



Thioredoxin Reductase: Signaling Pathways and Pharmacological Targets

Guest Editor:

Prof. Dr. Junmin Zhang

School of Pharmacy, State Key
Laboratory of Applied Organic
Chemistry, College of Chemistry
and Chemical Engineering,
Lanzhou University, Lanzhou
730000, China

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Message from the Guest Editor

Dear Colleagues,

Human thioredoxin reductase (TrxR) is a selenoprotein that utilizes highly reactive selenocysteine (Sec) residues in its active site. The pharmacological regulation of TrxR is typically presented in light of interactions with Sec residues and a few other mechanisms of action. The small-molecule-mediated regulation of TrxR signaling pathways has been investigated for its clinical significance and translational medicine in a variety of diseases. For example, the remarkable role of small molecules targeting the TrxR signaling pathway in anticancer and anti-infection, rheumatoid arthritis, and ischemia therapies has been explored in detail. Nonetheless, the rationale of the TrxR signaling pathway in therapeutic strategies for human diseases, the underlying molecular basis of small molecules targeting TrxR for therapeutic purposes, the downstream mechanisms of effect that may mediate therapeutic efficacy, and the diversity of therapeutic molecule structures and druggability are all topics that we will focus on in this Special Issue.

Keywords:

- thioredoxin
- thioredoxin reductase
- redox homeostasis
- redox regulation





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Prof. Dr. Maurizio Battino

Department of
Odontostomatologic and
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Sez-Biochimica, Faculty of
Medicine, Università Politecnica
delle Marche, Via Ranieri 65,
60100 Ancona, Italy

Message from the Editor-in-Chief

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MDPI, St. Alban-Anlage 66
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