

Ratiometric Singlet Oxygen Nano-optodes and Their Use for Monitoring Photodynamic Therapy Nanoplatfoms

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Received 18 May 2005; accepted 11 August 2005; published online 17 August 2005 DOI: 10.1562/2005-05-18-RA-532

ABSTRACT

Ratiometric photonic explorers for bioanalysis with biologically localized embedding (PEBBLE) nanoprobcs have been developed for singlet oxygen, using organically modified silicate (ORMOSIL) nanoparticles as the matrix. A crucial aspect of these ratiometric singlet-oxygen fluorescent probes is their minute size. The ORMOSIL nanoparticles are prepared via a sol-gel-based process and the average diameter of the resultant particles is about 160 nm. These sensors incorporate the singlet-oxygen-sensitive 9,10-dimethyl anthracene as an indicator dye and a singlet-oxygen-insensitive dye, octaethylporphine, as a reference dye for ratiometric fluorescence-based analysis. We have found experimentally that these nanoprobcs have much better sensitivity than does the conventional singlet-oxygen-free dye probe, anthracene-9,10-dipropionic acid disodium salt. The much longer lifetime of singlet oxygen in the ORMOSIL matrix, compared to aqueous solutions, in addition to the relatively high singlet oxygen solubility because of the highly permeable structure and the hydrophobic nature of the outer shell of the ORMOSIL nanoparticles, results in an excellent overall response to singlet oxygen. These nanoprobcs have been used to monitor the singlet oxygen produced by “dynamic nanoplatfoms” that were developed for photodynamic therapy. The singlet oxygen nanoprobcs could potentially be used to quantify the singlet oxygen produced by macrophages.

INTRODUCTION

There are two different lowest excited singlet states of oxygen, $^1\Sigma_g^+$ and $^1\Delta_g$. The $O_2(^1\Delta_g)$, an oxygen molecule at its lowest excited singlet state, is commonly called singlet oxygen (1O_2). 1O_2 is known as a highly reactive and metastable species that can

oxidize many chemical and biological substrates (1). Though not a radical (it does not have an unpaired electron) it is a reactive oxygen species (ROS) often associated with free radicals that have a strong oxidizing activity. 1O_2 is generated by input of energy such as light excitation or irradiation. Many organic dye molecules such as photofrin or methylene blue (MB) will produce singlet oxygen upon light excitation; these dye molecules are called photosensitizers and the process is called photosensitization (2).

1O_2 plays an important role in many natural photochemical and photobiological processes, such as photodegradation and aging processes. There is also much evidence that 1O_2 is a key element in the defense system of the mammalian body, being generated by phagocytes to kill invading microorganisms (3,4). It is also reported that 1O_2 plays a role as an activator of gene expression (5). Reactions of 1O_2 are used in several fields, including organic synthesis, bleaching processes and the photodynamic therapy (PDT) of cancer (6,7). PDT has emerged as a promising method for overcoming some of the inherent problems in classical cancer therapies (8–12). It involves the selective delivery of photosensitizers such as photofrin to specific tumors. Singlet oxygen is widely believed to be the primary cytotoxic agent for killing the tumor cells in PDT. The theory is that upon excitation the photosensitizer produces singlet oxygen and other ROS, resulting in the *in situ* initiation of apoptosis in many types of tumors, with minimal systemic toxicity (13).

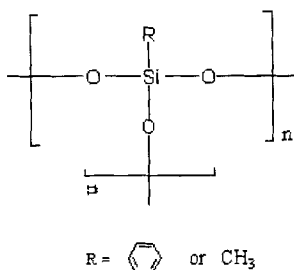
Because of the above-listed important roles of 1O_2 , the detection of 1O_2 has drawn lots of interest. So far, singlet oxygen detection methods can be sorted into two groups: 1) physical detection, given by the most direct measurement of near-infrared luminescence of 1O_2 at 1270 nm and 2) chemical 1O_2 probes, to be discussed in this article.

Although many studies have been reported on the subject, it is still difficult to detect 1O_2 in biological systems, mainly because of its relatively short lifetime in aqueous solutions or biological environments (14). Even though it has been recently reported that singlet oxygen can be quite long-lived and diffuse over appreciable distances contrary to common perception (15), the lifetime of singlet oxygen in water itself or in live cells is still quite short (about 2 μ s) and the solubility of singlet oxygen is very poor, the combination of which presents challenges to both the physical detection method and the naked molecular probes for singlet oxygen. The released 1O_2 produced in cells encounters an aqueous environment that results in the immediate quenching of a large fraction of the 1O_2 produced, hence reducing the emission signal below the sensitivity of detection methods. Several investigators

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Abbreviations: ADPA, anthracene-9,10-dipropionic acid disodium salt; DCM, dichloromethane; DMA, 9,10-dimethyl anthracene; DNP, dynamic nanoplatfoms; DPIBF, 1,3-diphenylisobenzofuran; MB, methylene blue; MTMS, methyltrimethoxysilane; OEP, octaethylporphine; ORMOSIL, organically modified silicate; PAA, poly(acrylamide); PDMA, poly(decyl methacrylate); PDT, photodynamic therapy; PEBBLE, photonic explorers for bioanalysis with biologically localized embedding; PTMS, phenyltrimethoxysilane; ROS, reactive oxygen species; SEM, scanning electron microscopy.

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Scheme 1. Chemical structure of ORMOSIL. R is the phenyl group in the inner layer of ORMOSIL and the methyl group in the outer layer of ORMOSIL.

have reported positive results from cells in suspension (16,17) or from red-cell ghosts (18). There is also a recent report on time-resolved measurements of $^1\text{O}_2$ luminescence that was generated by PDT in cells *in vitro*, or in tissues *in vivo* (19). However, the combination of the relatively short lifetime of singlet oxygen and the very low quantum yield of singlet oxygen luminescence has either required the use of deuterium oxide (D_2O) to increase the lifetime of $^1\text{O}_2$ and eliminate absorption of the 1270-nm luminescence by H_2O , or, alternatively, did not adequately distinguish between intracellular and extracellular $^1\text{O}_2$. This significantly limits the sensitivity of the physical method, thus making the detection very unreliable and irreproducible. Furthermore, it is extremely difficult for the physical method to reveal the accumulation of singlet oxygen production during the PDT process, which is one of the most important parameters to determine the efficiency of PDT.

In the chemical detection method, the $^1\text{O}_2$ chemical probes react with $^1\text{O}_2$ to produce endoperoxides, as shown in Scheme 2. In most cases the $^1\text{O}_2$ chemical probes show fluorescence or absorbance peaks at certain wavelength(s), however the endoperoxides do not. Thus, by monitoring the fluorescence emission decrease of the $^1\text{O}_2$ chemical probes, one can monitor the singlet oxygen quantitatively. The chemical trap method is useful even when the singlet oxygen generation is very low, because the concentration of endoperoxides is proportional to the cumulative amount of singlet oxygen generated. However, most of the well-known chemical probe molecules are hydrophobic dyes that are not suitable for use in an aqueous environment. Although water-soluble derivatives of these chemical probes have been developed (20–23), the problem associated with the relatively short lifetime of $^1\text{O}_2$ still remains, in addition to the possible dye toxicity problem.

The combination of the chemical detection method with the photonic explorers for bioanalysis with biologically localized embedding (PEBBLE) approach promises to tackle some of the problems mentioned above. Like the other chemical methods, it is suitable for the measurement of the minute amounts of singlet oxygen produced from cells, because the concentration of endoperoxides (product of reaction between singlet oxygen and chemical probe) formed is proportional to the cumulative $^1\text{O}_2$ generated, thus it allows one to monitor the accumulation of singlet oxygen during the natural or PDT process. Furthermore, the PEBBLE probes can be introduced inside cells by either gene-gun delivery or liposomal delivery with negligible physical perturbation (24–26). PEBBLE technology has been used especially for intracellular measurements of biologically important ions and gas molecules (24–26). These PEBBLE-type probes have several advantages over traditional molecular dye probes: (1) the dye-embedded nanoparticles can be used in aqueous solution regardless

of the hydrophobicity of the dye; (2) the dye molecule will be protected from the chemical and biological environment in the physiological system while, simultaneously, the physiological system is also protected from the potential toxicity of the dye molecules, thus enabling one to measure the concentration of the desired species in an unperturbed environment; and (3) by also loading a reference dye into the nanoparticles, ratiometric measurements can be made.

Here we developed singlet oxygen nanoprobe, designed to quantitatively monitor the singlet oxygen production during PDT, by adopting the PEBBLE technology. Organically modified silicate (ORMOSIL) nanoparticles incorporated with the hydrophobic singlet oxygen-detecting dye 9,10-dimethyl anthracene (DMA) were synthesized and tested for their feasibility as spectroscopic singlet oxygen detection probes, with very good results described below. This incorporation makes the detection take place inside the ORMOSIL particles, which significantly increases the lifetime of singlet oxygen molecules, thus allowing more time for them to react with DMA to be detected. It turns out that the incorporation of DMA inside ORMOSIL significantly enhances the sensitivity of these nanoprobe in comparison to that of the water-soluble molecular probe anthracene-9,10-dipropionic acid disodium salt (ADPA). In addition, these ORMOSIL nanoparticles are chosen as the matrix because they have been shown to provide a near-perfect environment for oxygen measurements (26) because of their good solubility, permeability and homogeneity to oxygen molecules. We thus expect that this matrix will provide similar advantages for singlet oxygen measurements, specifically the partition advantage for singlet oxygen in ORMOSIL rather than in water. Furthermore, these ORMOSIL nanoparticles can load either hydrophobic or hydrophilic dye molecules and this provides flexibility when it comes to which dyes should be chosen for optimal fabrications. We also incorporated DMA into poly (decyl methacrylate) (PDMA) nanoparticles and these nanoprobe demonstrated good responses to singlet oxygen. However, because of the severe aggregation of PDMA particles in aqueous solution, we concentrated our efforts on the ORMOSIL nanoparticles.

DMA was chosen as the chemical trap mainly because it has a relatively high quenching rate constant and unique selectivity for $^1\text{O}_2$ (k is $9.1 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ for DMA quenching of singlet oxygen) (27). In addition, the fluorescence of DMA does not change upon contact with hydrogen peroxide, nitric oxide or superoxide, confirming the specificity of DMA for $^1\text{O}_2$ (27). 1,3-Diphenylisobenzofuran (DPIBF) is traditionally considered to have the fastest reaction rate constant with singlet oxygen ($9.6 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$); however, it suffers from a severe auto-oxidation problem, especially in water (K. Gwangseong, Y. Cao, Y. E. Koo, W. Tang and R. Kopelman, in preparation), therefore it was not chosen as the probes in this paper. DMA reacts with $^1\text{O}_2$ to produce DMA-endoperoxide, as shown in Scheme 1. As for DMA endoperoxide, it is quite thermally stable; it must be heated to about 120°C to dissociate into DMA and oxygen molecules (28,29). Thus the reaction of DMA with singlet oxygen to produce DMA endoperoxide is irreversible under our experimental conditions, which is conducive to singlet oxygen detection. We also believe that the physical quenching of singlet oxygen by DMA is relatively small, as is the case for similar probes like ADPA or diphenyl anthracene (DPA), and that it is negligible when compared to the overall quenching by the solvent (30). DMA also has a strong fluorescence emission in the range of 400–450 nm, whereas DMA-endoperoxide has no emission in that range. Thus by monitoring the fluorescence

decrease of DMA, one can detect the amount of singlet oxygen produced over time. For ratiometric measurement, the octaethylporphine (OEP) dye was incorporated into the ORMOSIL nanoparticles, because of its relatively low reaction rate with $^1\text{O}_2$. Both reference and probe molecules are hydrophobic but the nanoprobe containing them are easily suspended in water, as required for the measurement of singlet oxygen production in an aqueous environment. The previously used ADPA molecules (30) are inferior in their kinetic and spectroscopic parameters compared to DMA, but were developed to have the advantage of being water soluble compared to DMA and other similar anthracene derivatives. The PEBBLE combines the advantages of both systems. In the work described below, ORMOSIL nanoprobe have been used to successfully determine the singlet oxygen production efficiency of photosensitizer-loaded PDT nanoplatforms designed and developed in our lab (31–33).

Intracellular measurements of $^1\text{O}_2$ produced during phagocytosis were reported by Steinbeck *et al.* (3), where they used micron-sized glass beads coated with 9,10-diphenyl anthracene, with perylene coated glass beads as an internal standard. Although they successfully demonstrated that $^1\text{O}_2$ is produced from the cells, there are still aspects to be improved: (1) Because the dye is coated onto the surface of the glass beads, and hence exposed to the cells directly, dye toxicity may affect the cells. (2) The glass bead size was $1.6\ \mu\text{M}$, which may be large enough to perturb the cells. We note that our singlet oxygen nanoprobe contain the dye inside the nanoparticles, avoiding the toxicity problem caused by the dye molecule; also, the probe size is reduced to 1/10 the size of the glass bead, *i.e.* the perturbing volume is reduced by a factor of 1000. We thus believe that these nanoprobe are promising candidates for intracellular singlet oxygen measurements, such as the singlet oxygen production inside live macrophages.

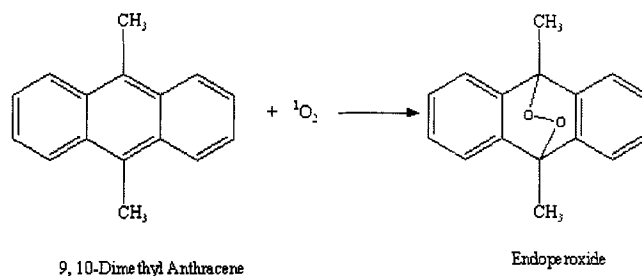
MATERIALS AND METHODS

Chemicals. Phenyltrimethoxysilane (PTMS), methyltrimethoxysilane (MTMS), ammonia, 9,10-dimethyl anthracene (DMA), methylene blue (MB), and dichloromethane (DCM) were obtained from Aldrich (Milwaukee, WI) and used without further purification. Octaethylporphine (OEP) was purchased from Frontier Scientific, Inc. (Logan, UT). Anthracene-9,10-dipropionic acid, disodium salt (ADPA) was purchased from Molecular Probes (Eugene, OR). A stock solution of $10\ \mu\text{M}$ ADPA was made in water and kept in the dark until use. The chemicals for the synthesis of photofrin-loaded Poly(acrylamide) (PAA) nanoparticles including hexane, Aerosol OT, and brij 30, acrylamide, 3-Aminopropyl-methacrylamide, N, N-Methylene(bis)acrylamide, 10% APS and TEMED were purchased from Aldrich (Milwaukee, WI). Photofrin was a gift from Axcan Pharmaceuticals (Mont Saint-Hilaire, Quebec, Canada). The $18\ \text{M}\Omega$ water used for all the solutions was purified by a Barnstead 1 Thermolyne Nanopure II system.

Gases. O_2 (99.6%, extra-dry grade), air (dry grade), N_2 (99.998%, prepurified) and Argon (99.7%, prepurified) were obtained from Cryogenic Gases (Detroit, MI). Nitric oxide (99.9%, CP grade) was obtained from Matheson Tri-Gas (Montgomeryville, PA).

Experimental sections

Preparation of ORMOSIL nanoparticles incorporated with DMA or OEP. Two different batches of ORMOSIL particles were prepared exactly as described in our previous work (25): one batch was loaded with DMA and the other was loaded with OEP. The dye amounts used in each batch were as follows: for DMA-loaded PEBBLE, 0.6 mL of DMA (1 mg/mL in DCM); and for OEP-loaded ORMOSIL PEBBLE, 0.5 mL of OEP (1 mg/mL in DCM). The resulting two batches of ORMOSIL PEBBLE were suction-filtered through a Fisher-brand glass microanalysis vacuum filter holder with a $0.1\text{-}\mu\text{m}$ Osmonics/MSI MAGNA nylon membrane filter. The PEBBLE were rinsed three times with water and then resuspended in the



Scheme 2. The reaction of DMA and singlet oxygen.

water-ethanol 1:2 mixture, sonicated for 5 min and then filtered through a $0.02\text{-}\mu\text{m}$ Whatman Anodisc filter membrane, washed with ethanol and allowed to air-dry. Emission spectra of $0.1\ \text{mg/mL}$ of these two batches of PEBBLE in 50:50 water/ethanol were taken on fluorimeter. Every one part of DMA-loaded ORMOSIL powder was mixed together with two parts of OEP-loaded ORMOSIL powder. This mixed powder was the ratiometric singlet oxygen nanoprobe and, unless noted otherwise, was used throughout all the experiments described in this paper.

Synthesis of amine-functionalized photofrin-incorporated PAA nanoparticles. A clean 250-mL round-bottom flask was charged with argon-purged hexanes. Aerosol OT (3.2 g) and brij 30 (6.4 mL) were added to the reaction flask and the contents were stirred under argon atmosphere until a uniform solution resulted. A 20-mL glass sample tube was charged with acrylamide (1.1 g) and $0.1\ \text{M}$ sodium phosphate buffer (4 mL, pH 7.25) and sonicated 10 min to yield a uniform solution. 3-Aminopropyl-methacrylamide (0.2 g) was added to the acrylamide solution and sonication was continued until a clear solution formed. N, N-Methylene(bis) acrylamide (0.36 g) was added to the acrylamide solution and sonicated again until a clear solution resulted. Photofrin (20 mg, Axcan Pharmaceuticals) was added to the acrylamide solution and sonicated for a period of 20 min at room temperature. The clear, dark red monomer solution was added to the hexane reaction mixture and stirred vigorously for 20 min at room temperature. Ten percent APS ($65\ \mu\text{L}$) and TEMED ($85\ \mu\text{L}$) were added to the reaction mixture under argon atmosphere. The reaction mixture was gently stirred at room temperature in the dark for 24 h. The reaction mixture was concentrated to a thick liquid residue and the residue was diluted with 100 mL of ethanol. The mixture was sonicated for 2 min and the separated particles were washed in an amicon stirred cell (Millipore, 200 mL) with ethanol ($5 \times 180\ \text{mL}$). The deep brown nanoparticles were further dried under nitrogen atmosphere and gently crushed to a fine powder. The material was covered with aluminum foil to protect it from light and stored in the refrigerator at 4°C .

Scanning electron microscopy (SEM) imaging. The above two individual batches of ORMOSIL PEBBLE were dispersed in 50:50 water/ethanol and sonicated for 20 min to prevent aggregation of particles. Then a drop of the PEBBLE solution was placed on the SEM specimen mount (aluminum) and dried gradually at room temperature. The sample was then sputter-coated with gold and the SEM images were taken on the Phillips XL30 Scanning Electron Microscope.

Optics. All fluorescence emission spectra are taken on a FluoroMax-2 spectrofluorimeter (ISA Jobin Yvon-Spex, Edison, NJ), slits set to 5 nm for excitation and 5 nm for emission respectively. If without any further specification, all emission spectra are taken with excitation at 380 nm. Without any further specification, all experiments and spectra were done in a darkroom with the lights off, because the photosensitizers used in this paper are sensitive to room light.

Detection of singlet oxygen produced by free MB in distilled water. To detect the singlet oxygen production by the photosensitizers (MB or photofrin), the ORMOSIL nanoprobe suspension was mixed with certain amount of photosensitizer solution. The mixture was then irradiated continuously at 650 nm (for MB) or 630 nm (for photofrin) for a fixed amount of time; after each irradiation, the emission spectrum of the mixture was taken. By monitoring the emission spectra over the irradiation time, it is possible to monitor the singlet oxygen production.

Singlet oxygen was produced using $10\ \mu\text{M}$ MB in distilled water. Based on the excitation and absorption spectra of MB in water, a 650-nm light source was used to irradiate the mixed suspension to produce singlet oxygen. Two milligrams of the above-mixed ORMOSIL PEBBLE was suspended in 10 mL of distilled water. Then, 0.5 mL of the suspension was mixed with 2 mL of $10\ \mu\text{M}$ MB in distilled water inside a quartz cell. Three

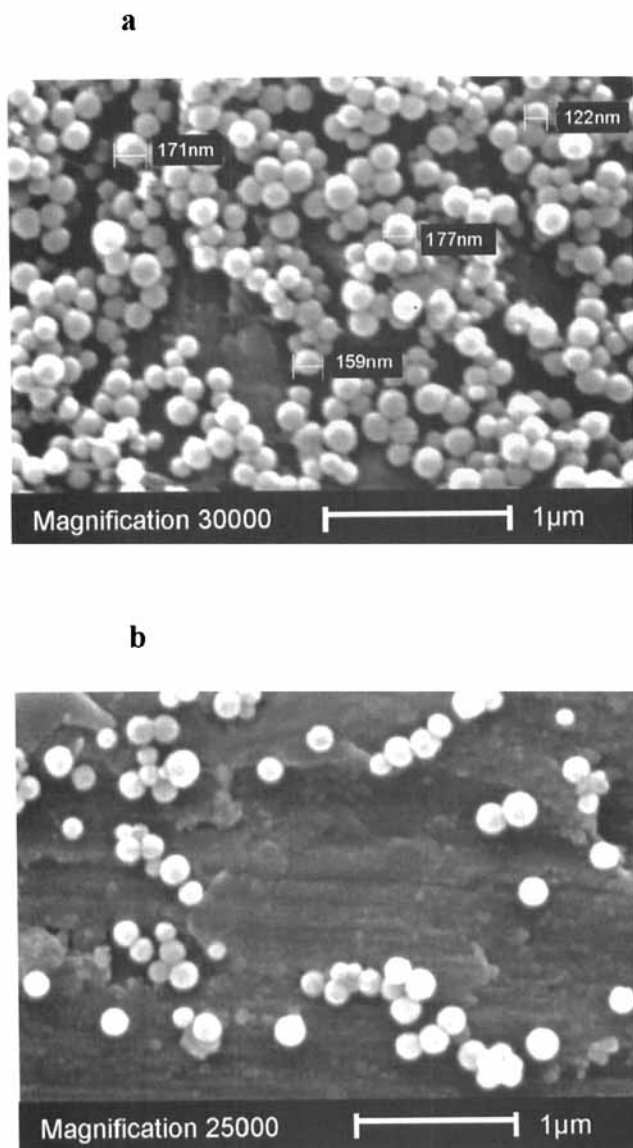


Figure 1. SEM images of ORMOSIL PEBBLE: (a) DMA-loaded particles; (b) OEP-loaded particles. The average diameter of the particles is about 160 nm.

emission spectra of this mixed sample were taken on the fluorometer immediately following the mixing. The sample was then illuminated at 650 nm (with five 5-nm excitation/emission slit widths) for 1 min, 2 min, 3 min, 10 min, 20 min and 40 min respectively; three emission spectra were taken right after each irradiation. The average of the three emission spectra represents each irradiation.

To check the photobleaching of MB in distilled water under the same experimental condition, 2.5 mL of 10 μ M MB in distilled water was continually irradiated at 650 nm for up to 40 min and emission spectra were taken to monitor the emission intensity changes.

Auto-oxidation and control tests. The above ORMOSIL PEBBLE suspension was measured into 0.5-mL samples and mixed with 2 mL of distilled water. The samples were irradiated for 0 min, 1 min, 3 min, 5 min, 10 min, 20 min, and 40 min respectively and three emission spectra were taken after each irradiation.

Another 0.5 mL of the above PEBBLE suspension was mixed with 2 mL of 10 μ M of MB in distilled water inside the quartz cell. The sample was then put in the darkness and three emission spectra were taken after 0 min, 1 min, 3 min, 5 min, 10 min, 20 min and 40 min, respectively.

Detection of singlet oxygen produced by free MB or MB incorporated inside PAA nanoparticles. To compare the performance between these ORMOSIL nanoprobe and free dye probes ADPA, samples A and B were prepared.

MB was incorporated inside PAA nanoparticles and the concentration of MB inside was measured by absorption calibration to be 1×10^{-4} mg/(30). Ten milligrams of this sample was suspended in 10 mL of distilled water. Two milliliters of this suspension was mixed with 0.5 mL of the above-described ORMOSIL nanoprobe suspension to form sample A.

To make sample B, 1×10^{-3} mg MB was dissolved in 10 mL of distilled water and mixed with 10 mg of blank PAA particles, thus making a final concentration of MB of 1×10^{-4} mg/mg. This mixture was sonicated for about 15 min. Two milliliters of this suspension was then mixed with 0.5 mL of above ORMOSIL singlet oxygen nanoprobe, forming sample B.

For both samples A and B, three emission spectra were taken after 0 min, 1 min, 2 min, 5 min and 20 min of irradiation at 650 nm respectively. The same experiment was done using 10 μ M ADPA in distilled water as the singlet oxygen detector, instead of the ratiometric ORMOSIL nanoprobe.

Detection of singlet oxygen produced by photofrin-incorporated PAA nanoparticles. In this experiment, the above-synthesized amine-functionalized photofrin-incorporated modified PAA particles were used as singlet oxygen source. Because the emission of photofrin overlaps with OEP, only DMA-loaded ORMOSIL nanoparticles were used as the singlet oxygen probe. Two milligrams of DMA ORMOSIL particles were suspended in 10 mL of distilled water. A 0.5-mL portion of this suspension was mixed with 2 mL of 0.1 mg/mL photofrin-incorporated PAA in distilled water. The mixed suspension was irradiated at 630 nm for 0 min, 1 min, 3 min, 5 min, 10 min, 15 min and 30 min, respectively. After each irradiation, three emission spectra were taken, with excitation and irradiation slit widths set as 5 nm and emission slit width set as 2 nm.

Exactly the same experiments were done using 10 μ M ADPA in distilled water as the singlet oxygen probe.

To correlate the relationship between the amount of singlet oxygen produced and the concentration of the photosensitizer, the following experiments were designed.

The above amine-functionalized photofrin-incorporated PAA nanoparticles were suspended in distilled water with two different concentrations: one at 1 mg/mL, the other at 0.25 mg/mL. A 2-mL portion of each of these two suspensions was mixed individually with 0.5 mL of the above singlet oxygen nanoprobe. Each mixed suspension was irradiated at 630 nm for up to 30 min. The emission spectra of the samples were monitored at different time intervals.

Leaching and photobleaching. The leaching test was performed by preparing a 0.2 mg/mL PEBBLE solution in distilled water and monitoring the fluorescence of the filtrate (filtered with a 20 nm Whatman cellulose membrane), every 2 h over a period of 3 days.

A photobleaching test of these PEBBLE samples was also performed to evaluate the accuracy and lifetime of the singlet oxygen nanoprobe. The same 0.2 mg/mL PEBBLE suspension in water was prepared and 3 mL of this suspension was placed in a quartz optical cell. The sample was continuously irradiated at 380 nm (excitation for DMA) and 630/650 nm (excitation for Photofrin/MB), with the excitation slit width set at 10 nm. The peak intensities at 424 nm (from DMA) and 621 nm (from OEP) were monitored over a period of 30 min.

Interference of oxygen and nitric oxide (NO). Two milligrams of the mixed ORMOSIL powder was suspended in 10 mL of distilled water and sonicated for about 30 min. A 3-mL portion of the suspension was then transferred into an adjustable airtight quartz cell capped with Teflon. A 4-inch hypodermic needle was inserted to transfer gas into the solution inside the quartz cell, and another 1.5-inch needle was used for ventilation. For the oxygen interference test, a combination of cylinders of pure nitrogen and oxygen and gas blender (Cole Parmer Instrument Co., Vernon Hills, IL) was used to achieve precise gas mixtures that were passed at selected flow rates. The interference from NO was tested by repeatedly purging the PEBBLE suspension first with nitrogen, second with NO and then with nitrogen again, and so on. The type of NO used was 99.9% NO. The time to bubble the mixed gas through the solution for each measurement was about 10 min, enough time to reach the distribution equilibrium. Emission spectra were taken immediately after each purging with air, oxygen, nitrogen and NO respectively.

RESULTS AND DISCUSSION

ORMOSIL particles

The morphology of the dye-incorporated MTMS/PTMS nanoparticles was determined by SEM. Figure 1 shows typical

monodispersed spherical ORMOSIL PEBBLE. They have an average diameter of about 160 nm.

Mixing DMA and OEP ORMOSIL PEBBLE for ratiometric analysis

Many chemical probes do not have reference signals. However, for previous PEBBLE nanosensors in this lab there was an attempt to introduce the reference dye into the PEBBLE so as to make it ratiometric, *i.e.* to minimize such effects as excitation light-intensity fluctuation or sample-size fluctuation, thus making the detection more reproducible and reliable. Therefore, historically each PEBBLE nanosensor/nanoprobe contained both sensing and reference dye molecules (24–26). However, attempts to load both dyes simultaneously resulted in loading inconsistency, presumably because of solubility and/or energy transfer. After many frustrating trials of mixing different molar ratios of OEP and DMA to get reasonably similar intensities of emission out of both DMA and OEP, a different approach was taken to overcome this problem. A batch of DMA-incorporated ORMOSIL PEBBLE was prepared, then a batch of OEP-incorporated ORMOSIL PEBBLE was prepared, then two different batches of PEBBLE were mixed at various ratios to reach the goal. The advantages of this approach are as follows:

1. It still fits the ratiometric scheme.
2. Instead of incorporating both sensing and reference dyes at the same time, which usually causes added complications for the synthesis of the nanoparticles, doping one at a time makes the synthesis easier.
3. It makes it easier to adjust the ratios between sensing and reference dyes. Instead of preparing new batches of PEBBLE for each ratio of sensing and reference dyes and repeatedly adjusting the ratios, which is usually very time-consuming (especially true for PEBBLE preparation, where the average time of preparing a batch of PEBBLE is 24 h) and expensive (especially if the dyes involved are expensive) one can just mix different amounts of two different batches of PEBBLE at one's will.
4. It minimizes energy transfer between indicator and reference dyes, which is usually a major concern in terms of choosing sensing and reference dyes that are shared by the same PEBBLE.

What should be noted here is that this approach is based on the assumption that two different batches of PEBBLE are monodispersed and homogeneous (when mixed and suspended in the solvent). This assumption is usually true if the two batches of PEBBLE have the same matrix, but it could also work with different matrixes. As stated, this new approach also provides more flexibility in terms of choosing sensing and reference dyes. However, one potential problem with this approach is that, for cell measurements, and whenever the local area of interest is small, it is still necessary for the mixed particles to be monodispersed and homogeneously distributed.

Based on the emission spectra of two different batches of DMA- (emission peaks: 408 nm, 430 nm and 456 nm) or OEP- (621 nm) loaded ORMOSIL particles, every one part of DMA ORMOSIL nanoparticles was mixed with two parts of OEP ORMOSIL nanoparticles so that the emission peak intensity of DMA would be about the same as that of OEP after the singlet oxygen measurement (a 40–50% signal decrease of DMA during the measurement was taken into account). Without any other specification, these

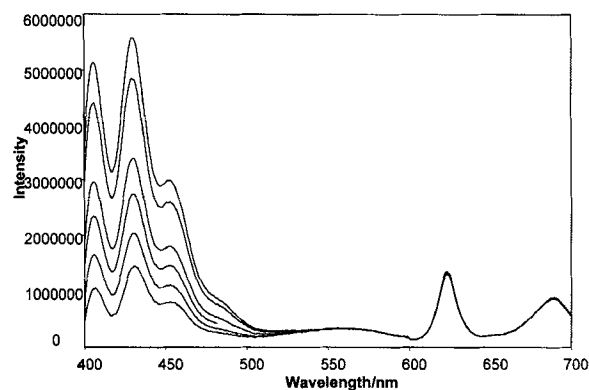


Figure 2. Detection of singlet oxygen produced by 10 μM MB in distilled water with DMA-loaded ORMOSIL singlet oxygen nanoprobres. Accumu-

same mixed DMA/OEP ORMOSIL nanoparticles were used for all of the singlet oxygen measurements.

Detection of singlet oxygen released by free MB in distilled water

One of the traditional photosensitizers that have been used for a variety of applications, including PDT, is MB; its high quantum yield for $^1\text{O}_2$ generation ($\Phi\Delta \sim 0.5$) in the therapeutic window (600–900 nm), coupled with its low toxicity, has led to the testing of MB as a promising candidate for PDT of cancer (34–35). It is also inexpensive and it is FDA-approved for ethemoglobinemia. Therefore MB has been chosen in this paper as one of the primary photosensitizers, *i.e.* as singlet oxygen source.

Regarding Fig. 2, it should be noted that the emission of DMA decreases more dramatically at the beginning of the irradiation. After 40 min irradiation, about 73% DMA emission signal reduction was observed, which means that about 73% of the DMA molecules had reacted with singlet oxygen to form DMA endoperoxides. On the other hand, the emission of OEP stays almost constant during the entire irradiation process, up to 40 min. Based on Fig. 2, one can figure out that $k_{\text{DMA}}/k_{\text{OEP}}$, the singlet oxygen quenching rate constant ratio between DMA and OEP, which is at least 100 (a lower order of magnitude limit). This confirms OEP's role as a very good reference dye for these nanoprobres.

To make sure that the emission signal decrease of DMA is indeed induced by singlet oxygen, control autooxidation and other blank tests were done and the results are shown in Figs. 3 and 4. Compared with Fig. 2's setup, Fig. 3's setup is the same, except that there is no MB inside the mixture. Fig. 3 shows that irradiation with 650-nm light causes very little signal change where there are no MB molecules in the mixture. In Fig. 4's setup, everything stays the same as in Fig. 2, (including MB), except that there is no 650-nm light irradiation.

A photobleaching test of MB under exactly the same experimental conditions reveals less than 1% signal loss over 40 min irradiation. Combining the messages of Figs. 2, 3 and 4, a reasonable conclusion is that the observed DMA signal decrease could only be caused by the simultaneous presence of both the 650-nm irradiation and the presence of MB. Neither irradiation nor MB by itself could be responsible for this behavior. Furthermore, the aromatic hydrocarbons have been known to have a specific

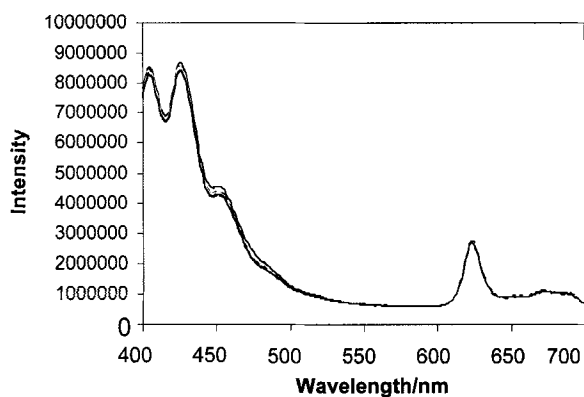


Figure 3. Control 1: Emission spectra (excited at 380 nm) of 0.5 mL of the above DMA ORMOSIL PEBBLE suspension, mixed with 2 mL of distilled water. No MB was added. The irradiation is at 650 nm for specified time. Accumulated irradiation time (from top to bottom): 0 min, 1 min, 3 min, 5 min, 10 min, 20 min and 40 min, respectively.

reactivity toward singlet oxygen because of the electrophilic nature of singlet oxygen and such selectivity over other chemical entities has been reported (36). In the case of some of the very reactive singlet oxygen acceptors such as DPIBF, however, auto-oxidation by ground state oxygen was observed. We did oxygen and nitrogen flow-through tests to check the contribution of O₂/N₂ on the emission signal decrease of DMA and it turns out to be negative. Additionally, our preliminary results (Y. Koo and R. Kopelman, in preparation) show that other ROS, such as hydroxyl radical or superoxide, barely diffuse into the ORMOSIL nanoparticles. The combination of the above points indicates that the DMA signal decrease is indeed caused by the production of singlet oxygen. This tends to confirm that the singlet oxygen produced by photo-irradiation is mainly responsible for the fluorescence emission decrease of DMA.

If one could figure out the concentrations of DMA inside the ORMOSIL matrix before and after the exposure to the singlet oxygen, the absolute amount/concentration of singlet oxygen could be determined, based on the molar ratio of 1:1 between endoperoxide and singlet oxygen and the assumption that the emission intensity of DMA is proportional to the concentration of DMA. This approach turns out to be ideal for the detection of intracellular singlet oxygen production (3,4).

Another analytical approach for singlet oxygen detection by chemical probe is to use the kinetics model proposed by Moreno *et al.* (30). Based on the model, one gets

$$[\text{DMA}] = [\text{DMA}]_{t=0} \cdot \exp(-k \cdot t)$$

And $k = \Phi^{1\text{O}_2} k_2 I^{\text{abs}}/k_1$, with the assumption of $k_1 \gg k_2 \cdot [\text{DMA}]$, where k_1 is the singlet oxygen solvent quenching rate constant and k_2 is the reaction rate constant of DMA with singlet oxygen in ORMOSIL. Both k_1 and k_2 may be affected by the ORMOSIL matrices. Because of the fact that the bulk of the suspension is water, with only a relatively small portion of the volume taken up by ORMOSIL nanoparticles, and the singlet oxygen molecules are mostly produced in water and have to diffuse through the water to get into the ORMOSIL matrices to be detected, we use the k_1 value for water, which is $3 \times 10^5 \text{ s}^{-1}$ (37). The k_2 value will be slightly different from that in water because the ORMOSIL matrices affect the lifetime and the solubility of ¹O₂ as described in the comparison with free ADPA part below. The matrix effects are under further study and preliminary results showed that the high k_2

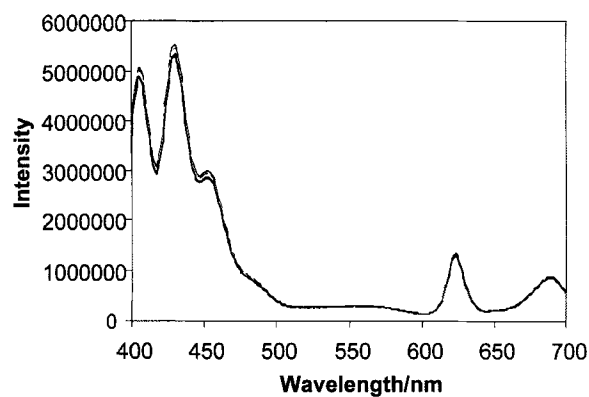


Figure 4. Control 2: Emission spectra (excited at 380 nm) of 0.5 mL of the ORMOSIL PEBBLE suspension mixed with 10 μM MB in distilled water, without 650 nm irradiation. Accumulated irradiation time (from top to bottom): 0 min, 1 min, 3 min, 5 min, 10 min, 20 min and 40 min, respectively.

value of DMA in water was maintained in ORMOSIL matrices (K. Gwangseong, Y. Cao, Y. E. Koo, W. Tang and R. Kopelman, in preparation). If we approximate the k_2 value as $9.1 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$, the value for DMA reacting with singlet oxygen in water (27) and the DMA concentration in the tested solution is estimated to be around 10^{-7} M , based on the number of dye molecules added during the synthesis, the experimental condition for this work meets the assumption made above.

With the caveat that the fluorescence intensity of DMA is proportional to the concentration of DMA, one can plot the semilog of the peak ratios of DMA/OEP against time and thus obtain the k value for each irradiation case. We note that indeed the fluorescence of the DMA free-dye was found to vary linearly with that of the concentration below 10^{-6} M (data not shown). These k values can be used for quality control of singlet oxygen production by the same PDT agents when tested with the same amount of PS. Still, it cannot be applied for comparing different PS values. However, one needs to note here that the k value by itself cannot be directly used to determine the absolute amount of singlet oxygen detected or produced. Therefore our approach in this work is to compare the relative k values for different experimental situations, where only one parameter is varied. This approach has been used to compare the singlet oxygen production efficiency by MB when the same amount of MB is either dissolved in distilled water or incorporated in PAA nanoparticles.

Comparisons between ratiometric ORMOSIL nanoprobe and free dye probe ADPA

As described in the experimental sections, two sets of independent experiments have been done to compare the performances of these ORMOSIL nanoprobe and free-dye ADPA probes. ADPA and DMA are quite similar to each other in terms of chromophore structures, fluorescence quantum yield, excitation/absorption/emission spectra and the reaction rate constant (k is $1 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ for ADPA in water) (30), which makes the comparison easier and more straightforward.

In the experimental setups of Figs. 5 and 6, all conditions are the same except that in Fig. 5, $1 \times 10^{-4} \text{ mg/mL}$ free MB molecules in distilled water are irradiated at 650 nm to release singlet oxygen whereas in Fig. 6, the same number of MB molecules are incorporated inside PAA particles. When Figs. 5 and 6 are

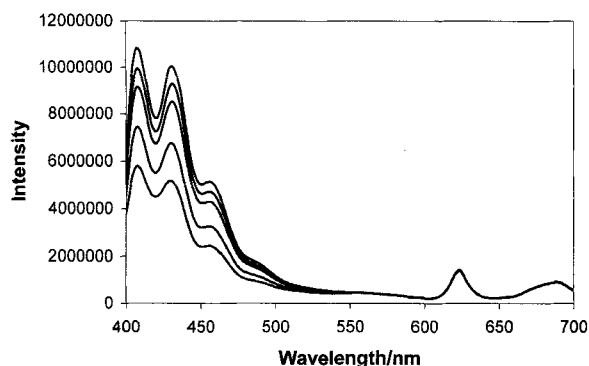


Figure 5. Emission spectra (excited at 380 nm) of 1×10^{-4} mg/mL of free MB in distilled water and DMA ORMOSIL nanoprobe upon light illumination at 650 nm. The same quantity of blank PAA nanoparticles (as in Fig. 6) was added to the free-dye mixture to minimize the light scattering interference caused by the PAA nanoparticles. Accumulated irradiation time (from top to bottom): 0 min, 1 min, 2 min, 5 min and 20 min, respectively.

compared, as shown in Fig. 7, the k value for the free-MB dye case is $0.0005 \text{ (s}^{-1}\text{)}$ whereas it is $0.0002 \text{ (s}^{-1}\text{)}$ for the same amount of MB dyes incorporated inside PAA particles. This tells us that about 150% more singlet oxygen is detected by the DMA ORMOSIL nanoprobe in the first 5 min for the sample of free MB molecules than when the same number of dye molecules are incorporated inside PAA particles (both samples are irradiated at 650 nm).

Similar experiments were done by Tang *et al.* using free ADPA singlet oxygen probes (34). However, they reported quite different results: 20% more singlet oxygen was detected for a free-MB sample than for the MB-loaded PAA sample, compared to 150% in this paper. We believe that this large difference (20% vs 150%) is caused by the diffusion of ADPA into and through the PAA nanoparticles. ADPA is very soluble in water and PAA is very hydrophilic. Thus we believe that ADPA can readily diffuse through the PAA matrix and therefore can have direct contacts with the MB molecules trapped inside, which means that ADPA can detect most of the singlet oxygen released by the MB molecules entrapped inside PAA. Therefore one only observes a minor change when the same amount of MB is incorporated inside PAA.

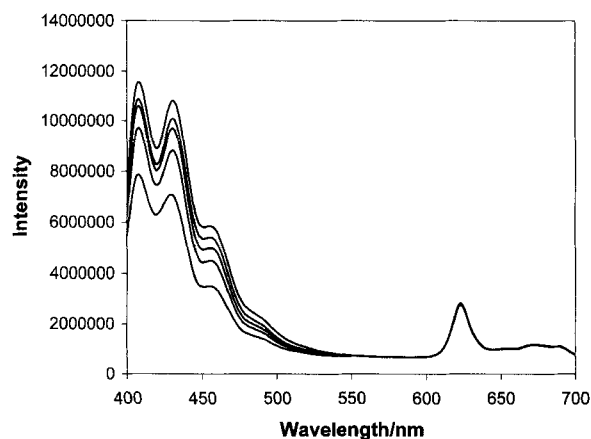


Figure 6. Emission spectra (excited at 380 nm) of 1 mg/mL MB-incorporated PAA in distilled water upon light irradiation at 650 nm. The overall concentration of MB is 1×10^{-4} mg/mL in distilled water. Accumulated irradiation time (from top to bottom): 0 min, 1 min, 2 min, 5 min and 20 min, respectively.

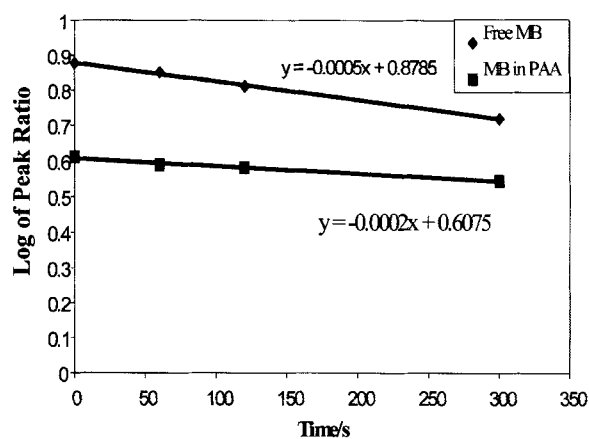


Figure 7. The semilog plot of peak ratio between DMA and OEP against time and the linear fits, used to figure out the k values for the singlet oxygen detection cases for free MB in water (top line, R^2 value is 0.997) and MB incorporated inside PAA nanoparticles (bottom line, R^2 value is 0.988).

However, the above described fabricated ratiometric ORMOSIL nanoprobe can only detect the singlet oxygen coming out of the PAA matrix, which is only a small portion of the singlet oxygen produced by MB incorporated inside PAA, upon light illumination. The above comparison provides some evidence that these new ratiometric nanoprobe can differentiate between the singlet oxygen coming out of the nanoparticles and the overall singlet oxygen production. This is extremely important, because in PDT what really matters is the amount of singlet oxygen coming out of the matrix, not the overall production of singlet oxygen. Our lab has been actively developing photosensitizer-loaded nanoprobe for PDT (31–33,38) and the singlet oxygen nanoprobe described here could become a very useful tool for determining the singlet oxygen production efficiency of the above PDT nanoprobe.

The conditions of the experimental setups depicted in Figs. 8 and 9 are identical except that, in Fig. 8's setup, the ORMOSIL DMA nanoprobe are used as singlet oxygen probes, whereas in Fig. 9's setup, free ADPA dye probes are used to detect the singlet oxygen. Again, one assumption here is that the concentration of DMA inside ORMOSIL is very close to that of the free ADPA

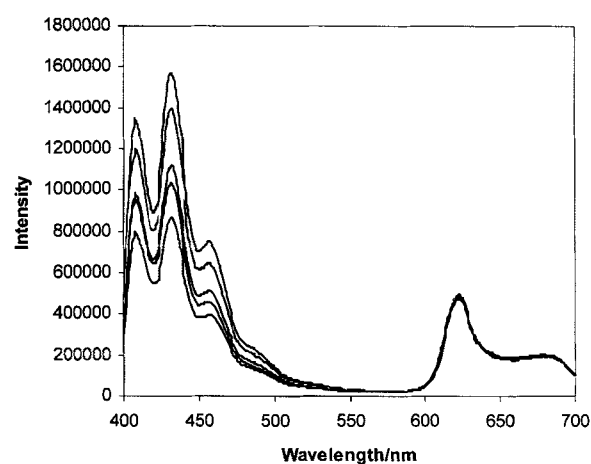


Figure 8. Detection of singlet oxygen produced by amine-functionalized photofrin-incorporated PAA particles via DMA ORMOSIL nanoprobe. Accumulated irradiation time (from top to bottom): 0 min, 1 min, 5 min, 10 min, 20 min, 30 min and 120 min, respectively.

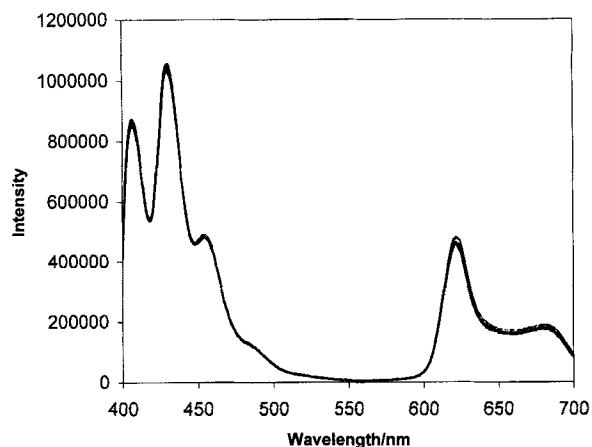


Figure 9. Detection of singlet oxygen produced by amine-functionalized photofrin-incorporated PAA particles via 10 μM ADPA in water. Accumulated irradiation time (from top to bottom): 0 min, 1 min, 3 min, 5 min, 10 min, 20 min and 30 min, respectively.

probes. This assumption can be roughly justified by the emission intensity comparison of both DMA and ADPA in Figs. 8 and 9.

What is striking, after comparing Figs. 8 and 9, is that the DMA signal decreases about 50% after 20 min of irradiation at 630 nm, whereas the ADPA's emission stays almost constant during the whole process, which means that almost no singlet oxygen is detected by the ADPA probes in this case.

It should be noted here that the amine-functionalized photofrin-loaded PAA particles are more hydrophobic than are ordinary PAA nanoparticles, which is necessary to retain the photofrin molecules; thus it will be very difficult for ADPA molecules to get into the PAA matrix in this scenario. We believe that the much longer lifetime of singlet oxygen inside ORMOSIL, compared with distilled water, singlet oxygen's higher solubility in ORMOSIL than in water and ORMOSIL's high permeability to singlet oxygen, and the higher reaction rate constant (about eight times higher) for DMA, compared to ADPA, play significant roles in explaining this result. The lifetime of singlet oxygen varies significantly from one solvent to another (39). It has also been demonstrated that the lifetime of singlet oxygen in a homogeneous polymer is practically the same as in a liquid solution consisting of analogous molecular structures (1). The lifetime of singlet oxygen in silicone rubber is estimated to be about 51 μs (1). As

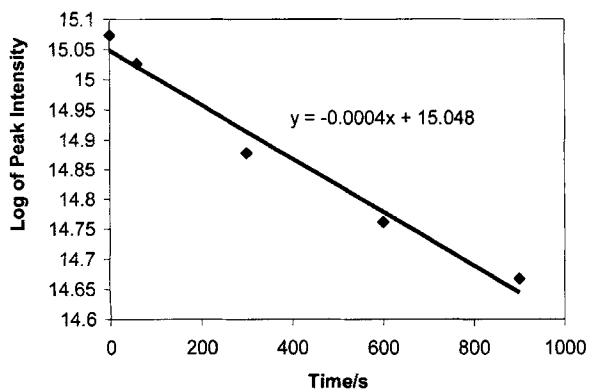


Figure 10. The semilog plot of DMA peak at 432 nm against time and the linear fit (R^2 value is 0.987) used to figure out the k value for 0.25 mg/mL amine-functionalized photofrin-PAA particles in water.

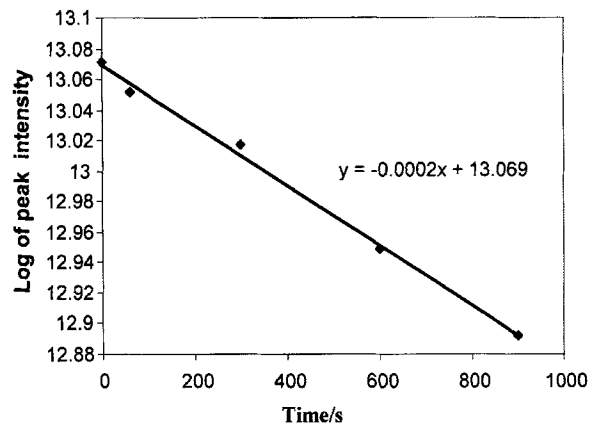


Figure 11. The semilog plot of DMA peak at 432 nm against time and the linear fit (R^2 value is 0.996) used to figure out the k value for 1.0 mg/mL amine-functionalized photofrin-PAA particles in water.

demonstrated in Scheme 1, the chemical structures of ORMOSIL and silicone rubber are quite similar to each other. Therefore we believe that it is quite reasonable to estimate the lifetime of singlet oxygen in ORMOSIL to be about 50 μs , which is about 25 times longer than in water. Furthermore, as demonstrated in our previous work (26), ORMOSIL is a very efficient oxygen-permeable polymer with a low diffusion barrier for oxygen, and we believe that the use of ORMOSIL as a matrix favors a scenario in which singlet oxygen molecules will partition in ORMOSIL more than in water and thus will move preferentially into this hydrophobic matrix, out from the aqueous solution and the hydrophilic PAA nanoparticles. Therefore both the relatively longer lifetime of $^1\text{O}_2$ in ORMOSIL than in water and the higher solubility of $^1\text{O}_2$ in ORMOSIL than in water significantly enhance these singlet oxygen nanoprobe's sensitivity.

The more photosensitizers, the more singlet oxygen?

The amount of singlet oxygen produced is dependent upon the type of photosensitizers, the concentration of the photosensitizers, the irradiation light intensity and the nature of the solvents. Also, in many cases it is true that the amount of singlet oxygen produced is proportional to the concentration of the photosensitizers. However in some rare situations, this may not be true, as demonstrated below.

When 100 μM MB in water is used as the singlet oxygen source, almost no signal decrease (K is almost 0) is observed for both ratiometric ORMOSIL nanoprobe and ADPA free-dye probes, even after about 60 min of irradiation at 650 nm. However, as presented above, the DMA's signal decreases about 73% after 40 min of irradiation when 10^{-4} mg/mL (0.34 μM) MB in water is used to produce singlet oxygen.

For amine-functionalized photofrin-incorporated PAA particles, as shown in Figs. 10 and 11, the K value is 0.0004 (S^{-1}) for the 0.25 mg/mL photofrin PAA sample; however it is 0.0002 (S^{-1}) for a 1.0 mg/mL sample, which means more singlet oxygen production for a 0.25 mg/mL sample than for a 1.0 mg/mL sample after 30 min of irradiation. Again this appears to be counterintuitive.

Therefore, we have observed for both the MB and photofrin samples that higher concentrations of photosensitizers do not necessarily produce more singlet oxygen, but rather the opposite. We believe that this is caused by the fact that the higher concentrations of MB and photofrin favor the formation of aggregates, *e.g.* dimers or higher order aggregates, which do not produce singlet oxygen

upon illumination (40, K. Gwangseong, Y. Cao, Y. E. Koo, W. Tang and R. Kopelman, in preparation).

Interference by oxygen and NO

Experiments have been done to check the interference of oxygen molecules and NO on these ORMOSIL nanoprobe. Our data show that the peak ratios between DMA and OEP are little affected by a continuous flow of oxygen, nitrogen or NO through the aqueous ORMOSIL suspensions.

Leaching

Leaching is a major concern for PEBBLE nanosensors. The performance, accuracy and lifetime of our nanosensors are directly related to the degree of leaching. Factors such as the molecular size of the dye (smaller molecules more readily leach out through the pores) and the pore size of the matrix (bigger pores favor leaching), and especially the relative solubility of the dyes in the matrix and in water, play very important roles in defining the degree of leaching. In terms of solubility, the solubility of DMA and OEP is much larger in ORMOSIL than in water, as they are hydrophobic and are almost insoluble in water, and this should minimize the leaching. These factors have of course been taken into consideration during the design of these nanosensors. In the tests for leached-out dyes, no leaching could be detected, as none of the spectra taken on these samples show any fluorescence signal being three times higher than the background noise at the emission wavelengths of both dyes entrapped inside the ORMOSIL matrix. This result shows the excellent stability of these ORMOSIL singlet oxygen nanoprobe, in terms of dye leaching, over a period of 3 days.

Photostability

Our results show that there is almost no emission intensity change for DMA after continuous illumination for 30 min. Thus photobleaching should not be a problem for DMA. The same experiment was carried out for monitoring the emission of OEP, and there is about a 10% signal decrease after continuous illumination at 380 nm for 30 min, but only about a 2% signal decrease after continuous illumination at 650 nm for 30 min. It should be noted that, during a typical run, using pulsed illumination, the total light exposure time is less than 30 s. Thus, in summary, these ORMOSIL singlet oxygen nanoprobe are sufficiently photostable over their operational time scale.

CONCLUSIONS

Because most singlet oxygen molecular probes with high reaction rate constants are hydrophobic, they have been conventionally modified through synthesis to be hydrophilic (such as ADPA) so that they could be used for singlet oxygen detection in aqueous solutions or biological environments. However, the performance of these modified naked molecular probes is often limited by the relatively short lifetime and poor solubility of $^1\text{O}_2$ in aqueous solutions, in addition to the possible dye toxicity problem. In this paper, the highly sensitive singlet oxygen molecular DMA probes are encapsulated inside ORMOSIL polymer matrix, thus making the detection of $^1\text{O}_2$ take place inside ORMOSIL. This approach has demonstrated significant advantages over the naked dye probes such as ADPA, and we believe that this is because of the following conditions: (1) the much longer lifetime of singlet oxygen inside the ORMOSIL matrix than in water; (2) singlet oxygen's higher

solubility in ORMOSIL than in water and ORMOSIL's high permeability to singlet oxygen; and (3) the protection of the dyes by the matrix. Therefore, the introduction of ORMOSIL as the matrix not only protects the dye molecules from direct contact with the outside environment but also significantly enhances its sensitivity to singlet oxygen. These nano-PEBBLE probes appear to be the first ratiometric singlet oxygen nanoprobe ever developed. These nanoprobe have been used to monitor the singlet oxygen production by photosensitizer-loaded PDT nanoprobe, with significant success, and they have revealed the dimmer- or higher-order aggregation effects of the PDT platforms. Another unique feature about these nanoprobe is that they only detect the singlet oxygen coming out of the photodynamic nanoprobe, which is critical in determining the singlet oxygen production efficiency of the nanoprobe. Thus these new nanoprobe could become ideal tools for quality control in PDT, for both qualitative and quantitative analysis and monitoring. Because of their high sensitivity (resulting from a combination of the highly sensitive DMA dye molecules with the more favorable singlet oxygen detection environment inside the ORMOSIL matrix) as well as their good biocompatibility, these nanoprobe are quite promising for intracellular singlet oxygen detection.

Acknowledgements—This work was sponsored by NIH/NCI contract N01-CO-07013 and by NIH grant EB00250. We also thank the University of Michigan Electron Microbeam Analysis Laboratory for use of the TEM and SEM.

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