Novel Oxidative stress responses Biosensor: sodA::luxCDABE fusion Escherichia coli

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Abstract

A recombinant bioluminescent *Escherichia coli* strain, EBHJ, (*sodA::luxCDABE*), containing the promoter for the manganese superoxide dismutase (*sodA*) gene fused to the *Vibrio fischeri luxCDABE* operon, was successfully constructed and characterized. Redox-cycling agents, such as paraquat and chromium, strongly induced a *sodA*- regulated response in dose-dependent manners, resulting in an increase of the bioluminescence. In a comparison with an existing oxidative stress responsive strain, DPD2511 (*katG::luxCDABE*), which is sensitive to H 2O2, the mechanism of chemicals that cause oxidative damage was elucidated via the key transcriptional factors involved in induction of the *sodA* and *katG* promoters, i.e. SoxRS and OxyR, respectively. It was found that responses from the *katG*- and *sodA*-based strains were significantly different dependent upon the chemicals being tested. Therefore, EBHJ, alone or in parallel with DPD2511, can be used to characterize and monitor chemicals that cause oxidative damage.

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