

SUMMARY OF PRODUCT CHARACTERISTICS

Fluorouracil “EBEWE” 50mg/ml Concentrate for Solution for Injection / Infusion

1. NAME OF THE MEDICINAL PRODUCT

Fluorouracil “EBEWE” 50mg/ml

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

One vial containing 20 mL contains 1,000 mg of fluorouracil.
One vial containing 50 mL contains 2,500 mg of fluorouracil.
One vial containing 100 mL contains 5,000 mg of fluorouracil.

Excipient with known effect:
Each ml contains 7.58-9.31mg sodium.

For the full list of excipients, see Section 6.1.

3. PHARMACEUTICAL FORM

Concentrate for solution for injection/infusion.

Clear and colourless to slightly yellowish solution.

4. CLINICAL PARTICULARS

4.1 Therapeutic indications

Palliative management of carcinoma of the colon, rectum, breast, stomach and pancreas, in selected patients considered incurable by surgery or other means.
As leucovorin-fluorouracil chemotherapy combination for cancer treatment.

4.2 Posology and method of administration

Note: when using leucovorin (calcium folinate) - fluorouracil chemotherapy combination, strict caution should be exercised not to mix the 2 drugs in the same administration set because of incompatibility (see section 6.2).

Selection of an appropriate dose and treatment regime will depend upon the condition of the patient, the type of carcinoma being treated and whether fluorouracil is to be administered alone or in combination with other therapy. Initial treatment should be given in hospital and the total daily dose should not exceed 1 gram. It is customary to calculate the dose in accordance with patient's actual weight unless there is obesity, oedema or some other form of abnormal fluid retention such as ascites. In this case, ideal weight should be used as the basis for the calculation. Reduction of the dose is advisable in patients with any of the following:

- 1) Cachexia
- 2) Major surgery within preceding 30 days
- 3) Reduced bone marrow function
- 4) Impaired hepatic or renal function

Fluorouracil injection can be given by intravenous injection or infusion. Fluorouracil injection should not be mixed directly, in the same container, with other chemotherapeutic agents or intravenous additives.

Fluorouracil is applied alone and in combination with other cytostatic drugs. The fluorouracil dosage depends on the schedule opted for, the use of other cytostatic drugs, the application of radiotherapy and the method of administration. The total daily dosage will usually not exceed 1 gram.

Colorectal tumours:

The initial therapy may be given as intravenous injections or intravenous infusion. The toxicity of fluorouracil is usually higher after injection than after infusion.

As intravenous infusion, 600 mg/m² daily (with a maximum of 1 g each time) in 300-500 ml 5% glucose solution may be given during 4 hours. This dosage is repeated daily until the first side effects occur. Therapy should then be interrupted. After disappearance of the haematological and gastrointestinal side effects, a maintenance therapy is given.

Fluorouracil is also given as a continuous infusion. The dosage and duration of the infusion depends on the schedule chosen, the use of other cytostatic drugs and the application of radiotherapy. In a dosage up to 300 mg/m² daily for 30-60 consecutive days, toxicity will rarely occur. In higher dosages stomatitis will be the dose-limiting side effect. A common dosage is 350 mg/m² daily.

As injection, 480 mg/m² daily is given intravenously on 3 consecutive days. If toxic side effects do not appear, 240 mg/m² is given intravenously on days 5, 7 and 9, followed by a maintenance therapy. The maintenance therapy consists of injections: once a week 200-400 mg/m² in intravenous injection.

Breast cancer:

For the treatment of breast cancer fluorouracil is given, for example, in combination with methotrexate and cyclophosphamide or in combination with doxorubicin and cyclophosphamide. The usual fluorouracil dosage in these schedules is 400-600 mg/m², intravenously administered on days 1 and 8, in a 28-day cycle.

In some schedules fluorouracil is administered as a continuous infusion. A common dosage is 350 mg/m²/day.

Other types of administration:

Fluorouracil is also used as a continuous infusion. The dosage and duration of the infusion depends on the regimen chosen, the use of other cytostatic drugs and the application of radiotherapy. A common dosage is 350 mg/m²/day.

When fluorouracil concentrate for solution for injection is used for continuous infusion, the fact that the solution for injection has not been preserved should be taken into account.

Dosage adjustment:

The fluorouracil dose should be adjusted in accordance with the schedule below if leukocytes or thrombocytes are reduced on the first day of therapy; the lowest value determines the height of the dose.

% of dose	Leukocytes	Thrombocytes
100	> 3,500	> 125,000
50	2,500-3,500	75,000-125,000
0	< 2,500	< 75,000

If the number of leukocytes is 2500-3500 /mm³ and/or the number of thrombocytes 75,000-125,000 /mm³, it is better not to administer cytostatic drugs for one week. When the blood count has been restored the course may be continued; if not, dose reduction may be carried out.

The administration of fluorouracil should be discontinued if a bilirubin plasma concentration over 85 micromol/L is reached. If the patient has undergone major surgery within 30 days prior to administration, the recommended dosage should be reduced by a third to a half from the very beginning.

Children

No recommendations are made regarding the use of fluorouracil in children.

Elderly

Fluorouracil should be used in the elderly with similar considerations as in younger adult dosages, notwithstanding that incidence of concomitant medical illness is higher in the former group.

4.3 Contraindications

Fluorouracil should not be used in the case of:

- Hypersensitivity to the active substance or to any of the excipients listed in Section 6.1
- Bone marrow suppression (myelosuppression)
- Marked changes in blood counts
- Severely impaired liver function
- Acute infections
- Patients in poor general health
- Patients with known complete absence of dihydropyrimidine dehydrogenase (DPD) activity (see Section 4.4)
- During Pregnancy and breast-feeding (see Section 4.6)
- Recent or concomitant treatment with brivudine (see Sections 4.4 and 4.5 on drug interactions)

In patients with dihydropyrimidine dehydrogenase (DPD) deficiency, the commonly used fluorouracil doses cause increased side effects. If severe adverse effects occur, control of DPD activity may be indicated. Patients with a DPD deficiency should not be treated with fluorouracil.

Active vaccinations should not be performed in temporal association with fluorouracil therapy. Contact with individuals who received a polio vaccine should be avoided.

4.4 Special warnings and precautions for use

Precautions to be taken when handling and using fluorouracil

Because of the potentially mutagenic and carcinogenic effects, increased safety regulations apply to nursing staff and physicians. When handling fluorouracil, any contact with the skin and mucous membranes should be avoided. The preparation must be carried out by means of an absolutely aseptic procedure. The use of a vertical laminar airflow workbench (LAF) is recommended. Protective clothing should be worn when handling fluorouracil. Pregnant personnel should not handle fluorouracil.

Cardiotoxicity

Treatment with fluoropyrimidines was associated with cardiotoxicity, including myocardial infarction, angina, arrhythmias, myocarditis, cardiogenic shock, sudden death, and ECG changes (including, in very rare cases, prolongation of the QT interval). These adverse events are more common in patients who receive a continuous infusion of fluorouracil than in the recipients of a bolus injection. A history of coronary artery disease may be a risk factor for cardiac side effects. Caution should therefore be exercised in the treatment of patients who experienced chest pain during treatment cycles, and in patients with known heart disease. During the treatment with fluorouracil, heart function should be monitored regularly. In the case of severe cardiotoxicity, therapy should be discontinued.

Encephalopathy

During post-marketing surveillance, cases of fluorouracil treatment-related encephalopathy (including hyperammonemic encephalopathy and leukoencephalopathy) were reported. Signs and symptoms of encephalopathy include mental state changes, confusion, disorientation, coma and ataxia. If any of these symptoms occur, treatment should be discontinued immediately, and serum ammonia levels should be determined. If serum ammonia levels are elevated, ammonia-lowering treatment should be initiated.

Caution should be exercised when administering fluorouracil to patients with impaired renal and/or hepatic function. Patients with impaired renal and/or hepatic function may be at increased risk for hyperammonemia and hyperammonemic encephalopathy.

Dihydropyrimidine dehydrogenase (DPD) deficiency

A rare, unexpected and severe toxicity associated with the use of 5-FU, manifesting as stomatitis, diarrhea, mucositis, neutropenia and neurotoxicity, was explained with impaired DPD activity. Patients with a low or no DPD activity (an enzyme involved in the degradation of fluorouracil) are at increased risk for serious, life-threatening or fatal side effects caused by fluorouracil. Although a DPD deficiency cannot be clearly defined, it is known that patients with certain homozygous or certain complex heterozygous mutations in the DPYD locus (e.g. DPYD*2A, c.1679T>G, c.2846A>T and c.1236G>A/HapB3 variants), which may result in a complete or nearly complete absence of enzymatic DPD activity (as determined in laboratory experiments), are at greatest risk for life-threatening or fatal adverse events and should not be treated with fluorouracil (see Section 4.3). There is no proven dosage that is safe for patients with a complete absence of DPD activity.

Patients with certain heterozygous DPYD variants (including DPYD*2A, c.1679T>G, c.2846A>T and c.1236G>A/HapB3 variants) were at increased risk for severe toxicities when treated with fluorouracil.

The frequency of heterozygous DPYD*2A genotypes in the DPYD gene in Caucasian patients is about 1%, 1.1% for c.2846A>T, 2.6% to 6.3% for c.1236G>A/HapB3 variants, and 0.07% to 0.1% for c.1679T>G. Genotyping for these alleles is recommended to identify patients at increased risk for severe toxicities. Information on the frequency of these DPYD variants in populations other than Caucasians is limited. It cannot be ruled out that other rare variants may also be associated with an increased risk of severe toxicities.

Special care must be exercised in patients with partial DPD deficiency (such as those with heterozygous mutations in the DPYD locus), where the benefit of fluorouracil outweighs the risk, taking into account the suitability of an alternative non-fluoropyrimidine chemotherapy regimen. Regular follow-up with dose adjustment according to toxicity must be performed. In these patients, a reduced initial dose should be considered to avoid severe toxicity. There is not enough data to recommend a specific dose in patients with partial DPD activity measured with a specific test. It was reported that the DPYD*2A, c.1679T>G variants resulted in a greater reduction in enzymatic activity than the other variants, which was associated with a higher risk of side effects. The effects of a reduced dose on efficacy are currently uncertain. Therefore, if there are no severe toxicities, the dose could be increased while the patient is being monitored carefully. Patients who tested negative for the allele mentioned above may still be at high risk for serious adverse events.

Patients with an undiagnosed DPD deficiency, who are treated with fluorouracil, as well as patients who tested negative for specific DPYD variants, may experience life-threatening toxicities that clinically manifest as an acute overdose (see Section 4.9). In case of acute toxicity grade 2 – 4, therapy should be discontinued immediately. A permanent discontinuation should be considered based on the clinical assessment of the onset, duration and severity of the observed toxicity.

Other notes

Brivudine should not be used together with fluorouracil. Deaths due to this drug interaction have been reported. After the end of treatment with brivudine and before the start of therapy with fluorouracil, an

interval of at least 4 weeks is required. Brivudine treatment may be initiated 24 hours after the last dose of fluorouracil (see Sections 4.3 and 4.5),

In the case of an accidental administration of brivudine to patients treated with fluorouracil, effective measures must be taken to reduce the toxicity of fluorouracil. Immediate hospitalization is recommended. All measures should be taken to prevent systemic infections and dehydration.

Patients taking phenytoin concomitantly with fluorouracil should be regularly evaluated for a possible increase in phenytoin plasma levels.

Damage to the intestinal wall requires symptomatic treatment commensurate with the severity, e.g. fluid replacement. Mild diarrhea may respond to antidiarrheal agents. In moderate to severe diarrhea, however, such agents do not suffice.

Before and during therapy with fluorouracil, the following follow-up examinations are recommended:

- Daily inspection of the oral cavity and pharynx for mucosal changes
- Blood count including the differential count, and platelet count, before each fluorouracil administration
- Kidney parameters
- Liver function tests (LFTs)

When co-administering fluorouracil and oral anticoagulants, the prothrombin time (Quick test) should be monitored closely.

Patients should be specifically alerted to the possibility of experiencing stomatitis/mucositis, diarrhea and bleeding (especially from the gastrointestinal tract). Patients should be instructed to consult their attending physician as soon as they have the first symptoms.

Patients should also be alerted to the possibility of experiencing hair loss, which is usually reversible, and skin changes (see also Section 4.8).

Pediatric population

There is insufficient experience on the efficacy and safety of fluorouracil in children.

4.5 Interaction with other medicinal products and other forms of interaction

Brivudine: A clinically significant interaction between brivudine and fluoropyrimidines (e.g. capecitabine, fluorouracil, Tegafur) was reported, which is based on an inhibition of dihydropyrimidine dehydrogenase by brivudine. This interaction, which leads to increased fluoropyrimidine toxicity, is potentially fatal. For this reason, brivudine should not be co-administered with fluorouracil (see Sections 4.3 and 4.4). After the end of therapy with brivudine, at least 4 weeks must pass before therapy with fluorouracil may be initiated. Therapy with brivudine may be initiated 24 hours after the last dose of fluorouracil.

Any treatment measure that worsens the patient's general condition or interferes with bone marrow function (e.g. other cytostatic agents) may increase the toxicity of fluorouracil. Fluorouracil may increase the cutaneous toxicity of radiation therapy.

Calcium folinate enhances the action of fluorouracil. Serious, including sometimes lethal, diarrhea may occur as a clinical consequence of this interaction.

An accumulation of such deaths has been reported particularly in the case of the administration regimen of a single weekly I.V. bolus injection of 600 mg/m² body surface area of fluorouracil in combination with calcium folinate.

Concomitant administration of phenytoin and fluorouracil has been reported to increase the plasma levels of phenytoin, which resulted in symptoms of phenytoin intoxication (see Section 4.4).

Cimetidine, metronidazole and interferons may increase the plasma levels of fluorouracil. This can increase the toxic effects of fluorouracil.

In female patients who received a thiazide diuretic in addition to cyclophosphamide, methotrexate and fluorouracil, the granulocyte count decreased more than after the same cytostatic cycles not containing thiazide.

In isolated cases, a drop in Quick-type PT was observed in patients treated with warfarin who also received fluorouracil alone or in combination with levamisole.

When treated with fluorouracil and levamisole, hepatotoxic effects (elevated serum levels of alkaline phosphatase, transaminases or bilirubin) are frequently observed.

Female patients with breast cancer who received combination treatment with cyclophosphamide, methotrexate, fluorouracil and tamoxifen displayed an increased risk of developing thromboembolic events.

Co-administration of vinorelbine with fluorouracil/folinic acid may cause severe mucositis leading to death.

The detection methods for bilirubin and 5-hydroxyindoleacetic acid in the urine may reveal elevated or false-positive values.

General comments:

Cytostatic agents may reduce antibody production after influenza vaccination.

Cytostatic agents may increase the risk of infection after the administration of live vaccines.

4.6 Fertility, pregnancy and breast-feeding

Pregnancy

Fluorouracil may have mutagenic effects and should not be used during pregnancy. Women of childbearing potential should use effective contraception during chemotherapy and up to 6 months thereafter. If pregnancy occurs during treatment, the possibility of genetic counseling should be used.

Breast-feeding

Women should not breast-feed during treatment.

Fertility

Fluorouracil may have mutagenic effects. Therefore, men treated with fluorouracil are advised not to father a child during treatment and up to 6 months thereafter.

4.7 Effects on ability to drive and use machines

Fluorouracil may indirectly affect the ability to drive or use machines by causing nausea and vomiting. Therefore, during treatment with fluorouracil, driving a car and using machines should be avoided.

4.8 Undesirable effects

The most common and significant side effects of fluorouracil are bone marrow toxicity and gastrointestinal symptoms.

The frequency of adverse reactions is defined using the following convention:

<i>very common</i>	$\geq 1/10$
<i>common</i>	$\geq 1/100$ to $< 1/10$
<i>uncommon</i>	$\geq 1/1,000$ to $< 1/100$
<i>rare</i>	$\geq 1/10,000$ to $< 1/1,000$
<i>very rare</i>	$< 1/10,000$
<i>not known</i>	<i>Frequency cannot be estimated from available data</i>

Infections and infestations

Very common: Infections

Common: Immunosuppression with an increased infection rate.

Rare: Sepsis.

Blood and lymphatic system disorders

Common: Myelosuppression, neutropenia, thrombocytopenia and anemia, febrile neutropenia.

Rare: Agranulocytosis, pancytopenia.

Myelosuppression is common and is one of the dose-limiting adverse events (see Sections 4.2 and 4.4). Mild to extremely severe cases of neutropenia and thrombocytopenia, as well as agranulocytosis, anemia and pancytopenia have all been reported.

The degree of myelosuppression (NCI grades 1 to 4) depends on the method of administration (I.V. bolus injection or continuous intravenous infusion) and the dosage regimen.

Neutropenia occurs after each course of treatment as I.V. bolus injections in adequate doses (nadir: 9th to 14th [to 20th] day of treatment; normal values usually achieved after the 30th day).

Immune system disorders

Rare: Generalized allergic reactions including anaphylactic shock.

Endocrine disorders

Not known: Increases in serum levels of total thyroxine (T4) and total triiodothyronine (T3), with no increase in free T4 and TSH, and no clinical signs of hyperthyroidism.

Metabolism and nutrition disorders

Common: Hyperuricemia.

Nervous system disorders

Rare: Nystagmus, headache, dizziness, Parkinson's symptoms, pyramidal signs and euphoria. Peripheral neuropathy (in combination regimens with radiation therapy).

- Very rare: Dysgeusia.
(Leuko-)encephalopathy with symptoms such as ataxia, speech disorders, confusion, disorientation, muscle weakness, aphasia, seizures or coma.
- Not known: Hyperammonemic encephalopathy.

Eye disorders

- Rare: Excessive flow of tears, blurred vision, disorders of ocular motility, optic neuritis, diplopia, loss of vision, photophobia, conjunctivitis, blepharitis, scar-related ectropion, and tear-duct fibrosis.

Cardiac disorders

- Common: Ischemic changes in ECG.
- Uncommon: Chest pain resembling angina pectoris.
- Rare: Arrhythmias, myocardial infarction, myocarditis, heart failure, dilated cardiomyopathy and cardiogenic shock.
- Very rare: Cardiac arrest and sudden cardiac death.
- Not known: Pericarditis.

Cardiotoxic side effects usually occur during or a few hours after the first administration cycle. Patients with pre-existing coronary artery disease or cardiomyopathy are at an increased risk of developing cardiotoxic side effects.

Vascular disorders

- Rare: The occurrence of thrombophlebitis has been reported.
- Not known: Cerebral, intestinal and peripheral ischemia, Raynaud's syndrome and thromboembolism.

Respiratory, thoracic and mediastinal disorders

- Common: Bronchospasm, epistaxis.

Gastrointestinal disorders

- Common: Mucositis (stomatitis, esophagitis, proctitis), watery diarrhea, nausea and vomiting.
- Rare: Dehydration as well as ulcers and bleeding in the gastrointestinal tract.

Gastrointestinal side effects are common and may be life-threatening. Mucositis (stomatitis, esophagitis, proctitis), watery diarrhea, nausea and vomiting (ranging from mild to severe), and a calculous cholecystitis have been reported (see also Section 4.4). The severity (NCI grades 1 through 4) of gastrointestinal adverse effects depends on the dosage regimen and method of administration. In administration by continuous intravenous infusion, stomatitis rather than myelosuppression is the dose-limiting factor.

As long as inflammations, ulcers or diarrhea persist, the use of fluorouracil should be avoided.

Hepatobiliary disorders

Rarely, liver cell damage and, in isolated cases, liver necrosis were observed, some of which were lethal.

Skin and subcutaneous tissue disorders

The so-called "hand-foot syndrome" with dysesthesia as well as redness, swelling, pain and peeling of the skin on the palms and soles occurs more frequently after administration as a continuous I.V. infusion than after I.V. bolus injections.

- Common: Alopecia (usually reversible).

Rare: Exanthema, dry skin with fissures, dermatitis, urticaria, photosensitivity, hyperpigmentation of the skin and streaky hyperpigmentation or pigment loss along the vein course.
Nail changes (e.g., diffuse superficial bluish pigmentation, hyperpigmentation, nail dystrophy, pain and thickening of the nail bed, paronychia) and onycholysis.

General disorders and administration site conditions

Common: Delayed wound healing, fatigue, general weakness, fatigue and listlessness.

Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Any suspected adverse events should be reported to the Ministry of Health according to the National Regulation by using an online form: <https://sideeffects.health.gov.il>

4.9 Overdose

Symptoms of intoxication

As a result of an overdose, the following side effects usually occur more frequently:
Nausea, vomiting, diarrhea, severe mucosal inflammation, ulcers and bleeding in the gastrointestinal tract, bone marrow suppression (thrombocytopenia, leukocytopenia, agranulocytosis).

Handling intoxication

In the case of intoxication, the administration of fluorouracil should be discontinued immediately. Symptomatic treatment should be initiated. Treatment of a marked myelosuppression should be administered on an inpatient basis. It may consist of the substitution of missing blood components, and antibiotic treatment. The transfer of the patient into a germ-free room may become necessary. Follow-ups of blood counts should be performed up to 4 weeks after the overdose.

5. PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

Pharmacotherapeutic group: Antineoplastic agents, antimetabolites, ATC code: L01BC02

5-Fluorouracil fluorinated pyrimidine derivative, itself without antineoplastic effects. The inhibition of cell division occurs after metabolism by the active metabolites 5-fluorouridine triphosphate (FUTP) and 5-fluorodeoxyuridine monophosphate (FdUMP).

Known mechanisms of action:

- Blockage of DNA synthesis (inhibition of thymidilate synthetase by FdUMP).
- Inhibition of RNA synthesis (formation of a defective RNA by incorporation of FUTP).
- DNA strand breaks after incorporation of fluorodeoxyuridine triphosphate (phosphorylated FdUMP) into the DNA.

The inhibitory effects are particularly evident in cells that grow rapidly and thus absorb fluorouracil to a greater extent.

5.2 Pharmacokinetic properties

Half-life

After intravenous administration of fluorouracil (5-FU), the (monophasic) elimination half-life is 10–20 minutes and is dose-dependent; a biphasic half-life of 8 or 40 minutes has been reported. 3 hours after administration, 5-FU plasma levels are no longer measurable.

Distribution

The distribution corresponds to the total body fluid. Fluorouracil crosses the blood-cerebrospinal fluid barrier.

Biotransformation

Approximately 85% of the administered dose is metabolized. Active metabolites are the intracellularly formed 5-fluorouridine triphosphate (FUTP) and 5-fluorodeoxyuridine monophosphate (FdUMP). In addition to the active metabolites, 5-FU is mainly metabolized in the liver to inactive metabolites (major metabolites: 5-fluorouridine, 5-fluorodeoxyuridine) and catabolized to uracil. Carbon dioxide, urea and other metabolites are also produced.

Elimination

15% of the administered amount is excreted unchanged within 6 hours by the kidneys, of which about 90% within the first hour.

Fluorouracil is further catabolized by the enzyme dihydropyrimidine dehydrogenase (DPD) to the much less toxic 5,6-dihydro-fluorouracil (FUH₂). The enzyme dihydropyrimidinase cleaves the pyrimidine ring to 5-fluoro-ureido-propionic acid (FUPA). Finally, β -ureido-propionase splits FUPA to α -fluoro- β -alanine (FBAL), which is excreted in the urine. The activity of dihydropyrimidine dehydrogenase (DPD) is rate-limiting. DPD deficiency may increase the toxicity of fluorouracil (see Sections 4.3 and 4.4).

5.3 Preclinical safety data

Acute toxicity

See Section 4.9 Overdose

Chronic toxicity

Fluorouracil has a preferential effect on proliferating cells, which is why bone marrow depression and damage to the mucous membrane of the gastrointestinal tract are the principal consequences.

Mutagenic and carcinogenic potential

Fluorouracil has been shown to be mutagenic in several *in vitro* and *in vivo* mutagenicity tests. There is a suspicion of a mutagenic effect in humans.

Animal experiments with fluorouracil have not provided any indications of a carcinogenic effect. However, fluorouracil belongs to a substance class that promises tumorigenic effects.

Reproductive toxicity

Fertility and teratogenicity studies in various animal species have suggested an embryotoxic and teratogenic potential, as well as impaired fertility and reproductive behavior.

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Sodium hydroxide, Water for injections.

6.2 Incompatibilities

Fluorouracil should only be diluted with physiological saline or 5 % glucose solution. Fluorouracil must not be diluted with strongly buffered solutions with pH <8, since fluorouracil precipitates in this environment. Do not mix with other chemotherapeutic solutions.

Incompatibilities with following substances have been reported:

Folinic acid, carboplatin, cisplatin, cytarabine, diazepam, doxorubicin, droperidol, filgrastim, gallium nitrate, leucovorin, methotrexate, metoclopramide, morphine, ondansetron, parenteral nutrition solutions, vinorelbine and other anthracyclines.

Calcium folinate

Fluorouracil must not be mixed in the same infusion with calcium folinate because a precipitate may form. Fluorouracil 50 mg/ml with calcium folinate 20 mg/ml, with or without dextrose 5% in water, has been shown to be incompatible when mixed in different amounts and stored at 4° C, 23° C or 32° C in polyvinyl chloride containers.

5-Fluorouracil Ebewe solution for injection/infusion must not be mixed with other medicinal products, including oxaliplatin or irinotecan.

6.3 Shelf life

The expiry date of the product is indicated on the packaging materials.

6.4 Special precautions for storage

Store below 25°C. Do not refrigerate or freeze.
Store in the original packaging in order to protect the content from light.

For single use only.
Only use clear and colourless to slightly yellowish solutions.

If a precipitate has formed as a result of exposure to low temperatures, redissolve by careful heating to 60°C accompanied by shaking. Allow to cool prior to use.

Shelf life after first opening

After first opening, can be used for 28 days at a temperature of 20 - 25°C.

Shelf life after dilution

Dilution can be done with 0.9% sodium chloride solution or 5% glucose solution. Stability data in concentrations of 0.35 mg/ml and 15 mg/ml have shown that the maximum shelf life of ready-made fluorouracil infusion solution is 28 days.

This shelf life refers both to refrigerator storage (2 - 8°C) including protection from light and storage at ambient temperature (20 - 25°C) with or without protection from light.

The prepared infusate demonstrated chemical and physical stability for 28 days. From a microbiological point of view, however, the product should be used immediately. If not used immediately, storage times and conditions of the prepared infusate prior to use are the responsibility of the user and would normally not be longer than 24 hours at 2 - 8°C, unless dilution has taken place under controlled and validated aseptic conditions.

6.5 Nature and contents of container

Glass vial containing 20ml
Glass vial containing 50ml
Glass vial containing 100ml

Type I amber glass vials with grey bromobutyl rubber stopper with aluminium crimp caps.

Not all pack sizes may be marketed.

6.6 Special precautions for disposal and other instructions for handling

Due to the possible mutagenic and carcinogenic effects, enhanced safety precautions for hospital staff and physicians apply. While handling fluorouracil any contact with skin and mucous membranes should be avoided. Failing this, immediately wash with soap and water.

In the event of contact with the eyes, the affected area should be washed with copious amounts of water, and medical attention should be sought. All provisions must be made to allow for absolutely aseptic work. The use of a workspace with laminar flow is recommended. Protective clothing must be worn while handling fluorouracil.

Pregnant staff must not work with Fluorouracil.

Inactivation:

700 °C

Sodium hypochlorite (liquor natrii hypochlorosi) diluted with 10 parts of water
Concentrated NaOH over several hours.

The ready solution should be used immediately after preparation.

If a precipitate has formed as a result of exposure to low temperatures, redissolve by careful heating to 60°C accompanied by shaking. Allow to cool prior to use.

Loss of efficacy due to the adsorption of fluorouracil in the glass infusion container has been described in the literature.

The handling and disposal regulations for cytostatic agents must be met.

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

7. LICENCE HOLDER AND MANUFACTURER

Licence holder:

Novartis Israel Ltd.,
P.O.Box 7126,
Tel Aviv

Manufacturer:

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A-4866 Unterach, Austria

8. REGISTRATION NUMBER

130 30 30866

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