

Escalated BEACOPDac in Hodgkin lymphoma

The possible benefits of this treatment vary; for some people this chemotherapy may reduce the risk of the cancer coming back, for others it may control the cancer and its symptoms. Your treatment will be fully explained by your doctor or nurse who will be happy to answer any questions you have. You will find it useful to refer to the booklet 'Chemotherapy, a guide' which gives general information on chemotherapy and side effects.

Your treatment

Your doctor or nurse clinician has prescribed for you a treatment with doxorubicin, cyclophosphamide, etoposide, dacarbazine, bleomycin, vincristine and prednisolone (Escalated BEACOPDac):

Day 1

Doxorubicin by short infusion Cyclophosphamide by short infusion Etoposide by short infusion

Days 2 and 3

Etoposide by short infusion Dacarbazine by short infusion

Day 8

Bleomycin by slow injection Vincristine by short infusion

Days 1 - 14

Prednisolone tablets every day for 14 days

Days 9 - 15

G-CSF injections under the skin

The treatment will repeat every 21 days (a 3-weekly cycle) Most patients will receive either 4 or 6 cycles

You will have a routine blood test and medical review every week whilst on treatment with escalated BEACOPDac. This is so your team can monitor and manage any side effects as well as assess your response to treatment.

After Cycle 2 of treatment, if clinically possible you will have your treatment at a location closer to home. The Christie are committed to providing treatment closer to home as part of the Outreach and Christie at Home service.



This treatment can have serious or possibly life-threatening side effects. It is very important that you report side effects straight away. Don't delay, if you feel unwell, please ring The Christie Hotline on **0161 446 3658**. The lines are open 24 hours a day.

• Take your **prednisolone** tablets with breakfast in the morning, not on an empty stomach as they may cause indigestion. It is also better to take them earlier in the day since they can make you feel more alert and prevent sleep.

Flu vaccinations

Is it alright for me to have a flu jab during the time I'm having chemotherapy?

It's safe to have a flu jab, but depending on the sort of chemotherapy you have had, it may not give quite as much protection against infection as usual. Some patients may need two vaccinations. However, if you you're thinking of having any other vaccinations, do check with your Christie doctor first, because some vaccines should be avoided.

Possible side effects

Chemotherapy can cause many different side effects. Some are more likely to occur than others. Everyone is different and not everyone gets all the side effects. Most side effects are usually temporary, but in some rare cases they can be life-threatening. It is important to tell your hospital doctor or nurse about any side effects so they can be monitored and, where possible, treated.

Common side effects (more than 1 in 10)

Increased risk of serious infection

You are vulnerable to infection while you are having chemotherapy. Minor infections can become life-threatening in a matter of hours if left untreated. Symptoms of infection include fever, shivering, sweats, sore throat, diarrhoea, discomfort when you pass urine, cough or breathlessness. We recommend that you use a digital thermometer so you can check your temperature. You can buy one from your local pharmacy.

If you feel unwell, you have symptoms of an infection or your temperature is 37.5°C or above, or below 36°C contact The Christie Hotline straight away.

Anaemia (low number of red blood cells)

While having this treatment you may become anaemic. This may make you feel tired and breathless. Let your doctor or nurse know if these symptoms are a problem. You may need a blood transfusion.

Bruising or bleeding

This treatment can reduce the production of platelets which help the blood clot. Let your doctor know if you have any unexplained bruising or bleeding, such as nosebleeds, bloodspots or rashes on the skin, or bleeding gums. You may need a platelet transfusion.

Lung function (Warning!)

Bleomycin may cause some changes to your lung tissue. Report any cough or breathlessness to your doctor without delay.

• Flu-like symptoms (Warning!)

Bleomycin may cause flu-like symptoms such as fever, aches and pains and shivering about 3 to 5 hours after it is given. These symptoms are temporary and should go within 12 to 24 hours. Paracetamol will help. If your symptoms persist or are particularly severe, please contact The Christie Hotline on 0161 446 3658 as these symptoms can also be a sign of infection or sepsis. Inform them when you had the chemotherapy and they will provide you with the appropriate advice.

• Changes to bowel habits

Vincristine as well as some anti-sickness tablets can cause constipation. Try to drink plenty of fluids and eat foods high in fibre. Report this to your hospital doctor who may prescribe a suitable laxative. Ask the

staff for a copy of The Christie booklet 'Eating - help yourself' which has useful ideas about diet when you are having treatment.

This treatment can also cause diarrhoea. Report this to your hospital doctor who may prescribe suitable medication once an infection has been ruled out.

Mood changes

Prednisolone tablets can affect your mood. This may result in mood swings and irritability. You may also have difficulty in sleeping. You may feel very energetic and have a good appetite when you take the tablets, and then lethargic and low in mood when you stop. If you experience this side effect and find it hard to tolerate, you should discuss it with your doctor.

Nausea and vomiting (sickness)

The severity of this varies from person to person. Anti-sickness medication will be given along with your chemotherapy to prevent this. You will also be given anti-sickness tablets to take at home. If you continue to feel or be sick, contact your GP or this hospital, because your anti-sickness medication may need to be changed or increased.

Hair loss

Hair loss is usually total but this depends on the number of cycles of chemotherapy you have. The hair falls out gradually 10 to 14 days following your first course of treatment. The time scale varies from person to person. Please remember that this is a temporary side effect and your hair will grow back when your treatment is completed. Very rarely, hair loss may be permanent.

The cancer information centre offers a coping with hair loss service to all patients where support, information and_advice will be given. Drop in, contact **0161 446 8100** or email **informationcentre@christie.nhs.uk**. Information about the wig service can also be found here and vouchers for wigs can also be obtained for eligible patients. The wig room provides a drop in service, please see The Christie leaflet 'The wig fitting service' for further information.

The Maggie's Centre runs a Talking Heads hair loss support workshop for anyone who is experiencing hair loss (both men and women). These sessions cover the practicalities of hair loss as well as offering support with its emotional impact. Contact Maggie's on **0161 446 4848** or email manchester@maggiescentres.org.

Lethargy

Some chemotherapy may make you feel tired and lacking in energy. It can be frustrating when you feel unable to cope with routine tasks. If you do feel tired, take rest and get help with household chores. If necessary, take time off work. Gentle exercise such as walking can be beneficial.

Skin and nails

This treatment can sometimes cause a rash, skin sensitivity and/or blistering to the hands and feet. Report this to your doctor if this happens. Sometimes as a result of the chemotherapy, your skin may appear darker in colour or lightly tanned, especially around the joints. This is known as hyperpigmentation. Asian and African-Caribbean people may develop noticeable light patches on their skin. Your nails may also discolour or become brittle during treatment. The skin and nails will usually return to normal when treatment is finished.

• Tingling and numbness

This is typically in the fingers and/or toes. Muscle weakness may also occur. This is common but is usually only mild and temporary. It can sometimes last for some time or become permanent. Sometimes you may also have difficulty controlling the muscles in your arms and legs, or you balance. Please report these symptoms to your doctor on your next hospital visit.

• Eye irritation

We will give you eye drops to try to prevent this. Please let your doctor or nurse know if your eyes feel uncomfortable.

Urine discolouration

Doxorubicin because of its red colour, may discolour your urine red or pink for up to 24 hours following treatment. This is perfectly normal and nothing to worry about.

• Bone pain

We will give you injections to boost your white blood cell count (G-CSF). These can sometimes cause pain, particularly in the lower back or pelvis. This usually disappears after the injections have been stopped. Paracetamol may help, but if the pain is severe then you may need to reduce or stop the injections. Please discuss this with your nurse specialist or your doctor.

Uncommon side effects (less than 1 in 10)

• Blood sugar

Prednisolone may cause your blood sugar to rise. If you are diabetic then you may need to increase the dose of any diabetic medication you take (insulin or tablets). You should discuss this with your doctor before starting the chemotherapy. Prednisolone can also induce diabetes in people not known to have this disease. This is normally reversible although you may need to take some diabetic medication while you are having chemotherapy.

• Sore mouth and altered taste

Your mouth may become sore or dry, or you may notice small mouth ulcers during this treatment. Drinking plenty of fluids and cleaning your teeth regularly and gently with a soft toothbrush can help to reduce the risk of this happening. We can prescribe a mouthwash for you to use during treatment. You can dilute this with water if your mouth is sore. Ask your doctor or nurse for further advice. There is also general mouth care information in the chemotherapy booklet. If you continue to have a sore mouth, please contact The Christie Hotline. You may also develop a strange taste sometimes described as metallic or bitter. A strongly flavoured sweet or mint will help to disguise this. Normal taste will usually return after the treatment finishes.

Bone changes

Prednisolone can occasionally cause weakness in your bones (in about 1 in 100 patients). Symptoms can develop several months or even years after chemotherapy. If you have persistent hip pain (not related to G-CSF) injections) then let your doctor know.

Heart function

Doxorubicin can occasionally affect your heart function. This is more common in people with other heart problems, diabetes or high blood pressure, but is much less common in other patients. Sometimes symptoms can develop months or years after treatment has finished. Your doctor can discuss whether any heart scans or monitoring is needed.

• Bladder irritation

Cyclophosphamide can occasionally cause irritation of the bladder. Let your doctor know if it becomes painful to pass urine or you pass blood clots. This usually happens within the first few days of each treatment only.

Rare side effects (less than 1 in 100)

Extravasation

Extravasation is when chemotherapy leaks outside the vein. If you develop redness, soreness or pain at the injection site **at any time**, please let us know straight away.

Serious and potentially life threatening side effects

In a small proportion of patients chemotherapy can result in very serious side effect which may rarely result in death. The team caring for you will discuss the risk of these side effects with you.

Sex, contraception and fertility

Protecting your partner and contraception: We recommend that you or your partner use a condom during sexual intercourse while you are having the course of chemotherapy. Chemotherapy is dangerous to unborn babies and this will also protect you and your partner from any chemotherapy drugs that may be present in semen and in the vagina. If you suspect that you may be pregnant please tell your doctor immediately.

Fertility: This chemotherapy may affect your ability to have children. Your doctor or nurse should have discussed this with you. If not, please ask them before you start treatment.

Late side effects

Some side effects may become evident only after a number of years. In reaching any decision with you about treatment, the potential benefit you receive from treatment will be weighed against the risks of serious long term side effects to the heart, lungs, kidneys and bone marrow. With some drugs there is also a small but definite risk of developing another cancer. If any of these problems specifically applies to you, the doctor will discuss these with you and note this on your consent form.

Contacts

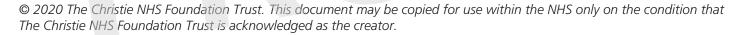
If you have any general questions or concerns about your treatment, please ring the area where you are having treatment:

Lymphoma clinical nurse specialists - 0161 446 8573

Lymphoma secretaries - 0161 446 3753 or 0161 918 7246

HTDU (department 26) - **0161 446 3924** Young Oncology Day Unit - **0161 446 3953**

Your consultant is:
Your hospital number is:
Your key worker is:



If you need information in a different format, such as easy read, large print, BSL, braille, email, SMS text or other communication support, please tell your ward or clinic nurse.

The Christie is committed to producing high quality, evidence based information for patients. Our patient information adheres to the principles and quality statements of the Information Standard. If you would like to have details about the sources used please contact **the-christie.patient.information@nhs.net**

For information and advice visit the cancer information centres at Withington, Oldham or Salford. Opening times can vary, please check before making a special journey.



Contact The Christie Hotline for urgent support and specialist advice

The Christie Hotline: 0161 446 3658

Open 24 hours a day, 7 days a week

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The Christie Patient Information Service Tel: 0161 446 3000 www.christie.nhs.uk