Patient agreement to systemic anti-cancer therapy (SACT):

Ponatinib

Patient details

Patient's surname/family name: _____

_____

Patient's first name(s): _____

_____

Date of birth: _____

NHS number: _____
(or other identifier)

Special requirements:
(e.g. other language/other communication method)

_____

_____

_____

_____

_____

_____

Hospital/NHS Trust/NHS Board: _____

_____

_____

Responsible consultant:

Name: _____

Job title: _____

Name of proposed course of treatment (include brief explanation if medical term not clear)

☐ Ponatinib for the treatment of chronic myeloid leukaemia (CML).

☐ Tablets are taken orally once a day. Treatment is continued until disease progression or unacceptable toxicity.

Where the treatment will be given

☐ Outpatient ☐ Day unit/case ☐ Inpatient ☐ Other: _____

Statement of health professional

(to be filled in by health professional with appropriate knowledge of proposed procedure, as specified in the hospital/Trust/NHS board's consent policy)

☑ Tick all relevant boxes

☐ I confirm the patient has capacity to give consent.

I have explained the course of treatment and intended benefit to the patient.

The intended benefits (there are no guarantees about outcome)

☐ To obtain or maintain remission of your leukaemia, in order to improve both quality and quantity of life.

Statement of health professional

(continued)

Patient identifier/label

Significant, unavoidable or frequently occurring risks

Common side effects:

More than 10 in every 100 (>10%) people have one or more of the side effects listed:

- ☐ Anaemia (low red blood cells), bleeding or bruising (due to low platelets), low white blood cell counts.
- ☐ Diarrhoea, feeling sick (nausea) and being sick (vomiting), constipation, abdominal (tummy) pain, lack of appetite.
- ☐ Skin changes (rash, dryness, itching).
- ☐ Tiredness and feeling weak (fatigue), pain in the muscles/joints/bones/back, muscle spasms.
- ☐ Fluid build-up in ankles and legs, shortness of breath, cough.
- ☐ High blood pressure, changes in liver function tests.
- ☐ Headache, dizziness, difficulty sleeping.

Occasional side effects:

Between 1 and 10 in every 100 (1-10%) people have one or more of these effects:

- ☐ If you have a low white blood cell count, you have an increased risk of getting an infection.
- ☐ **If you have a severe infection this can be life-threatening. Contact your doctor or hospital straight away if:**
  - **your temperature goes over 37.5°C or over 38°C, depending on the advice given by your chemotherapy team**
  - **you suddenly feel unwell (even with a normal temperature)**
- ☐ Thinning of the hair or hair loss, skin changes (thickening, redness, colour change), excessive sweating, night sweats, flushing.
- ☐ Eye problems (dryness, watery, swelling, change in vision).
- ☐ Sore mouth and ulcers, dry mouth, indigestion, weight loss, inflammation of the pancreas (pancreatitis) causing severe pain in the centre of your tummy.
- ☐ Numbness/tingling in the hands/feet and/or loss of sensation.
- ☐ Abnormal electrolyte levels (low calcium, potassium, sodium and phosphate), dehydration, underactive thyroid.
- ☐ Ponatinib can increase the risk of developing a blood clot (thrombosis), causing pain, redness, swelling in a leg or breathlessness and chest pain – you must tell your doctor straight away if you have any of these symptoms.
- ☐ Ponatinib can raise your blood sugar. If you have diabetes, it may lead to higher blood sugar levels.

Occasional side effects continued:

- ☐ Ponatinib can increase your risk of vascular occlusive events. You may experience pain in the leg muscles when walking, and less commonly stroke, heart attack or short term loss of blood supply to the brain.

Other risks:

- ☐ A very rare side-effect is changes in kidney function tests.
- ☐ A very rare risk of tumour lysis syndrome. This occurs when the treatment destroys cancer cells too quickly for the kidneys to cope leading to changes in blood tests. Rarely, dialysis may be needed. High risk patients are prescribed medicines for prevention.
- ☐ Allopurinol may be prescribed to prevent gout if you are at risk, and taking Ponatinib soon after diagnosis.
- ☐ Before treatment you might have blood tests to check for viruses such as Hepatitis B, Hepatitis C, HIV or more unusual infections. Treatment for cancer could weaken your natural defence (immune) system, so infections like this could worsen or become active again if you've had them in the past. You may have medicines to prevent or treat infection. At present there is no evidence that this occurs with Ponatinib.
- ☐ Some anti-cancer medicines can damage ovaries and sperm. At present there is no evidence that this occurs with Ponatinib.
- ☐ Ponatinib may damage the development of a baby in the womb. It is important not to become pregnant while you are having treatment. It is important to use effective contraception during treatment. You can talk to your doctor or nurse.
- ☐ If you are due to have surgery for reasons other than CML, it is important to consult your doctor about the suitability of taking Ponatinib, before, during and after your procedure.
- ☐ You may notice changes in your memory, concentration, or your ability to think clearly. There can be many causes of this including your treatment, diagnosis, or both.
- ☐ The side-effects of Ponatinib are usually mild or moderate. Complications of treatment can very occasionally be life threatening and may result in death. The risks are different for every individual. Potentially life threatening complications include those listed on this form, but, other, exceedingly rare side-effects may also be life threatening.

Statement of health professional

(continued)

Patient identifier/label

Any other risks and information

_____

_____

_____

☐ I have discussed the intended benefit and risks of the recommended treatment, and of any available alternative treatments (including no treatment).

☐ I have discussed the side effects of the recommended treatment, which could affect the patient straight away or in the future, and that there may be some side effects not listed because they are rare or have not yet been reported. Each patient may experience side effects differently.

☐ I have discussed what the treatment is likely to involve (including inpatient / outpatient treatment, timing of the treatment, blood and any additional tests, follow-up appointments etc) and location.

☐ I have explained to the patient, that they have the right to stop this treatment at any time and should contact the responsible consultant or team if they wish to do so.

☐ I have discussed concerns of particular importance to the patient in regard to treatment

(please write details here): _____

_____

_____

Clinical management guideline/Protocol compliant (please tick):

☐ Yes ☐ No ☐ Not available

If No please document reason here: _____

_____

The following written information has been provided:

☐ Information leaflet for Ponatinib

☐ 24 hour alert card or SACT advice service contact details

☐ SACT treatment record (cruk.org/treatment-record)

☐ Other, please state: _____

_____

Health professional details:

Signed: _____

Date: _____

Name (PRINT): _____

_____

Job title: _____

_____

Statement of interpreter (where appropriate)

Interpreter booking reference (if applicable): _____

I have interpreted the information above to the patient to the best of my ability and in a way in which I believe they can understand.

Signed: _____ Date: _____

Name (PRINT): _____

Job title: _____

Statement of patient

Patient identifier/label

Please read this form carefully. If your treatment has been planned in advance, you should already have your own copy of the form which describes the benefits and risks of the proposed treatment. If not, you will be offered a copy now. If you have any further questions, do ask – we are here to help you. You have the right to change your mind at any time, including after you have signed this form.

☐ I have **had enough time to consider** my options and make a decision about treatment.

☐ **I agree** to the course of treatment described on this form.

A witness should sign below if the patient is unable to sign but has indicated their consent. Young people/children may also like a parent to sign here (see notes).

Patient's signature: _____ Date: _____

Name (PRINT): _____

Parent's/Witness' signature: _____ Date: _____

Name (PRINT): _____

Copy accepted by patient: yes / no (please circle)

Confirmation of consent

(health professional to complete when the patient attends for treatment, if the patient has signed the form in advance)

On behalf of the team treating the patient, I have confirmed that the patient has no further questions and wishes the course of treatment/procedures to go ahead.

Signed: _____

Date: _____

Name (PRINT): _____

Job title: _____

Important notes: (tick if applicable)

☐ See also advance decision to refuse treatment

☐ Patient has withdrawn consent (ask patient to sign /date here)

Signed:_____

Date: _____

Further information for patients

Contact details (if patient wishes to discuss options later):

_____

Contact your hospital team if you have any questions about cancer and its treatment.

Cancer Research UK can also help answer your questions about cancer and treatment. If you want to talk in confidence, call our information nurses on freephone **0808 800 4040**, Monday to Friday, 9am to 5pm. Alternatively visit **www.cruk.org** for more information.

These forms have been produced by Guy's and St. Thomas' NHS Foundation Trust as part of a national project to support clinicians in ensuring all patients are fully informed when consenting to SACT. The project is supported by Cancer Research UK. This does not mean you are taking part in a clinical trial.

CANCER RESEARCH UK

Guidance for health professionals (to be read in conjunction with the hospital's consent policy)

Patient identifier/label

What a consent form is for

This form documents the patient's agreement to go ahead with the treatment you have proposed. It is not a legal waiver – if patients, for example, do not receive enough information on which to base their decision, then the consent may not be valid, even though the form has been signed. Patients are also entitled to change their mind after signing the form, if they retain capacity to do so. The form should act as an aide-memoir to health professionals and patients, by providing a check-list of the kind of information patients should be offered, and by enabling the patient to have a written record of the main points discussed. In no way should the written information provided for the patient be regarded as a substitute for face-to-face discussions with the patient.

The law on consent

See the following publications for a comprehensive summary of the law on consent. Consent: Patients and doctors making decisions together, GMC 2020 (available at **www.gmc-uk.org/guidance**), and Reference guide to consent for examination or treatment, Department of Health, 2nd edition 2009 (available at **www.doh.gov.uk**).

Who can give consent

Everyone aged 16 or over is presumed to have the capacity to give consent for themselves, unless the opposite is demonstrated. If a child under the age of 16 has "sufficient understanding and intelligence to enable him or her to understand fully what is proposed", then the child will have capacity to give consent for himself or herself.
Young people aged 16 and 17, and younger children with capacity, may therefore sign this form for themselves, but may like a parent to countersign as well. If the child is not able to give consent, someone with parental responsibility may do so on their behalf and a separate form is available for this purpose. Even where children are able to give consent for themselves, you should always involve those with parental responsibility in the child's care, unless the child specifically asks you not to do so. If a patient has the capacity to give consent but is physically unable to sign a form, you should complete this form as usual, and ask an independent witness to confirm that the patient has given consent orally or non-verbally.

When NOT to use this form

If the patient is 18 or over and lacks the capacity to give consent, you should use an alternative form (form for adults who lack the capacity to consent to investigation or treatment). A patient lacks capacity if they have an impairment or disturbance of the brain, affecting the way their mind works. For example, if they cannot do one of the following:

- understand information about the decision to be made
- retain that information in their mind
- use or weigh this information as a part of their decision making process, or
- communicate their decision (by talking, using sign language or any other means)

You should always take all reasonable steps (for example involving more specialist colleagues) to support a patient in making their own decision, before concluding that they are unable to do so.

Relatives cannot be asked to sign a form on behalf of an adult who lacks capacity to consent for themselves, unless they have been given the authority to do so under a Lasting Power of Attorney or as a court deputy.

Information

Information about what the treatment will involve, its benefits and risks (including side-effects and complications) and the alternatives to the particular procedure proposed, is crucial for patients when making up their minds. The courts have stated that patients should be told about 'significant risks which would affect the judgement of a reasonable patient'. 'Significant' has not been legally defined, but the GMC requires doctors to tell patients about 'significant, unavoidable or frequently occurring' risks. In addition if patients make clear they have particular concerns about certain kinds of risk, you should make sure they are informed about these risks, even if they are very small or rare. You should always answer questions honestly. Sometimes, patients may make it clear that they do not want to have any information about the options, but want you to decide on their behalf. In such circumstances, you should do your best to ensure that the patient receives at least very basic information about what is proposed. Where information is refused, you should document this on the consent form or in the patient's notes.

NHS Scotland

NHS Scotland staff should refer to Healthcare Improvement Scotland. Guidance on consent for SACT and local NHS Board guidance on consent aligned to the Scottish legal framework.

References

1. Summary of Product Characteristics (SmPCs) for individual drugs: https://www.medicines.org.uk/emc
2. Cancer Research UK: https://www.cancerresearchuk.org/about-cancer/cancer-in-general/treatment/cancer-drugs
3. Macmillan Cancer Support: https://www.macmillan.org.uk/information-and-support/treating/chemotherapy/drugs-and-combination-regimens
4. Guy's and St. Thomas' NHS Foundation Trust, Chemotherapy consent form