A real-world experience of Desidustat in Chronic Kidney Disease-anemia patients: A single centre experience

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Introduction:

Desidustat, an oral HIF-PHI agent which helps in endogenous erythropoietin production approved for management anemia associated with CKD.

■This study was aimed to see the effects of Desidustat (Oxemia[™]) therapy in real-world setup.

Material and Method:

A real world, single centered, retrospective study

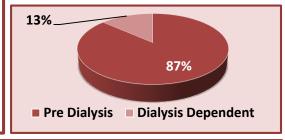
■52 Patients with CKD anemia, with or without erythropoietin therapy were given Desidustat (Oxemia[™]).

■As part of CKD Anemia treatment, patients were also receiving IV Iron (n=20, 38.46%), and Erythropoietin (n=13, 25%).

Monthly follow up was done to track the Hb changes.

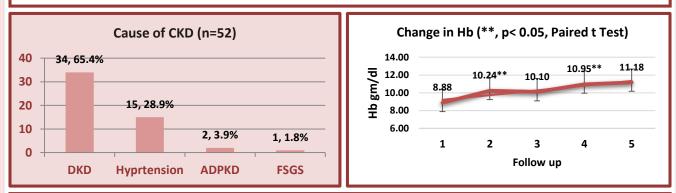
Dose adjustment was done according to the Hb level

Desidustat Dose (thrice weekly dosing)	Subjects (n, %)
50 mg	10, 19.23%
100 mg	41, 78.85%
150 mg	1, 1.92 %



Result:

- Study shown significant and stable improvement in Hb level post baseline 1.19 gm/dl (p<0.05, paired t Test), with Desidustat.
- •Majority of patients remain stable with 100 mg dose, 17.3% required dose adjustments.
- •Tolerability to the Desidustat therapy is very good (n=49, 94.23%),
- No adverse events observed, except 1 had constipation and throat pain.
- Lack of rigorous follow-up is a limitation of this study.



Conclusion:

Desidustat significantly increase the Hb in CKD Anemia without causing severe adverse drug reaction.
More studies are required assess the long term efficacy and safety of Desidustat in DD- CKD patients.

References:

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