ELLIOTTS B- sodium cation, sodium bicarbonate, anhydrous dextrose, magnesium sulfate, potassium chloride, calcium chloride, sodium phosphate injection Lukare Medical, LLC

ELLIOTTS B^{\circledR} SOLUTION (buffered intrathecal electrolyte/dextrose injection) $Rx\ Only$

DESCRIPTION

Elliotts $B^{\mathbb{B}}$ Solution is a sterile, nonpyrogenic, isotonic solution containing \underline{no} bacteriostatic preservatives. Elliotts B Solution is a diluent for intrathecal administration of methotrexate sodium and cytarabine.

Each 10 mL of Elliotts B Solution contains:

Sodium Chloride, USP	73 mg
Sodium Bicarbonate, USP	19 mg
Dextrose, USP	8 mg
Magnesium Sulfate • 7H ₂ O, USP	3 mg
Potassium Chloride, USP	3 mg
Calcium Chloride • 2H ₂ O, USP	2 mg
Sodium Phosphate, dibasic • 7H ₂ O, USP	2 mg
Water for Injection, USP	qs 10 mL

Concentration of Electrolytes:

Sodium	149 mEq/liter	Bicarbonate	22.6 mEq/liter
Potassium	4.0 mEq/liter	Chloride	132 mEq/liter
Calcium	2.7 mEq/liter	Sulfate	2.4 mEq/liter
Magnesium	2.4 mEq/liter	Phosphate	1.5 mEq/liter

The formulae and molecular weights of the ingredients are:

INGREDIENT	MOLECULAR FORMULA	MOLECULAR WEIGHT
Sodium Chloride	NaCl	58.44
Sodium Bicarbonate	NaHCO ₃	84.01
Dextrose	$C_6H_{12}O_6$	180.16
Magnesium Sulfate • 7H ₂ O	$Mg_2SO_4 \cdot 7H_2O$	246.48
Potassium Chloride	KCl	74.55
Calcium Chloride • 2H ₂ O	CaCl ₂ • 2H ₂ O	147.01
Sodium Phosphate, dibasic • 7H ₂ O	$Na_2HPO_4 \cdot 7H_2O$	268.07

The pH of Elliotts B Solution is 6.0-7.5, and the osmolarity is 288 mOsmol per liter (calculated).

CLINICAL PHARMACOLOGY

Elliotts B Solution provides a buffered salt solution for use as a diluent for the intrathecal administration of methotrexate sodium and cytarabine. It has been demonstrated that Elliotts B Solution is comparable to cerebrospinal fluid in pH, electrolyte composition, glucose content, and osmolarity:

Comparison of Electrolyte Composition, pH and Nonelectrolytic Constituents of Elliotts B Solution and CSF

Solution	Na ⁺ mEq/L	K ⁺ mEq/L	Ca ⁺⁺ mEq/L	0	HCO ₃ - mEq/L	Cl ⁻ mEq/L	pН	Phosphorus mg/dL	Glucose mg/dL
Cerebrospinal Fluid	117-137	2.3-4.6	2.2	2.2	22.9	113-127	7.31	1.2-2.1	45-80
Elliotts B Solution	149	4.0	2.7	2.4	22.6	132	6.0- 7.5	2.3	80

The approximate buffer capacity of Elliotts B Solution is 1.1×10^{-2} equivalents when the challenge solution is 0.01 N HCl and 7.8×10^{-3} equivalents when the challenge solution is 0.01 N NaOH.¹

Compatibility studies with methotrexate sodium and cytarabine indicate these drugs are physically compatible with Elliotts B Solution.

INDICATIONS AND USAGE

Elliotts B Solution is indicated as a diluent for the intrathecal administration of methotrexate sodium and cytarabine for the prevention or treatment of meningeal leukemia or lymphocytic lymphoma.

CONTRAINDICATIONS

None known.

WARNINGS

Intrathecal administration of drugs such as methotrexate sodium and cytarabine should be performed by personnel skilled in the technique of lumbar puncture under the supervision of a physician who is experienced in the use of cancer chemotherapeutic agents. The labeling for methotrexate sodium and cytarabine should be consulted.

PRECAUTIONS

General

Particular attention should be taken to assure the maintenance of sterile technique throughout the procedure. (See DOSAGE AND ADMINISTRATION.)

Carcinogenesis, Mutagenesis, Impairment of Fertility

No standard mutagenicity or carcinogenicity studies have been conducted with Elliotts B Solution.

Usage in Pregnancy

All components of Elliotts B Solution are normal body constituents. Animal reproduction studies have not been conducted with Elliotts B Solution.

ADVERSE REACTIONS

Adverse reactions may occur with any given intrathecal injection due to the chemotherapy or the technique of intrathecal administration. (See product labeling for methotrexate sodium and cytarabine.)

Preservative-free methotrexate sodium and cytarabine should be used to minimize adverse reactions due to preservatives.

If an adverse reaction does occur, discontinue the administration, evaluate the patient, institute appropriate therapeutic countermeasures and, if possible, save the remainder of the unused solution(s) for examination.

DRUG ABUSE AND DEPENDENCE

There is no potential for drug abuse or drug dependence.

OVERDOSAGE

Elliotts B Solution is a diluent. In the event of a drug, fluid or solute overload following administration, evaluate the patient's condition, and institute appropriate corrective treatment. (See product labeling for methotrexate sodium and cytarabine.)

DOSAGE AND ADMINISTRATION

See product labeling for methotrexate sodium and cytarabine.

Elliotts B Solution is intended for intrathecal administration only. Elliotts B Solution does not contain antibacterial preservatives and introduction of contaminated solutions into the cerebrospinal fluid may have extremely serious consequences. Therefore, administration of intrathecal solutions should be accomplished as soon as possible after preparation.

A sterile filter-needle should be used to withdraw the contents of the ampule.

Intrathecal drug products should be inspected visually for particulate matter and discoloration prior to administration.

Preparation and Administration Precautions

Elliotts B Solution is a diluent for the cytotoxic anticancer agents, methotrexate sodium and cytarabine. Care should be exercised in the handling and preparation of infusion solutions with these products. (See product labeling for methotrexate sodium and cytarabine.)

HOW SUPPLIED		
NDC	SIZE	

Elliotts B Solution is available in single-use clear glass ampules, packaged 10 ampules per box.

Store at controlled room temperature, 20°-25°C (68°-77°F) [See USP].

Preservative Free. Discard unused portion. Use only if solution is clear and ampule is intact.

Distributed by:

Lukare Medical, LLC Scotch Plains, NJ 07076 1-855-752-9317 www.elliottsbsolution.com

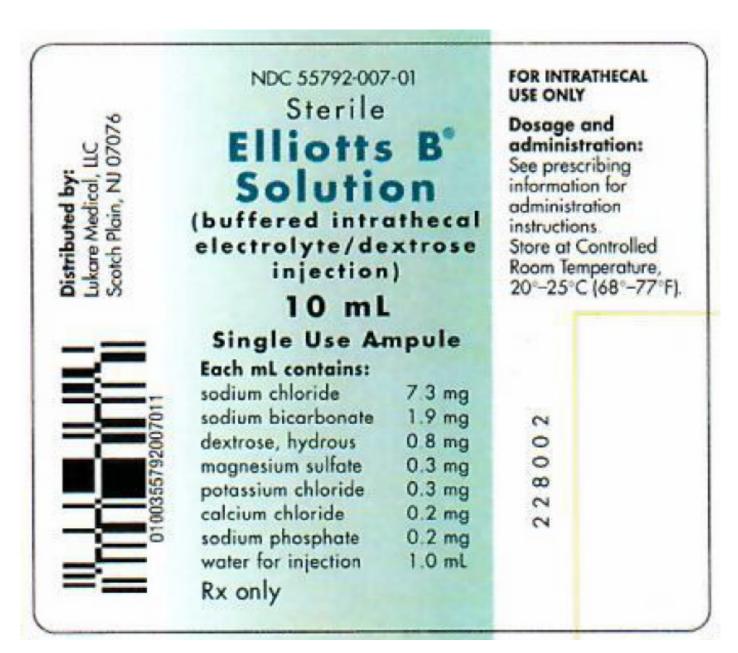
REFERENCES:

1. Cradock JC, et al. Evaluation of some pharmaceutical aspects of intrathecal methotrexate sodium, cytarabine and hydrocortisone sodium succinate. American Journal of Hospital Pharmacy (1978); 35:402.

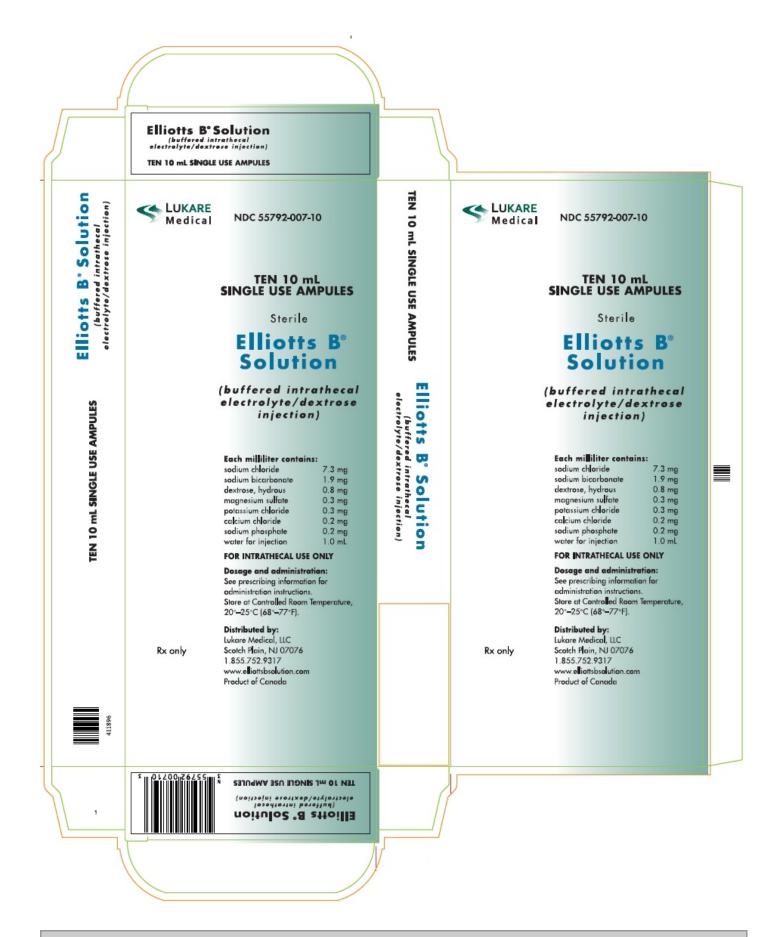
Rev. 10/2015

PACKAGING

Ampule labeling:



Box labeling:



ELLIOTTS B

sodium cation, sodium bicarbonate, anhydrous dextrose, magnesium sulfate, potassium chloride, calcium chloride, sodium phosphate injection

Product Information					
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:55792-007		
Route of Administration	INTRATHECAL				

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
$ \begin{array}{l} \textbf{so dium chloride} \ (UNII: 451W47IQ8X) \ (so dium \ cation - UNII:LYR4M0NH37, \ chloride \ ion - UNII:Q32ZN48698) \end{array} $	sodium chloride	73 mg in 10 mL			
sodium bicarbonate (UNII: 8MDF5V39QO) (bicarbonate ion - UNII:HN1ZRA3Q20)	sodium bicarbonate	19 mg in 10 mL			
anhydrous dextrose (UNII: 5SL0G7R0OK) (anhydrous dextrose - UNII:5SL0G7R0OK)	anhydrous dextrose	8 mg in 10 mL			
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB) (MAGNESIUM CATION - UNII: T6 V3LHY838)	MAGNESIUM SULFATE, UNSPECIFIED	3 mg in 10 mL			
potassium chloride (UNII: 660 YQ 98 I10) (potassium cation - UNII:295O53K152)	potassium chloride	3 mg in 10 mL			
calcium chloride (UNII: M4I0 D6 VV5M) (calcium cation - UNII:2M8 3C4R6 ZB)	calcium chloride	2 mg in 10 mL			
sodium phosphate (UNII: SE337SVY37) (phosphate ion - UNII:NK08V8K8HR)	sodium phosphate	2 mg in 10 mL			

Inactive Ingredients				
Ingredient N	Jame Strength			
Water (UNII: 059QF0KO0R)	10 mL in 10 mL			

Packaging						
# Item Code	Package Description	Marketing Start Date	Marketing End Date			
NDC:55792-007-10	10 in 1 BOX	06/04/2013				
NDC:55792-007-01	10 mL in 1 AMPULE; Type 0: Not a Combination Product					

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
NDA	NDA020577	06/04/2013			

Labeler - Lukare Medical, LLC (062862393)

Registrant - Lukare Medical, LLC (062862393)

Revised: 11/2018 Lukare Medical, LLC