skin. The chamber of the port is inserted into this pocket and is stitched in place. The operator will then use an ultrasound scan to look for a suitable vein in the neck. Once they have found one, they will inject the area around the vein with local anaesthetic to make it numb for a few hours.

The catheter section of the port is then tunnelled under the skin on the chest wall until it is near the neck vein. They then make a small puncture into the neck vein and thread the catheter into it until it reaches the right side of the heart. The operator makes a final check that the implantable port is working correctly by taking an x-ray. They will then confirm this by taking a small blood sample from the port, flush the port with a salt water solution (saline) and then fill it with heparin solution to stop blood clots forming.

The chest incision is closed with dissolvable stitches and then covered with sticky paper stitches. These should
stay in place for five days or so. The small puncture wound in the neck is covered with a dressing. The port can be used immediately by inserting a special needle into the reservoir. If a needle is inserted, the entire area will be covered with a see-through dressing to keep the needle in place and help prevent infection.

**Are there any risks associated with the insertion procedure?**

The implantable port is inserted while your child is under general anaesthetic. Although every anaesthetic carries a risk, this is extremely small. There is a small risk of bleeding when the incisions are made but this is unlikely as they are very small.

As the port gives direct access to the bloodstream, infection can be a risk after the procedure. You will need to look after the port very carefully but we will teach you everything you need to know before you leave GOSH. The nurses will also give you a written information booklet to remind you about what we have taught you.

**What happens afterwards?**

Your child will return to the ward after they have recovered from the general anaesthetic. Some children feel sick and vomit after a general anaesthetic. Your child may have a headache or sore throat or feel dizzy, but these side effects are usually short-lived and not severe. The area where the implantable port was inserted will feel uncomfortable for a while afterwards but this soon passes. The nurses will give your child pain relief as needed. Your child can start eating and drinking as normal once they feel like it. The implantable port can be used immediately if required. The sticky paper stitches should stay in place for five days or so but will then start to drop off – this is normal. The see-through dressing may need to be changed after 24 hours, particularly if the incision used to create the pocket for the port has oozed after the operation.

**Are there any alternatives to insertion of an implantable port?**

The doctors will only suggest that your child needs an implantable port or other central venous access device if they are having frequent or long term treatment needing access to the bloodstream. If your child only needs intravenous medicines or blood samples occasionally, a cannula (thin plastic tube) inserted into a vein on the hand or foot each time might be a better option.

There are various types of central venous access device. However, depending on your child’s age and the reason why they need one, some options may not be appropriate. The doctors and nurses will explain the benefits and drawbacks of each available option before you and your child make a decision.
Going home
If your child does not need to stay in hospital for treatment, you can return home once you are confident in caring for the port. Your nurse will provide you with instructions on how to look after the port at home. We will also make contact with your local children’s community nursing team who will help you to look after the port, for instance by changing dressings or by taking blood samples.

In the first few days after insertion, you should call the hospital if:

- The neck and chest incisions look red, swollen and feel hotter than the surrounding skin
- The incisions are oozing
- Your child is in a lot of pain and pain relief does not seem to help
- Your child has a temperature of 38°C or higher

If you have any questions, please telephone 020 7405 9200 and ask for the ward from which your child was discharged

Notes

Compiled by the Interventional Radiology and Intravenous Therapy teams in collaboration with the Child and Family Information Group

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