

Darbepoetin treatment algorithm in anemia of chronic renal failure with Hb < 11 g/dl

EVALUATE ETIOLOGY OF ANEMIA/TREAT CONDITION

- Check iron studies¹; supplement orally or IV as indicated
- Check guaiac stool for occult bleeding
- Check folic acid and B₁₂ level if MCV is increased

Initiate darbepoetin therapy

*If financial assistance is needed call:
Reimbursement Connection 1-800-273-9376*

Epoetin alfa patients *converting* to darbepoetin alfa

Patients *new* to erythropoietic therapy

Epoetin alfa (Total 2 wk dose)	darbepoetin alfa (Q2W dose, s.c.)
5,000-11,999 units	25 µg
12,000-19,999 units	40 µg
>20,000 units	60 µg

**Target Hb
not to exceed
13 g/dl**

Patient weight	darbepoetin alfa (Q2W dose s.c.)
< 60 kg	25 µg
61 – 80 kg	40 µg
> 80 kg	60 µg

If Hb increases
< 1.0 g/dl in 4 wks

Hb target
achieved

If Hb increases > 1.0 g/dl in 2 wks or if Hb
is increasing and approaching 13 g/dl

If Hb >13 g/dl

Increase dose² to the
next higher vial
strength⁺⁺ and evaluate
for hyporesponsive
factors³

Maintain/adjust dose
to maintain target Hb

Increase dosing interval
to every 3-4 wks

Hold until Hb < 13
g/dl and start at next
lower vial strength⁺⁺

⁺⁺Vial strengths:

- 25 µg
- 40 µg
- 60 µg
- 100 µg

Footnotes:

1. Monitor iron studies every 2 months
2. If Hb increased < 1.0 g/dl in 4 wks and iron stores are adequate.
3. Darbepoetin alfa dose increases should not be made more frequently than once a month.