DOSAGE ADJUSTMENT OF EPOETIN β AND DARBEPOETIN α IN CHRONIC KIDNEY DISEASE.

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OBJECTIVES

Erythropoietin (EPO) is indicated in treatment of anaemia associated with chronic kidney disease (CKD). Determination of epoetin β and darbepoetin α average dose required to achieve adequate levels of hemoglobin (Hb) in patients with CKD not yet undergoing dialysis and rate conversion factor between both EPOs.

STUDY DESIGN

Retrospective study.

Patients included: stage 3 to 5 CKD who started treatment with EPO between January 2012-December 2012.

Follow-up period: 6 months.

Hb target range: 10.0 g/dL-12.5 g/dL.

Data collected: sex, age, CKD stage; baseline, 3 and 6 months data analysis; type of EPO and posology.

Data: medical and pharmacotherapeutic history (Farmatools®).

Patients included (n = 81)] [Type of EPO and dosage		
Sex		59.3%			Average weekly dose	
Age		73.8 years (30-88)				Average weekly uose
Stage	3 CKD	29.7%		Epoetin ß	40.7%	7.718,18±6.155,72UI 20.55±10.30μg
	4 CKD	57.8%]	Darbepoetin α	59.3%	
	5 CKD	12.5%				
Basal Hb		10.13±1.16 g/dL		Patients who changed EPO treatment	46.9%	
Serum ferritin values>100 ng/mL		63.0%	1	treatment		

Hb and Hematocrit levels									
Average increase	After 3 r	nonths	After 6 months						
	Average increase	р	Average increase	р					
Hb level	1.52 g/dL	<0.001	1.60 g/dL	<0.001					
Hematocrit level	4.82 g/dL	<0.001	5.08 g/dL	<0.001					

After 3 months, 53.1% of patients had Hb 10.0-12.5 g/dL. Average weekly dose to achieve Hb target range: 6.875,0 UI of epoetin β and 20.4 μ g of darbepoetin α , which represent a relationship between both doses of EPOs: 337:1.

CONCLUSIONS

EPO increase statistically significantly Hb and hematocrit baseline levels after 3 and 6 months of treatment. The relationship between both doses of EPOs to achieve Hb target range (epoetin β : darbepoetin α) found in our study is different from the relationship described in the Summary of Product Characteristics (337:1 vs 200:1 respectively).





RESULTS