

NEW ZEALAND DATA SHEET

1. PRODUCT NAME

DBL™ Dacarbazine 200 mg Powder for Injection.

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

DBL Dacarbazine for Injection is a sterile parenteral dosage form for reconstitution. When reconstituted as directed each mL of the solution in the 200 milligram vial contains dacarbazine 10 milligrams, citric acid monohydrate 10 milligrams and mannitol 3.75 milligrams.

For the full list of excipients, see section 6.1.

3. PHARMACEUTICAL FORM

Powder for injection.

White or very pale yellow powder or plug.

The pH of the reconstituted solution is 3.0 – 4.0.

4. CLINICAL PARTICULARS

4.1 Therapeutic indications

Chemotherapy of metastatic malignant melanoma and various sarcomas. In other cancers, the available evidence shows dacarbazine to be ineffective or less effective than established regimens.

Note: The use of dacarbazine is restricted to hospitals with an oncology service.

4.2 Dose and method of administration

Adult

There are two commonly used dose regimens:

1. 4.5 milligrams/kg/day for 10 days; the 10 day course may be repeated every 4 weeks.

Note: 2 milligrams/kg/day for 10 days has been used by one investigator and found to be equally as effective as the higher dose.

2. 250 milligrams/m²/day for 5 days; the 5 day course may be repeated every 3 weeks.

In general, effectiveness is likely to be evident after the second course of dacarbazine. Of the 1427 patients with metastatic malignant melanoma treated with dacarbazine, 81 patients (5.7%) had complete remissions and 208 patients (14.6%) had partial remissions with a total response rate of 20.3%. The duration of remissions (partial and complete combined) varied from 5 to

100 weeks. The median remission duration obtained by three principle investigators was about 6 months. Once a patient has relapsed it is unlikely that subsequent courses of dacarbazine will be effective.

Combination therapy

Combinations of cancer chemotherapeutic agents have often shown an improved response over the use of single agents. This has not been the case in metastatic malignant melanoma except at a very high and toxic dosage of the combinations in small numbers of patients. However, in treatment of various soft tissue sarcomata combinations with doxorubicin and/or vincristine have increased the remission rates. The user should be familiar with the current cancer chemotherapeutic literature.

Geriatric

As for paediatric use.

With impaired hepatic function

As the drug partly undergoes metabolism in the liver impairment of liver function is likely to necessitate a variation in dosage (See section 5.2).

With impaired renal function

As the drug is excreted 50% unchanged in the urine by tubular secretion, impairment of renal function is likely to necessitate a variation in dosage (See section 5.2).

Paediatric population

No special information submitted to indicate whether or not children require a different dosage range or whether they metabolise the drug differently or react differently to the drug.

Method of Administration

Administration is by the intravenous route only.

Reconstitute vial contents by adding 19.7 mL of Water for Injections to the 200 milligrams vial.

The resulting solution is hypotonic and will contain 10 milligrams/mL of dacarbazine with a pH of 3 to 4.

Intravenous injection may be given over about one minute. Extravasation of the drug into surrounding tissue during intravenous administration may result in tissue damage and severe pain.

4.3 Contraindications

- Patients who are pregnant or are breast feeding.
- Patients with known hypersensitivity to dacarbazine or any of the excipients.
- Patients who have previously had severe myelosuppression.

4.4 Special warnings and precautions for use

Toxicity: In the treatment of each patient, the physician must weigh carefully the possibility of achieving therapeutic benefit against the risk of toxicity (See below and section 4.8).

The drug can produce severe and possibly fatal, haematologic or hepatic toxicity. Administer only to patients in hospitals with an oncology service so that they can be observed carefully and frequently during and after therapy particularly for haematopoietic toxicity.

Haematopoietic depression is the most serious form of toxicity and involves primarily the leucocytes and megakaryocytes causing depression of platelets, but also other blood forming elements. Leucopenia and thrombocytopenia may be severe enough to cause death.

Long-term therapy can cause cumulative bone marrow toxicity. Possible bone marrow depression requires careful monitoring of red blood cells, white blood cells and platelet level. Haematotoxicity may warrant temporary suspension or termination of therapy. The bone marrow depressant effects of dacarbazine may result in an increased incidence of microbial infection, delayed healing and gingival bleeding. Dental work, whenever possible, should be completed prior to initiation of dacarbazine therapy, or deferred until blood counts have returned to normal. Patients should be instructed on proper oral hygiene during treatment, including caution in the use of toothbrushes, dental floss and toothpicks.

Hepatic toxicity accompanied by hepatic vein thrombosis and hepatocellular necrosis (Budd-Chiari Syndrome) resulting in death has been reported. The incidence of such reactions has been low, approximately 0.01% of patients treated. This toxicity has been observed when dacarbazine has been administered concomitantly with other antineoplastic drugs; however, it has also been reported in some patients treated with dacarbazine alone. Therefore, frequent monitoring of liver size, function and blood counts (especially eosinophils) is required (see section 4.8).

Restriction of food intake for 4 to 6 hours prior to treatment may reduce the severity of nausea and vomiting which occurs in most patients particularly during the first 2 days of treatment. In rare cases, intractable nausea and vomiting have necessitated discontinuation of therapy. Administration of an antiemetic may also reduce the severity of these effects.

Impairment of liver and renal function: See section 4.2.

Care should be taken to avoid contact with the skin and eyes when reconstituting or administering dacarbazine.

Immunisation with live virus vaccines should only be undertaken with extreme caution (See section 4.5). Administration of live or live-attenuated vaccines in patients immunocompromised by chemotherapeutic agents including dacarbazine, may result in serious or fatal infections. Vaccination with a live vaccine should be avoided in patients receiving dacarbazine. Immunisation with oral poliovirus vaccines should be postponed in people in close contact with the patient, especially family members. Killed or inactivated vaccines may be administered; however, the response to such vaccines may be diminished (see section 4.5).

4.5 Interaction with other medicines and other forms of interaction

Dacarbazine is metabolised by cytochrome P450 (CYP1A1, CYP1A2, and CYP2E1). This has to be taken into account if other drugs are co-administered which are metabolised by the same hepatic enzymes. Microsomal liver enzyme inducers eg barbiturates, rifampicin, phenytoin may theoretically hasten the activation of dacarbazine to aminoimidazole-carboxamide (AIC).

The concomitant use of dacarbazine with phenytoin is not recommended due to the risk of exacerbation of convulsions resulting from the decrease of phenytoin digestive absorption by cytotoxic drug or risk of toxicity enhancement or loss of efficacy of the cytotoxic drug due to increased hepatic metabolism by phenytoin. It may be advisable to administer an anticonvulsant benzodiazepine.

Mercaptopurine, azathioprine, allopurinol: Dacarbazine inhibits xanthine oxidase and may theoretically potentiate the activity of these medicines.

The incidence or severity of side effects may be altered when dacarbazine is used in combination with other antineoplastic agents. The leucopenic and/or thrombocytopenic effects of dacarbazine may be increased with concurrent or recent therapy with other medications which cause these effects. Additive bone marrow depression may occur if dacarbazine is administered with other bone-marrow depressants or with radiation therapy. Dosage adjustment of dacarbazine may be necessary.

Fotemustine and dacarbazine should not be used concomitantly. Sequential administration of dacarbazine (400 – 1000 milligrams/m²) and fotemustine (100 milligrams/m²) has been associated with acute lung toxicity, in the form of adult respiratory distress syndrome. Dacarbazine should be administered over one week after fotemustine administration.

It has been reported that dacarbazine reduced the response to levodopa in a patient with Parkinson's disease. The mechanism of this interaction is unclear, but since the plasma levels of levodopa were unchanged, it is unlikely to be due to pharmacokinetic changes.

Administration of dacarbazine may potentiate the replication of live virus vaccines, increase the adverse effects of the vaccine, or decrease the antibody response to the vaccine. Dacarbazine may also suppress the antibody response to killed virus vaccines. Viral vaccines should not be administered for 3 to 12 months after discontinuing immunosuppressive drug treatment (see section 4.4).

Ciclosporin (and by extrapolation tacrolimus) given with dacarbazine may cause excessive immunosuppression with risk of lymphoproliferation.

It has been noted that administration of dacarbazine and interleukin-2 may result in alterations to the pharmacokinetics of dacarbazine. An increased clearance (of approximately 38%) and volume of distribution (of approximately 36%) have been reported when doses of 2 to 4×10⁶ U/m² interleukin-2 were used. This alteration appears to correlate with the dose of interleukin-2 and should be taken into account, particularly when high doses of interleukin-2 are combined with dacarbazine.

4.6 Fertility, pregnancy and lactation

Fertility

No data available.

Pregnancy

Category D: *This category specifies drugs which have caused, are suspected to have caused or may be expected to cause, an increased incidence of human foetal malformations or irreversible damage. These drugs may also have adverse pharmacological effects.*

The drug is teratogenic and carcinogenic when used in animals. When administered intraperitoneally to rats at doses of 50 or 70 milligrams/kg/day (approximately 11 times the human dose), teratogenic effects have been observed, including anomalies of the skeletal system, eyes, cardiovascular system and abdominal wall. Teratogenic effects have also been observed in rabbits administered 10 milligrams/kg intraperitoneally. No adequate and well controlled studies have been performed in pregnant women.

Dacarbazine for injection is therefore contraindicated in patients who are pregnant.

Advice on contraception in males and females

Based on reproductive toxicity and genetic toxicity findings, women of childbearing potential should be advised to use effective contraception during treatment with dacarbazine and for at least 6 months after the last dose.

Based on genetic toxicity findings, male patients should be advised to use effective contraception during treatment with dacarbazine and for at least 3 months after the last dose.

Lactation

It is not known whether dacarbazine is distributed into breast milk. Due to the potential risk to the infant, dacarbazine is contraindicated in breast-feeding mothers.

4.7 Effects on ability to drive and use machinery

No data available. Caution should be taken when driving or using machinery while on this medication as it may cause some undesirable effects (eg, nausea, vomiting or neurological side effects).

4.8 Undesirable effects

More common reactions

Gastrointestinal: 90% of patients experience nausea and vomiting in the first two days of treatment. Diarrhoea may also occur. A degree of tolerance may develop to these effects after about 2 days of treatment Vomiting lasts 1 to 12 hours Prophylactic antiemetic therapy with a 5HT₃ blocker or dexamethasone is usually required. Rarely, intractable nausea and vomiting have necessitated discontinuance of dacarbazine therapy.

Haematological: bone marrow depression (25%) (See section 4.8 - Life threatening reactions).

Leucocytopenia was usually seen 14 days after commencement of therapy but was noted as early as day 10 and, in 10% of patients, as late as day 30 (ie. 25 days after completion of therapy). The average length of duration was 1 week and the longest, 3 weeks.

Thrombocytopenia was most frequently seen 18 days after commencement of therapy but in 43% of patients it was noted by day 12 and in 10% not until after day 30. The average length of duration was 1 week and the longest 3 weeks.

Eosinophilia has been reported in one patient receiving dacarbazine.

Pancytopenia and anaemia have been reported in patients receiving dacarbazine, frequency unknown.

Less common reactions

Cardiovascular: facial flushing, ECG abnormalities, orthostatic hypotension. Hypotension appears to be associated with high doses ($> 850 \text{ mg/m}^2$) of dacarbazine and may be dose-limiting.

Dermatological: (1%, usually transient) rash, alopecia. Photosensitivity reactions have occurred rarely.

General: (3%) flu-like syndrome with fever to 39°C , severe myalgias and malaise. This syndrome usually occurs after large single doses approximately 7 days after treatment with dacarbazine and lasts 7 to 21 days. It may recur with successive treatments.

Hepatic: (5%, usually transient). Increases in transaminases (AST and ALT), alkaline phosphatase, LDH. Levels usually return to normal within 2 weeks. Hepatic toxicity accompanied by hepatic vein thrombosis and hepatocellular necrosis (Budd-Chiari Syndrome) resulting in death has been reported (See section 4.4) (0.01%). A case of acute hepatitis has been reported during the first course of dacarbazine. Granulomatous hepatitis has also occurred.

Nervous System: (3% usually transient) blurred vision, seizures, headache, paraesthesia, confusion, malaise, lethargy.

Local reactions: Injection of concentrated dacarbazine solutions may cause severe pain along the vein. Extravasation of the drug into surrounding tissue may cause severe pain, tissue damage and cellulitis.

Hypersensitivity: Anaphylaxis has occurred occasionally.

Dental effects: Dacarbazine may adversely affect dental procedures (See section 4.4).

Other: Dacarbazine may rarely cause stomatitis.

Life threatening reactions

Bone marrow depression: Death due to agranulocytosis or thrombocytopenia occurred in about 0.4% of patients in clinical trials.

Fatal hepatotoxicity: Budd-Chiari Syndrome (See section 4.4).

Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicine is important. It allows continued monitoring of the benefit/risk balance of the medicine. Healthcare professionals are asked to report any suspected adverse reactions <https://nzphvc.otago.ac.nz/reporting/>.

4.9 Overdose

Symptoms: Severe bone marrow depression and gastrointestinal effects such as nausea, vomiting and diarrhoea may be expected.

Treatment: There is no specific antidote to dacarbazine poisoning. Cease dacarbazine administration and institute supportive measures, e.g. appropriate transfusions, for bone marrow depression (See section 4.8 - Haematological).

For advice on the management of overdose please contact the National Poisons Centre on 0800 POISON (0800 764766).

5. PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

Mechanism of action

Relationship to other drugs: Dacarbazine is a structural analogue of 5-amino-imidazole-4-carboxamide which is an intermediate in purine biosynthesis.

Site and mode of action: This drug inhibits cell replication by an unknown mechanism. However, three possible mechanisms have been postulated:

1. Since dacarbazine is an analogue of 5-amino-imidazole-4-carboxamide, an intermediate in the de novo biosynthesis of purine, it might interfere with purine biosynthesis and hence DNA bio-synthesis. This appears to be true at high concentrations of the drug, but low concentrations appear to enhance DNA, RNA and protein biosynthesis.
2. One metabolite of dacarbazine, diazomethane, is an alkylating agent and may act in the same way as the nitrogen mustards.
3. The drug might act as a sulphhydryl reagent since the inhibition of bacterial growth by dacarbazine can be prevented by glutathione.

5.2 Pharmacokinetic properties

Absorption

Dacarbazine is poorly and erratically absorbed from the gastrointestinal tract, which could result in unpredictable tumour responses and possible increased toxicity. Therefore the drug is recommended for intravenous administration only. Peak plasma concentrations of about 8

micrograms per mL are reached immediately following administration of dacarbazine 4.5 milligrams/kg by intravenous push.

Distribution

The volume of distribution of dacarbazine exceeds total body water content, suggesting localisation in some body tissue, probably the liver. The drug is only slightly bound to plasma proteins. Dacarbazine crosses the blood-brain barrier to a limited extent; CSF concentrations are reported to be about 14% of plasma concentrations. It is not known if dacarbazine crosses the human placenta or distributes into milk.

Biotransformation/ Elimination

Dacarbazine is N-demethylated by liver microsomal enzymes to yield CO₂ which is excreted in expired air and aminoimidazole-carboxamide (AIC) which is excreted in the urine. About half the drug remains unchanged and is rapidly excreted by tubular secretion.

As dacarbazine is approximately 50% metabolised by the liver and the remaining unchanged drug and metabolites are excreted in the urine, impairment of hepatic or renal function may require a reduction in dosage to avoid toxicity.

Plasma concentrations of dacarbazine appear to decline in a biphasic manner. The initial phase half life ($t_{1/2\alpha}$) is very short, with one study reporting $t_{1/2\alpha}$ as 2.9 minutes. The terminal phase half life ($t_{1/2\beta}$) is consistently longer 41.4 to 75 minutes. In one patient with renal and hepatic dysfunction, the $t_{1/2\alpha}$ was 55 minutes and the $t_{1/2\beta}$ was 7.2 hours.

5.3 Preclinical safety data

Genotoxicity

Dacarbazine has been shown to be mutagenic in bacteria and produced evidence of both clastogenicity and mutagenicity in mammalian cells *in vitro*. In addition, dacarbazine has shown clastogenicity to mouse bone marrow *in vivo*. These results are consistent with the pharmacological activity of dacarbazine, which is a methylating cytostatic drug.

Carcinogenicity

The drug is carcinogenic when used in animals. See section 4.6 - Pregnancy.

Reproductive and developmental toxicity

See section 4.6 - Pregnancy.

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

- Citric acid monohydrate
- Mannitol

6.2 Incompatibilities

No data available.

6.3 Shelf life

24 months from date of manufacture stored at 2° to 8°C (Refrigerate, do not freeze). Protect from light.

8 hours reconstituted stored at or below 25°C. Protect from light.

24 hours reconstituted stored at 2° to 8°C (Refrigerate, do not freeze). Protect from light.

6.4 Special precautions for storage

Store between 2°C to 8°C. Refrigerate. Do not freeze. Protect from light.

Reconstituted vials are chemically stable for up to 8 hours if stored at 25°C and up to 24 hours if stored at 2°C to 8°C, protected from light. However, in order to reduce microbiological hazard use as soon as practicable after reconstitution/preparation. If storage is necessary, hold at 2°C - 8°C, for not more than 24 hours.

6.5 Nature and contents of container

DBL Dacarbazine for Injection is available in size 20 mL amber glass vial, single packs.

6.6 Special precautions for disposal and other handling

Intravenous admixtures should be inspected visually for clarity, particulate matter, precipitate, discolouration and leakage prior to administration. Solution showing haziness, particulate matter, precipitate, discolouration or leakage should not be used. Discard unused portion.

HANDLING PRECAUTIONS

As with all antineoplastic agents, trained personnel should prepare DBL Dacarbazine for Injection. This should be performed in a designated area (preferably a cytotoxic laminar flow cabinet). Protective gown, mask, gloves and appropriate eye protection should be worn when handling dacarbazine. Where solution accidentally contacts skin or mucosa, the affected area should be immediately washed, thoroughly with soap and water. It is recommended that pregnant personnel not handle cytotoxic agents such as dacarbazine.

Luer-Lock fitting syringes are recommended. Large bore needles are recommended to minimise pressure and possible formation of aerosols. Aerosols may also be reduced by using a venting needle during preparation.

Items used to prepare DBL Dacarbazine for Injection, or articles associated with body waste should be disposed off by placing in a double sealed polythene bag and incinerated at 1100°C.

SPILLS AND DISPOSAL

If spill occurs, restrict access to the affected area. Wear two pairs of latex rubber gloves, a suitable mask, a protective gown and safety glasses. Limit the spread of the spill by covering with a suitable material such as absorbent towels or adsorbent granules. Spills may also be treated with 5% sodium hypochlorite. Collect the absorbent/adsorbent and other debris from the spill and place in a leak proof plastic container and label accordingly. Cytotoxic waste should be regarded as toxic and hazardous and clearly labelled 'CYTOTOXIC WASTE FOR INCINERATION AT 1100°C'. Waste material should be incinerated at 1100°C for at least 1 second. Clean the remaining spill area with copious amounts of water.

Any unused product or waste material should be disposed of in accordance with local requirements.

7. MEDICINE SCHEDULE

Prescription Medicine

8. SPONSOR

Pfizer New Zealand Limited

P O Box 3998

Auckland, New Zealand

Toll Free Number: 0800 736 363

www.pfizermedicalinformation.co.nz

9. DATE OF FIRST APPROVAL

11 May 1995

10. DATE OF REVISION OF THE TEXT

14 November 2022

TM Trademark

Summary table of changes

Section changed	Summary of new information
3	Add visual description of the appearance of product.
6.3	Typographical correction to registered shelf-life.
All	Minor editorial changes.