

Universitatea de Stat de Medicină și Farmacie "Nicolae Testemitanu, din Republica Moldova

Bazele culturii informatiei

Biblioteca Științifică Medicală a Universității de Stat de Medicină și Farmacie "Nicolae Testemitanu,



Medicina bazată pe dovezi

- Acesta este un domeniu de cunoaștere, care include rezultatele cercetării de încredere, experiența clinică a medicului și interesele pacientului.
- metode de medicina bazata pe dovezi + set de reguli de cercetare şi analiză ulterioară, care asigură fiabilitatea rezultatelor.



Aplicarea medicinei bazată pe dovezi necesită timp, competență și experiență, fiind necesară parcurgerea mai multor etape:

- transformarea problemei medicale in întrebare clinică,
- căutarea evidențelor relevante pentru răspunsul la întrebarea respectivă,
- •evaluarea validității (conformitatea cu adevărul), a impactului (mărimea efectului) și a relevanței (utilitatea) metodei,
- integrarea evidențelor medicale relevante în practică,
- revaluarea eficienței și eficacității acestora în teren.

Actualmente există o serie de facilități care permit aplicarea metodologiei medicale bazată pe evidențe, între care se pot menționa:

- •existența sistemelor informatice de asistare a deciziei medicale,
- •realizarea sintezelor informatice și a metaanalizelor,
- •apariția a numeroase publicații științifice de practică a medicinei bazatăpe dovezi (Evidence-Medicina bazată pe dovezi Based Medicine, Evidence-Based Healthcare& Public Health etc.),
- dezvoltarea informaticii medicale, crearea de strategii de educație medicală continuă.

4

Logistica medicinii bazată pe dovezi este structurată pe trei nivele:

Nivelul organizatoric. In anul 1979 Archie Cochrane critica societatea medicală pentru lipsa unei modalități prin care să se pună la dispoziția medicilor rezultatele cercetărilor recente, lipsa cărora poate conduce la aplicarea cu întârziere a acestora în practica medicală. Au luat naștere în acest sens, o serie de organizații:

Colaborarea Cochrane are scopul de a crea și de a aduce la zi sinteze sistematice cu ultimele actualități din diverse domenii medicale.



Această organizație este constituită dintr-un grup metodologic care creează metodologii și îndrumă organizația în îmbunătățirea validității și preciziei sintezelor sistematice, precum și o rețea internațională de centre naționale care promovează obiectivele colaborării la nivele naționale.

Colaborarea AGREE este o organizație internațională care se ocupă cu implementarea ghidurilor de practică medicală



Nivelul tehnologic și financiar este asigurat din granturi de cercetare, cheltuielile fiind necesare pentru crearea și menținerea la zi a evidențelor primare și secundare, implementarea și administrarea bazelor de date, accesul la bazele de date, instruirea personalului etc.

Resursele umane sunt reprezentate de *specialiști în domeniu* care au rolul de a elabora evidențele medicale; *informaticieni* care se ocupă de suportul informatic pentru crearea și managementul evidențelor, respectiv pentru decizia medicală asistată pe calculator; persoane cu rol în dezvoltarea de politici sanitare (normative și legislative);

medicii practicieni, care integrează evidențele valide în practica curentă, alături de experiența individuală.



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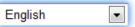


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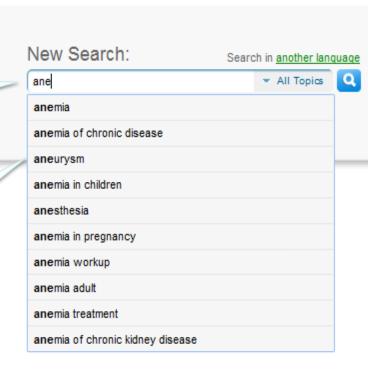




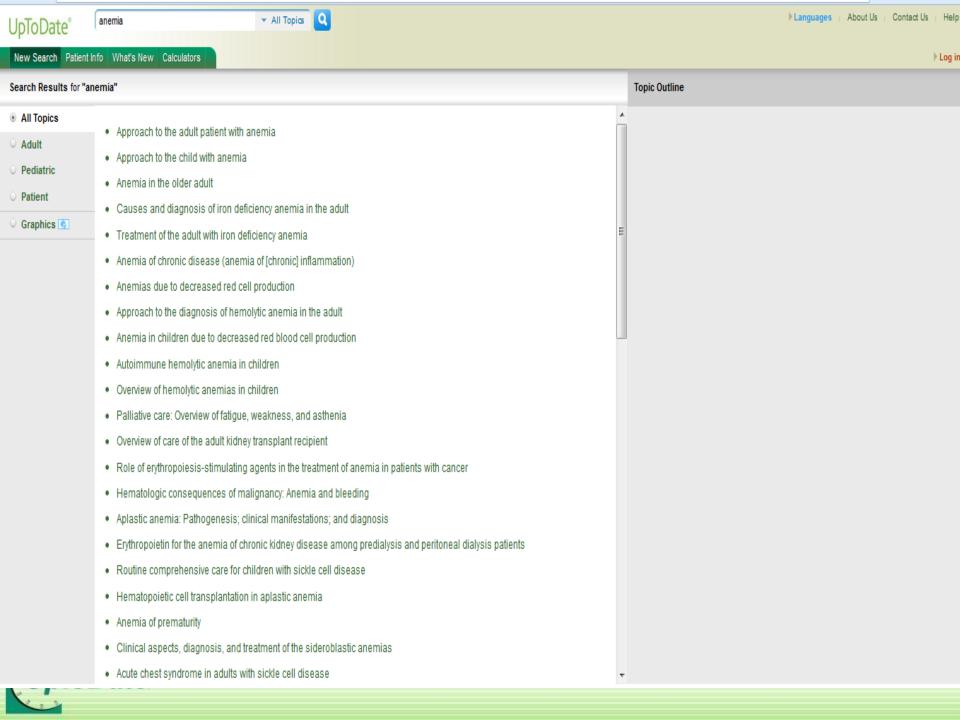


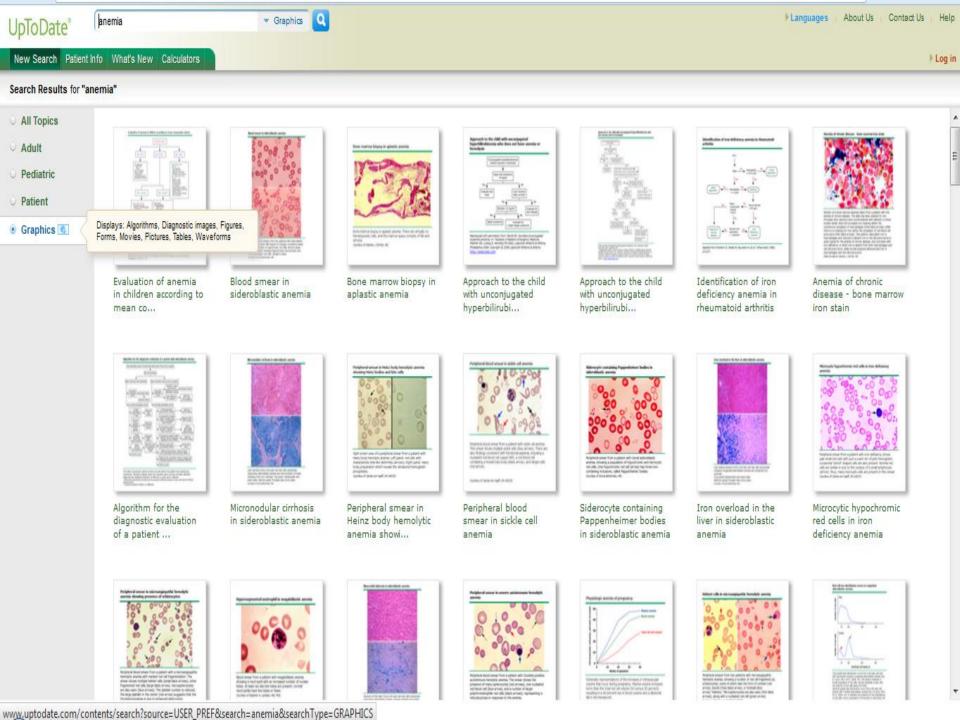
Introduceți cuvîntul de căutare în limba engleză (unul sau mai multe)

După ce ați întrodus trei simboluri, UpToDate afișează lista corespondențelor posibile a termenului.



Faceți clic pe elementul din lista de corespondențe posibile pentru rezultatele căutării a elementului necesar





anemia

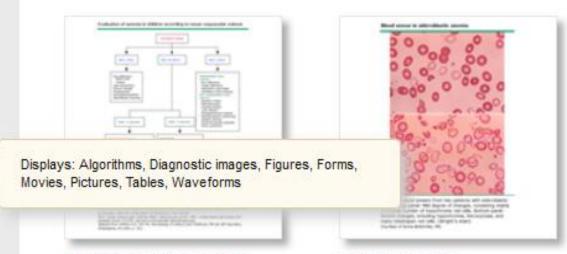




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Search Results for "anemia"

- O All Topics
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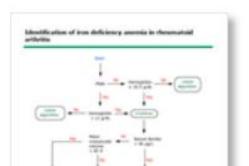


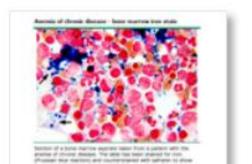
Evaluation of anemia in children according to mean co... Blood smear in sideroblastic anemia



Bank marrow biogray in aplantic promi

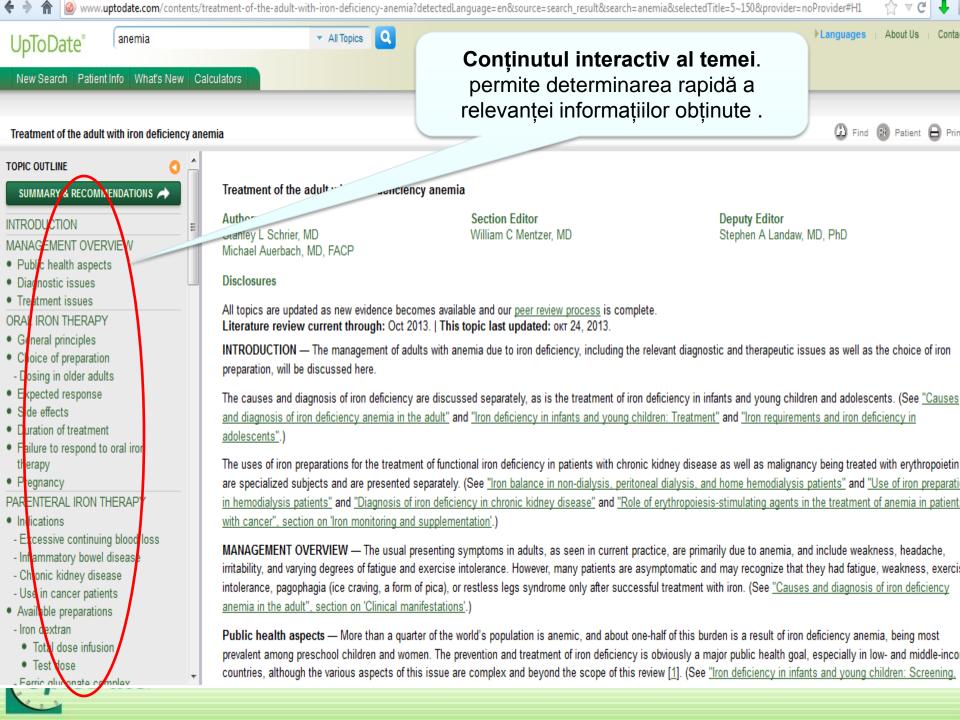
Bone marrow biop aplastic anemia













ailable and our peer review process is complete.

his topic last updated: окт 24, 2013.

anemia due to iron deficiency, including the relevant diagnostic and therapeutic issues as well as the choice of iron

cussed separately, as is the treatment of iron deficiency in infants and young children and adolescents. (See <u>"Causes"</u> and <u>"Iron deficiency in infants and young children: Treatment"</u> and <u>"Iron requirements and iron deficiency in</u>

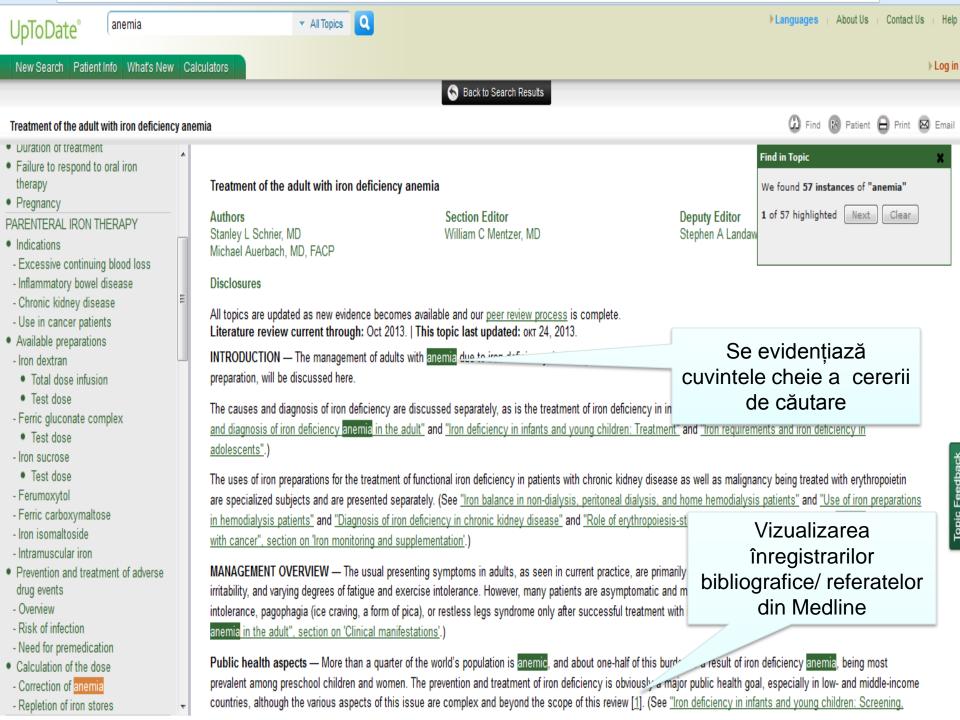
nctional iron deficiency in patients with chronic kidney disease as well as malignancy being treated with erythropoietin ely. (See "Iron balance in non-dialysis, peritoneal dialysis, and home hemodialysis patients" and "Use of iron preparations iciency in chronic kidney disease" and "Role of erythropoiesis-stimulating agents in the treatment of anemia in patients mentation.

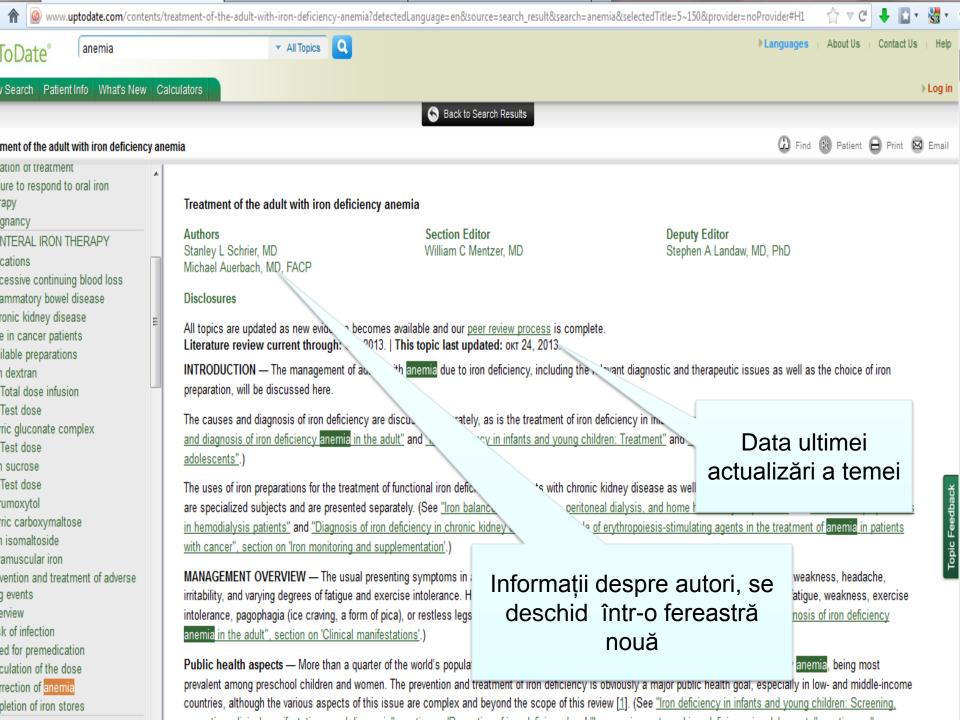
ng symptoms in adults, as seen in current practice, are primarily due to anemia, and include weakness, headache, se intolerance. However, many patients are asymptomatic and may recognize that they had fatigue, weakness, exercise

Căutarea în cadrul temei

Topic Feed







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Treatment of the adult with iron deficiency anemia

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INFORMATION FOR PATIENTS

NONANEMIC SUBJECTS

SUMMARY AND

RECOMMENDATIONS

- Indications for treatment
- Initial treatment
- · Resistant or relapsed disease

REFERENCES

TABLES

GRAPHICS View Ali-

Iron bioavailability

- · Failure to respond to iron
- Drug ratings in pregnancy
- IV iron preparations
- Parenteral iron dose

RELATED TOPICS

Allergic reactions to vaccines

Anemia of chronic disease (anemia of [chronic] inflammation)

Causes and diagnosis of iron deficiency anemia in the adult

Clinical manifestations of hereditary hemochromatosis

Diagnosis of iron deficiency in chronic kidney disease

Hereditary hemorrhagic telangiectasia (Osler-Weber-Rendu syndrome)

Prevention and treatment of adverse drug events — The following sections should be required reading for all clinicians contemplating the use of an intravenous iron preparation.

Overview — Each of the available formulations of IV iron has been associated with reports of life-threatening adverse drug events (ADEs) [39,60,61]. While the Food and Drug Administration's Adverse Event Reporting System (AERS) has been employed for comparing such rates [39.62], it is not possible to draw valid conclusions about comparative risks from the AERS database and other voluntarily submitted reports [60,63].

There exist only a small number of randomized, prospective trials comparing the relative safety of the various intravenous iron preparations [64]: three have shown relatively equal safety for low molecular weight iron dextran and iron sucrose [61,65,66]; one has shown relatively equal safety for iron sucrose and ferric gluconate complex [67]; while one, in abstract form, has shown comparate

Patients with a history of drug anergy equipment and personnel trained in the However, while we have seen such ana weight iron dextran preparations [59],

Systemic effects of IV iron includ.

benefit is obtained from the use of non-

-leader iron devtran preparations are

Tabelele și graficele pot fi vizualizate prin deschiderea lor de la text făcând clic pe linkurile din paranteze, precum și grafice în conținutul subiectelor.

- A flare of arthritis in patients with inflammatory arthritis, such as rheumatoid arthritis, commonly occurs. This can usually be abated by the use of 125 mg of intravenous methylprednisolone immediately before the iron infusion and 1 mg/kg per day of oral prednisone for four days thereafter. The cause of these reactions is unknown as tryptase levels, a marker of mast cell degranulation, have been normal following these reactions. (See "Allergic reactions to vaccines", section on 'Dextran' and "Perioperative anaphylaxis: Evaluation and prevention of recurrent reactions", section on 'Colloids'.)
- Approximately 0.5 to 1 percent of patients may complain of arthralgias, myalgias, or flushing without associated hypotension, tachypnea, tachycardia, wheezing, stridor, or periorbital edema [30,59]. After symptoms abate, rechallenge is appropriate and symptoms rarely recur. The empiric use of steroids prior to rechallenge may be beneficial, but is untested. In the rare patient who reacts in this manner twice, the IV formulation should be changed. No intervention for these complaints is necessary. Specifically, the use of antihistamines for prevention or treatment is inappropriate. (See 'Need for premedication' below.)

Risk of infection — Bacteria and other infectious agents require iron as a growth factor, and patients with hereditary hemochromatosis and iron overload are known to be

Vizualizarea tabelelor și a graficelor



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Radiation biodosimetry

Dose (Gy)	Vomiting (%)	Time to vomiting (hours)	ALC day 1 (/microL)	Lymphocyte fall rate constant (k)*	Lymphocyte dicentrics (per 1000)
0	0	-	2450	-	1-2
1	19	-	2160	0.126	88
2	35	4.6	1900	0.252	234
3	54	2.6	1680	0.378	439
4	72	1.7	1480	0.504	703
5	86	1.3	1310	0.63	1000
6	94	1.0	1150	0.756	
7	98	0.8	1010	0.881	
8	99	0.7	890	1.01	
9	100	0.6	790	1.13	
10	100	0.5	700	1.26	

Gy: absorbed whole body dose in Grey units; ALC: absolute lymphocyte count per microL.

Modified with permission from: Waselenko, JK, MacVittie, TJ, Blakely, WF, et al. Medical management of the acute radiation syndrome: Recommendations of the strategic national stockpile radiation working group. Ann Int Med 2004; 140:1039. Copyright @ 2004 American College of Physicians.

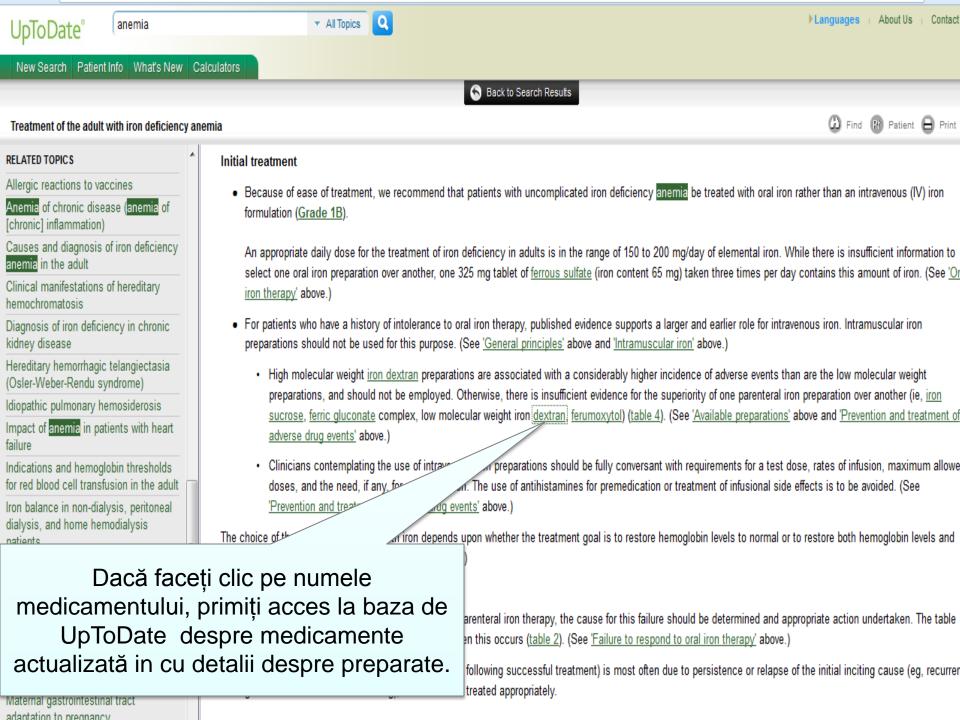


^{*} The lymphocyte fall rate constant is derived from a semilogarithmic plot of the absolute lymphocyte count (ALC) versus time in days, in the form of 2450 x e(-kt). The time (in days) for the ALC to fall to one-half of its original value [half-time, T(1/2)] can be obtained from the following equation: T(1/2) = 0.693/k.

concentration. Blood 2011: 118:3222

Maternal gastrointestinal tract

adaptation to pregnancy



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Dextran: Drug information

TOPIC OUTLINE 0 Brand Names: U.S.

Pharmacologic Category Dosing: Adult Dosing: Pediatric

Dosing: Geriatric Dosing: Renal Impairment

Dosing: Hepatic Impairment Dosage Forms: U.S.

Generic Equivalent Available: U.S. Administration Compatibility

Use Medication Safety Issues Adverse Reactions Significant

Contraindications Warnings/Precautions Metabolism/Transport Effects

Drug Interactions

Lactation

Topic Feedback

(For additional information see "Dextran: Pediatric drug information")

Treatment of shock or impending shock (when blood or blood products are not available): I.V.: Dextran 40: Infuse 10 mL/kg as rapidly as possible (maximum: 20 mL/kg/day for the first 24 hours; 10 mL/kg/day thereafter); therapy should not be continued beyond 5 days

Dosing: Geriatric Refer to adult dosing.

Dosing: Pediatric

Pregnancy Implications

Pregnancy Risk Factor

Breast-Feeding Considerations

Pricing: U.S. (Medi-Span®)

International Brand Names

Monitoring Parameters

Mechanism of Action

Volume expansion/shock: I.V.: Dextran 40: Infuse 500-1000 mL (~10 mL/kg) as rapidly as possible (maximum: 20 mL/kg/day for first 24 hours; 10 mL/kg/day thereafter);

Pump prime: Dextran 40: Varies with the volume of the pump oxygenator; generally, the solution is added in a dose of 10-20 mL/kg (or 1-2 g/kg); usual maximum total

Postoperative prophylaxis of venous thrombosis/pulmonary embolism (Dextran 40): Begin during surgical procedure and give 500-1000 mL (~10 mL/kg); an

additional 50 g (500 mL) should be administered every 2-3 days during the period of risk (up to 2 weeks postoperatively); usual maximum infusion rate for nonemergency

dose: 20 mL/kg (or 2 g/kg)

use: 4 mL/minute

Dosing: Adult

Dextran: Drug information Lexicomp®

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(For additional information see "Dextran: Pediatric drug information")

For abbreviations and symbols that may be used in Lexicomp (show table)

Brand Names: U.S. LMD in D5W: LMD in NaCl

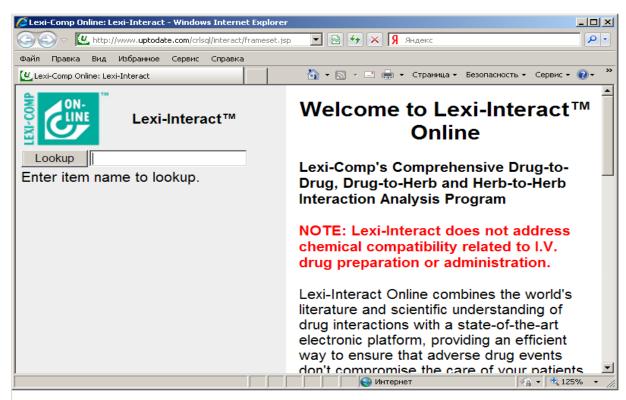
Pharmacologic Category Plasma Volume Expander, Colloid

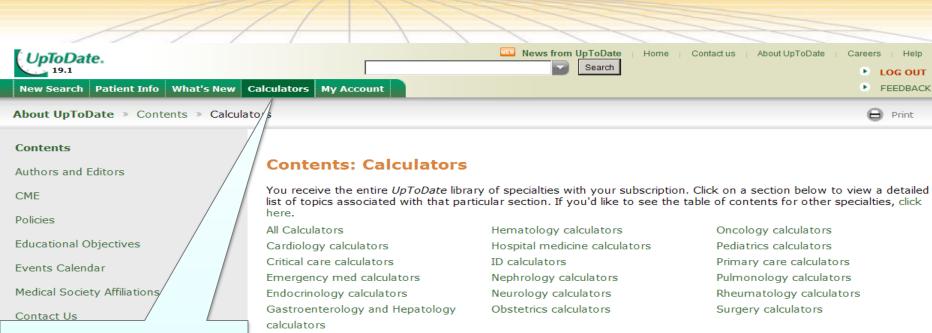
therapy should not be continued beyond 5 days



Informații despre medicamente. Baza de date Lexi-Interact







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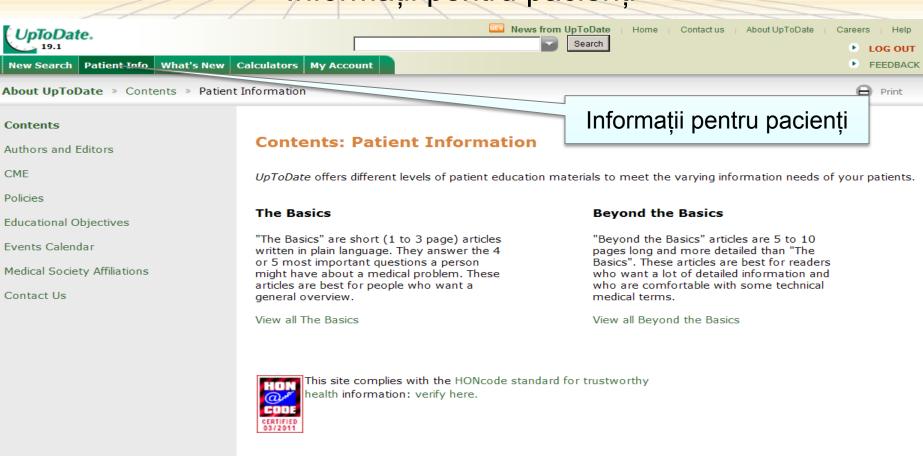
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