



THOMSON REUTERS™

PRUSSIAN BLUE

MICROMEDEX® DRUGPOINTS SUMMARY

DrugPoints Summaries provide information on dosage, pharmacokinetics, cautions, interactions, clinical applications, adverse effects, comparative efficacy, and drug of choice information.

For more information about Thomson Reuters Micromedex, visit www.micromedex.com.

Information valid as of March 17, 2011.

The information contained in the Thomson Reuters (Healthcare) Inc. products is intended as an educational aid only. All Treatments or procedures are intended to serve as an information resource for physicians or other competent healthcare professionals performing the consultation or evaluation of patients and must be interpreted in view of all attendant circumstances, indications and contraindications.

The use of the Thomson Reuters (Healthcare) Inc. products is at your sole risk. These products are provided "AS IS" and "as available" for use, without warranties of any kind, either express or implied. Thomson Reuters (Healthcare) Inc. makes no representation or warranty as to the accuracy, reliability, timeliness, usefulness or completeness of any of the information contained in the products. Additionally, THOMSON REUTERS (HEALTHCARE) INC. MAKES NO REPRESENTATION OR WARRANTIES AS TO THE OPINIONS OR OTHER SERVICE OR DATA YOU MAY ACCESS, DOWNLOAD OR USE AS A RESULT OF USE OF THE THOMSON REUTERS (HEALTHCARE) INC. PRODUCTS. ALL IMPLIED WARRANTIES OF MERCHANTABILITY AND FITNESS FOR A PARTICULAR PURPOSE OR USE ARE HEREBY EXCLUDED. THOMSON REUTERS (HEALTHCARE) INC. DOES NOT ASSUME ANY RESPONSIBILITY OR RISK FOR YOUR USE OF THE THOMSON REUTERS (HEALTHCARE) INC. PRODUCTS."

Prussian blue

DrugPoint® Summary

DOSING & INDICATIONS

Adult Dosing

[View Detailed information in DRUGDEX](#)

- Toxic effect of cesium, Radioactive: 3 g ORALLY 3 times daily, minimum of 30 days; after internal radioactivity is substantially reduced, may reduce dose to 1 to 2 g ORALLY 3 times daily to improve gastrointestinal tolerance [1]
- Toxic effect of cesium, Radioactive: [1]
- Toxic effect of thallium: 3 g ORALLY 3 times daily [1]
- Toxic effect of thallium: [1]

Pediatric Dosing

[View Detailed information in DRUGDEX](#)

- Toxic effect of cesium, Radioactive: 12 y of age and older, 3 g ORALLY 3 times daily, minimum of 30 days; after internal radioactivity is substantially reduced, may reduce dose to 1 to 2 g ORALLY 3 times daily to improve gastrointestinal tolerance [1]
- Toxic effect of cesium, Radioactive: 2 to 12 y of age, 1 g ORALLY 3 times daily, minimum of 30 days [1]
- Toxic effect of thallium: 12 y of age and older, 3 g ORALLY 3 times daily [1]
- Toxic effect of thallium: 2 to 12 y of age, 1 g ORALLY 3 times daily [1]

FDA-Labeled Indications

[View Detailed information in DRUGDEX](#)

- Toxic effect of cesium, Radioactive
- Toxic effect of thallium

CONTRAINDICATIONS/WARNINGS

Contraindications

[View Detailed information in DRUGDEX](#)

- none

The information contained in the Thomson Reuters (Healthcare) Inc. products is intended as an educational aid only. All Treatments or procedures are intended to serve as an information resource for physicians or other competent healthcare professionals performing the consultation or evaluation of patients and must be interpreted in view of all attendant circumstances, indications and contraindications. Copyright © 2011 Thomson Reuters (Healthcare) Inc. All rights reserved. Information is for individual use only and may not be sold, redistributed or otherwise used for commercial purposes.

Precautions

[View Detailed information in DRUGDEX](#)

- impaired intestinal motility
- preexisting electrolyte imbalances

Pregnancy Category

[View Detailed information in DRUGDEX](#)

- [C\[14\]](#)(FDA)

Breast Feeding

[View Detailed information in DRUGDEX](#)

- [Thomson: Infant risk is minimal.](#)

ADVERSE EFFECTS

[View Detailed information in DRUGDEX](#)





Common




- **Endocrine metabolic:** Hypokalemia
- **Gastrointestinal:** Constipation, High-fiber laxatives and/or diet advised

IV COMPATIBILITY (SINGLE)

Solution

Common Solutions

Common Solutions	
D5W (D5W-Dextrose 5%)	 Not Tested
D10W (Dextrose 10%)	 Not Tested
D5LR (Dextrose 5% in lactated Ringers)	 Not Tested
D5NS (Dextrose 5% in sodium chloride 0.9%)	 Not Tested

D5W - 1/2 NS (Dextrose 5% in sodium chloride 0.45%)		Not Tested
NS (Normal saline- Sodium chloride 0.9%)		Not Tested
1/2 NS (Sodium chloride 0.45%)		Not Tested

Other Solutions

No other drug-solution combinations have been tested.

Y-Site

"Not Tested"

Admixture

"Not Tested"

Syringe

"Not Tested"

TPN/TNA

TPN (2-in-1)

There are no TPN Results.

TNA (3-in-1)

There are no TNA Results.

The information contained in the Thomson Reuters (Healthcare) Inc. products is intended as an educational aid only. All Treatments or procedures are intended to serve as an information resource for physicians or other competent healthcare professionals performing the consultation or evaluation of patients and must be interpreted in view of all attendant circumstances, indications and contraindications. Copyright © 2011 Thomson Reuters (Healthcare) Inc. All rights reserved. Information is for individual use only and may not be sold, redistributed or otherwise used for commercial purposes.

Definitions



Compatible IV compatibility is compatible.



Incompatible IV compatibility is incompatible.



Caution: Variable Caution: IV compatibility is variable.



Uncertain IV compatibility is uncertain.



Not Tested IV compatibility is not tested.

NAME INFO

US Trade Names

- Radiogardase

[All Trade Names](#)

Class

[View Detailed information in DRUGDEX](#)

- Pigmentation Agent

Regulatory Status

- RX

Generic Availability

- No

MECHANISM OF ACTION/PHARMACOKINETICS

Mechanism of Action

[View Detailed information in DRUGDEX](#)

- Oral-Local: Chelating agent—Prussian blue has a strong affinity for thallium and cesium [2] and binds with them in the intestine [2][3][4][5][6][7][8][9] to form a complex, [10][2][11][5] which is eliminated via the feces. [2][12][3][11][5]

Pharmacokinetics

[View Detailed information in DRUGDEX](#)

Absorption

- Oral-Local: Poorly absorbed [\[2\]\[12\]\[111\]\[4\]\[6\]\[7\]](#)

Metabolism

- Oral-Local: Metabolite: cyanoferrate 7% [\[11\]](#)

Excretion

- Oral-Local: Prussian blue—Fecal [\[10\]\[2\]\[12\]\[111\]\[13\]\[6\]](#)
- Cyanoferrate—Renal [\[11\]](#)

ADMINISTRATION/MONITORING

Administration

[View Detailed information in DRUGDEX](#)

Oral

- capsules may be opened and mixed with bland foods or liquids [\[1\]](#)
- administration with food is preferred to stimulate the excretion of cesium or thallium [\[1\]](#)

Monitoring

[View Detailed information in DRUGDEX](#)

- reduction in signs and symptoms of thallium/cesium toxicity and/or radiation poisoning
- CBC and electrolytes; weekly [\[1\]](#)
- constipation [\[1\]](#)
- systemic radiation levels
- radioactivity counts in urine and fecal samples; recorded weekly [\[1\]](#)

HOW SUPPLIED

Radiogardase

- Oral Capsule: 0.5 GM

CLINICAL TEACHING

- Advise patient that stools may be blue-tinted during therapy.
The information contained in the Thomson Reuters (Healthcare) Inc. products is intended as an educational aid only. All Treatments or procedures are intended to serve as an information resource for physicians or other competent healthcare professionals performing the consultation or evaluation of patients and must be interpreted in view of all attendant circumstances, indications and contraindications. Copyright © 2011 Thomson Reuters (Healthcare) Inc. All rights reserved. Information is for individual use only and may not be sold, redistributed or otherwise used for commercial purposes.

- Instruct patient on proper handling and disposal of bodily fluids/waste during drug therapy to minimize radiation exposure to others.
- Whenever possible, patient should use a toilet instead of a urinal. Flush several times after each use [1].
- This drug may cause constipation.
- Patient should take drug with food to stimulate the excretion of cesium or thallium.

REFERENCES

1. Product Information: RADIOGARDASE (TM) oral capsules, insoluble prussian blue capsules. HEYL Chemisch-pharmazeutische Fabrik GmbH & Co., Berlin, Germany, 2006.
2. Product Information: Antidotum Thallii-Heyl, Radiogardase-Cs. Heyl, Germany, 94.
3. Pearce J: Studies of any toxicological effects of prussian blue compounds in mammals-a review.. *Food Chem Toxicol* 1994; 32:577-82.
4. Ghezzi R & Marrubini M: Prussian blue in the treatment of thallium intoxication.. *Vet Hum Toxicol* 1979; 21 Suppl:64-6.
5. Pai V: Acute thallium poisoning. Prussian blue therapy in 9 cases.. *West Indian Med J* 1987; 36:256-8.
6. Kamerbeek HH, Rauws AG, Ham M, et al: Prussian blue in therapy of thallotoxicosis. An experimental and clinical investigation.. *Acta Med Scand* 1971; 189:321-4.
7. Farina R & Mello CE: Medical aspects of 137Cs decorporation: the Goiânia radiological accident.. *Health Phys* 1991; 60:63-6.
8. Gilman AG, Rall TW, & Nies AS: Goodman and Gilman's the pharmacological basis of therapeutics. 8th ed, Pergamon Press, New York, 1990, pp 1633.
9. Haddad LM & Winchester JF: Clinical management of poisoning and drug overdose. 2nd ed., W.B. Saunders Company, Philadelphia, 1990, pp 1032.
10. Reynolds JEF: Martindale, the extra pharmacopeia. 30th ed, The Pharmaceutical Press, London, 1993, pp 693.
11. Mulkey JP & Oehme FW: A review of thallium toxicity.. *Vet Hum Toxicol* 1993; 35:445-53.
12. Ellenhorn MJ, Barceloux DG, & 1988: Diagnosis and treatment of human poisoning. *Medical toxicology* 1988; 83:1060-2.
13. Stevens W, Peteghem C, Heyndrickx A, et al: Eleven cases of thallium intoxication treated with prussian blue.. *Int J Clin Pharmacol* 1974; 10:1-22.
14. Product Information: Radiogardase(TM), Insoluble Prussian blue capsules. Haupt Pharma Berlin GmbH, Berlin, Germany, 2003.

The information contained in the Thomson Reuters (Healthcare) Inc. products is intended as an educational aid only. All Treatments or procedures are intended to serve as an information resource for physicians or other competent healthcare professionals performing the consultation or evaluation of patients and must be interpreted in view of all attendant circumstances, indications and contraindications. Copyright © 2011 Thomson Reuters (Healthcare) Inc. All rights reserved. Information is for individual use only and may not be sold, redistributed or otherwise used for commercial purposes.