

Video Article

Production and Detection of Reactive Oxygen Species (ROS) in Cancers

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Abstract

Reactive oxygen species include a number of molecules that damage DNA and RNA and oxidize proteins and lipids (lipid peroxidation). These reactive molecules contain an oxygen and include H₂O₂ (hydrogen peroxide), NO (nitric oxide), O₂⁻ (oxide anion), peroxyntrite (ONOO⁻), hydrochlorous acid (HOCl), and hydroxyl radical (OH⁻).

Oxidative species are produced not only under pathological situations (cancers, ischemic/reperfusion, neurologic and cardiovascular pathologies, infectious diseases, inflammatory diseases¹, autoimmune diseases², etc...) but also during physiological (non-pathological) situations such as cellular metabolism^{3,4}. Indeed, ROS play important roles in many cellular signaling pathways (proliferation, cell activation^{5,6}, migration⁷ etc...). ROS can be detrimental (it is then referred to as "oxidative and nitrosative stress") when produced in high amounts in the intracellular compartments and cells generally respond to ROS by upregulating antioxidants such as superoxide dismutase (SOD) and catalase (CAT), glutathione peroxidase (GPx) and glutathione (GSH) that protects them by converting dangerous free radicals to harmless molecules (i.e. water). Vitamins C and E have also been described as ROS scavengers (antioxidants).

Free radicals are beneficial in low amounts³. Macrophage and neutrophils-mediated immune responses involve the production and release of NO, which inhibits viruses, pathogens and tumor proliferation⁸. NO also reacts with other ROS and thus, also has a role as a detoxifier (ROS scavenger). Finally NO acts on vessels to regulate blood flow which is important for the adaptation of muscle to prolonged exercise^{9,10}. Several publications have also demonstrated that ROS are involved in insulin sensitivity^{11,12}.

Numerous methods to evaluate ROS production are available. In this article we propose several simple, fast, and affordable assays; these assays have been validated by many publications and are routinely used to detect ROS or its effects in mammalian cells. While some of these assays detect multiple ROS, others detect only a single ROS.

Video Link

The video component of this article can be found at <http://www.jove.com/video/3357/>

Protocol

1. Detection of ROS using carboxy-H₂DCFDA

Carboxy-H₂DCFDA is non-fluorescent but in the presence of ROS, when this reagent is oxidized, it becomes green fluorescent.

1. Immediately prior to use, prepare a fresh stock solution of carboxy-H₂DCFDA in sterile dimethylsulfoxide (DMSO) or 100% ethanol. Avoid multiple thaw/freeze cycles of your dye.
2. Wash the cells with HEPES buffered salt solution (HBSS) or phosphate-buffered saline (PBS) to remove traces of the original medium.
3. Load the cells with the dye (and the control dye, cf note section). In this assay we used Jurkat, a human leukemia cell line. Use carboxy-H₂DCFDA at a final concentration of 1 μM in regular culture medium with reduced serum (2%).
4. Incubate the cultures for 30 minutes in the dark, in a conventional incubator (37°C, 5% CO₂). Discard all unused dye solutions.
5. Remove carboxy-H₂DCFDA containing medium and wash twice with HBSS or PBS. From this step forward, protect your cells from light.
6. Add fresh medium containing your drug of choice to the carboxy-H₂DCFDA-loaded cells and incubate as desired. For this example we use H₂O₂ (0.03%) for 1 hour.
7. Assess ROS by immediately analyzing your cells by flow cytometry using the FL1 channel (green fluorescence), or by fluorescence plate reader, or by fluorescence microscopy. The fluorescence can be detected by using excitation and emission wavelengths that are appropriate for green fluorescence.

Note: Controls should include carboxy-H₂DCFDA-loaded untreated cells and unstained untreated cells. Carboxy-H₂DCFDA is known to detect peroxides but could also be oxidized by other ROS. This reagent can also be modified by other means, oxidation insensitive control dye such as 5-(and-6)-carboxy-2',7'-dichlorofluorescein diacetate (carboxy-DCFDA) should therefore be included in the test.

2. Measurement of nitric oxide (NO) production

You will need Sulfanilamide and N-1-naphthylethylenediamine dihydrochloride (NED) solutions, and Nitrite standard. This assay is called the Griess assay.

NED solution : Make a 0.1% solution of N-1-naphthylethylenediamine dihydrochloride diluted in sterile water. Sulfanilamide solution : Make a 1% solution of sulfanilamide diluted in 5% phosphoric acid. Nitrite Standard: Dilute the 0.1M standard stock sodium nitrite to 100µM in sterile medium, do a serial dilution in the same medium.

Storage Conditions: Store chemicals as directed by manufacturer at room temperature. When reconstituted, NED and Sulfanilamide solutions are stored immediately after use at 4°C, in the dark, and for a maximum of 3 months.

1. Culture cells in a 96 well plate, use triplicates for each condition, and include proper controls according to your experiments.
2. Treat cells to induce NO production. In our experiment we use lipopolysaccharide (LPS) (100 ng/ml) and recombinant IL-4 to treat our cells. In this protocol, we used RAW 264.7, a mouse macrophage cell line (Mouse leukaemic monocyte macrophage cell line).
3. On the day of the assay, bring both reagents to room temperature.
4. Spin your plate, collect cell supernatants and transfer 50µl to a new 96 well plate. Prepare limiting dilution wells of your standard stock solution to make a standard curve.
5. Add 50µl of Sulfanilamide Solution to each sample and control well and mix well.
6. Incubate at room temperature for 10 minutes in the dark.
7. Add 50µl of N-1-naphthylethylenediamine dihydrochloride solution to each sample and control well and mix well.
8. Incubate at room temperature for 10 minutes in the dark.
9. Measure absorbance immediately using a plate reader with filter of wavelengths between 520nm and 550nm.

If you utilize different plate/dish sizes, use 1/1/1 volume for each solution and sample supernatant.

A violet color will appear in the positive wells. Results obtained with the standard will help you check the stability of your solutions.

3. Detection of ROS action: oxidized proteins

A different method to identify the production of ROS is to look at the end results by detecting the oxidation of proteins. Indeed, ROS modify glutathione, an antioxidant that is expressed in most cells and plays a protective role against ROS. Following oxidation by ROS, modification of the reduced glutathione (GSH) results in the sulfhydryl (thiol) group of its cysteine being linked to a second glutathione via a disulfide bridge. This leads to the formation of a dimerized protein (oxidized protein GSSG). GSH can be restored via modification of GSSG by the enzyme glutathione reductase. The increase in the GSSG/GSH ratio reflects oxidative stress. The following assay is based on the detection and quantification of these oxidized proteins. This method is not selective for specific ROS but rather detects the effects of NO, H₂O₂, O₂⁻ and other ROS. Here we measure the total amount of oxidized (GSSG) and reduced (GSH) glutathione using bioluminescent signals.

1. Seed your cells in a 96 well plate (white/transparent, flat bottoms) as usual. For our experiment, we used 1x10⁴ cells per well for adherent Raw 264.7 and A549 cells and suspension Jurkat cells. Depending on the size of your cells these numbers could be adjusted. We recommend to plate adherent cells the day before the test to allow them to attach to the plate. Enough wells should be prepared for oxidized and reduced protein detection as well as for "medium only" and untreated cells as controls. Use triplicates for all conditions.
2. Treat your cells with your drug for the required time. Here we treated Jurkat cells and A549 cells for 1h with H₂O₂ (at 5mM and 2.5mM respectively). Raw 264.7 cells were treated with 200ng/ml of LPS for 16 h. During this incubation time, bring all your reagents to room temperature (RT). Make all reagents no more than 30 minutes before performing the test. The table below shows the volumes of reagents per well. When possible, it is important to remove the media containing the reducing agent before proceeding with the test.

	Adherent cells		Suspension cells	
Total Glu Lysis	Oxidized Glu Lysis	Total Glu Lysis	Oxidized Glu Lysis	
NEM, 25mM	none	0.5µl	none	0.5µl
Luciferin-NT	1 µl	1 µl	1 µl	1 µl
Passive Lysis Buffer, 5X	10µl	10µl	10µl	10µl
Distilled water	39.0µl	38.5µl	14µl	13.5µl
Final volume per well	50µl	50µl	25µl	25µl

3. Remove medium from wells with adherent cells. Do not remove medium in wells with suspension cells.
4. Add reduced glutathione lysis reagent or oxidized glutathione lysis reagent to the corresponding wells and shake for 5 minutes at RT.
5. Add the Luciferin generation reagent and incubate for 30 minutes at RT.
6. Add the Luciferin detection reagent and incubate for 15 minutes at RT.
7. Read the bioluminescent signal at integration time of 0.25-1 second per well using a plate reader luminometer.

	Adherent cells	Suspension cells
Cell number per well	1x10 ⁴	1x10 ⁴
Cell suspension per well+ drug	100 µl, to be removed before 3.4	25µl, not to be removed
Reduced Glutathione Lysis Reagent	50 µl	25µl
Oxidized Glutathione Lysis Reagent	50 µl	25µl
Luciferin Generation Reagent	50 µl	50 µl
Luciferin Detection Reagent	100 µl	100 µl

4. Representative Results:

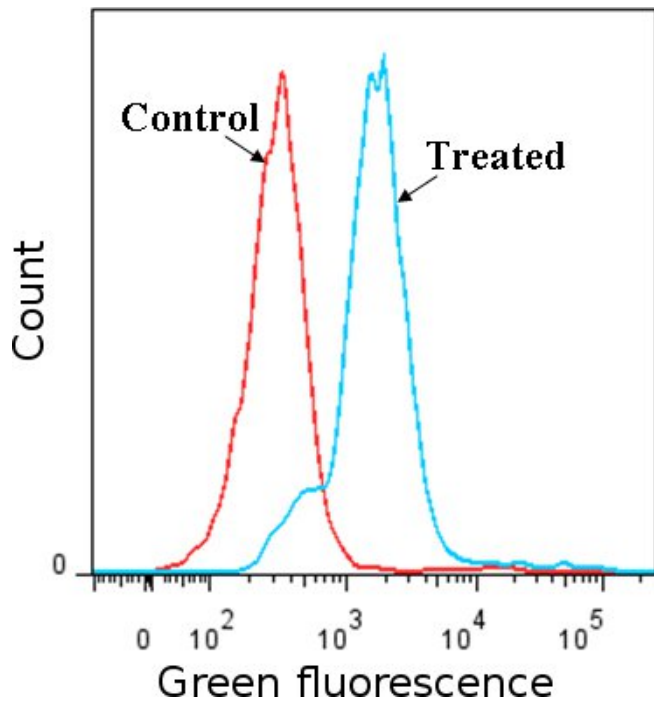


Figure 1 Detection of ROS using carboxy-H2DCFDA dye. Jurkat cells (human leukemia cell line) treated with H₂O₂ were compared to non-treated cells. ROS induces the modification of carboxy-H₂DCFDA that fluoresces green as detected by flow cytometry, the fluorescent peak in H₂O₂ treated cells shift compared to the peaks in controls (H₂O₂ treated cells stained with oxidation insensitive dye and non-treated cells stained with carboxy-H₂DCFDA). Results confirm the presence of ROS in treated cells.

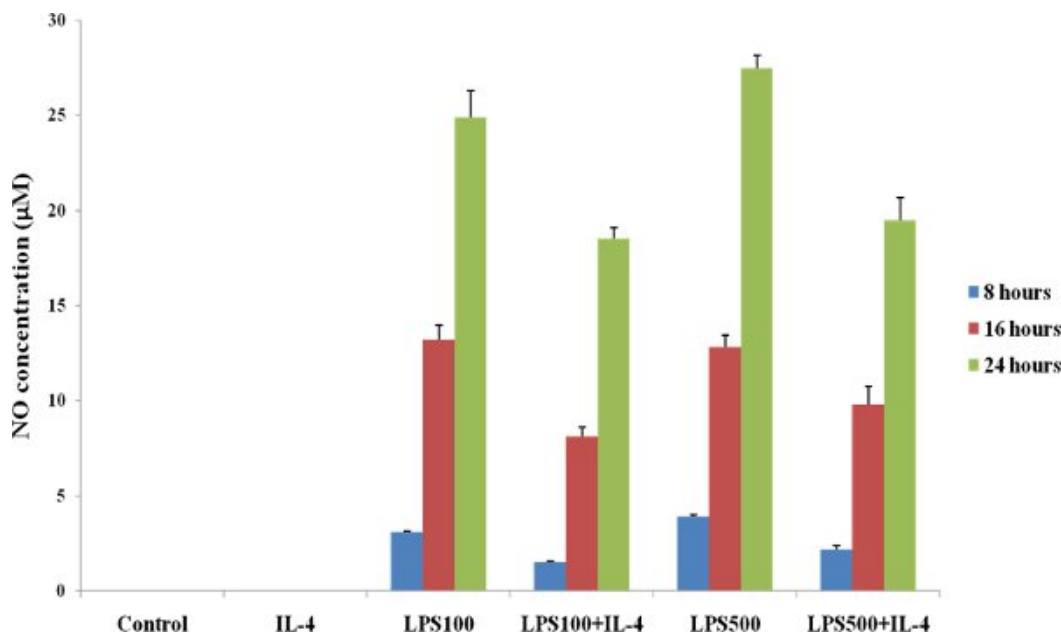


Figure 2 Detection of NO using Griess reagents. RAW 264.7 cells (mouse macrophage) were treated with LPS and IL-4. A significant increased in NO production was detected in treated cells compared to control untreated cells.

Cell lines	Non-treated	Treated
Raw 264.7	13.0	8.3
A549	21.6	10.5
Jurkat	5.2	2.8

Table 1 Detection of ROS mediated oxidation of proteins. RAW 264.7 cells were treated with LPS, Jurkat and A549 (human lung cancer) cells were treated with H₂O₂. Results are expressed as the ratio reduced (GSH)/oxidized (GSSG) glutathione. Lower ratios of glutathione (GSH)/(GSSG) were detected in treated cells compared to control untreated cells, revealing that proteins were more oxidized in treated cells.

Discussion

Several pathological situations such as inflammatory diseases, cancers, ischemia/reperfusion, and also treatments such as radiation or chemotherapy (i.e cisplatin) induce ROS overproduction. Thus, detecting and measuring ROS levels is important in many basic, pre-clinical and

clinical studies. However, ROS have very short half lives and could be complicated to detect. Here, we propose simple tests that are routinely used and widely accepted for the detection of free radical production in mammalian cells.

Detection protocols using dyes allow the experimenter to use various widely available methods (such as microscopy, flow cytometry, plate reader etc....) to assess the presence of ROS. Flow cytometry and microscopy provide information on cell viability and the percent of null, low, and high ROS producing cells. Microscopy also provides information on the effects of ROS on cell morphology and cell adhesion. Detection using plate readers allows mostly quantitative information. NO detection by Griess is quantitative but does not provide any information about the shape or adhesion of the cells studied. NO detection is relatively straightforward in mouse cells. However, lower amounts of NO are found in human samples, it is therefore important to be even more careful at each step of the detection procedure and to include controls. Detection of protein oxidation investigates the total effects of all ROS produced by the studied cells; here the intensity of the signal obtained represents the amount of ROS produced. The detection of H₂O₂ effects is trickier as it is very rapidly eliminated by catalase. The GSH-GSSG-Glo assay has the ability to detect GSSG directly, this assay minimize the issues of oxidation because it is performed directly on samples with no extraction, and with fewer cells. To improve GSH/GSSG readings, HBSS solution or glutathione-free media (such as DMEM) should be preferred; working with serum free media will further improve the reading. It is important to keep in mind that several factors including the nature of the cells (tissue type, normal versus pathologic tissues), the confluency of the monolayer cell culture, the activation and stress status, the age (number of passages), as well as the proliferation rate of your cells will have an effect on the production of ROS.

Most reagents used for ROS detection are susceptible to light, oxygen, and temperature changes; it is therefore crucial that the experimenter uses special care for storage and usage of all reagents and samples to be tested, and perform the work quickly.

Each experiment using dyes should include proper controls such as untreated (no drug), unloaded (no dye), loaded untreated, and unloaded treated cells as controls for ROS production/detection. Using medium without phenol red will reduce interference with fluorescent dyes (i.e. carboxy-H₂DCFDA, CM-H₂DCFDA). Before starting these experiments, it is important to check the natural fluorescence of your drug/treatment and of your untreated cells to adapt to this protocol. Loading your cells with the fluorescent dye should be done prior to or after treatments depending on the duration (minutes/hours versus several days) and type of treatment (drugs, ischemia/reperfusion, mechanical stress) that will be applied to the cells. The concentration of dye required to detect ROS can vary depending on the cell type and on the activation status of the cells. Our advice is to start with dye concentrations of 10µM and 100µM, check for toxicity and efficacy of the staining, and then reduce the concentration to determine the best dose to use for your experiments.

Disclosures

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References

1. Guzik, T. J., Korb, R., & Adamek-Guzik, T. Nitric oxide and superoxide in inflammation and immune regulation. *J Physiol Pharmacol.* **54**, 469-487 (2003).
2. Perl, A., Gergely, P., Jr., & Banki, K. Mitochondrial dysfunction in T cells of patients with systemic lupus erythematosus. *Int Rev Immunol.* **23**, 293-313 (2004).
3. Valko, M., Leibfritz, D., Moncol, J., Cronin, M. T., Mazur, M., & Telser, J. Free radicals and antioxidants in normal physiological functions and human disease. *Int J Biochem Cell Biol.* **39**, 44-84 (2007).
4. Droge, W. Free radicals in the physiological control of cell function. *Physiol Rev.* **82**, 47-95 (2002).
5. Nakamura, K., Yube, K., Miyatake, A., Cambier, J. C., & Hirashima, M. Involvement of CD4 D3-D4 membrane proximal extracellular domain for the inhibitory effect of oxidative stress on activation-induced CD4 down-regulation and its possible role for T cell activation. *Mol Immunol.* **39**, 909-921 (2003).
6. Los, M., Droge, W., Stricker, K., Baeuerle, P. A., & Schulze-Osthoff, K. Hydrogen peroxide as a potent activator of T lymphocyte functions. *Eur J Immunol.* **25**, 159-165 (1995).
7. Deem, T. L., & Cook-Mills, J. M. Vascular cell adhesion molecule 1 (VCAM-1) activation of endothelial cell matrix metalloproteinases: role of reactive oxygen species. *Blood.* **104**, 2385-2393 (2004).
8. Pacher, P., Beckman, J. S., & Liaudet, L. Nitric oxide and peroxynitrite in health and disease. *Physiol Rev.* **87**, 315-424 (2007).
9. Griendling, K. K., Sorescu, D., Lassegue, B., & Ushio-Fukai, M. Modulation of protein kinase activity and gene expression by reactive oxygen species and their role in vascular physiology and pathophysiology. *Arterioscler Thromb Vasc Biol.* **20**, 2175-2183 (2000).
10. Powers, S. K., Duarte, J., Kavazis, A. N., & Talbert, E. E. Reactive oxygen species are signalling molecules for skeletal muscle adaptation. *Exp Physiol.* **95**, 1-9.
11. Loh, K., Deng, H., Fukushima, A., Cai, X., Boivin, B., Galic, S., Bruce, C., Shields, B. J., Skiba, B., Ooms, L. M., Stepto, N., Wu, B., Mitchell, C. A., Tonks, N. K., Watt, M. J., Febbraio, M. A., Crack, P. J., Andrikopoulos, S., & Tiganis, T. Reactive oxygen species enhance insulin sensitivity. *Cell Metab.* **10**, 260-272 (2009).
12. Goldstein, B. J., Mahadev, K., & Wu, X. Redox paradox: insulin action is facilitated by insulin-stimulated reactive oxygen species with multiple potential signaling targets. *Diabetes.* **54**, 311-321 (2005).