Veno-occlusive disease (VOD) - Information for young people and families



Introduction

This factsheet has been written to help you, as a young person, parent or carer to understand veno-occlusive disease (VOD) - one of the less common but still serious complications which can occur during stem cell transplant (SCT). Veno-occlusive disease is not another illness, but a complication that affects the liver. The information given in this factsheet explains what VOD is and why it happens, how it is diagnosed, what the impact of it is, and how it is treated. It should give you a little more background and help you when you are asking questions to the SCT team. This factsheet provides details about:

- Young people/children at greater risk of developing VOD.
- Signs and symptoms.
- Accurate diagnosis.
- Managing the care of a young person/ child with VOD.

Veno-occlusive disease may also be referred to as sinusoidal obstruction syndrome (SOS). VOD can range from mild to severe, and occurs in approximately 10 - 30% of children/young people who have received total body irradiation (TBI), Busulfan, or high dose Melphalan as part of their conditioning treatment before a stem cell transplant. Although it is serious it is usually a temporary problem, but it can be more complicated or even cause long-term problems. Your SCT team will discuss this in more detail with you.

In VOD the chemotherapy causes damage to cells in the liver, so the veins within the liver become increasingly occluded (obstructed) by cellular debris (dead or damaged cells), causing the blood flow from the liver to back up. The protein-rich fluid content of the blood leaks out into the peritoneal cavity (a potential space between the two membranes that separate the organs in the abdominal cavity from the abdominal wall). There will be an increase in weight as the fluid collects in this space. As the tummy gets bigger it will become uncomfortable and may be quite painful requiring pain relief medicine.

The role of the liver and the impact of VOD

The liver is a large organ in the right upper part of the abdomen, you can't usually feel it as it is under your ribs. It is made up of two lobes, each lobe consisting of 50,000 to 100,000 lobules. Lobules are a series of liver cells in rows and beyond each row is a tiny channel or blood vessel, called a sinusoid. Blood flows through the sinusoids, bringing nutrients to be stored in the liver cells, and collecting products that need to be removed from the body. The sinusoids are lined with a type of cell called endothelial cells and these form the linings of the blood vessels. It is the endothelial cells that are damaged and cause VOD, and this damage stops the liver from working properly.

The liver helps the body in many ways, some of which are described below. If the cells of the liver are damaged these functions may be slower or not as effective as usual.

Digestion

The liver helps in the breakdown of carbohydrates, proteins, vitamins, minerals and fats gained from food. You may not be able to eat or drink enough and may need support with this.

Detoxifies (removing/changing)

Liver cells help to break down drugs so they can be used by the body. They also remove some drugs from the body once their job is done.

Breakdown of red blood cells

Red blood cells live for around 120 days and when they die, the liver is involved in their breakdown. Bilirubin (a yellowish pigment found in bile, a fluid made by the liver) is produced when red blood cells break down. Bilirubin is usually removed from the body in bile. In VOD it may not be removed from the body in this way. It may stay in the blood, and as your body tries to balance this, you start to look jaundiced (whites of eyes and skin become yellow).

Production

The liver produces vital blood products, including factors essential for stopping you from bleeding (clotting).

Risk factors

We know that some young people/children are more likely than others to develop VOD. If you fall into any of the categories listed below it does not mean that you will get VOD, but that you may be more likely to.

Before the transplant, your SCT team will discuss your care in more detail and will explain why this happens. Below is a list of some of the circumstances which may trigger VOD:

- · Children with Osteopetrosis.
- Children with Thalassaemia.
- Children with Adrenoleukodystrophy (ALD).
- Children with Macrophage Activating Syndromes (MAS, i.e.Haemophagocytic Lymphohistiocytosis HLH, Griscelli).
- Evidence of previous liver damage (including prior abdominal radiotherapy).
- Previous treatment with certain cytotoxic (anti-cancer) drugs. (This generally affects patients aged over 15, but can also be seen in babies or children of any age.)
- Any infection before or during the conditioning phase of SCT.
- Long-term treatment drugs that may harm the liver.
- Unrelated donor transplants.
- Second stem cell transplant.
- The use of Busulfan, high dose Melphalan and/or total body irradiation in the conditioning regimen.

Signs and symptoms of VOD

Sudden weight gain

May take place over a few days or, in extreme cases, hours. Due to fluid collecting in the peritoneum rather than staying in the blood system. The fluid has collected in the wrong place, so it cannot be passed by the kidneys.

Ascites (fluid collecting in the abdomen)

As the liver swells, fluid is squeezed out of the liver and collects in the abdomen (tummy). This fluid contains a lot of protein which encourages more fluid to collect in the abdomen. As a result, the abdomen gets bigger and it may be uncomfortable or difficult to breathe especially when lying down.

Abnormal clotting and requirement for more frequent platelet transfusions and blood products

Abnormal clotting may require plasma or other clotting factors, e.g. cryoprecipitate or vitamin K. More frequent platelet transfusion may be required as platelets do not live as long in patients who have VOD.

Pain

The liver is surrounded by a capsule. As it enlarges, due to VOD, the liver presses on the capsule and this can be painful. Older children and young people may complain of pain in their right upper abdomen. On examination of the abdomen the liver may feel bigger than usual.

Abnormal bilirubin

In the breakdown of dead red blood cells bilirubin is formed. The liver helps to eliminate this from the body. In VOD the liver is unable to do this as well as normal. Blood tests, (liver function tests LFTs) will also show that the liver is not working as it should be

Jaundice

If the level of bilirubin in the blood becomes very high the skin and whites of the eyes may look yellow. This is referred to as jaundice, which may also make your skin dry and itchy.

Nausea and vomiting

Nausea and vomiting occur for many reasons in VOD: pain, ascites, or an enlarged liver pressing on the stomach are all possible causes.

Lethargy

The symptoms described above will all make you feel tired and generally unwell.

How is the diagnosis of VOD made?

Your SCT team will be familiar with the signs and symptoms of VOD. Various investigations may be needed to make an accurate diagnosis. Some of the symptoms listed may be signs of other problems with your SCT, such as graft versus host disease or infection. However, it is only in VOD that there is unexplained weight gain and fluid retention. There are two techniques available to confirm the diagnosis of VOD. The first is a procedure using ultrasound, which shows reversal of blood flow through the liver. This is done using a probe that is gently run over the skin to give a picture of what is happening in the liver. It shouldn't hurt but may be uncomfortable if your abdomen is already painful. A liver biopsy may be needed to confirm the diagnosis. Your SCT team will discuss which is the best option for you.

Treatment of VOD

Monitoring of Busulfan levels

In recent years monitoring of chemotherapy levels has become routine in SCT. Monitoring of Busulfan is aimed at reducing the risk of VOD.

Prophylaxis (preventative)

Where there is a risk of VOD, prophylactic or preventative treatment is often given. An oral medicine called Ursodeoxycholic Acid or an intravenous (IV) medicine, Defibrotide, may be given from the start of the chemotherapy conditioning,

Defibrotide

Defibrotide is commonly used for the prevention of VOD in children and young people undergoing SCT that are considered to be at high risk of developing the disease, or as treatment of suspected or established VOD. Defibrotide is usually given as an intravenous infusion (over 2 hours, four times a day, directly into a vein). In severe cases of VOD, or poorly responding cases, the dose may be increased. Defibrotide works in a number of ways; antithrombotic (anti-clotting), anti-inflammatory (reduce swelling) and anti-ischaemic (reduce the restriction of blood flow). Defibrotide appears to have a protective effect against endothelial cell injury caused by chemotherapy drugs.

The management of medicines

This can become a very difficult time as all the therapies required to support a child/ young person through transplant have an impact on the liver, including IV feeding (TPN - total parenteral nutrition), analgesia (pain killers), antibiotic or antimicrobial treatments or blood products. The SCT team will plan your care on a daily basis to consider how to best manage your medication. Your care will be discussed with you but remember you can always ask questions if you are finding something difficult to understand.

Supportive care

The management of fluid retention and ascites are an essential part of the treatment of VOD. The aim is to reduce the amount of fluid that has collected in the abdomen and elsewhere within the body. This is achieved by giving diuretics (drug treatment) to encourage the passing of urine, and restricting the amount of oral fluids allowed within a day. Restricting the amount you are allowed to drink within a day can often be very distressing if you are thirsty. Small drinks or ice lollies/cubes can occasionally relieve the distress.

To monitor the effects of this treatment regular blood tests will be done. The accurate monitoring of fluid input and output is essential, as well as weighing you and measuring your tummy once or twice a day. In severe cases of VOD, fluid retention and a distended abdomen can have an effect on your kidney function. Occasional support may be required from the renal team (kidney specialists). Your SCT team will discuss this with you should it be required.

VOD can cause you to be very uncomfortable, occasionally leading to a painful tummy. The use of pillows and wearing loose clothing may help you get comfortable. Your skin may also become dry and itchy so you should continue to bath every day and use emollients (moisturisers) to stop your skin from itching. Drug treatments with antihistamines may also help relieve the itching.

Do ask any members of the SCT team looking after you if there is anything you don't understand or if you have any questions. Written by Helen Webster on behalf of the UK Paediatric/Adolescent Stem Cell Transplant Nurses Group in conjunction with the CCLG Publications Committee, comprising multiprofessional experts in the field of children's cancer.

Production of this factsheet was funded by CCLG Designed and published by CCLG March 2014 Review date March 2017 © CCLG 2014

If you have any comments on this factsheet, please contact us at the address below. CCLG booklets are available to download from our website.

CCLG makes every effort to ensure that information provided is accurate and up-to-date at time of printing. We do not accept responsibility for information provided by third parties. Information in this factsheet should be used to supplement appropriate professional or other advice specific to your circumstances.





The CCLG supports the 1,700 children who develop cancer each year in Britain and Ireland. As an association for healthcare professionals involved in their care, it works to benefit children through development of the highest standards of care. CCLG is a major provider of accredited information for patients and families.

Children's Cancer and Leukaemia Group

3rd Floor, Hearts of Oak House 9 Princess Road West Leicester. LE1 6TH Tel: 0116 2494460 Fax: 0116 2494470 Email: info@cclg.org.uk Website: www.cclg.org.uk Registered Charity No: 286669



