Alert	Folinic acid is not the same as folic acid but does have an equivalent vitamin activity.
Indication	Also known as calcular formate of Leucovorm.
indication	Eclinic acid dependent seizures and cerebral folate deficiency ^{3,4}
Action	Folinic acid dependent seizeres and cerebra rolate denciency.
	R Group Vitamin
Trade Name	DRI Laucovarin Calcium Injection, Dfizer Laucovarin Calcium Solution for Injection
ITade Name	DBL Leucovorin Calcium Tablets, Folinic Acid Cansules (FIT-Rioceuticals)
Presentation	DBL Leucovorin Calcium injection $_{-}$ 15 mg/2 ml $_{-}$ 50 mg/5 ml $_{-}$ 300 mg/20 ml
Fresentation	Pfizer (Perth) Leucovorin Calcium Injection LISP 50 mg (folinic acid) in 5 mL (sterile) and 100 mg (folinic
	acid) in 10 ml. Plastic Vial
	Pfizer (Perth) Leucovorin Calcium Injection LISP 50 mg (folinic acid) in 5 mL (sterile) and 100 mg (folinic
	acid) in 10 ml Steriluer amnoule
	DBL Leucovorin Calcium tablet 15 mg folinic acid
	Folinic Acid Capsules (FIT-Bioceuticals) 500 microg (Not on the NSW State Formulary).
Dose	Concurrent therapy with dihydrofolate reductase inhibitors ^{1,2}
	10 mg three times per week .
	Folinic acid responsive seizures ^{3, 5}
	2.5 mg twice a day (doses up to 8 mg/kg/day have been used)
Dose adjustment	Therapeutic hypothermia – Not applicable.
-	ECMO – No information.
	Hepatic impairment – No dose adjustment.
	Renal impairment – No dose adjustment.
Maximum Daily	Not established.
Dose	
Route	Oral
Preparation	Liquid injection solution: ¹⁶⁻¹⁸
	Measure the dose and give undiluted orally.
	Folinic acid tablet:
	1. Dispense one tablet into 15 mL of water for injection to make a concentration of 1 mg/mL
	2. Shake well to ensure even dispersion
	3. Administer required dose immediately, discard any remaining solution
Administration	ORALLY or via gastric tube, ideally administer on an empty stomach (i.e. at least one hour before food or
•• •	two nours after food). ²²
Monitoring	No specific monitoring required.
Contraindications	Little information. Not effective in methylenetetrahydrotolate reductase deficiency.
Precautions Drug Internetions	Avoid use with folic acid antagonists unless under a specialist's advice."
Drug interactions	Antieplieptics – folinic acid may counteract the antieplieptic effect of phenobarbital (phenobarbitone),
	phenytolli, phimuone, and succimmus and increase the requency of seizures.
	Fluorouracii – foimic aciu may effiance the toxicity of hubrouracii.
	Folic acid antagonists – when folinic acid is given in conjunction with a folic acid antagonist (e.g.
	cotrimovazala, pyrimothamina) the officacy of the folic acid antagonist may either be reduced or
	cotrimoxazole, pyrimethamine) the efficacy of the folic acid antagonist may either be reduced or completely neutralised ⁶
	cotrimoxazole, pyrimethamine) the efficacy of the folic acid antagonist may either be reduced or completely neutralised. ⁶
	cotrimoxazole, pyrimethamine) the efficacy of the folic acid antagonist may either be reduced or completely neutralised. ⁶ Chloramphenicol - Concurrent administration of chloramphenicol and folinic acid in folate deficient patients may result in antagonism of haematopoietic response to folinic acid
Adverse	cotrimoxazole, pyrimethamine) the efficacy of the folic acid antagonist may either be reduced or completely neutralised. ⁶ Chloramphenicol - Concurrent administration of chloramphenicol and folinic acid in folate deficient patients may result in antagonism of haematopoietic response to folinic acid.
Adverse Reactions	cotrimoxazole, pyrimethamine) the efficacy of the folic acid antagonist may either be reduced or completely neutralised. ⁶ Chloramphenicol - Concurrent administration of chloramphenicol and folinic acid in folate deficient patients may result in antagonism of haematopoietic response to folinic acid. Allergic sensitisation, including anaphylactic reactions, and urticarial rash. ⁶ Nausea and vomiting with high doses
Adverse Reactions Overdose	cotrimoxazole, pyrimethamine) the efficacy of the folic acid antagonist may either be reduced or completely neutralised. ⁶ Chloramphenicol - Concurrent administration of chloramphenicol and folinic acid in folate deficient patients may result in antagonism of haematopoietic response to folinic acid. Allergic sensitisation, including anaphylactic reactions, and urticarial rash. ⁶ Nausea and vomiting with high doses.
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Adverse Reactions Overdose	cotrimoxazole, pyrimethamine) the efficacy of the folic acid antagonist may either be reduced or completely neutralised. ⁶ Chloramphenicol - Concurrent administration of chloramphenicol and folinic acid in folate deficient patients may result in antagonism of haematopoietic response to folinic acid. Allergic sensitisation, including anaphylactic reactions, and urticarial rash. ⁶ Nausea and vomiting with high doses. AUSTRALIA Contact the Poisons Information Centre on 13 11 26 for information on the management of overdose. NEW ZEALAND
Adverse Reactions Overdose	cotrimoxazole, pyrimethamine) the efficacy of the folic acid antagonist may either be reduced or completely neutralised. ⁶ Chloramphenicol - Concurrent administration of chloramphenicol and folinic acid in folate deficient patients may result in antagonism of haematopoietic response to folinic acid. Allergic sensitisation, including anaphylactic reactions, and urticarial rash. ⁶ Nausea and vomiting with high doses. AUSTRALIA Contact the Poisons Information Centre on 13 11 26 for information on the management of overdose. NEW ZEALAND Contact the National Poisons Centre on 0800 764 766 for information on the management of overdose.

Incompatibility	Not applicable.
Stability	Use solution prepared from tablets immediately. Discard remaining
Storage	Solution for injection
	Store at 2 to 8°C. (Refrigerate. Do not freeze). Protect from light.
	Tablets
	Store below 25°C.
Excipients	Solution for injection.
	DBL Brand: Sodium chloride and water for injections. Sodium hydroxide and/or hydrochloric acid (used to
	adjust pH of 300 mg/30 mL only).
	Pfizer Brand: Sodium chloride in water for injections.
	Tablets.
	Lactose monohydrate, microcrystalline cellulose, magnesium stearate.
Special	
Comments	
Evidence	Efficacy
	Concurrent therapy with dihydrofolate reductase inhibitors:
	Pyrimethamine/sulfadiazine: Current guidelines for treatment of the infant with congenital
	toxoplasmosis are for use of pyrimethamine and sulfadiazine plus folinic acid. ^{1, 2} Folinic acid 10 mg three
	times a week is recommended until 1 week following cessation of pyrimethamine treatment. It is advised
	not to use folic acid as a substitute for folinic acid $\frac{1}{2}$ levels of folinic acid in the cerebrospinal fluid (CSE)
	from folinic acid supplemented infants treated with pyrimethamine for congenital toxoplasmosis are
	thought to be too low to inhibit the effect of pyrimethamine. ⁷ However, there are no clinical trials
	comparing folate or folinic acid versus placebo in infants with toxoplasmosis.
	Methotrexate: Folate and folinic acid have a protective and probably similar effect against methotrexate-
	related adverse effects (including a reduction in gastrointestinal side effects, henatic dysfunction, and
	discontinuation of MTX treatment for any reason) in patients with inflammatory disease ^{8,9}
	Trimethonrim/sulfamethoxazole: There are no clinical trials comparing folate or folinic acid versus
	nlacebo in infants with treated with trimethonrim/sulfamethoxazole
	Folinic acid responsive seizures
	Folinic acid responsive seizures
	in the CSE. Genetic or autoimmune mechanisms cause cerebral folate deficiency and delayed treatment
	my lead to encephalonathy with severe learning disabilities. An EEG may show abnormal background
	activity with multifocal spike-wave complexes but typically has no diagnostic features. Neuroimaging
	results are also usually normal. Datients either do not respond to pyridovine at all or exhibit only a
	temporary improvement. However, such nationts show a marked neurological recovery including
	cossistion of solitures upon folinic acid treatment ^{3, 5,10} In infants, folinic acid responsive solitures typically
	ressation of seizures upon formic acid treatment. If infiniarits, formic acid responsive seizures typically
	present within days after birth as epileptic spashis – myocionic, absence, or generalized tonic cionic
	seizures. Identified gene abhormalities include ALDH/AL, SEC40AL, FOERL, MITHER, and MITHES. Folling
	with gradual increase of the doce over 14 menths to 45 mg twice a day $^{4.5}$ $^{10-15}$. Recommanded treatment
	includes initial treatment with folinic acid or E methyltatrahydrofolato 2. E ma/kg and long torm
	treatment with falinia acid or 5 methyltatrahydrofalata 2 5 mg/kg dailu iong-term
	Sofoty
	Salely Na paodiatria data ara availabla
Defenences	No paeulatific data are available.
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