

1.3.1.1: Proposed New Zealand Datasheet

Data Sheet

EVOLTRA®

NAME OF THE MEDICINE

Evoltra® clofarabine 20 mg/20 mL concentrated solution for infusion vial

PRESENTATION

Clofarabine is a purine nucleoside anti-metabolite structurally related to marketed substances like fludarabine and cladribine. It is a white to off-white solid which is soluble in saline up to 1.5 mg/mL at room temperature. The optical rotation is + 39.93° (average; $c = 5$ mg/mL, 25°C) in dimethylformamide solvent. The alcohol pKa is 16, the amine pKa is 12, and the protonated base pKa is approximately 2.

Evoltra® is a sterile, clear, practically colourless solution containing clofarabine 20 mg/20 mL. Each 20 mL vial contains 20 mg of clofarabine and 180 mg of sodium chloride. This is equivalent to 3.08 mmol (or 70.77 mg) of sodium and should be taken into consideration for patients on a controlled sodium diet. The solution also contains Water for Injections. The pH of the solution ranges between 4.5 and 7.5 and it has an osmolarity of 270 to 310 mOsm/L.

USES

Actions

Antineoplastic agents

Mechanism of action

Clofarabine is a purine nucleoside anti-metabolite. Its antitumour activity is believed to be due to 3 mechanisms:

- DNA polymerase α inhibition resulting in termination of DNA chain elongation and/or inhibition of DNA synthesis / repair
- Ribonucleotide reductase inhibition with reduction of intracellular deoxynucleotide triphosphate (dNTP) pools
- Disruption of mitochondrial integrity with the release of cytochrome C and other proapoptotic factors leading to programmed cell death even in non-dividing cells

In addition, halogenation of the adenine ring at the 2 position increases resistance to cellular degradation by adenosine deaminase, and substitution of a fluorine at the C-2' position in the arabino configuration decreases the susceptibility to phosphorylic cleavage by acid and bacterial purine nucleoside phosphorylase.

Pharmacodynamic effects

In vitro studies have demonstrated that clofarabine inhibits cell growth and is cytotoxic to a variety of rapidly proliferating haematological and solid tumour cell lines (clofarabine concentrations inhibiting 50% of cell growth were $\leq 1\mu\text{M}$ for solid tumours and mostly $> 1\mu\text{M}$ for leukaemia cell lines). It was also active against quiescent lymphocytes and macrophages (50% growth inhibition at $>0.45\mu\text{M}$). In addition, clofarabine delayed tumour

growth and, in some cases, caused transient tumour regression in an assortment of human tumour xenografts implanted in mice.

Pharmacokinetics

The population pharmacokinetics of Evoltra® were studied in 40 paediatric patients aged 2 to 19 years (21 males/19 females) with relapsed or refractory acute lymphocytic leukaemia (ALL) or acute myelogenous leukaemia (AML). At the given 52 mg/m² dose, similar concentrations were obtained over a wide range of body surface areas (BSAs). Clofarabine was 47% bound to plasma proteins, predominantly to albumin (27% binding). Based on non-compartmental analysis, systemic clearance and volume of distribution at steady-state were estimated to be 28.8 L/h/m² and 172 L/m², respectively. The terminal half-life was estimated to be 5.2 hours. No apparent difference in pharmacokinetics was observed between patients with ALL and AML or between males and females.

No relationship between clofarabine or clofarabine triphosphate exposure and toxicity or response was found in this population.

Based on 24-hour urine collections in the paediatric studies, 49 - 60% of the dose is excreted in the urine unchanged. *In vitro* studies using isolated human hepatocytes indicate very limited metabolism (0.2%). The pathways of non-hepatic elimination remain unknown.

CLINICAL TRIALS

Seventy-eight (78) paediatric patients with Acute Lymphocytic Leukaemia (ALL) were exposed to Evoltra®. Seventy (70) of the patients received the recommended paediatric dose of 52 mg/m² daily x 5 as an intravenous (IV) infusion.

Phase I Study in Paediatric Patients with Haematologic Malignancies

The safety and efficacy of Evoltra® were evaluated in paediatric patients with refractory or relapsed haematologic malignancies in an open-label, dose-escalation, noncomparative study. The starting dose was 11.25 mg/m²/day IV infusion daily x 5 and escalated to 70 mg/m²/day IV infusion daily x 5. This dosing schedule was repeated every 2 to 6 weeks depending on toxicity and response. Nine of 17 ALL patients were treated with 52 mg/m² daily x 5. In the 17 ALL patients there were 2 complete remissions (12%) and 2 partial remissions (12%) at varying doses. Dose-limiting toxicities (DLTs) in this study were reversible hyperbilirubinaemia and elevated transaminase levels and skin rash, experienced at 70 mg/m². As a result of this study, the recommended dose for subsequent study in paediatric patients was determined to be 52 mg/m²/day for 5 days.

Phase II Single Arm Study in Paediatric ALL

Evoltra® was evaluated in an open-label, single arm study of 61 paediatric patients with relapsed/refractory ALL. Patients received a dose of 52 mg/m² over 2 hours for 5 consecutive days repeated every 2 to 6 weeks for up to 12 cycles. There was no dose escalation in this study.

All patients that had disease relapsed after and/or were refractory to two or more prior therapies. Most patients, 38/61 (62%) received > 2 prior regimens and 18/61 (30%) of the patients had undergone at least 1 prior transplant. The median age of the treated patients was 12 years (range 1-20 years old), 61% were male, 39% were female, 44% were Caucasian, 38% were Hispanic, 12% were African-American, 2% were Asian and 5% were of Other race.

The overall remission (OR) rate (Complete Remission [CR] + CR in the absence of total platelet recovery [CRp]), defined as no evidence of circulating blasts or extramedullary disease, an M1 bone marrow (≤ 5% blasts), and recovery of peripheral counts [platelets ≥ 100

$\times 10^9/L$ and absolute neutrophil count (ANC) $\geq 1.0 \times 10^9/L$] and CRp, defined as meeting all criteria for CR except for recovery of platelet counts to $> 100 \times 10^9/L$ was evaluated. Partial Response (PR) was also determined, defined as complete disappearance of circulating blasts, an M2 bone marrow ($\geq 5\%$ and $\leq 25\%$ blasts), and appearance of normal progenitor cells or an M1 marrow that did not qualify for CR or CRp. The remission criteria are based on those of the Children's Oncology Group. Duration of remission and overall survival was also evaluated. Transplantation rate was not a study endpoint.

Response rates for these studies were determined by an unblinded Independent Response Review Panel (IRRP).

Table 1 summarizes results for the paediatric ALL study. Responses were seen in both pre-B and T-cell immunophenotypes of ALL. The median cumulative dose was 530 mg (range 29-2815 mg) in 1 (41%), 2 (44%) or 3 or more (15%) cycles. The median number of cycles was 2 (range 1-12). The median time between cycles was 28 days with a range of 12 to 55 days.

Table 1: Response Rates in Paediatric ALL Study

N = 61			
Responses	n	%	95% CI
CR	7	11.5	4.7 to 22.2
CRp	5	8.2	2.7 to 18.1
PR	6	9.8	3.7 to 20.2
CR + CRp	12	19.7	10.6 to 31.8

Of 35 patients who were refractory to their immediately preceding induction regimen, 6 (17.1%) achieved a CR or CRp. Of 18 patients who had at least 1 prior haematopoietic stem cell transplant (HSCT), 5 (27.8%) achieved a CR or CRp.

Among the 18 patients who achieved at least a PR, 9 patients achieved the best response after 1 cycle of clofarabine, 8 patients required 2 courses and 1 patient achieved a CR after 3 cycles of therapy. Of these 18 responding patients, 9 had post-clofarabine bone marrow transplantation (3 CR, 3 CRp, 3 PR). Four of these patients who achieved CR or CRp received HSCT while in continued remission, one patient proceeded to transplant following relapse. One additional patient who achieved a CRp proceeded to transplant following alternative therapy. Duration of response was censored at the time of transplant.

The effect of clofarabine on remission duration and overall survival can only be accurately assessed for patients who did not undergo transplantation because of the confounding effect of transplantation. Of the six subjects with complete remission who did not undergo transplant, the median duration of remission was only 9 weeks (range 4 to 59). In the six who did receive transplant, the duration of remission ranged from 10 to 108+ weeks, some of the effect likely drug-related since longer remission may have enabled transplant.

Median overall survival was 63 weeks (range 9 to 72) for non-transplanted responding subjects. Overall survival ranged from 42 to 145+ weeks for transplanted responding subjects. The transplanted result reflects the effects of both clofarabine and the transplant.

Achievement of complete remission appears to have translated into opportunity for transplant and longer survival, but this needs confirmation.

Table 2 lists the individual remission duration and survival data for patients who achieved a CR or CRp.

Table 2: Duration of Remission and Survival for Patients who achieved CR or CRp in Paediatric ALL Study

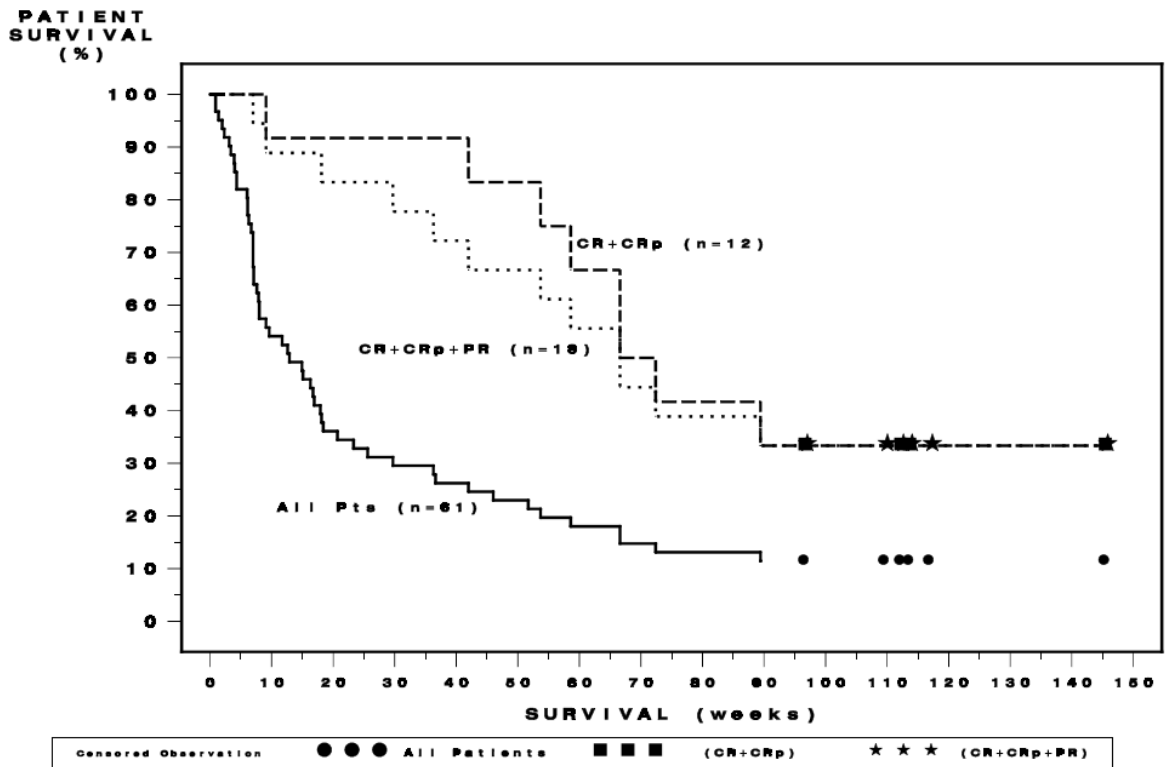
Best Response	Time to OR (weeks)	Duration of Remission (weeks)	Overall Survival (weeks)
Patients who did not undergo transplant			
CR	5.7	4.3	66.6
CR	14.3	6.1	58.6
CR	8.3	47.9	66.6
CRp	4.6	4.6	9.1
CR	3.3	58.6	72.4
CRp	3.7	11.7	53.7
Patients who underwent transplant while in continued remission*			
CRp	8.4	11.6+	145.1+
CR	4.1	9.0+	111.9+
CRp	3.7	5.6+	42.0
CR	7.6	3.7+	96.3+
Patients who underwent transplant after alternative therapy or relapse*			
CRp	4.0	35.4	113.3+**
CR	4.0	9.7	89.4***

* Duration of remission censored at the time of transplant

** Patient received a transplant following alternate therapy

*** Patient received a transplant following relapse

Figure 1:



Median overall survival for CR, CR + CRp, or CR + CRp + PR was 72.4, 69.5, 66.6 weeks respectively (Table 3). Median overall survival for all patients (n=61) was 12.9 weeks. Seven of 61 patients were alive at the time of last follow up (including 2 patients who achieved a PR).

Table 3: Overall Survival in Paediatric ALL Study

Response Category (IRRP)	N	Kaplan-Meier Median	Lower Limit of 95% CI	Upper Limit of 95% CI	Minimum	Maximum	% Censored
Complete Remission	7	72.4	66.6	-	58.6	111.9	28.6
Complete Remission w/o Platelet Recovery	5	53.7	9.1	-	9.1	145.1	40.0
Partial Remission	6	33.0	18.1	-	7.0	116.6	33.3
Treatment Failure/Not Evaluable	43	7.6	6.7	12.6	0.9	150.3	2.3
Overall Remission (CR+CRp)	12	69.5	58.6	-	9.1	145.1	33.3
Complete or Partial Remission (CR+CRp+PR)	18	66.6	42.0	-	7.0	145.1	33.3
All Patients	61	12.9	7.9	18.1	0.9	150.3	11.5

INDICATIONS

Evoltra® is indicated for treatment of acute lymphocytic leukaemia (ALL) in paediatric patients who have relapsed or are refractory after receiving at least two prior regimens. This use is based on the induction of complete responses (see **CLINICAL TRIALS**).

CONTRAINDICATIONS

Clofarabine is contraindicated:

- In patients with hypersensitivity to clofarabine or to any of the excipients (see **PRESENTATION**);
- In patients with severe renal insufficiency or severe hepatic impairment;
- In breast-feeding women.

Breastfeeding should be discontinued prior to, during and following treatment with Evoltra® (see **WARNINGS AND PRECAUTIONS: Use in lactation**).

WARNINGS AND PRECAUTIONS

Evoltra® should be administered under the supervision of a qualified physician experienced in the use of antineoplastic therapy.

Haematologic Toxicity

Obtain complete blood counts and platelet counts at regular intervals during Evoltra® therapy. Suppression of bone marrow function should be anticipated. This is usually reversible and appears to be dose dependent. Severe bone marrow suppression, including neutropenia, anaemia, and thrombocytopenia, has been observed in patients treated with Evoltra®. At initiation of treatment, most patients in the clinical studies had haematological impairment as

a manifestation of leukaemia. Because of the pre-existing immunocompromised condition of these patients and prolonged neutropenia that can result from treatment with Evoltra®, patients are at increased risk for severe opportunistic infections.

The most frequently reported haematologic adverse effects in paediatric patients have included febrile neutropenia (55%) and neutropenia (10%). One (1) report of grade 4 bone marrow depression, and 1 report of grade 4 thrombocytopenia were considered related to the study drug.

Infections

The use of Evoltra® is likely to increase the risk of infection, including severe sepsis, as a result of bone marrow suppression.

At baseline, 48% of the paediatric patients had 1 or more concurrent infections. A total of 83% of patients experienced at least 1 infection after Evoltra® treatment, including fungal, viral and bacterial infections. Twenty-one (18.3%) infection events were considered to be related to clofarabine of which catheter related infection (1 event), sepsis (2 events) and septic shock (2 events; 1 patient died) were considered to be serious.

Hyperuricaemia (Tumour Lysis), and Capillary Leak Syndrome / SIRS

Administration of Evoltra® results in a rapid reduction in peripheral leukaemia cells. Evaluate and monitor patients undergoing treatment with Evoltra® for signs and symptoms of tumour lysis syndrome, as well as signs and symptoms of cytokine release (e.g., tachypnoea, tachycardia, hypotension, pulmonary oedema) that could develop into systemic inflammatory response syndrome (SIRS)/capillary leak syndrome and organ dysfunction. Early intervention is recommended. Provide IV infusion fluids throughout the five days of Evoltra® administration to reduce the effects of tumour lysis and other adverse effects. Administer Allopurinol if hyperuricaemia (tumour lysis) is expected. Discontinue Evoltra® immediately in the event of clinically significant signs or symptoms of SIRS or capillary leak syndrome, either of which can be fatal, and consider use of steroids, diuretics, and albumin. Re-institute Evoltra® when the patient is stable and organ function has returned to baseline, generally with a 25% dose reduction. The use of prophylactic steroids (e.g., 100 mg/m² hydrocortisone on days 1 through 3) may be of benefit in preventing signs and symptoms of SIRS/Capillary Leak Syndrome or cytokine release.

Capillary leak syndrome or SIRS occurred in 6 paediatric patients overall (5 ALL, 1 AML). Several patients developed rapid onset of respiratory distress, hypotension, capillary leak (pleural and pericardial effusions), and multi-organ failure. Other concurrent medical conditions, including sepsis, may also have contributed to the incidence of capillary leak syndrome. In addition, prior therapies and/or disease progression may also make these patients more susceptible to capillary leak syndrome.

Hepatic and Renal Impairment

Evoltra® has not been studied in patients with hepatic or renal dysfunction. In severe dysfunction, Evoltra® is contraindicated. In mild to moderate dysfunction, Evoltra® should be used with the greatest caution [see **DOSAGE AND ADMINISTRATION**].

Hepatobiliary Disorders

Patients who have previously received a hematopoietic stem cell transplant (HSCT) may be at higher risk for hepatotoxicity, suggestive of veno-occlusive disease (VOD) following treatment with clofarabine (40 mg/m²) when used in combination with etoposide (100 mg/m²) and cyclophosphamide (440 mg/m²). Veno-occlusive disease has also been observed in patients who had not received HSCT and were treated with clofarabine (52mg/m²) as the single chemotherapeutic agent. Severe hepatotoxic events have been reported in an ongoing

Phase I / II combination study of clofarabine in paediatric patients with relapsed or refractory acute leukaemia.

Hepatobiliary toxicities were frequently observed in paediatric patients during treatment with Evoltra[®]. Grade 3 or 4 elevated aspartate aminotransferase (AST) occurred in 36% of patients and grade 3 or 4 elevated alanine aminotransferase (ALT) occurred in 43% of patients. Grade 3 or 4 elevated bilirubin occurred in 13% of patients, with 2 events reported as grade 4 hyperbilirubinaemia (2%), one of which resulted in treatment discontinuation, one patient had multi-organ failure and died. One report (1%) of veno-occlusive disease (VOD) was considered related to the study drug.

For patients with follow-up data, elevations in AST and ALT were transient and typically \leq 15 days duration. The majority of AST and ALT elevations occurred within 10 days of Evoltra[®] administration and returned to \leq grade 2 within 15 days. Where follow-up data are available, the majority of bilirubin elevations returned to \leq grade 2 within 10 days. Eight patients had grade 3 or 4 elevations in serum bilirubin at the last time point measured, these patients died due to sepsis and/or multi-organ failure.

Renal and Urinary Disorders

The most prevalent renal toxicity in paediatric patients was elevated creatinine. Grade 3 or 4 elevated creatinine occurred in 8% of patients. Nephrotoxic medications, tumour lysis, and tumour lysis with hyperuricaemia may contribute to renal toxicity. Haematuria was observed in 13% of patients overall. This event likely reflects thrombocytopenia that is a pre-existing condition in these patients.

Vascular Disorders

Sixty-four patients of 115 (55.7%) experienced at least one vascular disorder adverse effect. Twenty-three patients out of 115 experienced a vascular disorder considered to be related to clofarabine, the most frequently reported being flushing (13 events; not serious) and hypotension (5 events; all of which were considered to be serious). However, the majority of these hypotensive events were reported in patients who had confounding severe infections.

Cardiac Disorders

The most frequently reported cardiac disorder was tachycardia (35%), which was, however, already present in 35% of patients at study entry. Most of the cardiac adverse effects were reported in the first 2 cycles. Pericardial effusion was a finding in these patients on post-treatment studies, [9/115 (8%)]. The effusion was almost always minimal to small and in no cases had haemodynamic significance.

Use in Cardiac Disease

Patients with cardiac disease and those taking medicinal products known to affect blood pressure or cardiac function should be closely monitored during treatment with clofarabine (see **WARNINGS AND PRECAUTIONS: Interactions with other medicines**).

Dehydration/Hypotension

Patients receiving Evoltra[®] may experience vomiting and diarrhea; they should therefore be advised regarding appropriate measures to avoid dehydration. Patients should be instructed to seek medical advice if they experience symptoms of dizziness, lightheadedness, fainting spells, or decreased urine output. Evoltra[®] administration should be stopped if the patient develops hypotension for any reason during the 5 days of administration. If hypotension is transient and resolves without pharmacological intervention, Evoltra[®] treatment can be re-instituted generally with a 25% dose reduction [see **DOSAGE AND ADMINISTRATION**].

Use in Pregnancy

Category D¹

1Drugs which have caused, are expected to have caused or may be expected to cause, an increased risk of human foetal malformations or irreversible damage. These drugs may also have adverse pharmacological effects. Accompanying texts should be consulted for further details.

Evoltra® (clofarabine) may cause foetal harm when administered to a pregnant woman.

Clofarabine was teratogenic in rats and rabbits. Developmental toxicity (reduced foetal body weight and increased post-implantation loss) and increased incidences of malformations and variations (gross external, soft tissue, skeletal and retarded ossification) were observed in rats receiving 1 mg/kg/day (approximately 40% of the recommended clinical dose on an extrapolated AUC basis), and in rabbits receiving 1 mg/kg/day (12 mg/m²/day; approximately 23% of the recommended clinical dose on a mg/m² basis).

There are no adequate and well-controlled studies in pregnant women using clofarabine. If this drug is used during pregnancy, or if the patient becomes pregnant while taking this drug, the patient should be apprised of the potential hazard to the foetus.

Women of childbearing potential and sexually active males must use effective methods of contraception during treatment.

Use in Lactation

It is unknown whether clofarabine or its metabolites are excreted in human breast milk. The excretion of clofarabine in milk has not been studied in animals. Because of the potential for tumorigenicity shown for clofarabine in animal studies and the potential for serious adverse effects, women treated with clofarabine should not nurse. Female patients should be advised to avoid breast-feeding during treatment with Evoltra® (see **CONTRAINDICATIONS**).

Adults (> 21 and < 65 years old): Safety and effectiveness have not been established in adults.

Elderly patients (≥ 65 years old): There are currently insufficient data to establish the safety and efficacy of clofarabine in elderly patients.

Carcinogenicity, Genotoxicity, Effects on Fertility

Clofarabine has not been tested for carcinogenic potential.

Clofarabine showed genotoxic activity in the *in vitro* mammalian cell chromosome aberration assay (CHO cells) and in the *in vivo* rat micronucleus assay. It did not show evidence of mutagenic activity in the bacterial mutation assay (Ames test).

Studies in mice, rats, and dogs have demonstrated dose-related adverse effects on male reproductive organs. Seminiferous tubule and testicular degeneration and atrophy were reported in male mice receiving intraperitoneal (IP) doses of 3 mg/kg/day (9 mg/m²/day, approximately 17% of clinical recommended dose on an AUC basis). The testes of rats receiving 25 mg/kg/day (approximately 5 times the recommended clinical dose on an AUC basis) in a 6-month IV study had bilateral degeneration of the seminiferous epithelium with retained spermatids and atrophy of interstitial cells. In a 6-month IV dog study, cell degeneration of the epididymis and degeneration of the seminiferous epithelium in the testes were observed in dogs receiving 0.375 mg/kg/day (approximately 30% of the clinical recommended dose on an AUC basis). Ovarian atrophy or degeneration and uterine mucosal apoptosis were observed in female mice at 75 mg/kg/day IP (225 mg/m²/day, approximately 4-fold of recommended human dose on a mg/m² basis), the only dose administered to female

mice. Atrophy of the ovaries, uterus and vagina occurred in female rats receiving 12.5 g/kg/day (approximately 4 times the recommended clinical dose on an AUC basis). As the effect of clofarabine treatment on human fertility is unknown, reproductive planning should be discussed with patients as appropriate.

Interactions with Other Medicines

Although no clinical drug-drug interaction studies have been conducted to date, on the basis of the *in vitro* studies, cytochrome p450 inhibitors and inducers are unlikely to affect the metabolism of clofarabine. Clofarabine did not significantly inhibit the activity of CYP3A4, CYP2D6, CYP2C9, CYP2C19 and CYP1A2 at clinically relevant concentrations *in vitro*. Weak induction of CYP1A2 and CYP3A4 occurred in human hepatocytes *in vitro* at clofarabine concentrations of 0.5-12 µg/mL, which was slightly higher than the clinical C_{max}.

Clofarabine is predominately excreted via the kidneys. Hence, the concomitant use of medicinal products that have been associated with renal toxicity and those eliminated by tubular secretion should be avoided particularly during the 5 day clofarabine administration period.

Since the liver is a potential target organ for toxicity, the concomitant use of medicinal products that have been associated with hepatic toxicity should be avoided wherever possible.

Patients taking medicinal products known to affect blood pressure or cardiac function should be closely monitored during treatment with clofarabine.

There are no known clinically significant interactions of Evoltra® with other medications or laboratory tests. No formal drug/laboratory test interaction studies have been conducted with Evoltra®.

Incompatibilities/Compatibilities

This medicinal product must not be mixed with other medicinal products except those mentioned in **DOSAGE AND ADMINISTRATION**.

Driving/Use of Machinery

No studies on the effects of clofarabine on the ability to drive and use machines have been performed. However, patients should be advised that they may experience undesirable effects such as dizziness, light-headedness or fainting spells during treatment and told not to drive or operate machines in such circumstances.

ADVERSE EFFECTS

Clinical Trials Experience

The information provided is based on data generated from clinical trials in which 115 patients (> 1 and ≤ 21 years old) with either ALL or acute myeloid leukaemia (AML) received at least one dose of clofarabine at the recommended dose of 52 mg/m² daily x 5.

Patients with advanced stages of ALL or AML may have confounding medical conditions that make causality of adverse effects difficult to assess due to the variety of symptoms related to the underlying disease, its progression and the co-administration of numerous medicinal products.

Nearly all patients (98%) experienced at least one adverse effect considered by the study investigator to be related to clofarabine. Those most frequently reported were nausea (61% of

patients), vomiting (59%), febrile neutropenia (35%), headache (24%), rash (21%), diarrhoea (20%), pruritus (20%), pyrexia (19%), palmar-plantar erythrodysesthesia syndrome (15%), fatigue (14%), anxiety (12%), mucosal inflammation (11%), and flushing (11%). Sixty-eight patients (59%) experienced at least one serious clofarabine-related adverse effect. One patient discontinued treatment due to grade 4 hyperbilirubinaemia considered as related to clofarabine after receiving 52 mg/m²/day clofarabine. 3 patients died of adverse effects considered by the study investigator to be related to treatment with clofarabine: one patient died from respiratory distress, hepatocellular damage, and capillary leak syndrome; one patient from VRE sepsis and multi-organ failure; and one patient from septic shock and multi-organ failure.

Table 4 lists adverse effects by System Organ Class, including severe or life-threatening events (NCI CTC grade 3 or grade 4), reported in ≥ 1% of the 115 patients in the 52 mg/m²/day dose group (pooled analysis of paediatric patients with ALL and AML).

Table 5 lists the incidence of treatment emergent laboratory abnormalities after Evoltra® administration at 52 mg/m² among paediatric patients with ALL and AML (n=115).

Table 4: Adverse effects considered to be related to clofarabine reported at frequencies ≥ 1/100 (i.e. in > 1/115 patients) in clinical trials (<i>Very common = ≥ 1/10; Common = ≥ 1/100 to < 1/10</i>)	
Blood and the lymphatic system disorders	<i>Very common:</i> Febrile neutropenia <i>Common:</i> Neutropenia
Cardiac disorders	<i>Common:</i> Pericardial effusion*, tachycardia*
Ear and labyrinth disorders	<i>Common:</i> Hypoacusis
Gastrointestinal disorders	<i>Very common:</i> Vomiting, diarrhoea, nausea <i>Common:</i> Haematemesis, mouth haemorrhage, abdominal pain, abdominal pain upper, gingival bleeding, mouth ulceration, proctalgia, stomatitis
General disorders and administration site conditions	<i>Very common:</i> Pyrexia, mucosal inflammation, fatigue <i>Common:</i> Multi-organ failure, pain, oedema, peripheral oedema, feeling hot, feeling abnormal, chills, systemic inflammatory response syndrome*, irritability
Hepato-biliary disorders	<i>Common:</i> Jaundice, veno-occlusive disease, alanine aminotransferase increased (ALT)* and aspartate (AST)* aminotransferase increased, hyperbilirubinaemia
Immune system disorders	<i>Common:</i> Hypersensitivity
Infections and infestations	<i>Common:</i> Septic shock*, sepsis, bacteraemia, pneumonia, herpes zoster, herpes simplex, oral candidiasis
Injury, poisoning and procedural complications	<i>Common:</i> Contusion
Investigations	<i>Common:</i> Weight decreased
Metabolism and nutrition disorders	<i>Common:</i> Dehydration, anorexia, decreased appetite
Musculoskeletal, connective tissue and bone disorders	<i>Common:</i> Musculoskeletal chest pain, bone pain, neck pain, back pain, pain in extremity, myalgia, arthralgia
Neoplasms benign and malignant (including cysts and polyps)	<i>Common:</i> Tumour lysis syndrome*
Nervous system disorders	<i>Very common:</i> Headache <i>Common:</i> Neuropathy peripheral, paraesthesia, somnolence, dizziness, tremor

Psychiatric disorders	<i>Very common:</i> Anxiety <i>Common:</i> Agitation, restlessness, mental status changes
Renal and urinary disorders	<i>Common:</i> Haematuria*
Respiratory, thoracic and mediastinal disorders	<i>Common:</i> Tachypnoea, epistaxis, dyspnoea, cough, respiratory distress
Skin and subcutaneous tissue disorders	<i>Very common:</i> Pruritus, palmar-plantar erythrodysesthesia syndrome <i>Common:</i> Petechiae, rash generalised, erythema, rash pruritic, alopecia, rash maculo-papular, generalised erythema, rash erythematous, skin hyperpigmentation, dry skin, hyperhidrosis, skin exfoliation
Vascular disorders	<i>Very common:</i> Flushing* <i>Common:</i> Hypotension*, haematoma, capillary leak syndrome

* see **WARNINGS AND PRECAUTIONS**

Note all adverse effects occurring at least twice (i.e., 2 or more events (1.7%)) are included in this table.

Uncommon Effects

Gastrointestinal Disorders: Pancreatitis, serum amylase and lipase elevation

Skin Disorders: Stevens-Johnson Syndrome (SJS), toxic epidermal necrolysis (TEN)

Table 5: Incidence of Treatment Emergent Laboratory Abnormalities After Evoltra® Administration

Parameter	Any Grade	Grade 3 or higher
Haematology		
Anaemia (N=114)	95 (83.3%)	86 (75.4%)
Leukopenia (N=114)	100 (87.7%)	100 (87.7%)
Lymphopenia (N=113)	93 (82.3%)	93 (82.3%)
Neutropenia (N=113)	72 (63.7%)	72 (63.7%)
Thrombocytopenia (N=114)	92 (80.7%)	91 (79.8%)
Blood creatinine increased (N=115)	57 (49.5%)	9 (7.8%)
Aspartate aminotransferase increased (N=100)	74 (74.0%)	36 (36.0%)
Alanine aminotransferase increased (N=113)	91 (80.5%)	49 (43.4%)
Blood bilirubin increased (N=114)	51 (44.7%)	15 (13.2%)

Post-marketing Experience

The following adverse effects have been identified during post approval use of Evoltra®. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure. Decisions to include these reactions in labeling are typically based on one or more of the following factors: (1) seriousness of the reaction, (2) reported frequency of the reaction, or (3) strength of causal connection to Evoltra®.

- Blood and lymphatic system disorders: Bone marrow failure and pancytopenia.
- Gastrointestinal disorders: Colitis.
- Hepatobiliary disorders: Serious hepatotoxic events of venoocclusive disease have been reported in patients who had previous HSCT and who received conditioning regimens that included busulfan, melphalan, and/or the combination of cyclophosphamide and total body irradiation.
- Infections and infestations:
 - *Systemic*: neutropenic sepsis, bacterial sepsis (Streptococcus, Enterococcus, Klebsiella), fungal sepsis (Candida), bacteraemia (Streptococcus).
 - *Respiratory*: bronchopulmonary aspergillosis, lung abscess, pneumonia bacterial (Klebsiella), lower respiratory tract infection, respiratory tract infection fungal.
 - *GI*: tonsillitis, hepatosplenic candidiasis.
 - *Neurological*: brain abscess, central nervous system infection.
 - *Musculoskeletal*: necrotizing fasciitis.
 - *Skin*: wound infection bacterial (Staphylococcus).
 - *Unspecified*: Cytomegalovirus, Fusarium, Pseudomonas, central line infection, vaginal cellulitis.
- Nervous system disorders: Syncope.
- Psychiatric disorders: Confusional state.

DOSAGE AND ADMINISTRATION

Dosage

Paediatric patients: Administer the recommended paediatric dose of 52 mg/m² as an IV infusion over 2 hours daily for 5 consecutive days. Treatment cycles are repeated following recovery or return to baseline organ function, approximately every 2 to 6 weeks. The dosage is based on the patient's body surface area (BSA), calculated using the actual height and weight before the start of each cycle.

In clinical trials with clofarabine in paediatric acute leukaemias, the median duration between cycles was 28 days (range of 12-55 days) [see **CLINICAL TRIALS**].

Provide IV infusion fluids throughout the 5 days of Evoltra[®] administration to reduce the effects of tumour lysis and other adverse effects. Consider prophylactic anti-emetic medications as Evoltra[®] is moderately emetogenic. The use of prophylactic steroids may be of benefit in preventing signs or symptoms of SIRS or capillary leak (e.g., hypotension, tachycardia, tachypnoea, and pulmonary oedema).

Most patients who respond to clofarabine do so within 1 to 2 treatment cycles (see **CLINICAL TRIALS**). The potential benefits and risks of continued therapy should be considered in those not responding after 2 cycles.

Children (weighing < 20 kg): An infusion time of > 2 hours should be considered to help reduce symptoms of anxiety and irritability, and to avoid unduly high maximum concentrations of clofarabine (see **USES: Actions**).

Children (< 1 year old): There are no data on the pharmacokinetics, safety or efficacy of clofarabine in infants. Therefore, a safe and effective dosage recommendation for patients (< 1 year old) has yet to be established.

Dose reduction for patients experiencing haematological toxicities:

Administer subsequent cycles no sooner than 14 days from the starting day of the previous cycle provided the patient's ANC is $\geq 0.75 \times 10^9/L$.

If a patient experiences a Grade 4 neutropenia (ANC $< 0.5 \times 10^9/L$) lasting ≥ 4 weeks, reduce dose by 25% for the next cycle.

Dose reduction for patients experiencing non-haematological toxicities:

Withhold Evoltra[®] if a patient develops a clinically significant infection, until the infection is clinically controlled and then restart at the full dose. **In the event of a second clinically significant infection, clofarabine treatment should be withheld until the infection is clinically controlled and may be reinitiated at a 25% dose reduction.**

Withhold Evoltra[®] if a Grade 3 non-infectious non-haematologic toxicity (excluding transient elevations in serum transaminases and/or serum bilirubin and/or nausea/vomiting that was controlled by antiemetic therapy) occurs. Re-institute Evoltra[®] administration at a 25% dose reduction upon resolution or return to baseline.

Should a patient experience the same severe toxicity on a second occasion, treatment should be delayed until the toxicity resolves to baseline parameters or to the point where it is no longer severe and the potential benefit of continued treatment with clofarabine outweighs the risk of such continuation. It is then recommended that clofarabine be administered at a further 25% dose reduction.

Any patient who experiences a severe toxicity on a third occasion, a severe toxicity that does not recover within 14 days (see above for exclusions), or a life-threatening or disabling toxicity (US NCI CTC Grade 4 toxicity) should be withdrawn from treatment with clofarabine (see **WARNINGS AND PRECAUTIONS**).

Evoltra[®] administration should be stopped if the patient develops hypotension for any reason during the 5 days of administration. If hypotension is transient and resolves without pharmacological intervention, Evoltra[®] treatment can be re-instituted generally with a 25% dose reduction.

Discontinue Evoltra[®] administration if a patient shows early signs or symptoms of SIRS or capillary leak (e.g., hypotension, tachycardia, tachypnoea, and pulmonary edema) occur and provide appropriate supportive measures.

Discontinue Evoltra[®] administration if substantial increases in creatinine or bilirubin are noted. Re-institute Evoltra[®] when the patient is stable and organ function has returned to baseline, possibly with a 25% dose reduction. If hyperuricaemia is anticipated (tumour lysis), prophylactically administer allopurinol.

Patients with renal insufficiency: There is no experience in patients with renal insufficiency (see **CONTRAINDICATIONS** and **WARNINGS AND PRECAUTIONS**). Close monitoring of renal function during the 5 days of Evoltra[®] administration is advised [see **WARNINGS AND PRECAUTIONS**]. Avoid drugs with known renal toxicity during the 5 days of Evoltra[®] administration.

Patients with hepatic impairment: There is no experience in patients with hepatic impairment (see **CONTRAINDICATIONS** and **WARNINGS AND PRECAUTIONS**).

Close monitoring of hepatic function during the 5 days of Evoltra[®] administration is advised [see **WARNINGS AND PRECAUTIONS**]. Avoid concomitant use of medications known to induce hepatic toxicity.

Administration

Evoltra[®] contains no antimicrobial preservative. Product is for single use in one patient only. Discard any residue.

Evoltra[®] must be diluted prior to administration (see below). The recommended dosage should be administered by intravenous infusion although it has been administered via a central venous catheter in ongoing clinical trials. Evoltra[®] must not be mixed with or concomitantly administered using the same intravenous line as other medicinal products (see **Incompatibilities/Compatibilities**).

Evoltra[®] should be filtered through a sterile 0.2 micrometre syringe filter and then diluted with sodium chloride 9 mg/mL (0.9%) intravenous infusion to produce a total volume according to the examples given in the tables below. However, the final dilution volume may vary depending on the patient's clinical status and physician discretion. (If the use of a 0.2 micrometre syringe filter is not feasible, the sterile concentrate should be pre-filtered with a 5 micrometre filter, diluted and then administered through a 0.22 micrometre in-line filter).

Body surface area (m²)	Sterile concentrate (ml)*	Total diluted volume
≤ 1.44	≤ 74.9	100 ml
1.45 to 2.40	75.4 to 124.8	150 ml
2.41 to 2.50	125.3 to 130.0	200 ml

*Each ml of concentrate contains 1 mg of clofarabine. Each 20 ml vial contains 20 mg of clofarabine. Therefore, for patients with a body surface area ≤ 0.38 m², the partial contents of a single vial will be required to produce the recommended daily dosage of clofarabine. However, for patients with a body surface area > 0.38 m², the contents of between 1 to 7 vials will be required to produce the recommended daily dosage of clofarabine.

The diluted sterile concentrate should be a clear, colourless solution. Visually inspect for particulate matter and discolouration prior to administration.

For stability of the diluted sterile solution and associated information, refer to **STABILITY** section.

Procedures for proper handling of antineoplastic agents should be observed. Cytotoxic medicinal products should be handled with caution.

The use of disposable gloves and protective garments is recommended when handling Evoltra[®]. If the product comes into contact with eyes, skin or mucous membranes, rinse immediately with copious amounts of water.

Evoltra[®] should not be handled by pregnant women.

Stability

The diluted sterile concentrate is chemically and physically stable for 3 days at 2 to 8°C and at room temperature. From a microbiological point of view, it should be used immediately. If not used immediately, in-use storage times and conditions prior to use are the responsibility of the user and would normally not be longer than 24 hours at 2 to 8°C unless dilution has taken place under controlled and validated aseptic conditions.

OVERDOSAGE

Symptoms

No case of overdose has been reported. However, possible symptoms of overdose may include nausea, vomiting, diarrhoea and severe bone marrow depression. To date, the highest daily dose administered to human beings is 70 mg/m² for 5 consecutive days (2 paediatric ALL patients). The toxicities observed in these patients included vomiting, hyperbilirubinaemia, elevated transaminase levels and maculo-papular rash.

Treatment

No specific antidotal therapy exists. Immediate discontinuation of therapy, careful observation and initiation of appropriate supportive measures are recommended.

In case of overdose, immediately contact the Poisons Information Centre for advice on management. (in Australia, call 13 11 26; in New Zealand call 0800 764 766).

PHARMACEUTICAL PRECAUTIONS

Store below 25 °C. Do not freeze.

For storage conditions of the diluted medicinal product, see **STABILITY**.

MEDICINE CLASSIFICATION

Prescription Medicine

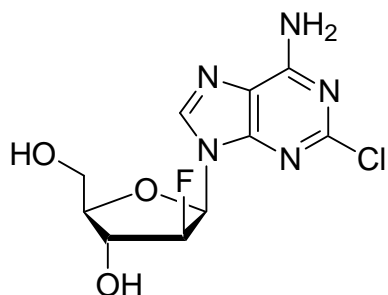
PACKAGING QUANTITIES

Strength	Pack*
20 milligram/20 mL	1 x 20 mL vial
	3 x 20 mL vials
	4 x 20 mL vials
	10 x 20 mL vials

* Not all pack sizes may be marketed.

FURTHER INFORMATION

Structure



Clofarabine

Chemical Name

2-chloro-9-(2'-deoxy-2'-fluoro-β-D-arabinofuranosyl)-9H-purine-6-amine.

Molecular Formula

C₁₀H₁₁ClFN₅O₃

CAS Number

123318-82-1

Molecular Weight

303.68

NAME AND ADDRESS

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31 August 2010

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