Cell-based bioassay for compounds with prooxidant activity

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Abstract

There is substantial evidence from epidemiological studies that the etiology of many ‘modern’ diseases is linked in part to environmental pollution. A number of observations suggest that an imbalance in cellular oxidants and antioxidants is a critical underlying factor. Many environmental pollutants, without being redox-active and capable of causing oxidation in vitro, are able to shift the oxidant/antioxidant balance of the cells, thus triggering pathological responses. It is difficult to assess the biological impact of mixtures of environmental pollutants because of their complexity. We propose the parallel use of normal (parental) and genetically engineered superoxide dismutase (SOD)-deficient E. coli strains, as a tool for the detection of prooxidant-acting environmental pollutants. Doubling time, viability, mutation rate, and induction of antioxidant regulons are used as parameters to assess the pollutants’ toxicity and prooxidant action. SOD-deficient E. coli is highly sensitive to agents causing oxidative stress directly or indirectly. This sensitivity can be measured and is highly reproducible. Using E. coli is advantageous because the organism is well studied, can be stored at low temperatures for a very long time, and can be grown under standard, well-defined conditions. The parallel use of SOD-deficient mutants and SOD-proficient parents permits discrimination between compounds that exert general toxicity and compounds that act as prooxidants in vivo. The bioassay was used for fast screening of potentially hazardous chemicals with pro-oxidative action.

Keywords: prooxidant, environmental pollutants, superoxide dismutase-deficient, oxidative stress, bioassay.
1 Introduction

Increasing awareness that the environment is an important determinant of both individual and community health stimulates the search for a better understanding of the mechanisms of action of various environmental pollutants. There is substantial evidence from epidemiological studies that the etiology of a number of ‘modern’ diseases is linked in part to environmental pollution (Hennig et al. [1]). Many environmental contaminants, especially persistent organic pollutants, are risk factors for chronic diseases because they may exacerbate an underlying disease by altering gene expression patterns. Many mechanisms and signaling pathways associated with the pathology of ‘modern’ diseases are modulated by environmental pollutants. Pollutants have been found to contribute to the development of a wide range of diseases such as atherosclerosis (Hennig et al. [1], cancer (Brody et al. [2], Rumchev et al. [3]), diabetes (Yoshida et al. [4], Carpenter [5]), renal failure (Bertin and Averbeck [6]), cardiovascular (Prüss-Ustün et al. [7], Bjerregaard [8]), neurological (Grandjean and Landrigan [9]), and liver diseases (Yoshida et al. [4], Carpenter [5]), and can cause immunosuppression (Bertin and Averbeck [6]). Many genes induced in such diseases are oxidative stress-sensitive (Grandjean and Landrigan [9], Calabrese et al. [10], Watkins et al. [11]), suggesting that an imbalance in cellular redox status is a critical underlying factor (Hennig et al. [1]). Many environmental pollutants including heavy metals and various organic contaminants, without being redox active and prooxidative in vitro, are able to shift the oxidant/antioxidant balance of the cells, thus triggering pathological responses.

It is difficult to assess the biological impact of environmental pollutants acting as mixtures of individual compounds with different chemical properties and reactivities. More than 15,000 high volume production manmade chemicals are in use and hundreds more are introduced each year. More than half of them have never been tested for human toxicity (Landrigan et al. [12]), because the capacity to produce new chemicals exceeds the ability to test them (Chalupka [13]). This poses a demand for fast, easy and reliable methods for screening of potentially toxic chemicals. Whole-cell biosensors are finding increasing use in the detection of environmental pollution and toxicity (Ron [14]). Here we describe how the parallel use of normal (parental) and mutant, superoxide dismutase (SOD)-deficient strains (where the genes for the cytoplasmic SODs are deleted), can be used as a tool for the detection of prooxidant-acting environmental pollutants. We and others have demonstrated that the SOD-deficient E. coli is highly sensitive to various agents contributing directly or indirectly to oxidative stress (Carlioz and Touati [15], Benov et al. [16, 17]). Such sensitivity is manifested as retarded growth, increased mutation rates, induction of specific genes, and loss of viability. These responses can be easily measured and were found to be highly reproducible. Using E. coli as a biosensor is advantageous because the organism is well studied, can be stored at low temperatures for a very long time, and can be grown under standard, well-defined conditions. The parallel use of SOD-deficient mutants and SOD-
proficient parents provides a basis for discrimination between compounds that exert general toxicity and compounds that act as prooxidants in vivo.

2 Materials and methods

2.1 Strains

The strains of *E. coli* used were: GC4468 = parental; QC1799 = GC4468 Δ sodA3 Δ sodB-kan Touati et al. [18]. These strains were prepared by D. Touati, Institute Jacques Monod, CNRS, University Paris, France. To ensure that the responses are not a consequence of the genetic background, the following strains were used in parallel: AB1157 = parental; JI132 = AB1157 plus (sodA::MmdPR13) 25 (sodB-kan) 1-Δ 2 Imlay et al. [19]; These strains were provided by J. Imlay, University of Illinois, Champaign-Urbana, Urbana, IL.

2.2 Growth media

Luria-Bertoni (LB) medium contained 10 g Bacto-tryptone, 5 g yeast extract, and 10 g NaCl per liter and was adjusted to pH 7.0 with ~ 1.5 g of K₂HPO₄. M9CA medium consisted of minimal A salts (6 g Na₂HPO₄, 3 g K₂HPO₄, 1 g NH₄Cl, and 0.5 g NaCl per liter (Maniatis et al. [20]); MgSO₄ and CaSO₄ were autoclaved separately and added to the cooled A salts to a final concentration of 20 mM and 100 µM respectively), 0.2% casamino acids, 0.2% glucose, 3 mg pantothenate and 5 mg of thiamine per liter. Minimal medium consisted of minimal A salts supplemented with 0.2% glucose.

Starter cultures were grown overnight at 37°C, with shaking in air, in LB medium containing 50 µg/ml kanamycin and/or 30 µg/ml chloramphenicol where indicated. For monitoring growth, the overnight cultures were diluted 200 fold into M9CA medium not containing antibiotics. Freshly prepared filter-sterilized solutions of chemicals to be tested were added and cultures were grown at 37°C, with shaking in air. Respective controls not containing additives or containing solvents only were run in parallel. Growth was followed turbidimetrically at 600 nm. For survival and MTT assays M9CA cultures were grown to A₆₀₀nm = 0.5 – 0.8 and the cells were resuspended to the same density in minimal medium.

2.3 MTT assay

MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) assay was performed using a modification of the original procedure of Mosmann [21]. Hundred µl aliquots of the cell suspensions were transferred into a 96-well plate, graded concentrations of the tested chemicals or progressively diluted samples were added and the plates were incubated for 1 h at 37°C on a shaker at 200 rpm. After the completion of the incubation 10 µl of MTT reagent (5 mg/ml) were added to all wells and the plates were incubated for 30 min at 37°C on a shaker. After 30 min, 100 µl of 10% SDS in 10 mM HCl were added and plates were
incubated for 1 h at room temperature. The solubilized product was assayed at 570 nm, and the absorbance at 700 nm was used as a background value.

2.4 Mutagenesis

Mutagenesis was monitored by assaying the frequency of thymine-negative (Thy\(^-\)) mutants. Thy\(^-\) mutants are resistant to the drug trimethoprim and can be selected from a Thy\(^+\) population (Miller [22]). The assay was performed as described by Farr et al. [23]. For counting Thy\(^-\) mutants, aliquots of the cultures were plated directly on LB plates containing thymine (50\(\mu\)g/ml) and trimethoprim (15 \(\mu\)g/ml). For enumeration of cells, cultures were suitably diluted and plated on LB plates containing thymine, but not trimethoprim.

Where indicated, exogenous metabolic activation was performed as described by Ames et al. [24] Hydrophobic organic compounds were dissolved in DMSO. All samples were tested at different concentrations in triplicate, both with and without microsomal activation. The S9 mixture contained the hepatic S9 fraction (1 mg/ml of microsomal protein), and mid-log bacterial culture in M9CA medium.

All experiments were repeated 3–5 times with three replicates. Student t-test was used to determine statistical significance.

3 Results and discussion

_E. coli_ contains three distinct SODs, CuZnSOD, FeSOD and MnSOD (Touati [25]). Genes coding for MnSOD are designated as _sodA_, for FeSOD – _sodB_, and for CuZnSOD – _sodC_ (Touati [26]). In _E. coli_ both FeSOD and MnSOD are cytoplasmic, while the CuZnSOD is periplasmic (Benov et al. [27]). The _sodA sodB_ strains are referred to as SOD-deficient even though they retain the periplasmic CuZnSOD, which is expressed in stationary phase and is practically not present during exponential growth (Benov and Fridovich [28]).

The _sodAsodB_ mutants grow as well as the SOD-proficient parents anaerobically, but exhibit poor aerobic growth (Al-Maghrebi and Benov [29]), auxotrophy for certain amino acids (Benov et al. [30], Benov and Fridovich [31]), increased aerobic mutation rates (Farr et al. [23], Benov and Fridovich [32], and hypersensitivity to H\(_2\)O\(_2\) and redox-cycling agents (Carlioz and Touati [15]).

The hypersensitivity of the SOD-deficient mutants to compounds causing oxidative stress suggested that those mutants could be used as sensors for potentially prooxidant environmental contaminants. To test this idea, parental and SOD mutants were subjected to treatment with various potentially toxic chemicals, including known prooxidants, and physiological responses such as changes in growth rate, decrease in viability, acceleration of mutation rate, and overall changes in metabolic rate were determined.
3.1 Effect of toxic compounds on *E. coli* growth and viability

Growth of parental and SOD-deficient strains was followed in the presence of chemicals with prooxidant activity and compared to the growth with chemicals which are known to be toxic, but without known prooxidant activity. Fig. 1 A shows growth curves of SOD-proficient and SOD-deficient cultures and the effect of Cd^{2+}. The effect of graded concentrations of CdCl\(_2\) on the growth of parental and mutant cultures is shown in fig. 1 B.

Figure 1: Effect of CdCl\(_2\) on proliferation of SOD-proficient and SOD-deficient strains in M9CA medium. Panel A, growth curves; Panel B, growth rates. Growth was monitored as absorbance at 600 nm. The growth rate of control cultures (without toxic additions) was calculated as 100%. Values significantly different from control (p<0.05) are indicated with a (*). Values are means of triplicates ± SE and are representative of at least three independent experiments.

Fig. 2 illustrates the higher sensitivity of the SOD-deficient mutants to some heavy metals (Hg, Cd), paraquat (methyl viologen), phenazine methosulfate (PMS), peroxides, short-chain sugars, dicarbonyls, and even to ascorbate, which is considered an antioxidant.

No differences between parental and *sodAsodB* mutants were observed with respect to sensitivity towards pyridine (fig. 2 A), and similar effect was observed for arsenate, dimethylamine, formaldehyde, and chlorhexidine digluconate (not shown). Other compounds, considered potentially toxic (acrylamide, nitrate) had no effect at the tested concentrations.

The hypersensitivity of the *sodAsodB* cells towards heavy metals such as Cd and Hg has its explanation. Even though those metals are not redox active and cannot act as prooxidants *in vitro*, *in vivo* Cd was shown to increase the production of superoxide by interfering with the respiratory electron transport chain (Belyaeva et al. [33], Wang et al. [34]), while Hg being a sulphydryl poison was shown to indirectly increase ROS production (Kovacic and Somanathan [35]) and to induce the O\(_2^-\) -sensing *soxRS* regulon in *E. coli* (Fuentes and Amábile-Cuevas [36]). The prooxidant effect of otherwise considered antioxidants, for example ascorbate and cysteine, can be explained by coupled
redox-cycling with transition metals (Park and Imlay [37]), and that of paraquat and PMS, to intracellular redox-cycling, producing superoxide (Hassan and Fridovich [38]). Fig. 2 demonstrates that compared to the SOD-proficient parent, less sodAsodB cells survived in the presence of common metabolites such as triose-phosphates. The reason lies in O$_2$-dependent oxidation to α,β-dicarbonyl compounds (Benov et al. [16], Benov and Fridovich [17]), and their lower rate of elimination by the SOD-deficient cells (Okado-Matsumoto and Fridovich [39]).

![Figure 2: Effect of toxic compounds on growth (Panel A) and viability (Panel B) of parental and sodAsodB strains. To compensate for the volume added by the tested sample, an equal volume of the carrier was added to control cultures. Values significantly different from control (p<0.05) are indicated with a (*). Means of triplicates ± SE are shown and are representative of at least three independent experiments. PQ - paraquat; PMS - phenazine methosulfate; TBHP – tert-Butyl hydroperoxide; Asc – sodium ascorbate; GA – glycolaldehyde; GX – glyoxal; Erythr – erythrose; AAM – acrylamide; Pyr – pyridine; GA3P – glyceraldehyde-3-phosphate; DHAP – dihydroxyacetone phosphate; MG – methylglyoxal; Cys – cysteine.](image)

### 3.2 MTT reduction test

Results shown above demonstrate that *E. coli* growth rate and viability can be used as relatively easily measurable parameters for assessment of toxicity and prooxidant effect. Growth, however, needs to be followed for relatively long time intervals and viability assessment requires plating and enumeration of colonies, which is time- and work-demanding. Methods based on reduction of tetrazolium salts to purplish-blue formazan products have become some of the most widely used tools for assessing cell viability and proliferation in cell biology. Since production of colored formazan depends on cell metabolism
(Berridge et al. [40]), the assay has been widely used for measuring the metabolic activity of cells ranging from mammalian to microbial. It has been proved suitable for assessing microbial cell proliferation (Tsukatani et al. [41]) and viability (Tsukatani et al. [42]), as well as bactericidal activity (Stevens et al. [43], Stevens and Olsen [44]). A comparison between the effects of prooxidant and non-prooxidant toxic compounds on cell proliferation and viability with MTT test, performed in a 96-well plate, demonstrated that MTT reduction can be used as a method for fast initial screening of a big number of samples (fig. 3).

![MTT reduction test](image)

**Figure 3:** MTT reduction test. GA3P – glyderaldehyde-3-phosphate; PQ – paraquat; GX – glyoxal; GA – glycolaldehyde; Pyr – pyridine; AAM – acrylamide.

### 3.3 Mutagenesis

It is known that in aerobic environment, the sodAsodB E. coli cells mutate faster than the parental cells, which has been attributed to ROS-mediated DNA damage. This finding implies that compounds, which increase ROS production, i.e. act as prooxidants either directly or indirectly, would eventually further increase mutation rates, and SOD-deficient cells would be more sensitive. This is illustrated by the bigger number of mutants in SOD-deficient cultures treated with short-chain aldehydes (fig. 4).

Activation by rat hepatic S9 fraction, as described in the Ames’ mutagenicity test, has been found to suppress the mutagenic effect of short-chain aldehydes, but potentiated the mutagenic effect of known mutagens such as benzpyrene (not shown). The assay was tested with samples of sea water, collected from different parts of Kuwait’s coastal area. Fig. 5 shows increased mutagenicity of the SOD-deficient cells by S9-activated seawater samples (SW2 and SW3) collected from areas polluted as a result of a spill of sewage water from a malfunctioning sewage-treatment plant.
Figure 4: Prooxidant-acting compounds are much stronger mutagens for the SOD-deficient than for the parental strain. Overnight LB cultures of SOD-proficient and SOD-deficient strains were diluted 200 fold in M9CA medium and were grown aerobically at 37°C and 200 rpm to a density of $A_{600\text{nm}} \sim 0.5$. At this point compounds to be tested were added and the cells were kept for 2 h on the shaker. After the incubation, the cells were diluted and plated on LB plates containing thymine for assessing cell number or aliquots were plated without dilution on LB plates containing thymine and trimethoprim for counting Thy$^-$ mutants. Bars represent mean ± S.E.M. (n=3). GA – glycolaldehyde; GX – glyoxal; GLA – glyceraldehyde; DHA – dihydroxyacetone.

Figure 5: Mutagenicity of seawater samples collected from areas polluted with raw sewage (SW2 and SW3). Non-polluted seawater (SW1) was tested for comparison. Mutagenicity test was performed after activation by rat hepatic S9 fraction.
4 Conclusions

In conclusion, the data presented here indicate that the parallel use of SOD-proficient and SOD-deficient E. coli strains is a promising cell-based system for detection of environmental pollutants with potentially prooxidant action. Detailed studies with a bigger number of chemicals will be needed in order to assess the practical usefulness and the limitations of the proposed biosensor.

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References


