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ORAL PRESENTATION

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Molecular aspects of sGC regulation

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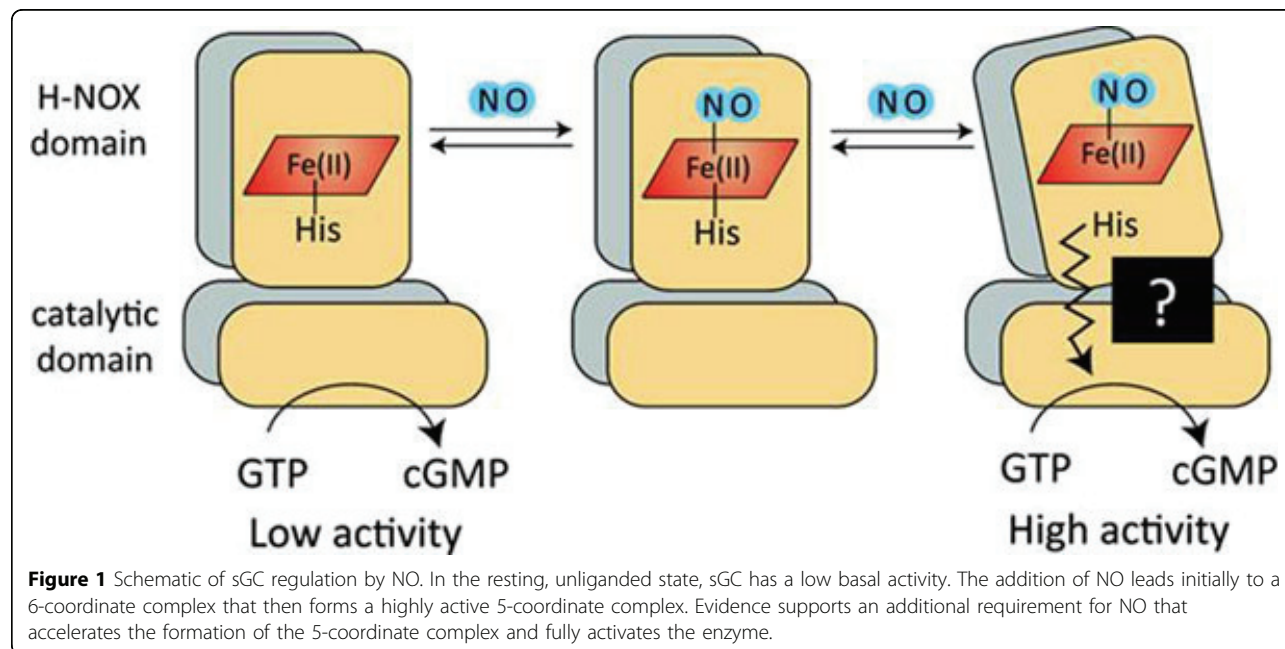
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Mammalian sGC is a heterodimer composed of α - and β -subunits (Figure 1) [1]. The C-terminus of each subunit contains a catalytic domain, and the active site is composed of residues from both subunits. Sequence analysis shows that each subunit also contains a well-defined PAS-like domain, and a predicted helical region. The N-termini of the α - and β -subunits are homologous to the H-NOX (Heme-Nitric oxide/Oxygen) family of proteins. The N-terminus of β -subunit contains a ferrous heme cofactor that serves a receptor for NO.

Ferric heme oxidized sGC has low activity, and the NO complex of the re-reduced heme generates a

desensitized, low-activity state of sGC. The molecular mechanism for this desensitization involves site specific S-nitrosation. The conformational changes associated with activation are both subtle and complex. Hydrogen-deuterium exchange mass spectrometry analysis can be used to probe conformational changes and protein-protein interactions. This method has been brought to bear on sGC, illuminating domain interactions within sGC and conformational changes induced by NO binding.

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Reference

1. Derbyshire ER, Marletta MA: Biochemistry of soluble guanylate cyclase. *Handb Exp Pharmacol* 2009, **191**:17-31.

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