

Influence of transition metals on stability of various s-nitrosothiols

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The analysis of the destroying action of copper and iron ions on S-nitrosocysteine (cys-NO) demonstrated that the amount of intrinsic copper in the solutions was too low ($0.2 \mu\text{M}$) to produce a destabilizing effect on cys-NO. Only admixed iron present in the solutions at the concentrations of about $1\text{--}2 \mu\text{M}$ was responsible for the process of cys-NO degradation, which occurred through the formation of dinitrosyl iron complexes (DNIC) resulting in free NO liberation. The conclusion was drawn from demonstration of the capacity of copper/iron chelator, bathophenanthroline disulfonic acid to selectively block the destructive effect of iron, but not of copper ions on cys-NO. Similar results were obtained in relation with the influence of intrinsic copper/iron on vasorelaxant properties of cys-NO on isolated rat aorta. By contrast, the concentrations of both intrinsic copper and iron ions in the solutions were proved to be not large enough to initiate the decomposition of S-nitrosoglutathione. The process was initiated when copper/iron ions were added to the solution at the concentration of $100 \mu\text{M}$ in the presence of reducing agents, ascorbate or GSH. S-dinitrosodithiotreitol (DTT-2NO) was decomposed spontaneously without any effect of intrinsic copper/iron ions in the solution. Nevertheless, the addition of copper/iron ($20 \mu\text{M}$) in the presence of ascorbate (1 mM) also accelerated the process of DTT-2NO decomposition. Such type of enhancing effect of copper/iron ions was suggested to be characteristics of enzymatic manner of RS-NO decomposition. The process of GS-NO or DTT-2NO decomposition catalyzed by iron ions ($100 \mu\text{M}$) in the presence of GSH or DTT was associated with formation of DNIC with GSH or DNIC with DTT, respectively. These results allow to propose that the process of GS-NO and DTT-2NO decomposition catalyzed by iron ions proceeded, as with cys-NO, through the formation of DNIC. The subsequent decomposition of DNIC ensures appearance of free NO molecules in the solution.