

عوارض ناشی از آهن زدایی و شیوه ی مدیریت آن نرگس بیگم میربهبهانی فوق تخصص هماتولوژی آنکولوژی کودکان استاد دانشگاه علوم پزشکی گلستان





* دفروکسامین(DFO)

* دفريپرون(DFP)

* دفرازیراکس(DFX)

Deferoxamine (DFO)

30 to 60 mg per kilogram of body weight per day, given over a period of 8 to 12 hours for 5 to 7 days per week

Common adverse events

lack of adherence to the treatment regimen

TIF2014

Defrasirox has been licensed as first-line monotherapy for thalassaemia major in over 100 countries worldwide, although the earliest age at which deferasirox qualifies as first-line treatment differs somewhat between the FDA and the EMEA

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Initiate Iron Chelation 2–6 yr of age: DFO (first-line therapy), DFX (first-line in U.S., second-line in E.U.) >6 yr of age: DFO (first-line), DFP (second-line), DFX (first-line), combination

NTDT

- Starting dose : DT 10 mg/kg/day
- FCT 7 mg/kg/day
- Maximum dose :DT 20mg/kg/day
- FCT 14 mg/kg/day
- Monitor iron overload:serum ferritin every 3 mo,LIC every 6-12 mo to tailor dose.
- Monitor and manage adverse events per local prescribing information

Deferoxamine (DFO) *ocular and auditory symptoms, * bone-growth retardation, *local reactions, *allergy

Defroxamine =DFO Ferritin Regularly (1-2 time yearly) Audiometry yearly Electroretinography yearly Regular monitoring of growth(children) every 3 month Bun, creatinin every 6 month U/A :every 3-6 month Alt, Ast: monthly (3m), then every 6 month

Defroxamine:

Local skin reaction : Adequate dilution(Not exceed10%) Ulceration Attention to sc (Not intradermal)

The deferiprone (DFP) given orally three times a day 75 to 100 mg per kilogram per day.

- * Common adverse events
- gastrointestinal symptoms,
- * arthralgia,
- * agranulocytosis,
- * neutropenia

Deferiprone=DFP CBC (weekly to monthly) ANC <1500/ml : interrupted treatment ANC<500/ml (Agranulocy tosis): stopped treatment ALT,AST : monthly (6month)then every 6 month Creatinin : every 6 month U/A : every 3 month Electroretinography : yearly Audiometry : yearly

Monitor and manage adverse

events:

per local prescribing information

Defriprone

GI symptoms : Nausea , vomiting , gasteric irritation change in appetite(loss or gain) Management : liquid preparation Arthropathy : Reduction of dose NSAID Stopped treatment Zinc defficiency :zinc supplementation Deferasirox (DFX)

given orally once a day 20 to 40 mg per kilogram per day (dispersible tablet [DT]) or 14 to 28 mg per kilogram per day (film-coated tablet [FCT]) Common adverse events *gastrointestinal symptoms, * increased creatinine levels *increased hepatic enzyme levels.

Defersirox= DFX

Creatinine weekly (4w), then monthly

Ferritin monthly <500u g /L : interrupting dosing

U/A (Proteinuria) monthly to every 3 month

Alt, Ast every 2 weeks or monthly (3 month)

then every 6 months.

LFT>5 fold : interrupting dosing

CBC monthly

Monitor and manage adverse

events:

per local prescribing information

Defersirox= DFX MRI or Echo T2 < 6ms : interrupted LVEF <55% ; interruptd Audiometry : yearly Electroretionyaxply : yearly



Defersirox= DFX

Skin rashes: if mod to severe : stopped and later restarted at low dose GI effect : diarrhoea , abdominal pain ,nausea , vomiting Management : adminstration in the evening or after food , film coated

- *Oral chelators have an established advantage over deferoxamine with respect to adherence to the treatment regimen,
- * New film-coated deferasirox tablet is associated with improved patient-reported outcomes, as compared with the dispersible form.

The choice of iron chelator should be based on

- *local guidelines,
- *clinical judgment,
- *Individual patient's iron overload profile

Successful treatment depends on

- * Dose adjustment according to ongoing iron intake,
- *Monitoring, attention to adherence issues,
- *Management of adverse events.

Continuous parenteral deferoxamine remains the first choice for patients who already have cardiac dysfunction,

and data on the benefit of deferoxamine combined with deferiprone are also available

