Structure-activity study of furyl aryloxazole fluorescent probes for the detection of singlet oxygen

Renzo P. Zanocco, Roger Bresoli-Obach, Santi Nonell, Else Lemp, Antonio L. Zanocco

1 Departamento de Química Orgánica y Fisicoquímica, Facultad de Ciencias Químicas y Farmacéuticas, Universidad de Chile, Santiago, Chile, 2 Institut Químic de Sarrià, Universitat Ramon Llull, Barcelona, Spain

* These authors contributed equally to this work.
* renzo.zanocco@ciq.uchile.cl (RPZ); azanocco@ciq.uchile.cl (ALZ)

Abstract

In this study, we report the synthesis and the photochemical behavior of a series of new "click-on" fluorescent probes designed to detect singlet oxygen. They include a highly fluorescent chemical structure, an aryloxazole ring, linked to a furan moiety operating as singlet oxygen trap. Their activity depends on both the structure of the aryloxazole fluorophore and the electron-donating and electron-accepting properties of the substituents attached to the C-5 of the furan ring. All probes are selectively oxidized by singlet oxygen to give a single fluorescent product in methanol and produce negligible amounts of singlet oxygen themselves by self-sensitization. The most promising dyad, (E)-2-(2-(5-methylfuran-2-yl)vinyl)naphtho[1,2-d]oxazole, FN-6, shows outstanding reactivity and sensitivity: it traps singlet oxygen with a rate constant \((5.8 \pm 0.1) \times 10^7 \text{ M}^{-1} \text{s}^{-1}\) and its fluorescence increases by a factor of 500 upon reaction. Analysis of the dyads reactivity in terms of linear free energy relationships using the modified Swain and Lupton parameter \(F\) and the Fukui condensed function for the electrophilic attack, suggests that cycloaddition of singlet oxygen to the furan ring is partially concerted and possibly involves an exciplex with a "more open" structure than could be expected for a concerted cycloaddition.

Introduction

Singlet oxygen, \((\text{O}_2(1\Delta_g))\), hereafter \(^1\text{O}_2\), the first electronic excited state of molecular oxygen, is a well-known reactive oxygen species (ROS), that can diffuse and oxidize several types of biomolecules such as proteins, nucleic acids and lipids constituent of cell membranes [1–8]. Due to this reactivity, \(^1\text{O}_2\) is recognized as an important ROS capable of promoting a large variety of cell responses [9–11]. On the other hand, \(^1\text{O}_2\) is one of the most central cytotoxic ROS produced by exposure of sensitizers to light in photodynamic therapy, PDT [12–15].

To understand in depth the mechanisms associated with cellular stimuli originated by the presence of \(^1\text{O}_2\) and/or to appropriately control the photooxidation process in PDT treatments, the detection and quantification of \(^1\text{O}_2\) is undoubtedly one the most relevant and
critical factors. Common $^{1}\text{O}_2$ detection methods include electron paramagnetic resonance [16–18], chemiluminescence [19–21] and fluorescence spectroscopy [22–29]. These methods are based on the observation of a signal produced by a probe attached to a $^{1}\text{O}_2$ chemical acceptor that responds to the oxidation stage of the acceptor. Typical chemical acceptors are electron-rich dienes, naphthalenes, anthracenes and furans.

Furan derivatives such as 2,5-diphenyl-3,4-isobenzofuran (DPBF) or 2,5-dimethylfuran, have been widely used as a $^{1}\text{O}_2$ scavengers since the 70s [30,31]. They are advantageous chemical traps of $^{1}\text{O}_2$ because they can be quantified by using routine analytical techniques such as spectrophotometry or gas chromatography, they react with $^{1}\text{O}_2$ mainly via a chemical channel to form endoperoxides with a minimal or null contribution of physical quenching [32], and the overall quenching rate constant, $k_q$, shows a very modest solvent dependence. In addition, furans react specifically with $^{1}\text{O}_2$ and thus are ideally suited to develop selective $^{1}\text{O}_2$ probes [33,34].

Our group has long been interested in the detection and quantification of $^{1}\text{O}_2$ using furan derivatives [27,35,36]. We recently started a program for the design, synthesis and study of new “click-on” furan-based fluorescent probes for $^{1}\text{O}_2$ sensing [27]. The novel molecular entities are dyads comprising an aryloxazole fluorescent moiety linked to a furan trap. In their native state, the inherently strong fluorescence of the aryloxazole moiety is quenched by the electron-rich furan. Upon reaction with $^{1}\text{O}_2$, the furan is oxidized and the quenching process ceases to operate, restoring the intrinsically strong fluorescence of the aryloxazole ring. Moreover, the absorption and fluorescence spectra of the native and oxidized forms of the probe are different, which allows the selective photoexcitation of either form, thereby further enhancing the fluorescence contrast to an unmatched level [22,27].

Having established the proof of concept for this new family of probes, we have now set out to explore the effect of structural modifications on the performance of aryloxazole-furan probes and we report herein the results of our studies. We aimed at rationalising the fluorescent response of the probes and their reactivity towards $^{1}\text{O}_2$ to understand the factors affecting their performance. Thus, we have been able to design an optimum candidate by modifying the structure of the aryloxazole moiety (Fig 1A) and by including different substituents at the C-5 position of the furan ring (Fig 1B).

**Materials and methods**

**Materials**

Perinaphthenone (Sigma-Aldrich), new methylene blue (NMB, Sigma-Aldrich), anthracene (Sigma-Aldrich), naphthalene (Sigma-Aldrich), 1-methyl-2-pyrrolidone, (Sigma-Aldrich), 2-methylbenzo[1,2-d]oxazole (Sigma-Aldrich), 2-methylnaphtho[1,2-d]oxazole (AK Scientific, Inc.), 2-methylnaphtho[3,2-d]oxazole (TCI), 2-aminophenol (Sigma-Aldrich), 1-aminomethyl-2-naphthol hydrochloride (Sigma-Aldrich), 2-amino-3-naphthol (Sigma-Aldrich), furfural (Sigma-Aldrich), 5-methylfurfural, (Sigma-Aldrich), 5-bromofurfural (Sigma-Aldrich), 5-phenylfurfural (Sigma-Aldrich) and furfuryl chloride (Sigma-Aldrich) were used as received. All solvents used were UV or HPLC grade.

**Spectroscopic measurements**

Absorption spectra were recorded on a UV4 spectrophotometer (Unicam). Fluorescence emission spectra were recorded on a Quattro III (OBB) or on a PC1 (ISS) spectrofluorometers. Fluorescence quantum yields ($\Phi_F$) were determined by the comparative method described by Eaton and Demas [37,38], using quinine sulphate in 0.1 N sulphuric acid ($\Phi_F = 0.55$) or naphthalene in ethanol ($\Phi_F = 0.21$) as references [39]. The absorbance of sample and reference
solutions was set below 0.1 at the excitation wavelength and the fluorescence emission spectra were corrected using rhodamine G as reference. Sample quantum yields were evaluated using Eq (1): 

$$\Phi_X = \left( \frac{Grad_X}{Grad_{Act}} \right) \left( \frac{\eta_X^2}{\eta_{Act}^2} \right) \Phi_{Act}$$  \hspace{1cm} (1)$$

where $Grad_X$ and $Grad_{Act}$ are the slope of integrated fluorescence vs. absorbance plots for the sample and the actinometer, respectively, and $\eta_X$ and $\eta_{Act}$ are the refractive index of sample and actinometer solutions, respectively. All measurements were carried out in nitrogen-purged solutions at $(20.0 \pm 0.5)^\circ C$. Fluorescence decays were recorded with a time-correlated single photon counting system (Fluotime 200; PicoQuant GmbH, Berlin, Germany) equipped with a red-sensitive photomultiplier. Excitation was achieved by means of a 375 nm picosecond diode laser working at 10 MHz repetition rate. The counting frequency was maintained always below 1%. Fluorescence lifetimes were analyzed using the PicoQuant FluoFit 4.0 software.

Light irradiation of NMB at 660 ± 5 nm in steady-state experiments, was performed in 1 cm spectrophotometer cuvettes using a ThorLabs M660L4 LED light source (15 mW cm−2). The same setup was employed to determine the chemical reaction rate constant, $k_r$, in methanol. The distance between the light source and the cell was set for each experiment so that the initial substrate concentration would diminish about 50% in 15 min. Dyad consumption was evaluated by observing the decrease of absorbance over time and $k_r$ values were derived from the slopes of the linear pseudo first-order photoconsumption plots using 9,10-dimethylanthracene ($k_r = 6.3 \times 10^7 \text{ M}^{-1} \text{s}^{-1} \text{ [40]}$) as reference. Photooxidation products for HPLC-MS analysis were prepared by long-term irradiation of the samples (> 90% conversion).
Singlet oxygen measurements

The phosphorescence of $^1$O$_2$ was detected by means of a customized PicoQuant Fluotime 200 system. A Diode-pumped Q-switched laser (Pulselas-A-660-50, AlphaLas) working at 2 kHz repetition rate was used for excitation of NMB at 660 nm. The luminescence exiting from the side of the sample was filtered by a 1100 nm cut-off filter (Edmund Optics) and a narrow bandpass filter at 1270 nm (NB-1270-010, Spectrogon) to remove any scattered laser radiation. A plane-convex lens (23 mm diam. X 75 mm) was used to focus the light emitted onto the photomultiplier window. A near-IR sensitive photomultiplier tube assembly (H1033A-45; Hamamatsu Photonics) was used as detector. Photon counting was achieved with a multichannel scaler (PicoQuant’s Nanoharp 250). Time-resolved emission signals $S_t$ were analyzed using the PicoQuant FluoFit 4.0 data analysis software to extract lifetime ($\tau_t$ and $\tau_\Delta$) and amplitude ($S_0$) values. Quantum yields for $^1$O$_2$ production ($\Phi_\Delta$) were calculated from the amplitudes using the following Eqs (2)–(4):

$$S_t = S_0 \left( \frac{\tau_\Delta}{\tau_\Delta} \right) \left( e^{\frac{t}{\tau_\Delta}} - e^{\frac{t}{\tau_\Delta}} \right)$$

$$S_0 \propto \Phi_\Delta$$

$$\Phi_\Delta(\text{sample}) = \Phi_\Delta(\text{ref}) \left( \frac{S_0(\text{sample})}{S_0(\text{ref})} \right)$$

Perinaphthenone was used as reference for which $\Phi_\Delta = 1$ was taken [41]. The rate constant for $^1$O$_2$ quenching by the dyads ($k_q$) was determined by measuring the $^1$O$_2$ lifetime as a function of the dyad concentration. $^1$O$_2$ was generated by photoexcitation of a 50 μM NMB solution at 665 nm and the concentration of the dyads was varied in the range (0.1–1 mM). A plot of the reciprocal lifetime, $1/\tau_\Delta$, vs. the concentration of the dyad afforded $k_q$ as the slope of the linear fit, Eq (5),

$$\frac{1}{\tau_\Delta} = \frac{1}{\tau_\Delta} + k_q[Dyad]$$

where $\tau_\Delta^0$ is the $^1$O$_2$ lifetime in the neat solvent.

Synthesis of furyl aryloxazoles

Furyl derivatives of aryloxazoles were synthesized using the method of El’chaninov et al [42]. Typically, 1.1 mmol of amino arylalcohol and 1 mmol of furoyl chloride in 4 mL of dry 1-methyl-2-pyrrolidone were stirred under nitrogen by 1 h. Addition of 6 mL of cold water gives a precipitate that was filtered and washed with 20 mL of cold acetonitrile. Recrystallization of the solid in acetonitrile affords the product with high purity.

2-(furan-2-yl)benzo[1,2-d]oxazole, FN-1. 1H-NMR (400 MHz, DMSO-d$_6$): δ: 8.06 (dd, 1H, J = 1.8; 0.8 Hz, Fu-H$_3$), 7.75 (m, 1H, Ar-H$_{6,9}$, Ar-H$_2$), 7.45 (dd, 1H, J = 3.5; 0.8 Hz, Fu-H$_2$), 7.41 (m, 1H, Ar-H$_4$, Ar-H$_2$), 6.80 (dd, 1H, J = 3.5; 1.8 Hz, Fu-H$_2$); 13C-NMR (100 MHz, DMSO-d$_6$): δ: 155.07; 149.98; 147.58; 142.06; 141.56; 126.04; 125.53; 115.58; 113.25; 111.32; IR(KBr) v(cm$^{-1}$): 3420, 3108, 1634, 1450, 1014, 955; Elem. Anal.: exp. C 73.97%, H 4.18%, N 6.75%; calc. C 71.35%, H 3.81%, N 7.56%; MS(ESI) m/z: 186.05.

2-(furan-2-yl)naphtho[2,3-d]oxazole, FN-3. 1H-NMR (400 MHz, DMSO-d$_6$): δ: 8.29 (s, 1H, Ar-H$_4$), 8.22 (s, 1H, Ar-H$_5$), 8.14 (dd, 1H, J = 1.7; 0.8 Hz, Fu-H$_1$), 8.06 (m, 2H, Ar-H$_2$), 7.58 (dd, 1H, J = 3.6, 0.8 Hz, Fu-H$_2$), 7.51 (m, 2H, Ar-H$_6$), 6.85 (dd, 1H, J = 3.6; 1.7 Hz, Fu-
H), 13C-NMR (100 MHz, DMSO-d6) δ: 156.83, 148.87; 148.29; 141.83; 141.50; 131.82; 131.69; 128.73; 128.36; 126.06; 125.38; 117.41; 116.99; 113.50; 106.95, IR(KBr) υ(cm−1): 3418, 3051, 1652, 1305, 1080, 1011, Elem. Anal.: exp.: 76.18%, H 4.12%, N 6.16% calc. C 76.59%, H 3.68%, N 6.95%; MS (ESI+) m/z 235,90.

Synthesis of furyl vinyl aryloxazoles

Furyl vinyl aryloxazoles were obtained using the method of Zajac et al. [43], employing dimethylsulfoxide as solvent and KOH as the base. Typically 1.1 mmol of 2-methylaryloxazole and 1 mmol of furfural were dissolved in 2 mL of dimethylsulfoxide. Then, were added 0.124 mL of an aqueous solution of KOH 50% and the mixture was stirred by 1 h at room temperature. Addition of 10 mL of water afforded a yellow precipitate, which was filtered, washed with cold water and cold methanol and recrystallized from acetonitrile giving the product in high purity with yields between 50–60%.

(E)-2-(2-(furan-2-yl)vinyl)benzo[1,2-d]oxazole, FN-2. 1H-NMR (400 MHz, DMSO-d6); δ: 7.85 (dd, 1H, J = 1.8; 0.5 Hz, Fu-H1), 7.68 (m, 1H, Ar-H7), 7.62 (d, 1H, J = 16.1 Hz, Ar-H5), 7.36 (m, 2H, Ar-H6), 6.95 (dd, 1H, J = 3.4; 0.5 Hz, Fu-H2), 6.87 (d, 1H, J = 16.1 Hz, Vi-H4), 6.65 (dd, 1H, J = 3.4; 1.8 Hz, Fu-H3); 13C-NMR (100 MHz, DMSO-d6); δ: 162.53; 151.27; 150.23; 146.02; 142.21; 126.81; 125.81; 125.19; 119.93; 115.20; 113.29; 111.01; 106.95, IR(KBr) υ(cm−1): 3429, 3131, 2928, 1636, 1452, 1085, 942; Elem. Anal.: exp. C 71.11%, H 3.87%, N 7.32%; calc. C 73.92%, H 4.29%, N 6.63%; MS (ESI+) m/z 212.21.

(E)-2-(2-(furan-2-yl)vinyl)naphtho[2,3-d]oxazole, FN-4. 1H-NMR (400 MHz, DMSO-d6); δ: 8.23 (s, 1H, Ar-H7), 8.14 (s, 1H, Ar-H6), 8.05 (m, 2H, Ar-H9), 7.90 (dd, 1H, J = 1.7; 0.4 Hz, Fu-H1), 7.75 (d, 1H, J = 16.1 Hz, Vi-H4), 7.50 (m, 2H, Ar-H8), 7.03 (dd, 1H, J = 3.4; 0.5 Hz, Fu-H3), 6.93 (d, 1H, J = 16.1 Hz, Vi-H6), 6.68 (dd, 1H, J = 3.4; 1.8 Hz, Fu-H2); 13C-NMR (100 MHz, DMSO-d6); δ: 164.60; 151.25; 149.16; 146.46; 142.24; 131.84; 131.57; 128.73; 128.31; 128.22; 125.97; 125.21; 117.07; 116.03; 113.47; 110.80; 106.45, IR(KBr) υ(cm−1): 3432, 3052, 2926, 1646, 1248, 1017; Elem. Anal.: exp. C 77.98%, H 4.18%, N 5.41%; calc. C 78.15%, H 4.24%, N 5.36%; MS (ESI+) m/z 261.82.

(E)-2-(2-(furan-2-yl)vinyl)naphtho[1,2-d]oxazole, FN-5. 1H-NMR (400 MHz, DMSO-d6); δ: 8.36 (d, 1H, J = 8.3 Hz, Ar-H8), 8.09 (d, 1H, J = 8.2 Hz, Ar-H11), 7.95 (d, 1H, J = 8.8 Hz, Ar-H7), 7.89 (d, 1H, J = 8.9 Hz, Ar-H6), 7.87 (d, 1H, J = 1.7 Hz, Fu-H1), 7.59 (dd, 1H, J = 8.2; 6.9 Hz, Ar-H5), 7.49 (dd, 1H, J = 3.3 Hz, Fu-H3), 6.97 (d, 1H, J = 3.3 Hz, Fu-H2); 13C-NMR (100 MHz, DMSO-d6); δ: 162.09; 151.43; 150.09; 147.67; 145.88; 137.44; 131.30; 129.30; 127.77; 126.79; 126.04; 125.94; 125.90; 121.96; 114.92; 113.29; 111.44; 111.13; IR(KBr) υ(cm−1): 3428, 3056, 2919, 1635, 1463, 1018; MS (ESI+) m/z 261.88.

(E)-2-(2-(5-methyl-furan-2-yl)vinyl)naphtho[1,2-d]oxazole, FN-6. 1H-NMR (400 MHz, DMSO-d6); δ: 8.35 (d, 1H, J = 8.3 Hz, Ar-H8), 8.08 (d, 1H, J = 8.2 Hz, Ar-H11), 7.95 (d, 1H, J = 8.9 Hz, Ar-H7), 7.89 (d, 1H, J = 8.9 Hz, Ar-H6), 7.69 (ddd, 1H, J = 8.2; 6.9; 1.2 Hz, Ar-H9), 7.66 (d, 1H, J = 16.1 Hz, Vi-H6), 7.59 (ddd, 1H, J = 8.2; 6.9; 1.3 Hz, Ar-H10), 6.99 (d, 1H, J = 16.2 Hz, Vi-H4), 6.97 (d, 1H, J = 3.3 Hz, Fu-H3), 6.66 (dd, 1H, J = 3.4; 1.8 Hz, Fu-H2); 13C-NMR (100 MHz, DMSO-d6); δ: 162.37; 155.42; 150.09; 147.59; 145.88; 137.44; 131.30; 129.30; 127.77; 126.79; 126.04; 125.94; 125.90; 121.96; 114.92; 113.29; 111.44; 111.13; IR(KBr) υ(cm−1): 3428, 3056, 2919, 1630, 1483, 1020; Elem. Anal.: exp. C 77.012%, H 4.63%, N 4.99%; calc. C 78.53%, H 4.76%, N 5.09%; MS (ESI+) m/z 276.17.
7.60 (d, 1H, J = 16.1 Hz, Vi-H3), 7.59 (ddd, 1H, J = 8.2; 6.9; 1.3 Hz, Ar-H9), 7.00 (d, 1H, J = 3.0 Hz, Fu-H2), 6.98 (d, 1H, J = 16.3 Hz, Vi-H4), 6.79 (d, 1H, J = 3.5 Hz, Fu-H1); $^{13}$C-NMR (100 MHz, DMSO-$_d^6$) δ: 199.78; 161.83; 153.62; 147.73; 131.31; 129.32; 127.82; 126.96; 126.09; 125.92; 124.71; 121.6; 117.08; 115.40; 114.53; 111.72; 111.45; IR (KBr) υ(cm$^{-1}$); 3432, 3061, 1632, 1460, 1085, 1006; Elem. Anal. exp. C 60.22%, H 3.15%, 4.05%; calc. C 61.04%, H 3.41%, N 3.95%; MS (ESI+) m/z 339.89.

(E)-2-(2-(5-phenyl-2-furan-2-yl)vinyl)naphtho[1,2-d]oxazole, FN-9. $^{13}$C-NMR (100 MHz, DMSO-$_d^6$) δ: 162.30; 155.33; 151.16; 147.69; 137.55; 131.31; 129.47; 129.45; 129.40; 129.32; 128.87; 127.77; 126.75; 125.93; 125.41; 124.56; 124.46; 124.43; 121.96; 117.42; 111.44; 111.13; 109.39; IR(KBr) υ(cm$^{-1}$); 3430, 3036, 1628, 1470, 1027, 1006; Elem. Anal.: exp. C 82.36%, H 4.59%, N 4.10%, calc. C 82.3%, H 4.88%, N 3.99%.

**Results and discussion**

**Photophysical characterization of unsubstituted furyl aryloxazoles**

Compounds FN-1 to FN-5 are dyads composed by an unsubstituted furan ring linked to an aryloxazole moiety through either a single C-C bond or a vinyl bridge. The aim of studying this series of compounds was to determine the most promising aryloxazole and link structures to detect and quantify $^1$O$_2$. Absorption spectra of dyads FN-1, FN-2, FN-4 and FN-5 in solvents representative of the polarity scale are shown in Fig 2.

As can be observed in Fig 2 and from the data in Table 1, the position of lowest-energy band is insensitive to the solvent polarity, a behavior previously observed for related compounds [27,44]. Furthermore, the large value of the experimental molar absorption coefficients and molecular-orbital analysis of the minimum energy structures obtained from DFT calculations (6311g+dp orbital base) with Gaussian 04W indicate that the lowest energy bands correspond to $\pi-\pi^*$ transitions. Also, comparison of wavelength maxima of FN-2 (naphthalene-like), FN-4 (anthracene-like) and FN-5 (phenanthrene-like), shows that the greater aromaticity of the dyad the smaller the transition energy [45–47].

![Fig 2](https://doi.org/10.1371/journal.pone.0200006.g002)
On the other hand, these results indicate a strong electronic coupling between the oxazole moiety and the furan ring as visualized from the shape of the HOMO orbitals for FN-1, FN-4 and FN-5 (S1 Fig). The spectra of 2-(2-(furan-2-yl)ethyl)naphtho[1,2-d]oxazole, compound that has a saturated bridge between the aryloxazole group and the furan ring, shows a 40 nm blue-shifted absorption maximum [27].

The fluorescence spectra dyads in the same solvent set show a similar trend (Fig 2) with the position of the emission maxima nearly independent of the solvent polarity. Fluorescence quantum yields, measured using the Eaton and Demas method [37,38], are included in Table 2. For comparison, aryloxazoles typically show fluorescence quantum yields in the range of 0.7 to 1 [44,48,49] when aromatic substituents, different of furyl, are linked to the vinyl bridge, e.g. (E)-2-(4-methylstyryl)naphtho[1,2-d]oxazole [27], or in (E)-4-(2-(naphtho[1,2-d]oxazole-2-yl)vinyl)benzonitrile [44].

The furyl derivatives clearly behave in a different way. Furyl vinyl aryloxazoles, FN-2, FN-4 and FN-5, show a very low fluorescence quantum yield that, in addition, is essentially independent of the solvent polarity, i.e. when the furyl group is linked to the aryloxazole moiety through a vinyl bridge, the intrinsic fluorescence of aryloxazole is quenched by the furyl substituent [27]. This is a very important piece of information because the biological cell contains microenvironments of very different polarity. On the contrary, when the furyl substituent is linked directly to the oxazole ring, e.g., in FN-1 or FN-3, the intramolecular quenching process does not operate and the fluorescence quantum yields are in the range of 0.2 to 0.7.

### Reactivity of unsubstituted furyl aryloxazoles towards singlet oxygen

To evaluate the reactivity of the probes with $^{1}\text{O}_2$ we employed steady-state experiments to observe the evolution of absorption and emission spectra, and time resolved methods to measure changes in the decay kinetics of $^{1}\text{O}_2$ luminescence.

Fig 3, shows the evolution of absorption spectra of FN-2—FN-5, observed over the course of 40 min irradiation of a solution in the presence of NMB as sensitizer.

All probes showed a strong absorbance decrease of the lowest energy band in a light-dose dependent fashion, a result that indicates that reaction occurs between $^{1}\text{O}_2$ and the dyad. A careful examination of the spectra corresponding to FN-2, FN-4 and FN-5, also reveal clear

### Table 1. Absorption maxima and molar absorption coefficient of furyl aryloxazoles in representative solvents of different polarity.

| Solvent | $\lambda_{\text{max}}$ nm (ε/M$^{-1}$ cm$^{-1}$) |
|---------|-----------------------------------|
| ACN     | FN-1 304 (28472)                  |
|         | FN-2 354 (29712)                  |
|         | FN-3 332 (35587)                  |
|         | FN-4 334 (53381)                  |
|         | FN-5 357 (32500)                  |
| Benzene | FN-1 307 (29166)                  |
|         | FN-2 356 (23613)                  |
|         | FN-3 337 (34665)                  |
|         | FN-4 340 (47996)                  |
|         | FN-5 362 (20000)                  |
| DMF     | FN-1 307 (30753)                  |
|         | FN-2 356 (27230)                  |
|         | FN-3 337 (37468)                  |
|         | FN-4 339 (51523)                  |
|         | FN-5 362 (19722)                  |
| Methanol| FN-1 307 (35138)                  |
|         | FN-2 356 (30064)                  |
|         | FN-3 337 (37058)                  |
|         | FN-4 340 (53992)                  |
|         | FN-5 360 (27500)                  |

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### Table 2. Fluorescence quantum yields for unsubstituted furyl- and furyl vinyl aryloxazoles.

| Solvent | FN-1 | FN-2 | FN-3 | FN-4 | FN-5 |
|---------|------|------|------|------|------|
| ACN     | 0.20 | < 0.01 | 0.43 | 0.05 | 0.02 |
| Benzene | 0.30 | < 0.01 | 0.73 | 0.08 | 0.09 |
| DMF     | 0.26 | < 0.01 | 0.46 | 0.04 | 0.02 |
| Methanol| 0.17 | < 0.01 | 0.33 | 0.03 | 0.09 |

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isosbestic points at 305, 289 and 298 nm, respectively, suggesting that a major photoproduct is formed when a furyl vinyl aryloxazole is the substrate. HPLC experiments confirm that only one product is formed in the photooxidation reaction (S2 Fig).

The modification of the absorption spectrum upon oxidation is a distinctive feature of this family of probes as compared to the most popular ones. The main benefit is that it allows to selectively exciting the fluorescence of the oxidized probe, which leads to very high fluorescent enhancement. The fluorescence changes upon reaction with $^1\text{O}_2$ are shown in Fig 4.

FN-2 and FN-5 show an important fluorescence increase as photooxidation proceeds, 25- and 70-fold, respectively, exciting at their optimum wavelength (290 nm and 330 nm, respectively). In contrast, FN-4 shows only minor changes and FN-3, on the other hand, shows a

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**Fig 3.** Evolution of the lower energy absorption band of dyads FN-2 (A), FN-3 (C), FN-4(B) y FN-5(D), due to reaction with $^1\text{O}_2$ produced by photosensitization. Solvent: methanol; Sensitizer: NMB; Irradiation: Led, 660 nm. [https://doi.org/10.1371/journal.pone.0200006.g003](https://doi.org/10.1371/journal.pone.0200006.g003)

**Fig 4.** Changes in the fluorescence spectra of dyads FN-2 (A), FN-3 (B), FN-4(C) y FN-5(D), due to reaction with $^1\text{O}_2$ produced by photosensitization. Solvent: methanol; Sensitizer: NMB; Irradiation: Led, 660 nm, irradiation time: 40 min. Inset: Increase of relative fluorescence with the irradiation time. [https://doi.org/10.1371/journal.pone.0200006.g004](https://doi.org/10.1371/journal.pone.0200006.g004)
3-fold decrease in fluorescence. The total (physical and reactive) $^1\text{O}_2$ quenching rate constant ($k_q$) for FN-2 and FN-5 were measured by observing singlet oxygen luminescence decay in time-resolved experiments. Lineal Stern-Volmer plots were obtained from which $k_q$ values equal to $(1.37 \pm 0.06) \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ and $(1.16 \pm 0.04) \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ in ACN and $(0.35 \pm 0.02) \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ and $(0.27 \pm 0.02) \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ in methanol, respectively, were determined. Notice that the $k_q$ values in methanol, are 25- and 36-fold lower, respectively, than that of 2-methylfuran in the same solvent ($k_q = 9.9 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$) [34]. This important decrease in the furan reactivity is consistent with a substantial electronic delocalization of the furan $\pi$-electrons through the vinyl bridge.

Data reported in this section indicate that the most promising dyads for $^1\text{O}_2$-detection are FN-2 and FN-5, in which the furan and aryloxazole rings are linked by a vinyl bridge. Although FN-5 is slightly less reactive than FN-2, it shows a larger fluorescence enhancement. We therefore selected FN-5 as a basis for further development of fluorogenic $^1\text{O}_2$ probes and proceeded to evaluate the effect of substituents in C-5 of the furan ring.

### Photophysical characterization of substituted furyl vinyl naphthoxazoles

Compounds FN-6 to FN-9 (Fig 1B) are dyads related to FN-5 that include electron-withdrawing and electron-donating groups in position C-5 of the furyl moiety. Their absorption spectra are shown in Fig 5. As for compounds belonging to the first series, they are nearly independent on the solvent polarity, the absorption coefficients are in the range 25,000–50,000 M$^{-1}$ cm$^{-1}$, and molecular calculation analysis showed that the lowest energy transition is also $\pi,\pi^*$. It is worth mentioning that the $\lambda_{\text{max}}$ value is red-shifted by 30 nm for FN-9.

Likewise, their emission spectra (Fig 5) are also independent of the solvent polarity. Emission quantum yields are very low, particularly in polar solvents such as methanol, consistent with intramolecular charge transfer quenching the fluorescence emission [27]. Fluorescence lifetimes, Table 3, are around of 6 ns, independent of solvent and similar to those measured for other naphthoxazole derivatives [44].
Reactivity of substituted furyl vinyl naphthoxazoles towards singlet oxygen

The ability of probes FN-6 –FN-9 to react with \( ^1\text{O}_2 \) was evaluated using steady-state and time-resolved methods. \(^1\text{O}_2\) was photogenerated by irradiation of the sensitizer NMB in methanol. Changes in the absorption spectra of the probes upon reaction with \(^1\text{O}_2\) are shown in Fig 6. All dyads showed a notable loss of absorbance upon reaction with \(^1\text{O}_2\) and the relative rates of photobleaching were FN-6 ≈ FN-9 > FN-5 > FN-7 ≈ FN-8. Clear isosbestic points were observed, indicating a clean reaction with a single major product, which was confirmed by HPLC.

The reaction of FN-6 –FN-9 with \(^1\text{O}_2\) produces likewise a notable increase on the fluorescence intensity, which is accompanied by a hypsochromic shift of the emission maxima (Fig 7). The rate of fluorescence growth is largest for FN-6 and the fluorescence intensity of its oxidized form is >500-fold larger than that of the native form at the optimum excitation wavelength (330 nm).

The rate constants for overall \(^1\text{O}_2\) quenching by the dyads (\( k_q \)) were determined by time-resolved luminescence spectroscopy (S3 Fig). Likewise, the reactive rate constants \( k_r \) were obtained by comparing the rate of consumption of the dyads to that of the reference compound dimethylanthracene. Both are collected in Table 4.

### Table 3. Fluorescence quantum yields for substituted furyl vinyl naphthoxazoles. In parenthesis fluorescence lifetime in ns.

| Solvent | \( \Phi_F \) (\( \tau /\text{ns} \)) |
|---------|----------------------------------|
|         | FN-6 | FN-7 | FN-8 | FN-9 |
| ACN     | 0.004 (4.6) | 0.014 (5.9) | 0.014 (6.5) | 0.014 (4.3) |
| Benzene | 0.025 (6.3) | 0.083 (5.9) | 0.088 (6.8) | 0.109 (7.2) |
| DMF     | 0.007 (5.6) | 0.020 (5.9) | 0.021 (6.4) | 0.029 (6.8) |
| Methanol| 0.003 (6.0) | 0.008 (6.2) | 0.010 (6.3) | 0.010 (6.0) |

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Fig 6. Absorption spectrum changes for FN-6 (A), FN-7 (B), FN-8 (C) and FN-9 (D) upon reaction with \(^1\text{O}_2\) generated by irradiation of NMB in methanol.

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Values of $k_q$ in methanol included in Table 4 follow the order $FN-6 > FN-9 >> FN-7 > FN-5 > FN-8$. Thus, FN-6 and to a lesser extent FN-9, are excellent $^1O_2$ quenchers, with efficiency comparable to that of dimethylfuran ($k_q = 2.4 \times 10^7 \text{M}^{-1} \text{s}^{-1}$) and methylfuran ($k_q = 10.1 \times 10^7 \text{M}^{-1} \text{s}^{-1}$) in methanol [31]. Regarding the $k_r$ values, some observations are worth highlighting: (i) Their reactivity with singlet oxygen follows the same trend as their overall quenching efficiency, FN-6 and FN-9 being the most reactive compounds; (ii) the trend is not affected by solvent polarity, as expected for a cycloaddition $[4 + 2]$ reaction [50]; (iii) compared to the unsubstituted homologous FN-5, the methyl- and phenyl-substituted compounds FN-6 and FN-9 are 26- and 8-fold, respectively, more reactive; (iv) the least reactive compound is the Br-substituted FN-8 as could possibly be expected by the electron-accepting character of the substituent, which decreases the electron density on the C-5 position of furan ring; (v) although it is not the compound with the highest singlet oxygen trapping efficiency ($k_r / k_q = 56\%$), FN-6 shows nevertheless the highest reactivity and therefore appears as the best candidate for being use as a singlet oxygen fluorescent probe.

**Structure-reactivity relationships**

Substituents effects on the reactive rate constant were analyzed using the Hammett free energy relationship. Thus, the measured rate constants were correlated with $\sigma_m$ and $\sigma_p$ Hammett

|  | $k_q / 10^7 \text{M}^{-1} \text{s}^{-1}$ | $k_r / 10^7 \text{M}^{-1} \text{s}^{-1}$ | $k_r / k_q$ | $\Phi_\lambda$ |
|---|---|---|---|---|
| FN-5 | 1.2 ± 0.1 | 0.27 ± 0.03 | 22% | 0.003 |
| FN-6 | 5.7 ± 0.4 | 7.4 ± 0.1 | 5.8 ± 1 | 56% | 0.003 |
| FN-7 | 0.18 ± 0.02 | 0.55 ± 0.06 | 0.25 ± 0.03 | 45% | 0.009 |
| FN-8 | 0.13 ± 0.01 | 0.16 ± 0.01 | 0.15 ± 0.02 | 94% | 0.010 |
| FN-9 | 1.2 ± 0.1 | 4.1 ± 0.3 | 1.8 ± 0.2 | 26% | 0.004 |

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parameters [51] and with \( F \) and \( R \), the Swain and Lupton modified parameters [52], that redefine the substituent \( \sigma \)-parameter in terms of field effects, \( F \) (inductive and pure field) and the resonance effects, \( R \). Correlations of \( k_q \) with \( \sigma_m \), \( \sigma_p \) and \( R \) did not show clear trends. Better correlations in both methanol and ACN were obtained with the field parameter \( F \). Fig 8A shows that there is a good correlation between the inductive donor effect of the substituent and the dyads reactivity. Additional insight on the substituent effect upon the reaction rate and the electron density on the C-5 of the furan was obtained from correlations of the reaction rate with local Fukui coefficients. Condensed Fukui functions, \( f_k^{+/-} \), account for physicochemical properties of atoms or functional groups in a molecule, such as the nucleophilicity and electrophilicity of different sites in molecule [53–55]. Fig 8B shows that there is a reasonable correlation between the values of the rate constants and the value of the Fukui condensed function for the C-5 position of the furan ring. In addition, the most reactive dyads, FN-6, with methyl and FN-9, with phenyl substituents, show the highest \( f_k^- \) values and the lowest \( F \) values (low \( F \) values are associated with groups that have the greatest inductive effect).

These results suggest that the [4 + 2]-cycloaddition of \( ^1O_2 \) to the furan ring does not occur through a concerted mechanism, as for other furan derivatives [56], but possibly the attack proceeds in a partially concerted manner with a primary interaction of \( ^1O_2 \) with the center of greater electronic density, on the C-5 of the furan ring, as has been suggested by Lemp et al. in the cycloadDITION of \( ^1O_2 \) to mono-substituted anthracenes [57], forming an exciplex of a "more open" structure with a large charge separation (Fig 9).

Moreover, mass spectra of main products obtained in the photooxidation of FN-5 and FN-6 (S4 Fig) shows the same fragmentation pattern, suggesting a common reaction mechanism. Both mass spectra are compatible with a classical photooxidation mechanism that involves a [4 + 2]-cycloaddition of \( ^1O_2 \) to the furan ring followed of methanolysis to give the final product (S5 Fig).

**Self-sensitization of \( ^1O_2 \) by the dyads and reactivity towards other ROS**

A drawback of common \( ^1O_2 \) fluorescent probes is the evolution of fluorescence due to self-sensitization of \( ^1O_2 \) [58]. Other probes suffer from poor selectivity towards \( ^1O_2 \) and are capable of reacting also with other ROS. We investigated whether the naphthoxazole dyads are affected by the same problems. All dyads sensitized the production of \( ^1O_2 \) in methanol, however for FN-6 and FN-9 quantum yields (\( \Phi_{\Delta} \), Table 4) were 3-fold smaller than for the most popular probe SOSG [58]. Reactivity towards other ROS was also tested for the series FN-5 –FN-9. We found negative results for all probes against superoxide (KO2) and H2O2 (S6 and S7 Figs).

**Fig 8.** Linear correlation between the total rate constant for reaction of furyl vinyl naphthoxazoles and \( ^1O_2 \) with the \( F \) parameter of Swain and Lupton (A) and the Fukui condensed function for the electrophilic attack of \( ^1O_2 \) on the C-5 of furan ring (B). Solvents: ACN (black circles), methanol (red squares). Blue inverted triangles correspond to the reactive rate constant in methanol.

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After 50 min. the extent of reaction is lower than 1.5% indicating a high degree of specificity for $^1$O$_2$.

The results described in previous sections suggest that dyads which include a polycyclic aromatic naphthoxazole [1,2-d] system with a vinyl bridge at the 2-position of the heterocycle linked to a substituted furan ring with an alkyl group at carbon-5 are the ones with the highest fluorescent response. Accordingly, this structure could be employed to engineer a “click-on” probe to detect and quantify singlet oxygen in biological interest media, a potential solution if a quick response is needed in a test system. However, various characteristics of the biological systems must be considered to planning a successful probe: i) aromatic fluorescent molecules, such as naphthoxazole—furan dyads, could form complexes with proteins [59,60], altering the expected fluorescent increases after the $^1$O$_2$ addition to the scavenger moiety and restricting substantively probe internalization in cells. Different approaches have been proposed to solve this problem, such as probe binding to nanoparticles, which maintains its reactivity towards $^1$O$_2$, reducing the interaction with proteins [61–63]; ii) in biological media, a complex redox system maintaining cell viability is always present, which may affect the fluorescent response of the dyad if oxidant and/or reducing biomolecules react with the products yielded by the reaction between the dyad and $^1$O$_2$. Nevertheless, various “off-on” fluorescent dyads based on polyaromatic systems have been employed to detect $^1$O$_2$ in a diversity of cellular environments by the use of bioimaging techniques [59, 64–70]. Independent of the advantages or drawbacks of each probe, all of them afford highly repetitive results showing that, even in systems under oxidative stress, primary products of reaction with $^1$O$_2$ (typically an endoperoxide), are stable.

Fig 9. Exciplex structure for the partially concerted cycloaddition of $^1$O$_2$ to furyl-substituted vinyl naphthoxazole.

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under oxidizing or reducing conditions in cellular systems, a behavior also expected for furyl vinyl
napthoxazole endoperoxides; iii) the solubilization locus of the dyad can also modify the
dyad fluorescent response due to the diversity of microenvironments in cells. However, we
previously show that furan derivatives are very appropriate probes to monitor singlet oxygen
dynamics in systems mimicking biological interest organizations [35,36]. In these systems,
furan reactivity is nearly independent of the solubilization site although different sensitivities
of the dyad towards $^1$O$_2$ are expected, since the fluorescence quantum yield slightly increases
in non-polar solvents; iv) at first, lasers employed to excite the dyad in cells after the reaction
with $^1$O$_2$ could involve high light intensities and/or focusing and it is reasonable to assume
that high heat release could occur under these experimental conditions. These thermal effects
open a new reaction pathway whereby thermally decomposition of endoperoxide could pro-
duce $^1$O$_2$ and/or $^3$O$_2$. Nevertheless, high molar absorptivity’s of dyads described herein can
help to avoid severe thermal loads.

Conclusions

Summing up, “click-on” dyads constructed by linking a furan ring to an aryloxazole at 2-posi-
tion of the heterocycle via a vinyl bridge possesses appropriate properties to monitor and
quantify singlet oxygen in solution. They show very low fluorescence quantum yields, are
chemically stable, produce negligible amounts $^1$O$_2$ by self-sensitization, and react selectively
with $^1$O$_2$ at high rate. Photooxidation yields a major product in which the structure of the fluo-
rescent fragment is maintained but the non-radiative deactivation channel is cancelled upon
oxidation, which leads to increasing the relative fluorescence up to an unprecedented 500-fold
value. Reaction proceeds via a partially concerted [4 + 2] cycloaddition involving the forma-
tion of a loose or "open" structure exciplex, in which the oxygen is weakly bound to the furan
ring. Dyads described herein are promising platforms to develop probes for monitoring singlet
oxygen behavior in biological systems.

Supporting information

S1 Fig. HOMO orbitals for the dyads FN-1 (A), FN-4 (B) y FN-5 (C).
(TIF)

S2 Fig. HPLC chromatograms of FN-5 before (solid line) and after (dotted line) the photo-
oxidation reaction using NMB as sensitizer in methanol (95% FN-5 consumption).
(TIF)

S3 Fig. Stern-Volmer plots for reactions of FN-7, FN-8 and FN-9 with $^1$O$_2$ in methanol. Sens-
itizer NMB.
(TIF)

S4 Fig. Low-resolution mass spectra of the main photooxidation product of FN-5 (A) and
FN-6 (B) in methanol. Sensitizer: NMB.
(TIF)

S5 Fig. Reaction mechanism of photooxidation of furyl vinyl napthoxazoles.
(TIF)

S6 Fig. Reaction of FN-6 (A), FN-7 (B), FN-8 (C) and FN-9 (D) with an excess of H$_2$O$_2$ in
methanol. Reaction time 50 min.
(TIF)
S7 Fig. Reaction of FN-6 (A), FN-7 (B), FN-8 (C) and FN-9(D) with an excess of KO$_2$ in methanol. Reaction time 50 min.

(TIF)

**Author Contributions**

**Conceptualization:** Santi Nonell, Antonio L. Zanocco.

**Investigation:** Renzo P. Zanocco, Roger Bresoli-Obach, Else Lemp.

**Writing – original draft:** Renzo P. Zanocco.

**Writing – review & editing:** Santi Nonell, Antonio L. Zanocco.

**References**

1. Davies MJ. Singlet oxygen-mediated damage to proteins and its consequences. Biochem Biophys Res Commun. 2003; 305: 761–770. PMID: 12763058

2. Kang P, Foote CS. Formation of transient intermediates in low-temperature photosensitized oxidation of an 8-$^{13}$C-guanosine derivative. J Am Chem Soc. 2002; 124: 4865–4873. PMID: 11971737

3. Martinez GR, Gasparutto D, Ravanat J-L, Cadet J, Medeiros MHG, Di Mascio P. Identification of the main oxidation products of 8-methoxy-2'-deoxyguanosine by singlet molecular oxygen. Free Radical Biol Med. 2005; 38: 1491–1500.

4. Redmond RW, Kocheva IE. Spatially resolved cellular responses to singlet oxygen. Photochem Photobiol. 2006; 82: 1178–1186. https://doi.org/10.1562/2006-04-14-IR-874 PMID: 16740059

5. Weishaupt KR, Gomer CJ, Dougherty TJ. Identification of singlet oxygen as the cytotoxic agent in photoactivation of a murine tumor. Cancer Res. 1976; 36: 2326–2329. PMID: 1277137

6. Girotti AW. Photosensitized oxidation of membrane lipids: reaction pathways, cytotoxic effects, and cytoprotective mechanisms. J Photochem Photobiol B Biol. 2001; 63: 103–113.

7. Ye Y, Muller JG, Luo W, Mayne CL, Shallop AJ, Jones RA et al., Formation of $^{13}$C-, $^{15}$N-, and $^{18}$O-labeled guanidinohydantoin from guanosine oxidation with singlet oxygen. Implications for structure and mechanism. J Am Chem Soc. 2003; 125: 13926–13927. https://doi.org/10.1021/ja0378660 PMID: 14611206

8. Cadet J, Ravanat J-L, Martinez GR, Medeiros MHG, Di Mascio P. Singlet oxygen oxidation of isolated and cellular DNA: product formation and mechanistic insights. Photochem Photobiol. 2006; 82: 1219–1225. https://doi.org/10.1562/2006-06-09-IR-914 PMID: 16808595

9. Babu PSS, Manu PM, Dhanya TJ, Tapas P, Meera RN, Surendran A, et al. Bis(3,5-diiodo-2,4,6 trihydroxyphenyl) squaraine photodynamic therapy disrupts redox homeostasis and induce mitochondria-mediated apoptosis in human breast cancer cells. Sci Rep. 2017; 7: 42126. https://doi.org/10.1038/srep42126 PMID: 28169351

10. Bauer G. The antitumor effect of singlet oxygen. Anticancer Res. 2016; 36: 5649–5663. https://doi.org/10.21873/anticancerres.11148 PMID: 27793886

11. Westberg M, Bregnhoj M, Banerjee CH, Biázquez-Castro A, Breitenbach T, Ogilby T. Exerting better control and specificity with singlet oxygen experiments in live mammalian cells. Methods 2016; 109: 81–91. https://doi.org/10.1016/j.ymeth.2016.07.001 PMID: 27389303

12. Sudhamani ChN, Naik HSB, Gowda KRS, Giridhar M, Girija D, Kumar PNP. Novel iron phenanthroline-based photosensitizers for antimicrobial PDT: synthesis, DNA binding and photo-induced DNA cleavage activity. Med Chem Res. 2017; 26:1160–1169.

13. Abrahamse H, Hamblin MR. New photosensitizers for photodynamic therapy. Biochem. J. 2016; 473: 347–364. https://doi.org/10.1042/BJ20150942 PMID: 26862179

14. Dolkmans DEJGJ, Fukumura D, Jain RK. Photodynamic therapy for cancer. Nat Rev Cancer 2003; 3: 380–387. https://doi.org/10.1038/nrc1071 PMID: 12724736

15. Castano AP, Mroz P, Hamblin MR. Photodynamic therapy and anti-tumour immunity. Nat Rev Cancer 2006; 6: 535–545. https://doi.org/10.1038/nrc1894 PMID: 16794636

16. Nardi G, Monti I, Monti S, Miranda MA, Lhiaubet-Vallet V. Scope and limitations of the TEMPO/EPR method for singlet oxygen detection: the misleading role of electron transfer. Free Radical Biol Med. 2014; 77: 64–70.
17. Wu H, Song Q, Ran G, Lu X, Xu B. Recent developments in the detection of singlet oxygen with molecular spectroscopic methods. Trends Anal Chem. 2011; 30: 133–141.

18. Hideg E, Deák Z, Hakala-Yatkin M, Karonen M, Rutherford AW, Tyystjärvi E, et al. Pure forms of the singlet oxygen sensors TEMP and TEMPD do not inhibit Photosystem II. Biochim Biophys Acta 2011; 1807: 1658–1661. https://doi.org/10.1016/j.bbapap.2011.09.009 PMID: 21982982

19. Adam W, Kazakov DV, Kazakov VP. Singlet-oxygen chemiluminescence in peroxide reactions. Chem Rev. 2005; 105: 3371–3387. https://doi.org/10.1021/cr0300035 PMID: 16159156

20. Zhu D, Xing D, Wei Y, Li X, Gao B. Evaluation of the degree of medical radiation damage with a highly sensitive chemiluminescence method. Luminescence 2004; 19: 278–282. https://doi.org/10.1002/bio.782 PMID: 15386798

21. MacManus-Spencer LA, Latch DE, Kroncke KM, Mcneill K. Stable dioxetane precursors as selective trap-and-trigger chemiluminescent probes for singlet oxygen. Anal Chem. 2005; 77: 1200–1205. PMID: 15859008

22. Ruiz-González R, Zanocco AL. Singlet oxygen fluorescent probes. In Nonell S, Flors C, editors. Singlet oxygen. Applications in biosciences and nanosciences. London: The Royal Society of Chemistry; 2016. pp. 103–120.

23. Liu H-W, Xu S, Wang P, Hu X, Zhang J, Yuan L, et al. Efficient two-photon fluorescent probe for monitoring mitochondrial singlet oxygen in tissues during photodynamic therapy. Chem Commun. 2016; 52: 12330–12333.

24. Garcia-Diaz M., Huang Y.-Y., Hamblin M. R., Use of fluorescent probes for ROS to tease apart Type I and Type II photochemical pathways in photodynamic therapy. Methods 2016; 109: 158–166. https://doi.org/10.1016/j.ymeth.2016.06.025 PMID: 27374076

25. Kim S, Fujitsuka M, Majima T. Photochemistry of singlet oxygen sensor green. J Phys Chem B. 2013; 117: 13985–13992. https://doi.org/10.1021/jp406638g PMID: 24111566

26. S. Pedersen K, Holmehave J, Blaikie FH, Gollmer A, Breitenbach T, Jensen HH, et al. Aarhus sensor green: a fluorescent probe for singlet oxygen. J Org Chem. 2014; 79: 3079–3087. https://doi.org/10.1021/jo500219y PMID: 24605923

27. Ruiz-González R, Zanocco R, Gidi Y, Zanocco AL, Nonell S, Lemp E. Naphthoxazole-based singlet oxygen fluorescent probes. Photochem Photobiol. 2013; 89: 1427–1432. https://doi.org/10.1111/php.12106 PMID: 23730728

28. Song D, Cho S, Han Y, You Y, Nam W. Ratiometric fluorescent probes for detection of intracellular singlet oxygen. Org Lett. 2013; 15: 3582–3585. https://doi.org/10.1021/ol401421r PMID: 23808624

29. Dai Z, Tian L, Xiao Y, Ye Z, Zhang R, Yuan J. A cell-membrane-permeable europium complex as an efficient luminescent probe for singlet oxygen. J Mater Chem B. 2013; 1: 924–927.

30. Matheson IBC, Lee J. Reaction of chemical acceptors with singlet oxygen produced by direct laser excitation. Chem Phys Lett. 1970; 7: 475–476.

31. Merkel PB, Keams DR. Direct measurement of the lifetime of $^1\Delta$ oxygen in solution. Chem Phys Lett. 1970; 12: 120–122.

32. Di Mascio P, Medeiros MHG, Sies H, Bertolotti S, Braslavsky SE, Veloso DP, et al. Quenching of singlet molecular oxygen by natural furan diterpenes. J Photochem Photobiol B Biol. 1997; 38: 169–173.

33. Okamoto M, Tanaka F, Teranishi H. Pressure effect on the lifetime of singlet oxygen in solutions. J Phys Chem. 1990; 94: 669–672.

34. Zanocco AL, Günther G, Lemp E, de la Fuente JR, Pizarro N, Kinetics and mechanism of the photosensitised oxidation of furosemide. Photochem Photobiol. 1998; 68: 487–493.

35. Castañeda F, Zanocco AL, Meléndez M, Günther G, Lemp E. Synthesis of 2-((N,N,N-trimethyl)-n-alkyl)-5-alkylfuryl halides: Useful probes for studying singlet oxygen dynamics and equilibria in micro-compartmentalized systems. J Photochem Photobiol A Chem. 2004; 168: 175–183.

36. Zanocco AL, Meléndez M, Günther G, Lemp E. Study of singlet oxygen equilibrium in dioctadecylmethylammonium chloride vesicles employing 2-(N,N,N-trimethylamine)-n-alkyl)-5-alkylfuryl halides. Photochem. Photobiol. 2007; 83: 584–591. https://doi.org/10.1562/2006-08-01-RA-991 PMID: 17115800

37. Demas JN, Crosby GA. The measurement of photoluminescence quantum yields. A review. J Phys Chem. 1971; 75: 991–1024.

38. Eaton DF. Reference materials for fluorescence measurement. Pure Appl Chem. 1988; 60: 1107–1114.

39. Montalti M, Credi A, Prodi L, Gandolfi MT. Handbook of Photochemistry. 3rd ed. Boca Raton: CRC Press; 2006.
40. Günther G, Lemp E, Zanocco AL. On the use of 9,10-dimethylanthracene as chemical rate constant actinometer in singlet molecular oxygen reactions. Bol Soc Chil Quím. 2000; 45: 637–644.

41. Schmidt R, Tanielian C, Dunsbach R, Wolff C. Phenalenone, a universal reference compound for the determination of quantum yields of singlet oxygen O\(_2^\text{(1Δg)}\) sensitization. J Photochem Photobiol A Chem. 1994; 79: 11–17.

42. Aleksandrov AA, Dedeneva AS, El'chaninov MM. Synthesis and some transformations of 2-(2-furyl)naphtho[1,2-d]oxazole. Russ J Org Chem. 2011; 47: 710–713.

43. Zajac M, Hrobářík P, Foltínová P, Zahradník P. Donor–π-acceptor benzothiazole derived dyes with an extended heteroaryl-containing conjugated system: synthesis, DFT study and antimicrobial activity. Tetrahedron. 2008; 64: 10605–10618.

44. Curitol M, Ragas X, Nonell S, Pizarro N, Encinas MV, Rojas P, et al. Solvent and media effects on the photophysics of naphthoxazole derivatives. Photochem Photobiol. 2013; 89: 1327–1334. https://doi.org/10.1111/php.12133 PMID: 23834078

45. Cyranński M, Krygowski TM. Separation of the energetic and geometric contributions to the aromaticity of π-electron carboyclics. Part V. Analysis of the aromatic character of aza-analogues of benzenoid hydrocarbons. Tetrahedron. 1995; 52: 13795–13802.

46. Krygowski TM, Cyranowski M. Separation of the energetic and geometric contributions to the aromaticity. Part IV. A general model for the π-electron systems. Tetrahedron. 1996; 52: 10255–10264.

47. Krygowski TM, Cyranowski M. Structural aspects of aromaticity. Chem Rev. 2001; 101: 1385–1420 PMID: 11710226

48. Fayed TA. Probing of micellar and biological systems using 2-(p-dimethylaminostyryl) benzoxazole. An intramolecular charge transfer fluorescent probe. Colloids and Surfaces A Physicochem Eng Aspects. 2004; 236: 171–177.

49. da Hora Machado AE, de Miranda JA, Guiliardi S, Nicodem DE, Severino D. Photophysics and spectroscopic properties of 3-benzoxazol-2-yl-chromen-2-one. Spectrochem Acta A 2003; 59: 345–355.

50. Lemp E, Zanocco AL, Lissi EA. Linear free energy relationship analysis of solvent effects on singlet oxygen reactions. Curr Org Chem. 2003; 7: 799–819.

51. Hansch C, Leo A, Taft RW. A survey of Hammett substituent constants and resonance and field parameters. Chem Rev. 1991; 97: 165–195.

52. Swain CG, Lupton EC. Field and resonance components of substituent effects. J Am Chem Soc. 1990; 90: 4328–4337.

53. Contreras R, Fuentealba P, Galván M, Pérez P. A direct evaluation of regional Fukui functions in molecules. Chem Phys Lett. 1999; 304: 405–413.

54. Tiznado W, Chamorro E, Contreras R, Fuentealba P. Comparison among four different ways to condense the Fukui function. J Phys Chem A. 2005; 109: 3220–3224. https://doi.org/10.1021/jp0450787 PMID: 16833651

55. Fukui K. Role of frontier orbitals chemical reactions. Science. 1982; 217: 747–784.

56. Clennan EL, Pace A. Advances in singlet oxygen chemistry. Tetrahedron. 2005; 61: 6665–6691

57. Castro-Olivares R, Günther G, Zanocco AL, Lemp E. Linear free energy relationship analysis of solvent effect on singlet oxygen reactions with mono and disubstituted anthracene derivatives. J Photochem Photobiol A Chem. 2009; 207: 160–166.

58. Koo Lee Y E, Kopelman R. Nanoparticle PEBBLE sensors in live cells. Methods Enzymol. 2012; 504: 419–470. https://doi.org/10.1016/B978-0-12-391857-4.00021-5 PMID: 22264547

59. Bresoli-Obach R, Nos J, Mora M, Sagristà ML, Ruiz-González R, Nonell S. Anthracene-based fluorescent nanoprobes for singlet oxygen detection in biological media. Methods 2016; 109: 64–72. https://doi.org/10.1016/j.ymeth.2016.06.007 PMID: 27302662
64. Flors C, Fryer MJ, Waring J, Reeder B, Bechtold U, Mullineaux PM, et al. Imaging the production of singlet oxygen in vivo using a new fluorescent sensor, singlet oxygen sensor green. J Exp Bot. 2006; 57: 1725–1734. https://doi.org/10.1093/jxb/erj181 PMID: 16595576

65. Shen Y, Lin H, Huang Z, Xiao L, Chen D, Li B, et al. Kinetic analysis of singlet oxygen generation in a living cell using singlet oxygen sensor green. Proc SPIE. 2010; 7845: 78451F1–78451F6;

66. Xiao L, Gu L, Howell S B, Sailor MJ. Porous silicon nanoparticle photosensitizers for singlet oxygen and their phototoxicity against cancer cells. ACS Nano. 2011; 5: 3651–3659. https://doi.org/10.1021/nn1035262 PMID: 21452822

67. Ragas X, Cooper LP, White JH, Nonell S, Flors C, Quantification of photosensitized singlet oxygen production by a fluorescent protein. ChemPhysChem 2011; 12: 161–165. https://doi.org/10.1002/cphc.201000919 PMID: 21226197

68. Westberg M, Bregnhøj M, Blázquez-Castro A, Breitenbach T, Etzerodt M, Ogilby P R. Control of singlet oxygen production in experiments performed on single mammalian cells. J Photochem Photobiol A Chem. 2016; 321: 297–308.

69. Kim S, Tachikawa T, Fujitsuka M, Majima T. Far-red fluorescence probe for monitoring singlet oxygen during photodynamic therapy. J Am Chem Soc. 2014; 136: 11707–11715. https://doi.org/10.1021/ja504279r PMID: 25075870

70. Lee JJ, Goncalves A, Smith BA, Palumbo R, White AG, Smith BD. In vivo imaging using polymeric nanoparticles stained with near-infrared chemiluminescent and fluorescent squaraine catenane endoperoxide. Chem. Commun. 2013; 49: 3016–3018.