Anemia in CKD not On HD

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Intruduction

- Anemia in CKD is associated with decreased quality of life
- Increased risk of cardiovascular disease and mortality

limori S, Naito S, Noda Y et al Nephrology 2015

Mechanisms of CKD-related anemia

- Including relative erythropoietin deficiency
- Decreased red cell life span
- Abnormal iron metabolism
- Chronic inflammation
- Metabolic abnormalities
- Effects of medications such as renin angiotensin system inhibitors

Babitt JL, Lin HY. Mechanisms of anemia in CKD. J Am Soc Nephrol 2012;

Clinical Kidney Journal, 2020, vol. 13, no. 4, 613–624

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Anemia and iron deficiency among chronic kidney disease Stages 3–5ND patients in the Chronic Kidney Disease Outcomes and Practice Patterns Study: often unmeasured, variably treated

CKDopps (Chronic Kidney Disease Outcomes and Practice Patterns Study)

- International prospective cohort study
- International variation in anemia assessment and management practices in CKD
- CKD clinics in Brazil, France, Germany and the USA.
- From 1 January 2013 until 13 April 2018

CKDopps (Chronic Kidney Disease Outcomes and Practice Patterns Study)

- The study sample was comprised of 6766 CKD Stages 3–5ND
- patients from 135 CKD clinics across the four countries
- Nondialysis patients with eGFR <60 mL/min/1.73 m2
- The hemoglobin measurement
- Iron status was assessed
- Anemia treatment with erythropoiesis stimulating agents





FIGURE 3: Hemoglobin distribution, by country and CKD stage. Patients could contribute once for each CKD stage experienced during the study for (a). First hemoglobin (Hgb) for each stage was taken if there were multiple measures.

(a) by CKD stage



(b) by first hemoglobin level



Conclusion

- Anemia management in patients with nondialysis CKD treated in nephrology clinics identified substantial clinic- and country-level variation in prevalence of anemia and iron deficiency, frequency of anemia Monitoring and treatment
- High proportion of patients without measurement of iron parameters
- IDA who are not treated with iron or ESA
- Especially in the USA and Brazil.
- Anemia monitoring and treatment are a ripe area for quality improvement in CKD care.

Original Report: Patient-Oriented, Translational Research



Am J Nephrol 2018;47:333-342 DOI: 10.1159/000489223 Received: February 8, 2018 Accepted: April 10, 2018 Published online: May 18, 2018

Association of Pre-End-Stage Renal Disease Hemoglobin with Early Dialysis Outcomes

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INTRUDUCTION

- Evaluation the impact of pre-ESRD hemoglobin (Hgb) and pre-ESRD Hgb slope on post-ESRD mortality and hospitalization outcomes
- Anemia management may impact early post-ESRD outcomes
- The study included 31,472
- 12-month post-ESRD all-cause and cardiovascular mortality and hospitalization







Fig. 2. Association of 6-month pre-end stage renal disease (ESRD) hemoglobin (Hgb) with 12-month hospitalization count in 31,472 veterans transitioning to ESRD. IRR, incidence rate ratio.

Result and coclosion

• Similarly, Hgb exhibited a U-shaped association with CV mortality

• Only lower Hgb was associated with a higher hospitalization rate

• Conclusions: Lower and higher pre-ESRD Hgb levels are associated with a higher risk of early post-ESRD mortality

DEFINITION of anemia

Anemia is defined by the World Health Organization (WHO)

1. Hb <1 3.0 g/dL for adult males and postmenopausal women

2. Hb <1 2.0 g/dL for premenopausal women

EPIDEMIOLOGY

- National Health and Nutrition Examination Survey (NHANES)
- The prevalence of anemia increased
 - 1 percent among patients with an estimated GFR (eGFR) of 60
 - 9 percent at an eGFR of 30 mL/min
 - 33 to 67 percent at an eGFR of 1 5 mL/min

Astor BC, Muntner P, Levin A, et al. Arch Intern Med 2002

The initial evaluation

- CBC
- RBC indices
- Reticulocyte count
- Serum iron
- Total iron-binding capacity (TIBC)
- Percent transferrin saturation (TSAT)
- Serum ferritin
- Serum folate and vitamin B1 2 levels
- Testing for occult blood in stool

Monitoring

- Patients without anemia
 - eGFR) ≥45 mL/min monitor annually
 - eGFR <45 mL/min/1.73 m least twice yearly
- Patients with anemia but not on an ESA
 - eGFR) ≥45 mL/min every 6 m if anemia is progressive every 3 m
 - eGFR <45 mL/min/1.73 m every 3 m

Monitoring

- Patients treated with an ESA
- On starting ESA or increasing dose
 - Measure the Hb every two to four weeks
 - Every three months thereafter
- Patients treated with iron
 - Iron assessment status every three months

TREATMENT

1. Iron

- 2. ESA
- 3. Rarely RBC transfusions

Erythropoiesis-stimulating agent

- Hb <1 0 g/dL
- TSAT is >25 percent
- Ferritin >200
- Dosing
 - The initial epoetin dose = 50 to 1 00 units/kg/week
 - 4000 or 1 0,000 units subcutaneously once weekly
 - 10,000 to 20,000 units subcutaneously every other week

Target hemoglobin value

- Nondialysis CKD patients Hb levels between 10 and 11.5 g/dL
- Do not target Hb concentration >1 3 g/dL

Thanks for your attention