



IRON POISONING

An adolescent with a pmh of anemia and depression took multiple iron tablets. She was having abdominal pain but now appears to be resting comfortably in bed. You should:

A) Call psychiatry for a full psychiatric assessment due to the risk of future more serious ingestions

B) Ask her when, how many and what type of iron pills she took

C) Check her Total Iron Binding Capacity (TIBC) and Serum Iron Concentration (SIC) to determine if she has successfully sequestered the ingestion

D) Perform a stat abdominal xray to localize the ingestion



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- Management of the clinical condition is critical in Iron poisonings, however, there noted phases of illness:

Gastrointestinal phase: (30 minutes to 6 hours) abdominal pain, vomiting, diarrhea, hematemesis, melena, lethargy, shock (from capillary leak and third spacing), metabolic acidosis

Latent: (6 to 24 hours) improvement in GI symptoms; may have poor perfusion, tachypnea, tachycardia

Shock and metabolic acidosis: (4 hours to 4 days) hypovolemic, distributive, or cardiogenic shock with profound metabolic acidosis, coagulopathy, renal insufficiency/failure, pulmonary dysfunction/failure, central nervous system dysfunction

Hepatotoxicity: (within 2 days) coma, coagulopathy, jaundice. Severity is dose dependent.

Bowel obstruction: (2 to 4 weeks) vomiting, dehydration, abdominal pain, usually gastric outlet obstruction

- This patient may have entered the latent phase. Time course and dosage help determine future management considerations
- >40mg/kg of elemental iron ingestion is typically the threshold of concern in patients without symptoms.
 - Birth control placebo pills may contain high doses of elemental iron
 - Ferrous gluconate 12% elemental iron
 - Ferrous sulfate 20% elemental iron
 - Ferrous fumarate 33% elemental iron



- These patients may benefit from x ray and labs:
- Serum iron concentration: measure serum iron concentration within 4 to 6 hours after ingestion (8 hours for extended release tablets)
- Arterial or venous pH
- Abdominal radiograph looking for radiopaque pills
- Other initial labs: electrolytes, BUN, creatinine, glucose, liver function tests, prothrombin, partial thromboplastin time, CBC with differential, type and cross match



- A) will be important after management of acute intoxication
- C)TIBC measurements are unreliable in Iron overdose.
- D)Abdominal x-ray may be valuable in identifying iron in the digestive tract which may be removed. This will only partially determine management decisions and may be more critical during the GI phase. Liquid formulations may not appear on xray.





<http://pedclerk.bsd.uchicago.edu/page/iron-toxicity>



A 3 yr old whose mother recently gave birth found some of his mother's prenatal candies. He presents with abdominal pain, persistent vomiting, diarrhea, and his stools are dark grey, green and black. You should:

- A) Reassure parents that as he is vomiting and stooling he will expel the ingestion and clinically improve in a couple hours
- B) Perform a stat abdominal film to localize the ingestion
- C) Perform a stat gastric lavage as he is having acute GI toxicity
- D) Start IV deferoxamine 15 mg/kg/hr
- E) Perform a stat measurement of serum iron concentration to determine whether there may be systemic effects, if greater than 500 mcg/ml begin treatment



A 3 yr old whose mother recently gave birth found some of his mother's prenatal candies. He presented with abdominal pain, and dark grey, green and black stools. He continues to have persistent vomiting, diarrhea. You should:

A) Reassure parents that as he is vomiting and stooling he will expel the ingestion and clinically improve in a couple hours

B) Perform a stat abdominal film to localize the ingestion

C) Perform a stat gastric lavage as he is having acute GI toxicity

D) Start IV deferoxamine 15 mg/kg/hr

E) Perform a stat measurement of serum iron concentration to determine whether there may be systemic effects, if greater than 500 mcg/ml begin treatment



- Iron causes damage through free radical formation and lipid peroxidation
- A) symptoms during the initial phase of illness are reflective of mucosal damage caused by this process and do not necessarily reflect expulsion of toxin
- B) abdominal radiographs may guide removal of iron from the GI tract, however if significant symptoms are noted it is critical to begin Deferoxamine as end organ damage may be irreversible



- C) Gastric lavage may be used after radiographic visualization in patients with large ingestions who can tolerate large bore orogastric tube. There is however risk of aspiration. A less dangerous option is Whole Bowel Irrigation using polyethylene glycol which has been shown to have some efficacy in decreasing toxicity.



- D) start Deferox. In all cases contacting poison control for recommendations is important, however general guidelines are that Deferox 15 to 35 mg/kg/hr should be started if there is any of:
 - Severe symptoms (hypovolemia/shock, lethargy/coma, persistent vomiting or diarrhea)
 - Anion gap metabolic acidosis
 - Peak serum iron concentration >500mcg/dl
 - Significant number of pills on radiograph



- E) While serum iron is a part of the overdose evaluation, iron is freely absorbed by the liver and rapidly cleared from serum. After 4-6hrs levels may be falsely decreased while end organ damage may be occurring. Enteric-coated tabs may reach peak concentration later than 4-6 hrs thereby also resulting in a reduced value depending on timing of evaluation.



Acute iron intoxication: Rapid overview

To obtain emergent consultation with a medical toxicologist, call the United States Poison Control Network at 1-800-222-1222, or access the World Health Organization's list of international poison centers (www.who.int/gho/phe/chemical_safety/poisons_centres/en/index.html).

History

What form of iron was ingested?

Ferrous gluconate (12 percent elemental iron)

Ferrous sulfate (20 percent elemental iron)

Ferrous fumarate (33 percent elemental iron)

How many mg/kg of elemental iron was ingested?

When did the ingestion occur?

Clinical features: overlapping phases of clinical manifestations

Gastrointestinal phase: (30 minutes to 6 hours) abdominal pain, vomiting, diarrhea, hematemesis, melena, lethargy, shock (from capillary leak and third spacing), metabolic acidosis

Latent: (6 to 24 hours) improvement in GI symptoms; may have poor perfusion, tachypnea, tachycardia

Shock and metabolic acidosis: (4 hours to 4 days) hypovolemic, distributive, or cardiogenic shock with profound metabolic acidosis, coagulopathy, renal insufficiency/failure, pulmonary dysfunction/failure, central nervous system dysfunction

Hepatotoxicity: (within 2 days) coma, coagulopathy, jaundice. Severity is dose dependent.

Bowel obstruction: (2 to 4 weeks) vomiting, dehydration, abdominal pain, usually gastric outlet obstruction

Diagnostic evaluation: for all patients with systemic symptoms, those who have ingested >40 mg/kg of elemental iron, and those for whom the amount of elemental iron ingested is unknown

Serum iron concentration: measure serum iron concentration within 4 to 6 hours after ingestion (8 hours for extended release tablets)

Arterial or venous pH

Abdominal radiograph looking for radiopaque pills

Other initial labs: electrolytes, BUN, creatinine, glucose, liver function tests, prothrombin, partial thromboplastin time, CBC with differential, type and cross match

Management

Secure airway and breathing

Treat volume depletion aggressively with isotonic infusion

Whole bowel irrigation: for all patients with a significant number of pills in stomach and small intestine on radiograph

Deferoxamine: continuous IV infusion (can cause hypotension). Begin at 15 mg/kg/hour. May increase to 35 mg/kg/hour during first 24 hours for severe ingestions. A toxicologist and/or regional poison control center should be consulted to determine the optimum dose of deferoxamine and duration of therapy. Treat in the following circumstances:

Severe symptoms: altered mental status, hemodynamic instability, persistent vomiting, diarrhea

Anion gap metabolic acidosis

Serum iron concentration >500 mcg/dL

Significant number of pills on x-ray



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