

Chemotherapy Protocol

SKIN CANCER

DACARBAZINE

Regimen

• Skin – Dacarbazine

Indication

- Dacarbazine is recommended as a first line option for unresectable stage IIIC or IV melanoma in those unsuitable for ipilimumab.
- Dacarbazine is recommended as an option for the second line treatment of unresectable stage IIIC or IV melanoma of cutaneous, ocular or mucosal origin following treatment with ipilimumab.
- Dacarbazine is recommended as an option for the third line treatment of unresectable stage IIIC or IV melanoma of cutaneous, ocular or mucosal origin.
- WHO performance status 0, 1

<u>Toxicity</u>

Drug	Adverse Effect
Dacarbazine	Influenza type syndrome starting up to 7 days after treatment and lasting up to 21 days, photosensitivity, anaphylaxis, vein pain, myelosuppression

The adverse effects listed are not exhaustive. Please refer to the relevant Summary of Product Characteristics for full details.

Monitoring

Regimen

• FBC, LFT's and U&E's prior to day one of each cycle

Dose Modifications

The dose modifications listed are for haematological, liver and renal function only. Dose adjustments may be necessary for other toxicities as well.

In principle all dose reductions due to adverse drug reactions should not be reescalated in subsequent cycles without consultant approval. It is also a general rule for chemotherapy that if a third dose reduction is necessary treatment should be stopped.



Please discuss all dose reductions / delays with the relevant consultant before prescribing if appropriate. The approach may be different depending on the clinical circumstances. The following is a general guide only.

Haematological

Prior to prescribing the following criteria must be met.

Criteria	Eligible Level
Neutrophil	equal to or more than 1.5x10 ⁹ /L
Platelets	equal to or more than 100x10 ⁹ /L

If the platelets are less than 100×10^9 /L and the neutrophils are less than 1.5×10^9 /L then delay treatment for seven days. Repeat the full blood count at this time. If the counts have returned to the eligible levels treatment may continue at 100% of the last dose.

Consider blood transfusion or erythropoietin if patient symptomatic of anaemia or has a haemoglobin of less than 8g/dL

Hepatic Impairment

Dacarbazine is activated and metabolised in the liver. It can be hepatotoxic. Consider a dose reduction or stopping therapy if the baseline liver function tests double during therapy

Renal Impairment

Drug	Creatinine Clearance (ml/min)	Dose (% of original dose)
	45 - 60	80%
Dacarbazine	30 - 45	75%
	less than 30	70%

Other

Dose reductions or interruptions in therapy are not necessary for those toxicities that are considered unlikely to be serious or life threatening. For example, alopecia, altered taste or nail changes.

Regimen

21 day cycle for 6 cycles

Drug	Dose	Days	Route
Dacarbazine	1000mg/m ²	1	Intravenous infusion in 500ml sodium chloride
	_		0.9% over 60 minutes

Dose Information

• Dacarbazine will be dose banded as per the CSCCN agreed bands



Administration Information

Extravasation

• Dacarbazine - vesicant

Other

• Dacarbazine is light sensitive. The infusion should be protected from light during administration. Do not use if the solution has a pink or red discolouration.

Additional Therapy

• Antiemetics

15-30 minutes before chemotherapy

- dexamethasone 8mg oral or intravenous
- ondansetron 8mg oral or intravenous

As take home medication

- dexamethasone 4mg twice a day for three days starting the day after chemotherapy
- metoclopramide 10mg three times a day for three days and then 10mg three times a day when required oral

Gastric protection with a proton pump inhibitor or a H_2 antagonist may be considered in patients considered at high risk of GI ulceration or bleed

Coding

- Procurement X70.1
- Delivery X72.3

References

1. Huncharek M, Caubert JF, McGarry R. Single agent DTIC versus combination chemotherapy with or without immunotherapy in metastatic melanoma: a meta-analysis of 3273 patients from 20 randomised trials. Melanoma Research 2001; 11 (1): 75-81.



REGIMEN SUMMARY

Dacarbazine

Day One

- 1. Dexamethasone 8mg oral or intravenous
- 2. Ondansetron 8mg oral or intravenous

3. Dacarbazine 1000mg/m 2 intravenous infusion in 500ml sodium chloride 0.9% over 60 minutes

Administration Instructions

Dacarbazine is light sensitive. The infusion should be protected from light during administration. Do not use if the solution has a pink or red discolouration

Take Home Medicines

4. Dexamethasone 4mg twice a day for 3 days starting the day after chemotherapy oral

5. Metoclopramide 10mg three times a day for three days starting on the day of chemotherapy and then 10mg three times a day when required oral



DOCUMENT CONTROL

Version	Date	Amendment	Written By	Approved By
1	February 2015	None	Dr Deborah Wright Pharmacist	Prof C Ottensmeier Consultant Medical Oncologist
				Dr M Wheater Consultant Medical Oncologist

This chemotherapy protocol has been developed as part of the chemotherapy electronic prescribing project. This was and remains a collaborative project that originated from the former CSCCN. These documents have been approved on behalf of the following Trusts;

Hampshire Hospitals NHS Foundation Trust NHS Isle of Wight Portsmouth Hospitals NHS Trust Salisbury Hospital NHS Foundation Trust University Hospital Southampton NHS Foundation Trust Western Sussex Hospitals NHS Foundation Trust

All actions have been taken to ensure these protocols are correct. However, no responsibility can be taken for errors which occur as a result of following these guidelines.