







Breaking photoswitch activation depth limit using ionising radiation stimuli adapted to clinical application

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Electromagnetic radiation-triggered therapeutic effect has attracted a great interest over the last 50 years. However, translation to clinical applications of photoactive molecular systems developed to date is dramatically limited, mainly because their activation requires excitation by low-energy photons from the ultraviolet to near infra-red range, preventing any activation deeper than few millimetres under the skin. Herein we conceive a strategy for photosensitive-system activation potentially adapted to biological tissues without any restriction in depth. High-energy stimuli, such as those employed for radiotherapy, are used to carry energy while molecular activation is provided by local energy conversion. This concept is applied to azobenzene, one of the most established photoswitches, to build a radioswitch. The radiation-responsive molecular system developed is used to trigger cytotoxic effect on cancer cells upon gamma-ray irradiation. This breakthrough activation concept is expected to expand the scope of applications of photosensitive systems and paves the way towards the development of original therapeutic approaches.

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Triggering process activation using stimuli is a promising approach to reach control and modulation of on-demand therapeutic actions in real time¹. Stimulus-triggered actions are mediated by endogenous or external stimuli. While the former restrict to specific local environments of the targeted zone, such as distinctive pH, redox potential, oxygen content or enzyme activity, the latter enable to escape from these local limitations thanks to an orthogonal actuation leading to high spatiotemporal control^{2,3}. Due to the promising benefits of this approach, light responsive systems have been developed for many decades and are now able to initiate specific and complex actions such as bond cleavage, switch, slide or rotation, which led to the research field of photoactivation⁴.

The first reported photosensitive systems were activated by ultraviolet (UV) photons carrying enough energy to induce bond cleavage or molecular motion. Because of the limited tissue penetration of such photons (<100 μm), many investigations were undertaken to reach deeper areas in the body by lowering the required energy of the activating photon⁵. New chemical entities, such as visible or near-infrared absorbing compounds and nanoparticles, were introduced and led to photoactivatable systems adapted to in vivo studies in small animals (zebrafish, rodent)⁶. Clinical applications relying on photodynamic therapy^{7,8} or photoimmunotherapy⁹ were also implemented to treat topical cancers or tumors accessible by endoscopic techniques. However, the photocontrolled systems developed until now are unable to trigger any action deeper than few millimetres in tissues because of the intrinsic low penetration of the required activating photons¹⁰. And this is probably today the main hurdle preventing any wide clinical application in spite of the extensive capacities of the reported systems such as drug delivery, protein and cell activity modulation, gene expression, molecular pump, slider and motor, and many more^{11–15}.

Herein we propose a strategy for the activation of light-sensitive systems by external stimuli without any restriction in depth. Our approach is to use high-energy waves/particles contained in ionising radiations (IRs) employed in cancer radiotherapy, such as X-ray (XR), gamma-ray (GR) or electron-beam (E) irradiations, to efficiently reach deep-tissues and then to benefit from the local conversion of carried energy into low-energy

particles and species to induce specific activation of photosensitive therapeutics (Fig. 1).

Radiotherapy is a cornerstone in cancer treatment since more than half of patients will benefit from it and many efforts have been made to improve it¹⁶. In the first place, enhancements of IR dose were reported in the presence of iodine-based radiopaque diagnostic agent¹⁷. Then, nanoparticles able to increase the effect of the IR dose by radiosensitizing and/or radioenhancement effects were described, which was more recently associated to immune system activation due to the intrinsic immunogenicity of IRs^{18,19}. In the last few years, several systems were reported to induce more complex and selective actions from IR stimuli^{20,21}. We could separate them into two main families depending on the type of activation based on down-conversion or oxidation by ROSs. The down-converting systems, such as nanoscintillators, are designed to convert incident XR photons into UV-vis light to induce the release of cytotoxic agents such as singlet oxygen (¹O₂) in the case of the widely studied photodynamic therapy approaches²². The other systems benefit from the generated ROSs (and/or from secondary electrons)²³ to induce specific bond cleavage (mainly diselenide, disulfide, C-N, C-O, S-N, coordination bonds with metal)^{24–27}, DNA break²⁸, atom oxidation (mainly sulphur, selenium and carbon from unsaturated lipids)^{29,30} leading to the disassembly of capsules, polymers or prodrugs, and the release of drug or gas (nitric oxide, carbon monoxide).

Our approach was to investigate if IR could be used in ways other than radioenhancement, bond break or atom oxidation to disassemble nanoparticle, and be employed to generate specific and non-destructive molecular actions such as molecular rearrangements described in photoactivation. For this purpose, a molecule containing both an IR-sensitive element and a photo-sensitive moiety was designed. High atomic number (Z) metals are known to efficiently interact with high-energy photons through their full or partial absorption and the local release of lower-energy secondary photons and electrons^{16,31}. These secondary particles then induce several types of interaction with the surrounding matter as well as with the neighbouring high-Z elements leading to a ripple effect. Thus, gadolinium (Gd)-chelate was selected for both efficient interaction with IRs and detection

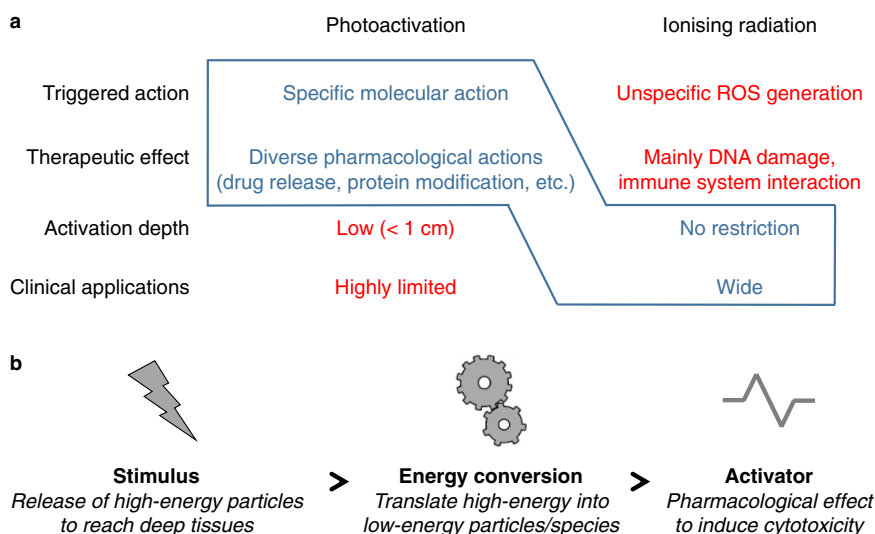


Fig. 1 Approach to tackle deep-tissue photoactivation to overcome the current limitation preventing wide clinical applications. **a** Highlight of advantages (blue) and limitations (red) of photoactivation and ionising radiation to exhibit the complementary properties of both domains which can be combined (blue delineation) for an original molecular-activation concept. **b** Our approach is based on using ionising-radiation stimulus to reach deep tissues and to benefit from local conversion of high-energy particles into low-energy particles and species suitable for molecular activation. In this way, specific pharmacological actions could be induced in biological tissues without any depth restriction.

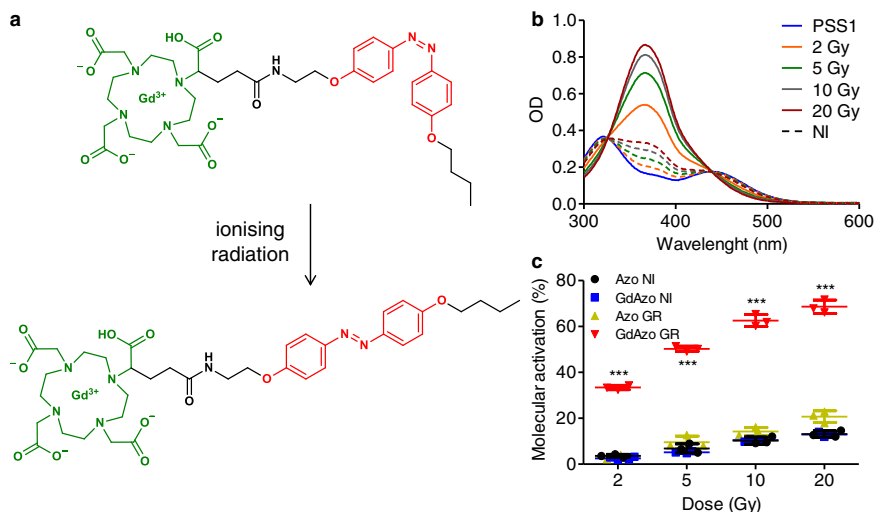


Fig. 2 Molecular activation upon IR. **a** Activation of *cis*-GdAzo (top) into *trans*-GdAzo (bottom) upon IR. **b** Absorbance spectra of *cis*-GdAzo compound (photostationary state 1, PSS1) recorded after GR irradiation at 2, 5, 10, and 20 Gy. The control absorbance spectra of non-irradiated (NI) *cis*-GdAzo compound analysed concurrently are drawn in dashed lines with corresponding color. **c** Molecular activation of *cis*-GdAzo and *cis*-Azo (control molecule without Gd atom) determined by HPLC and reported as the difference of *trans*-isomer proportion before and after GR irradiation ($n = 3$ independent experiments). The means \pm standard deviations are reported. Two-way Anova (Bonferroni post-test) was used for statistical analyses (All vs GdAzo GR, t -values = 20.78, 30.29, 34.99 and 37.43 for GdAzo NI vs GdAzo GR at doses 2, 5, 10, and 20 Gy). *** $P < 0.001$. OD optical density.

by magnetic resonance imaging (MRI)³². On the other hand, extensive investigations proved that the azobenzene photoswitch is adapted to many pharmacological applications thanks to photocontrol of the *cis-trans* configuration switch impacting both length and polarity of the moiety³³. Moreover, azobenzene compounds can be isomerised through different types of stimuli such as photon, heat and electron, which offers a great opportunity to implement triggering systems based on IRs.

In this study, Gd-chelate and azobenzene moiety are associated to design a special class of photosensitive system referred to as radioswitch (Fig. 2a). It is based on a photoswitch able to be activated by IRs through molecular rearrangement (isomerization) and to trigger cytotoxic effect. We investigate the underlying mechanism of the activation process and assess cancer-cell killing activity to highlight the therapeutic potential provided by this breakthrough photoswitch-activation approach.

Results and discussion

Design of radioswitch. An azobenzene moiety carrying a primary amine function and alkoxy chains at both opposite *para*-positions was prepared using standard procedures. The Gd-chelate was introduced through an anhydride opening reaction from a modified 1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetic acid (DOTA) chelate followed by complexation with Gd. The final product GdAzo was synthesised over 5 steps in 33% yield (Supplementary Section 2).

Azobenzene is known for reversible shift between *trans* and *cis* configurations through isomerisation of the azo double bond after excitation by photons from UV to near-infrared energies³⁴. The UV-vis absorption signatures of these two isomers significantly differ through the $\pi \rightarrow \pi^*$ electronic transition, detected at 322 and 367 nm for the *cis*-GdAzo and *trans*-GdAzo isomers respectively with much higher intensity for the latest. These spectrophotometric characteristics allow the monitoring of azobenzene isomerisation by absorbance measurement as we confirmed by proton nuclear magnetic resonance (¹H NMR) and high-performance liquid chromatography (HPLC) (Supplementary Section 3). The thermodynamically more stable *trans*-GdAzo isomer can be partially converted to the metastable isomer *cis*-GdAzo upon UV light

(photostationary state in PBS containing $90 \pm 2\%$ of the *cis*-isomer) and will then be recovered by thermal back relaxation (Supplementary Fig. 14, $t_{1/2} = 2.3$ h at 37 °C in PBS). It has to be noted that while the thermal half-life of *cis*-GdAzo is suitable for in vitro experiment, for any in vivo work, other systems need to be developed, with thermal half-lives of several days to weeks.

Molecular activation upon ionising radiation. Primary photons from IRs have energies above the kiloelectronvolt range (1 keV \sim 1.24 nm). They have energy theoretically much too high to induce intramolecular electronic transition (such as $\pi \rightarrow \pi^*$ or $n \rightarrow \pi^*$) and exhibit the same low probability to interact with azobenzene as with any other components constituted of carbon, nitrogen or oxygen (water molecules, proteins, DNA, etc.). In contrast, IRs deliver or generate photons which have higher probability of interaction with metals of high atomic number such as Gd ($Z = 64$) through the photoelectric effect (ejection of an electron from the inner shells) since the photoelectric cross-section scales as Z^3 - Z^4 depending on the incident-photon energy. This induces the release of low-energy particles and species in the very close vicinity (few nanometres) of the metal, mainly in the form of Auger electrons, lower-energy photons and reactive oxygen species (ROS)^{35,36}. Monte Carlo simulations showed that such interaction induces a nanoscale dose deposition with a local increase in the equivalent energy up to several orders of magnitude by generating a large amount of low-energy species, which highlights the potential of this energy reservoir^{37,38}. This increase in the radiation-dose effect has been validated for several IR sources (from kiloelectronvolt to megaelectronvolt) currently used in clinic^{36,39,40}.

GdAzo isomerisation has been assessed upon GR (662 keV, Supplementary Section 4) at irradiation doses compatible with clinical applications (2-20 Gy) (Fig. 2a). Using spectrophotometric and HPLC determinations, we discovered that *cis*-GdAzo was properly converted into *trans*-GdAzo (Fig. 2b, c and Supplementary Section 5). The activation efficacy was 33% to 69% at irradiation doses from 2 to 20 Gy (Figs. 2c, 3e). No other compound than *trans*-GdAzo was generated upon GR irradiation as confirmed by spectrophotometry (Fig. 2b and Supplementary

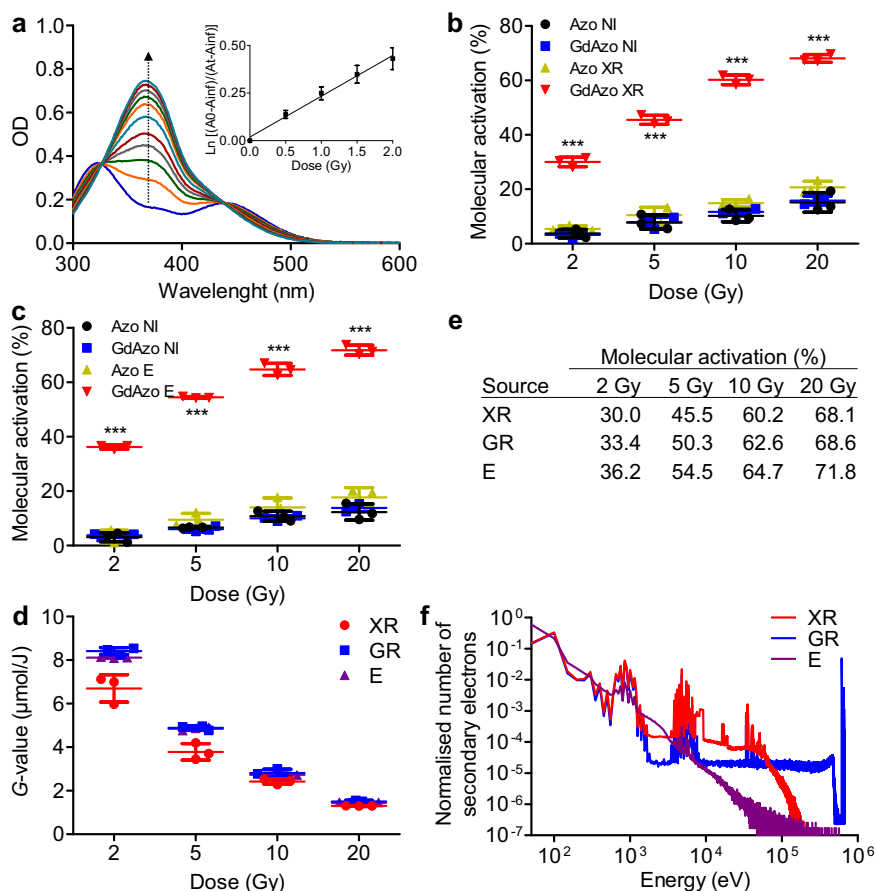


Fig. 3 Characterisation of *cis*-GdAzo activation upon different radiation sources. **a** Absorbance spectra of *cis*-GdAzo (PSS1 in blue) recorded after GR (0.5 Gy increment in the 0.5–2 Gy range and 1 Gy increment in the 2–8 Gy range). The graph $\text{Ln}[(A_0 - A_{\text{inf}})/(A_t - A_{\text{inf}})]$ by the irradiation dose (0.5–2 Gy range) is plotted on the top-right corner, with A_0 , A_{inf} and A_t the absorbances of PSS1, *trans*-GdAzo and the mixture at a specific irradiation dose respectively. Linear regression (least-squares method, r^2 : 0.9521, GraphPad Prism 5.00) were used to determine the activation constant $k = (2.1 \pm 0.2) \times 10^{-1} \text{ Gy}^{-1}$ (triplicate, $n = 3$ independent experiments, F -value = 258.3, $P < 0.0001$). **b**, **c** Molecular activation of *cis*-GdAzo and *cis*-Azo determined by HPLC and reported as the difference of *trans*-isomer proportion before and after XR (**b**) and E (**c**) irradiations ($n = 3$ independent experiments). Two-way Anova (Bonferroni post-test) was used for statistical analyses (All vs GdAzo GR, t -values = 16.93, 23.90, 30.65, 33.01 and 19.04, 28.43, 32.09, 34.01 for GdAzo NI vs GdAzo GR at doses 2, 5, 10, and 20 Gy upon XR and E respectively). $***P < 0.001$. **d** Radiochemical yields (G -values) of *trans*-GdAzo determined by HPLC (corrected from thermal back relaxation) upon XR, GR and E irradiations ($n = 3$ independent experiments). **e** Molecular activation of *cis*-GdAzo upon XR, GR and E irradiations determined by HPLC. **f** Monte Carlo simulation to quantify the energy of secondary electrons emitted by the interaction of the incident primary particle on the Gd atom upon XR, GR and E irradiations. The means \pm standard deviations are reported. OD optical density.

Figs. 15–17), HPLC (Supplementary Figs. 18–35) and LC-MS (Supplementary Figs. 37–44). Moreover, successive irradiations by GR and UV showed that GdAzo *trans*-*cis* isomerisation was reversible confirming that isomerisation is the molecular process happening upon IRs (Supplementary Fig. 45). The *cis*-Azo control compound (same structure as *cis*-GdAzo without Gd atom) as well as the non-irradiated *cis*-GdAzo and *cis*-Azo compounds were only slightly converted into the *trans*-isomer during the same time frame, a conversion mainly attributed to thermal back relaxation of the *cis*-isomer in the dark (Fig. 2b, c and Supplementary Section 5).

The *cis*-GdAzo activation upon GRs was characterised by a dose-related monoexponential increase in *trans*-GdAzo at low irradiation dose (< 2 Gy) and determination of a constant k to quantitatively characterise the activation process was then possible (Fig. 3a and Supplementary Fig. 49). The activation constants of *cis*-GdAzo and *cis*-Azo upon GRs are $k = (2.1 \pm 0.2) \times 10^{-1} \text{ Gy}^{-1}$ and $k = (1.83 \pm 0.07) \times 10^{-2} \text{ Gy}^{-1}$ respectively ($P < 0.0001$). Another approach to quantitatively characterise this activation method is to introduce the G -value which is the chemical yield currently used for IR-dosimetry calculation and

consisting in the number of molecules affected per 100 eV ($\sim 1.60 \times 10^{-17} \text{ J}$) of energy absorbed. *cis*-GdAzo activation at 2 Gy was characterised by $G = 81 \pm 2$ corresponding to a molecular activation of $8.4 \pm 0.2 \mu\text{mol/J}$ (corrected from thermal back relaxation). For comparison, hydroxyl radical (HO^*) and hydrogen peroxide (H_2O_2) are generated upon water radiolysis at $0.28 \mu\text{mol/J}$ and $0.073 \mu\text{mol/J}$ yield respectively (homogeneously distributed $\sim 10^{-7} \text{ s}$ after IR)⁴¹. Thus, the *cis*-GdAzo activation efficacy was particularly high considering the ROS released from IR sources. This type of molecular activation based on radioswitch appears very promising and open new avenues for the applications of photoswitch such as potential actuatable tools upon radiotherapy. In addition, considering the easy accessibility of radiation sources in hospitals for cancer treatment, it may turn into a very powerful technique for therapy applications.

Mechanism of activation upon ionising radiation. The GR photons (662 keV) used in our study mostly interact with matter through the Compton effect, which is the main physical effect occurring during clinical radiotherapy. It consists in the inelastic

scattering of the incident primary photon by a loosely-bound valence electron that is ejected. It is almost independent of the atomic number (cross-section only varies linearly with Z) and dominates in biological media for incident-photon energies ranging from about 50 keV to 20 MeV and 500 keV to 5 MeV for interaction with elements of Z at 10 and 70 respectively⁴². To gain insight into the physical process involved during the *cis*-**GdAzo** activation, we investigated the impact of the incident-particle energy and its type by using an XR generator (photons of 80 keV mean energy) and a linear accelerator (LINAC, electrons (E) of 4.5 MeV) respectively (Supplementary Section 4). Indeed, 80 keV photons mainly interact with high- Z elements through the photoelectric effect with an efficacy highly dependent on the atomic number of the interacting element, contrary to the Compton effect. This would suggest that *cis*-**GdAzo** (containing Gd, $Z = 64$) activation efficacy upon XRs could be increased compared to GR irradiation at 662 keV. On the other hand, LINAC provides high-energy charged particles mainly imparting energy through Coulomb interactions and inducing Čerenkov effect (release of UV photons), which could impact on *cis*-**GdAzo** activation efficacy. The obtained G -values and activation constants k upon XR and E irradiations were in the same order as those obtained upon GR irradiation (Fig. 3b–e and Supplementary Section 5). This clearly demonstrates that the activation of *cis*-**GdAzo** is independent from the energy and the type of the IR, meaning that primary particle-matter interaction is not significant and that secondary particles and species emitted after this first interaction lead to *cis*-**GdAzo** activation.

Monte Carlo simulations (PENELOPE code)⁴³, allowing to determine the energy of the secondary electrons released after the interaction of the three different IR sources with Gd atoms, confirmed that the relative amount of high-energy electrons drastically differed for the three sources, whereas a similar relative amount of low-energy electrons was released from them (Fig. 3f and Supplementary Section 6). These simulations support that *cis*-**GdAzo** activation cannot be due to high-energy particle-Gd interaction and it is mainly triggered by the low-energy particles and species generated by energy loss from the incident primary particles, explaining the similar efficacies obtained with the different IR sources.

To go further into the understanding of the mechanism of *cis*-**GdAzo** activation, GR irradiations were performed in media containing scavengers able to interact with different radicals generated during water radiolysis. Results highlighted the crucial role of oxidising species in the *cis*-**GdAzo** activation process (Fig. 4a and Supplementary Section 7). Indeed, species which quench the hydroxyl radical HO• (*tert*-butanol (Fig. 4b), mannitol and ethanol) abolished *cis*-**GdAzo** activation except when they converted HO• into other oxidant species (sodium azide, dimethylsulfoxide). On the contrary, electron-converting species (cadmium perchlorate) did not affect *cis*-**GdAzo** activation except when they converted electrons into oxidant species (sodium selenate).

Activation carried out in gas-saturated solutions validated the key role of HO• (Fig. 4c and Supplementary Section 8.1). Indeed, removing oxygen by nitrogen saturation was not affecting *cis*-**GdAzo** activation which bears out that hydrated electrons and hydrogen radicals (and thus the couple perhydroxyl radical/superoxide radical anion HO₂•/O₂•⁻ resulting from their reaction with oxygen) are not involved. It has to be noted that this independence from oxygen could be very valuable for hypoxic-tumor treatment. On the other end, nitrous oxide (N₂O) saturation led to an increase in *cis*-**GdAzo** activation which unambiguously implicates the HO• as N₂O converts all hydrated electrons into HO• upon irradiation of aqueous solution, resulting in doubling the production yield of HO• ($G(\text{HO}^\bullet) = 0.56 \mu\text{mol/J}$,

completed in ~14 ns). Furthermore, when the irradiation dose in N₂O saturated solution was brought back to the generated HO• amount, the molecular activation of *cis*-**GdAzo** was similar as in water (Fig. 4d). For instance, 5 Gy irradiation in N₂O-saturated solutions (equivalent to 10 Gy in non-saturated solutions with respect to generated HO• amount, and noted “eq 10” in Fig. 4d) resulted in similar activation efficacy as 10 Gy irradiation in non-saturated solutions. The central role of HO• led us to exclude some photon-mediated interactions in the *cis*-**GdAzo** activation process such as the Čerenkov effect (release of UV photons from accelerated charged particles such as electrons) and the scintillation effect (release of UV photons from gadolinium atoms). Finally, comparable quenching-effects were observed upon XR, GR and E irradiations (Supplementary Section 7), which revealed a similar *cis*-**GdAzo**-activation mechanism using sources with different primary-particle types and energies. This confirmed that *cis*-**GdAzo** activation was induced by the secondary low-energy particles and species generated upon IR, which is in line with the conclusions drawn from Monte Carlo simulations.

If the hydroxyl radicals HO• was the necessary and sufficient specie for *cis*-**GdAzo** activation, a chemical introduction of HO• would lead to the same process. This has been investigated using Fenton chemistry which generates HO• from the dismutation of H₂O₂ by ferrous-iron catalyst (Fig. 5a, b and Supplementary Section 8.2)^{44,45}. As expected, *cis*-**GdAzo** activation was induced by Fenton chemistry, while H₂O₂ alone was inefficient, even if some degradation was observed in these conditions (Supplementary Fig. 58).

These investigations demonstrate that *cis*-**GdAzo** activation is triggered by the oxidation of the azo double bond ($\text{N}=\text{N}^{+\bullet}$) by reaction with HO• generated by the low-energy particles released upon IR. Thus, the metastable *cis*-**GdAzo**^{+•} radical cation is instantaneously converted into the more stable *trans*-**GdAzo**^{+•} radical cation which then reduces into the final thermodynamically stable and neutral *trans*-**GdAzo** compound. Indeed, the isomerisation rate of *cis*-azobenzene-based radical cations is known to be several orders of magnitude faster than that of the corresponding neutral compound⁴⁶.

The three sources of radiation used in this study deliver low linear energy transfer radiations (LET, which is defined by the rate of energy loss per unit length of track of the particle). In low-LET radiations, the first events appear in small widely separated spurs (10⁻¹⁶–10⁻¹⁰ s time scale) and the generated radicals are homogeneously spread into water at about 10⁻⁷ s after irradiation^{47,48}. The HO• radiochemical yield ($G(\text{HO}^\bullet)$) is about 0.28 $\mu\text{mol/J}$ (at standard dose rate); however, it can be lowered when low-energy photons (1–100 keV) are used, which is the case of the XR generator used herein⁴⁹. Thus, $G(\text{HO}^\bullet)$ delivered by both the XR generator and the GR source has been determined (Supplementary Section 9.1, Supplementary Figs. 59–63). Indirect quantification of HO• was based on HO• scavenging by coumarin⁵⁰ and quantification of 7-hydroxycoumarin (7-OH-Coum) which is the only fluorescent product released from this scavenging reaction⁵¹. 7-OH-Coum is proportional to HO• concentration and specific for HO• among other reactive oxygen species⁵². In the conditions used, coumarin reacts with HO• ~10⁻⁷ s after the initial transfer of energy to water (rate constant $k = 1.05 \times 10^{10} \text{ L}\cdot\text{mol}^{-1}\cdot\text{s}^{-1}$)⁵³, which results in quantification of the homogeneously distributed HO• without interfering on the intratrack recombination of radicals occurring in the spurs. We quantified $G(\text{HO}^\bullet)$ of 0.200 and 0.280 $\mu\text{mol/J}$ upon XR and GR irradiations respectively (Fig. 5d when $[\text{Gd}^{3+}] = 0$), which is in line with the reported yields for similar radiation sources and dose rates^{48,49}. Thus, $G(\text{HO}^\bullet)$ ~10⁻⁷ s after irradiation were 0.200, 0.280, and 0.280 $\mu\text{mol/J}$ for the XR, GR and E sources respectively. Interestingly, the activation yield of *cis*-**GdAzo** was

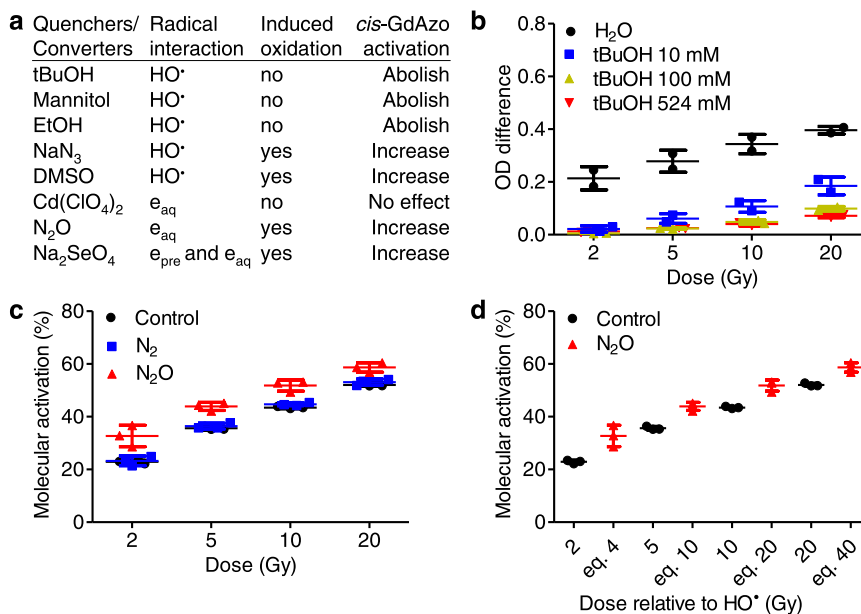


Fig. 4 Investigation on activation mechanism using scavengers. **a** Quenchers and converters used to highlight the impact of specific radicals on *cis*-GdAzo-activation efficacy upon XR, GR and E (cf. Supplementary Section 7 for details). **b** Absorbance difference (365 nm) of medium containing *cis*-GdAzo (50 μ M) in the presence of tBuOH (10, 100, 524 mM (~5% v/v), control in water) before and after GR irradiation ($n = 2$ independent experiments). **c** Molecular activation of *cis*-GdAzo (50 μ M, H₂O) upon GR irradiation in N₂ or N₂O-saturated solutions (Control without inerting) determined by HPLC and reported as the difference of *trans*-isomer proportion before and after GR irradiation ($n = 3$ independent experiments). **d** Comparison of *cis*-GdAzo activations in control (no inerting) and N₂O-saturated solutions in conjunction with the equivalent (eq.) amount of hydroxyl radicals HO• generated depending on doses (“eq 4”, “eq 10”, “eq 20” and “eq 40” relate to the doses of 2, 5, 10, and 20 Gy in N₂O saturated solutions) ($n = 3$ independent experiments). The means \pm standard deviations are reported. OD optical density.

also slightly lower upon XR compared to GR and E at 2–5 Gy (Fig. 3d and Supplementary Fig. 47), even if this difference is lower than what could be expected from the $G(\text{HO}^\bullet)$ difference. This could be explained by a different kinetic regime for HO• to react with coumarin and *cis*-GdAzo. Indeed, the non-homogeneous energy distribution at the first stage after irradiation leads to non-homogeneous kinetics stages in the spurs.

To gain insight into the role of Gd in the activation process, $G(\text{HO}^\bullet)$ was quantified at different concentrations of Gd³⁺ ions (Fig. 5d). We observed that $G(\text{HO}^\bullet)$ was gradually increased in solutions with Gd³⁺ concentrations up to about 10–200 μ M and then reached a plateau (25–500 μ M) before decreasing for higher concentrations (500–2000 μ M). The highest enhancement factors were 33% at 200 μ M [Gd³⁺] and 20% at 500 μ M [Gd³⁺] upon XR and GR respectively. It has to be noted that this study quantified the HO• that diffused into the bulk solution and not the HO• initially generated into the spurs. Thus, the saturation and decrease of $G(\text{HO}^\bullet)$ at high Gd³⁺ concentrations could be explained by the increase in the probability of HO• recombination⁵². Furthermore, addition of Gd³⁺ ions into a solution of *cis*-Azo led to *cis*-Azo activation upon GR irradiation, and the higher the Gd³⁺ concentration was, the more efficient the *cis*-Azo activation was (Supplementary Section 9.2, Supplementary Fig. 64). This confirmed that the presence of Gd³⁺ was required to induce activation by IR of this molecular system. Thus, the presence of Gd³⁺ in solution increased the radiochemical yield $G(\text{HO}^\bullet)$ and led to the activation of the control compound *cis*-Azo.

Finally, the impact of *cis*-GdAzo concentration on the activation efficacy was assessed to point out any cooperative effect in this process (Supplementary Section 10). The linear relation between the radiochemical yield of *trans*-GdAzo (G -value) and the initial concentration of *cis*-GdAzo showed there is no catalytic effect for *cis*-GdAzo activation, which resulted in the

same efficacy for low and high *cis*-GdAzo concentration (Fig. 5e). Thus, the catalytic pathway based on the activation of a substoichiometric amount of *cis*-GdAzo (Fig. 5c) was not involved, albeit already observed for the oxidative isomerisation of azobenzene in acetonitrile⁴⁶. Moreover, the higher $G(\text{trans-GdAzo})$ at lower doses revealed a larger loss of energy at higher doses (Fig. 5e) through non-specific reactions or recombination of HO• for instance. A predictive model to estimate $G(\text{trans-GdAzo})$ from the initial *cis*-GdAzo concentration and the radiation dose was established based on a logarithmic decrease in the 2–20 Gy dose range (Fig. 5f).

Pharmacological and cytotoxic effect. The isomerisation of azobenzene is known to highly impact its physicochemical properties by modifying both its length and dipole moment. Indeed, the *trans*-isomer is almost plane and has a dipole moment close to zero, whereas the *cis*-isomer exhibits an angular geometry and a much higher dipole moment³³. Azobenzene modification has already been introduced into many systems including biomolecules, liquid crystals, or polymers to make photosensitive devices adapted to control many actions, such as complex mechanical movement, enzyme-structure modification, ion-channel opening or gene expression using UV or visible-light irradiation^{5,54–60}. The *trans*-GdAzo has an amphiphilic structure, resulting from the hydrophilic Gd-chelate and the apolar *trans*-azobenzene moiety. We took advantage of this property to use *cis*-GdAzo as an IR-triggered prodrug and to demonstrate how this photochemical compound could be activated by deep-tissue penetrating lights to induce therapeutic outcomes. We, indeed, hypothesised that *trans*-GdAzo could act as a surfactant on cell membranes by inducing cell permeabilisation and eventually cell death. Moreover, GdAzo can be detected by MRI due to the presence of Gd in its structure,

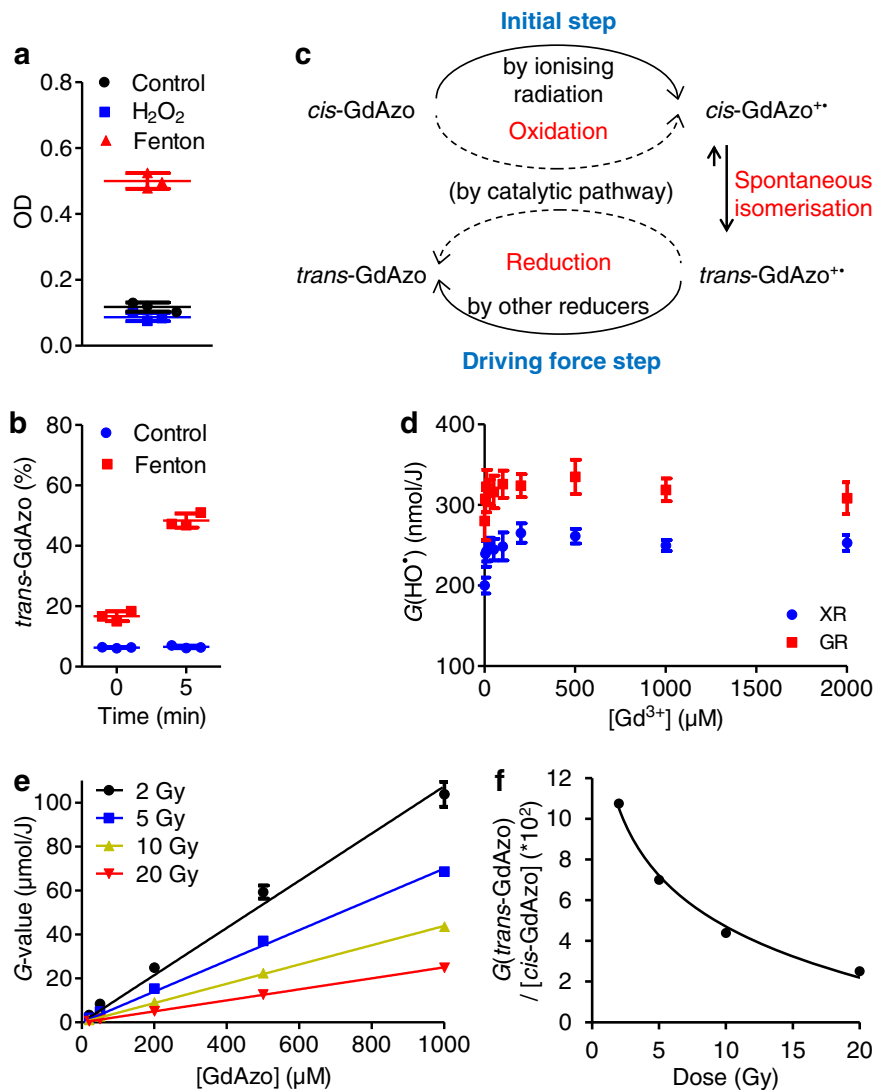


Fig. 5 Role of HO[•] radicals and proposition of mechanism. **a** Absorbance of *cis*-GdAzo (photostationary state at time 0) recorded 5 min after addition of H₂O₂ alone (bleu squares) or after running Fenton reaction (red triangle) (Control in water) ($n = 3$ independent experiments). **b** Proportion of *trans*-GdAzo after running Fenton reaction (Control without Fenton reagents) in *cis*-GdAzo-containing medium (photostationary state at time 0, just before addition of H₂O₂) determined by HPLC ($n = 3$ independent experiments). An increase in *trans*-GdAzo from 16.7% to 48.4% was observed 5 min after H₂O₂ introduction. **c** Proposal of a mechanism for the *cis*-GdAzo activation upon IRs. The initial step is the IR-induced oxidation of *cis*-GdAzo, leading to a radical cation on the azo double bond. The *cis*-GdAzo^{•+} isomer then spontaneously isomerises into the *trans*-GdAzo^{•+} which recovers its thermodynamic stable and neutral form through reduction. This last step could be induced by several reducer species in the media (hydrated electrons, hydrogen radicals, etc.) since the catalytic pathway has been rejected (*vide infra*). **d** Determination of $G(\text{HO}^{\bullet})$ upon XR and GR in the presence of different concentrations of Gd³⁺ by considering a conversion yield of coumarin into 7-OH-Coum of 3.1% (from ref. 52, $n = 5$ independent experiments). **e** Activation of *cis*-GdAzo upon GR (G -value) determined by HPLC (corrected from thermal back relaxation) at different initial concentration of *cis*-GdAzo ($n = 3$ independent experiments). **f** Linear regression to correlate $G(\text{trans-GdAzo})/[\text{cis-GdAzo}]$ to the irradiation doses. The relation $Y = -0.036 \cdot \ln(X) + 0.1304$ (with Y the slope of linear regressions from **e** in the unit G -value/ μM GdAzo and X the dose in Gy) was obtained with $r^2 = 0.9925$ (Excel 2016). The means \pm standard deviations are reported. OD optical density.

as experimentally confirmed by relaxivity measurements to determine MRI-detection efficacy (Supplementary Section 11.2). Such pharmacological and diagnostic properties could suit for a theranostic cancer-treatment approach.

The geometrical contrast between the *cis*-GdAzo and *trans*-GdAzo compounds was clearly revealed by calculations at the B3LYP/6-31 G* level of theory, respectively showing bended and planar structures of the azobenzene moiety (Fig. 6a). The electronic potentials at the surface correlate with the amphiphilic structure of *trans*-GdAzo and the theoretical dipole moment of the azobenzene moiety was decreased by 2.62 D through *cis-trans* conversion, which was expected to be sufficient to favour

cell-membrane disruption as compared to azobenzene-based polymers previously described⁶¹ (Supplementary Section 12).

The amphiphilic nature and geometry of *trans*-GdAzo led to the self-assembly of micelles at concentration above 0.42 mM in PBS at 37 °C, as shown by fluorescence and relaxivity measurements (Supplementary Section 11). These micelles were described by small angle X-ray scattering (SAXS) as ellipsoidal aggregates (the lengths for short and long semi-axes were 37 Å and 24.5 Å respectively, Supplementary Section 13.2).

Moreover, the insertion of *trans*-GdAzo within model phospholipid membranes of 1,2-dipalmitoyl-sn-glycero-3-phosphocholine (DPPC) was revealed by SAXS (Supplementary

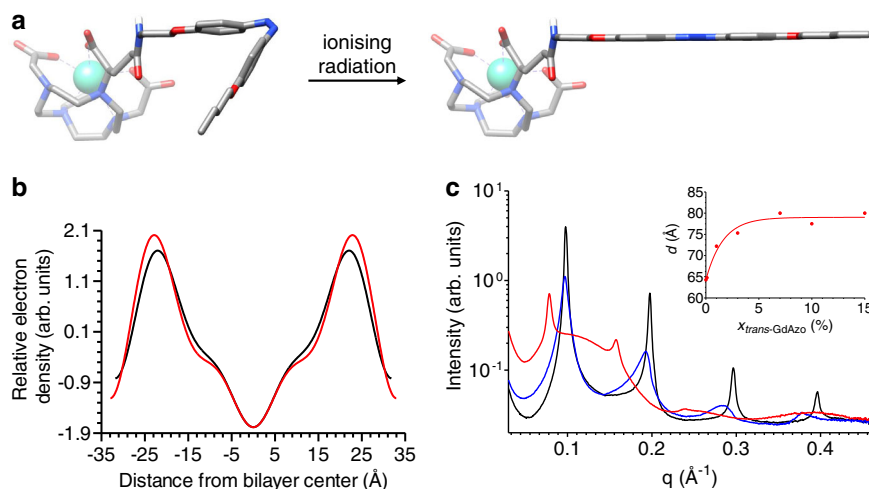


Fig. 6 Geometrical structure of GdAzo and insertion within model phospholipid membrane. **a** Structures of *cis*-GdAzo (left) and *trans*-GdAzo (right) calculated at the B3LYP/6-31G* level of theory. **b** Electronic density of the DPPC model phospholipid membrane in the presence of *trans*-GdAzo (0.1%) extracted from SAXS experiments (black, red: $x_{\text{trans-GdAzo}} = 0, 0.1\%$, respectively). **c** SAXS patterns of mixtures of DPPC and *trans*-GdAzo (black, blue, red: $x_{\text{trans-GdAzo}} = 0, 0.1, 7\%$, respectively). Inset: Variation of the d -spacing as a function of *trans*-GdAzo molar fraction.

Section 13). Indeed, the lamellar spacing (d -spacing, corresponding to the sum of the bilayer and water layer thicknesses) of the DPPC lamellar phase was increased from $63.7 \pm 0.3 \text{ \AA}$ to $65.9 \pm 0.8 \text{ \AA}$ upon addition of 0.1 mol% of *trans*-GdAzo. The location of *trans*-GdAzo was addressed by comparing the electron density profiles of bilayers of DPPC, on one hand, and DPPC with 0.1 mol% of *trans*-GdAzo on the other hand (Fig. 6b). The DPPC profile reflected the electron-rich phosphatidylcholine headgroups and electron-poor hydrocarbon chains, in agreement with previous studies⁶². The distance between the phosphate groups, d_{HH} , deduced from the position of the two maxima was 44.0 Å in pure DPPC. The addition of *trans*-GdAzo slightly shifted these maxima away from the bilayer center ($d_{\text{HH}}' = 46.4 \text{ \AA}$) and increased their intensity. These results are consistent with the insertion of *trans*-GdAzo between DPPC chains, with the electron-rich Gd-chelate moiety slightly protruding in water. Of note, the thickness of the bilayer hydrophobic core ($2D_{\text{C}} = 34.4$ and 28.5 \AA at 20 and 50 °C respectively)⁶² enables the insertion of the hydrophobic part of *trans*-GdAzo whose length can be estimated at 21.8–22.6 Å, according to the molecular model (cf. Supplementary Section 12). The addition of increasing amounts of *trans*-GdAzo to DPPC model membrane led to a progressive shift towards smaller scattering vectors q of the Bragg peaks ($q = 4\pi \sin(\theta)/\lambda$ where 2θ the scattering angle and λ the radiation wavelength; Fig. 6c). At room temperature, the d -spacing increased from $d = 64.3 \text{ \AA}$ in DPPC to $d = 80 \text{ \AA}$ in DPPC containing 7% of *trans*-GdAzo and then remained constant up to 15% of *trans*-GdAzo. This change in d -spacing was accompanied by a broadening and a decrease in intensity of the Bragg peaks, indicating a more disordered lamellar phase involving fewer bilayers. A broad maximum centred on $q \approx 0.12 \text{ \AA}^{-1}$, compatible with scattering from micelles, was also clearly observed for *trans*-GdAzo molar fraction $\geq 3\%$. These results suggest the coexistence of stacked bilayers and micelles, both of which would be mixed structures containing *trans*-GdAzo molecules. The interaction of *trans*-GdAzo with DPPC bilayers was further supported by differential scanning calorimetry (DSC) (Supplementary Section 13). The model emerging from all the SAXS and DSC results is that the partial solubilisation of phospholipid bilayers by *trans*-GdAzo at high concentration involved the formation of mixed micelles consisting of phospholipid nanodiscs whose hydrophobic edges were shielded from water by *trans*-GdAzo molecules.

The cell permeabilisation in the presence of either *cis*-GdAzo or *trans*-GdAzo was assessed by microscopic examination of cancer cells (PANC-1) incubated with propidium iodide (PI). While neither cell permeabilisation, nor cytotoxicity were observed with *cis*-GdAzo, the active *trans*-GdAzo isomer induced cell permeabilisation in few minutes after incubation (Fig. 7a, b and Supplementary Figs. 78–85). The cell permeabilisation of *cis*-GdAzo after activation into *trans*-GdAzo upon GR (2 Gy) was significantly higher compared to (i) exposure to *cis*-GdAzo without GR or (ii) GR in the absence of *cis*-GdAzo (Fig. 7c–e and Supplementary Figs. 86, 87). These observations revealed that *trans*-GdAzo could induce a loss of integrity for some cells leading to collapse (Fig. 7l and Supplementary Videos 1–3), which was attributed to the partial solubilisation of the phospholipid bilayers as shown by SAXS on model membrane (Supplementary Section 13). A similar collapse of cell structure due to the rapid breakdown of cell membrane has previously been reported for high concentration of the surfactant Triton X-100⁶³.

Electron energy-loss spectroscopy coupled to transmission electron microscopy (EELS-TEM) confirmed the presence of Gd in the permeabilised cells and revealed a heterogeneous distribution in the cell cytoplasm. The *trans*-GdAzo compound accumulated in specific cytoplasmic areas and was not homogeneously diffused (Fig. 7i and Supplementary Section 14.5). At this stage, we were wondering if *cis*-GdAzo activation could induce a lethal action by itself or favour drug penetration inside resistant cancer cells. The cytotoxicity induced by *cis*-GdAzo activation was assessed on a gemcitabine (Gem)-resistant cancer cell line (CCRF-CEM ARAC 8 C, human T lymphocytes) which does not express the hENT-1 membrane receptor required for the Gem to penetrate inside the cell, before phosphorylation and inhibition of DNA synthesis⁶⁴. First, the permeabilisation of the Gem-resistant cell membranes by *cis*-GdAzo upon GR (2 Gy) was confirmed using PI and flow cytometry quantification (Fig. 7f–h and Supplementary Section 14.4). To assess cytotoxicity, the cells were treated for 1 h with *cis*-GdAzo upon irradiation by GR (2 Gy) and the cell viability was assessed after four-day incubation in standard conditions. The treatment performed in the presence or absence of Gem confirmed the killing activity of *cis*-GdAzo upon IR and showed that the additional cytotoxicity effect of Gem was not significant (Fig. 7j, k and Supplementary Figs. 90, 91). The absence of any increase in Gem intracellular-accumulation in the resistant

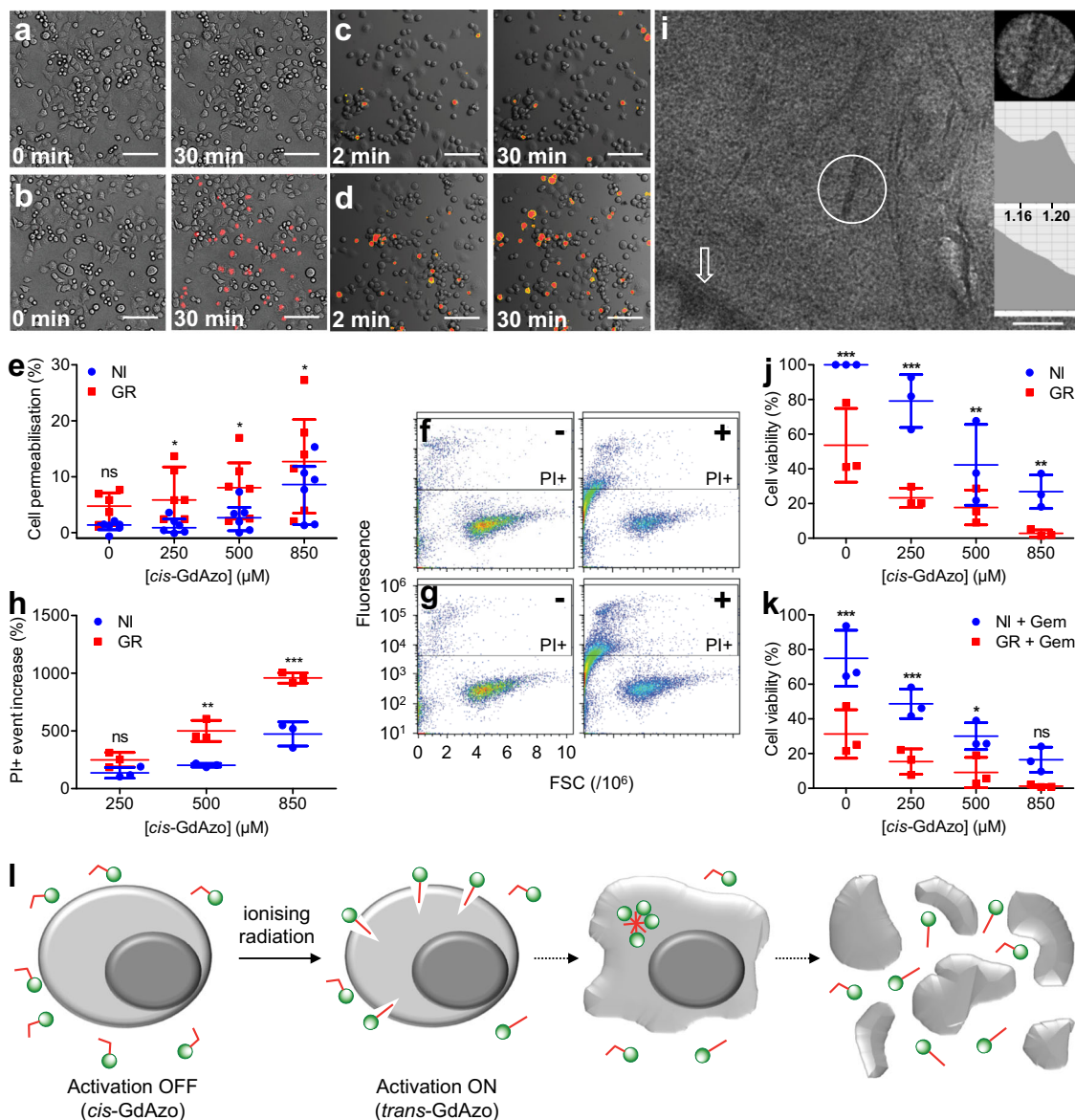


Fig. 7 Cell cytotoxicity triggered by *cis*-GdAzo activation upon IR. **a, b** Confocal microscopy of cancer cell (PANC-1) in the presence of PI before (0 min) and 30 min after incubation with either inactive *cis*-GdAzo (**a**) or active *trans*-GdAzo (**b**) (4 repetitions, permeabilised cells in red). **c, d** Confocal microscopy of cancer cell (PANC-1) in the presence of PI at 2 min and 30 min after *cis*-GdAzo introduction followed by GR (**d**) or no (**c**) irradiation (permeabilised cells in red). **e** Quantification of cell permeabilisation (PANC-1) by confocal microscopy after treatment with *cis*-GdAzo upon GR ($n = 6$ biologically independent samples, processed by a custom script, t -values = 3.110, 2.941, 2.815 at 250, 500, 850 μM). The medians \pm interquartile ranges are reported. **f, g** Flow cytometry of cancer cells (CCRF-CEM-ARAC-8C) in the presence of PI and *cis*-GdAzo (absence (-) or 500 μM (+)) upon GR (**g**) or without (**f**) irradiation at 30 min post-treatment. **h** Quantification of cell permeabilisation (CCRF-CEM-ARAC-8C) by flow cytometry after treatment with *cis*-GdAzo upon GR at 30 min post-irradiation ($n = 3$ biologically independent samples, t -values = 5.254, 8.565 at 500, 850 μM). The relative increase in PI-positive events compared to medium without *cis*-GdAzo is represented. **i**, EELS-TEM on cancer cell (PANC-1) in the presence of *trans*-GdAzo. Gd was detected in localised cytoplasmic areas (circle, top EELS spectrum, keV axis unit) and not in the surrounding cytoplasm (bottom EELS spectrum), cell membrane or nucleus (arrow). **j, k** Cell viability of Gem-resistant cancer cells (CCRF-CEM-ARAC 8 C) 4 days after treatment with *cis*-GdAzo upon GR in the absence (**j**) or presence (**k**) of Gem ($n = 3$ biologically independent samples, t -values = 6.854, 8.258, 3.636, 3.557 at 0, 250, 500, 850 μM in the absence of Gem and 6.454, 4.927, 3.099 at 0, 250, 500 μM in the presence of Gem). **l** Representation of the impact of *cis*-GdAzo on cancer cells upon IR as assumed from the microscopy experiments. The means \pm standard deviations are reported unless otherwise specified. Two-way Anova (Bonferroni post-test) was used for statistical analyses (NI vs GR). ns: not significant, $*P < 0.05$, $**P < 0.01$, $***P < 0.001$. Scale bars = 125 μm .

cancer cells in the presence of *trans*-GdAzo was confirmed by using radiolabelled Gem (Supplementary Fig. 93) and was attributed to dead-cell collapse. Even if the synergistic cytotoxic effect of *cis*-GdAzo and IR will need to be improved, this approach brought out that nearly complete cancer-cell death can be achieved at high concentration of *cis*-GdAzo upon IR (remaining of 2.8% living cells). It has to be noted that no

cytotoxicity was detected using a control compound containing the Gd-chelate without azobenzene modification (Dotarem[®]), thus confirming the need of the presence of the azobenzene moiety and refuting the idea that cytotoxicity could be due to the Gd-induced radiosensitisation only (Supplementary Fig. 92).

Photosensitive systems have been developed for many decades but their translation into clinic reveals to be very disappointing

mainly due to the non-penetrating lights they need for activation. We overcome here this limitation by designing a radioswitch, i.e. a photoswitch system adapted to IR at low clinical dose such as currently used for radiotherapy, and inducing a cell-permeabilising and destructuring effect. Thanks to the Gd-chelate moiety associated to the photosensitive system, theranostic approach should be possible by *in vivo* MRI detection of the prodrug before triggering a localised therapeutic action. Taking advantage of the various pharmacological actions able to be triggered by photoswitch systems, many cellular manipulations and therapeutic approaches could come out as the cell permeabilisation described in this work. This exciting new development in photochemistry opens the way towards novel opportunities in the translation of photoswitching molecular tools currently limited to research area into clinical applications.

Methods

The synthesis procedures and characterisation of the **Azo** and **GdAzo** compounds are described in the Supplementary Sections 1–3. The methods to assess activation upon IR are described in the Supplementary Sections 4, 5. The Monte Carlo simulation is reported in the Supplementary Section 6. The study on activation mechanism upon IR in the presence of various quenchers/converters is detailed in the Supplementary Section 7. Investigations on the role of hydroxyl radicals (experiments under gas saturation and Fenton chemistry), Gd (quantification of hydroxyl radicals and activation of *cis*-**Azo** in the presence of Gd) and the impact of *cis*-**GdAzo** concentration are reported in the Supplementary Sections 8–10. The physicochemical characterisation, DFT calculations and study of **GdAzo** aggregation (SAXS) and interactions with cell membrane model (SAXS and DSC) are described in the Supplementary Sections 11–13. The methods to assess *in vitro* cytotoxicity are reported in the Supplementary Section 14. The main methods are listed below even if more details are provided in the Supplementary Sections.

Quantification of *cis*-GdAzo activation upon IR. *trans*-**GdAzo** and the control compound without Gd (*trans*-**Azo**) (50 μM , 200 μL , PBS) were introduced in two 96-well microplates. Both microplates were irradiated by UV (365 nm, 0.817 $\text{mW}\cdot\text{cm}^{-2}$, 5 min) to obtain the PSS1 containing a majority of the *cis*-isomer (90 \pm 3%). One microplate (plate 1) was kept in the dark and used as control (non-irradiated by IR) whereas the second microplate (plate 2) was irradiated upon incremental doses of IRs (2, 3, 5 and 10 Gy). After each irradiation, absorbance and HPLC (method C) analyses were performed on the non-irradiated (plate 1) and IR-irradiated (plate 2) compounds. A time delay between the UV irradiation of plate 1 and plate 2 was introduced to analyse both the control (plate 1) and the IR-treated (plate 2) compounds concurrently. The relative amount of each isomer was obtained by running HPLC and the molecular activation (%) was determined by the difference in proportion of the *trans*-isomer in the media. The experiment was repeated 3 times independently.

For investigation on the activation mechanism using scavengers, the *cis*-**GdAzo** compound was dissolved in different media: water (control), aqueous solutions of *tert*-butanol (10, 100, 524 mM), mannitol (10, 100, 524 mM), ethanol (10, 100, 524 mM), sodium azide (1, 10 and 50 mM), dimethylsulfoxide (15% *v/v*), sodium selenate (1 and 25 mM) and cadmium perchlorate (20 mM). After each irradiation, absorbance analyses were performed on the non-irradiated (plate 1) and IR-irradiated (plate 2) compounds (triplicate).

***cis*-GdAzo activation under N_2 and N_2O gas saturation.** *trans*-**GdAzo** (50 μM , 200 μL , H_2O) was introduced in a 96-well microplate before UV irradiation (365 nm, 0.817 $\text{mW}\cdot\text{cm}^{-2}$, 10 min, *cis*-isomer at 90 \pm 3%). The compound was then introduced into two sealed glass tubes (700 μL) and the medium was saturated by N_2 or N_2O gas bubbling for 15 min. One of the glass tube was kept in the dark (control) whereas the second one was irradiated upon incremental doses of GRs (2, 3, 5 and 10 Gy). After each irradiation, HPLC analyses were performed (method C) and the molecular activation (%) was determined by the difference in proportion of the *trans*-isomer in the media. The experiment was repeated 3 times independently.

***cis*-GdAzo activation using Fenton chemistry.** *cis*-**GdAzo** (1.000 mL, final concentration 50 μM , H_2O), EDTA (9.00 μL , final concentration 75 μM , H_2O) and FeCl_2 (9.00 μL , final concentration 75 μM , H_2O) were introduced in an 1.5 mL Eppendorf tube. A first HPLC analysis was run before the introduction of H_2O_2 . The Fenton reaction was started by the addition of H_2O_2 (4.41 μL , final concentration 50 mM, H_2O) into the mixture which was stirred (orbital) in the dark. The proportion of *trans*-**GdAzo** was determined by HPLC (method C) 5 min after the addition of H_2O_2 . The experiment was repeated 3 times independently.

Quantification of hydroxyl radicals. Coumarin (600 μL , final concentration 0.5 mM, H_2O) and GdCl_3 (0–168 μL , final concentrations from 0 to 2000 μM , H_2O) were mixed and introduced in a 96-well microplate (200 μL , H_2O , duplicate). The

microplate was irradiated upon incremental doses of ionising radiations (final doses of 0, 2, 5, 10, 15, 20, 25, 30 and 40 Gy). After each irradiation, fluorescence (ex: 355 nm, em: 460 nm) was measured. The experiment was repeated 5 times independently. The standard curve of 7-OH-Coum led to quantify the production of 7-OH-Coum by increasing the radiation dose at different Gd^{3+} concentrations. The slope of these lines (related to $G(7\text{-OH-Coum})$) were plotted against the Gd^{3+} concentrations to assess the impact of Gd^{3+} on HO^\bullet production. Finally, $G(\text{HO}^\bullet)$ was determined by applying a conversion yield of Coum into 7-OH-Coum of 3.1% (cf. Supplementary Section 9).

Cell permeabilisation of *cis*-GdAzo upon GR using confocal microscopy. The cells (PANC-1) were seeded (10,000 cells in 200 μL /well) in imaging chamber and maintained in culture medium in a humid atmosphere at 37 $^\circ\text{C}$ with 5% CO_2 . 24 h postseeding, culture medium was replaced by PBS (95 μL) and propidium iodide (PI, 5 μL , final concentration 1 μM) was added in the medium. 15 min after addition, *cis*-**GdAzo** (100 μL , final concentration 0, 250, 500, or 850 μM , PBS) was introduced in the medium. A first set of images was acquired at this stage and then the imaging chamber was irradiated (GR, 2 Gy). The imaging chamber was kept in the dark at 37 $^\circ\text{C}$ using a stage heater and images were acquired each 5 min for 30 min. A similar procedure was used for the non-irradiated control experiment except that the imaging chamber was not irradiated upon GR. The cells were observed with an inverted Nikon microscope (10x dry objective lens). The red fluorescence emission of PI (ex: 561 nm, em: 598–672 nm) and transmission images were collected. Four images per well were acquired and two wells were used for each concentration of *cis*-**GdAzo**. The experiment was repeated 3 times independently and each well was considered as a biological independent replicate. The number of cells on images acquired before irradiation was manually counted and the number of fluorescent cells on images acquired before and after GR irradiation was automatically determined using an in-house script for ImageJ software⁶⁵ (Version 1.50i completed with Adjustable Watershed plugin).

Cell permeabilisation of *cis*-GdAzo upon GR using flow cytometry. Just before the experiment, cells (CCRF-CEM ARAC-8C) were dispersed in PBS and transferred in 96-well microplate (40,000 cells in 90 μL /well). *cis*-**GdAzo** (90 μL , final concentration 0, 250, 500, or 850 μM , PBS) was introduced before irradiation of the medium by GR (2 Gy). Then, PI (5 μL , final concentration 1 μM) was added and the medium was maintained in the dark at room temperature. Flow cytometry analyses were run at 15, 30, and 45 min after irradiation using a BD Accuri™ C6 Plus flow cytometer (runs of 30 μL , 100 $\mu\text{L}/\text{min}$, no threshold). A similar procedure was used for the non-irradiated control experiment except that the microplate was not irradiated upon GR. Cell membrane permeabilisation was quantified by numbering the PI-positive events (duplicate) and represented as the relative increase in PI-positive events compared to medium without *cis*-**GdAzo**. The experiment was repeated 3 times independently. The data were treated using the BD Accuri C6 Plus software (version 1.0.27.1) and the representative images were obtained using the Flowjo software (version 10.7.1).

Cytotoxicity of *cis*-GdAzo upon GR. Just before the experiment, cells (CCRF-CEM ARAC-8C) were dispersed in PBS and transferred in 48-well microplate (40,000 cells in 80 μL /well). Gemcitabine (Gem, 20 μL , final concentration 0.1 μM) or PBS (20 μL) and *cis*-**GdAzo** (100 μL , final concentration 0, 250, 500 or 850 μM , PBS) were added before irradiation of the medium by GR (2 Gy). The medium was maintained in the dark in a humid atmosphere at 37 $^\circ\text{C}$ with 5% CO_2 for 1 h. Then, culture medium (600 μL) was added in the wells and the cells were washed by 3 centrifugation cycles (300 g, 5 min). Cells were dispersed in culture medium (600 μL) containing Gem (0 or 0.1 μM final concentrations) and were maintained in a humid atmosphere at 37 $^\circ\text{C}$ with 5% CO_2 for 4 days. Living cell number was then determined by cell counting in presence of trypan blue 1:1 *v/v* (triplicate). The experiment was repeated 3 times independently. The cell viability was expressed as the ratio of living cell number after treatment to living cell number without any treatment (non-irradiated, no Gem and no *cis*-**GdAzo**).

Reporting summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability

All characterisation data and experimental protocols to evaluate the conclusions in the paper are available in the manuscript and/or the Supplementary Information. Moreover, source data is available for all the figures and Supplementary Figures in the associated Source Data file, and the main data generated in this study have been deposited in the Zenodo database⁶⁶ under accession code <https://doi.org/10.5281/zenodo.6379759>.

Code availability

The custom codes developed for this study are available at the Zenodo database⁶⁶ under accession code <https://doi.org/10.5281/zenodo.6379759>. They can be run with imageJ

(version 1.50i completed with Adjustable Watershed plugin, to determine cell permeabilisation from the optical microscopy data) and with UCSF Chimera (version 1.14, to classify and display polar and non-polar surfaces from the DFT calculations).

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References

- Aguado, B. A., Grim, J. C., Rosales, A. M., Watson-Capps, J. J. & Anseth, K. S. Engineering precision biomaterials for personalized medicine. *Sci. Transl. Med.* **10**, eaam8645 (2018).
- Qiao, Y. et al. Stimuli-responsive nanotherapeutics for precision drug delivery and cancer therapy. *WIREs Nanomed. Nanobiotechnol.* **11**, e1527 (2019).
- Liu, D., Yang, F., Xiong, F. & Gu, N. The smart drug delivery system and its clinical potential. *Theranostics* **6**, 1306–1323 (2016).
- Glusac, K. What has light ever done for chemistry? *Nat. Chem.* **8**, 734 (2016).
- Hüll, K., Morstein, J. & Trauner, D. In vivo photopharmacology. *Chem. Rev.* **118**, 10710–10747 (2018).
- Vickerman, B. M., Zywoot, E. M., Tarrant, T. K. & Lawrence, D. S. Taking phototherapeutics from concept to clinical launch. *Nat. Rev. Chem.* **6**, 1–19 (2021).
- Lan, M. et al. Photosensitizers for photodynamic therapy. *Adv. Healthc. Mater.* **8**, 1900132 (2019).
- Fisher, C. J. et al. Photodynamic therapy for the treatment of vertebral metastases: A phase I clinical trial. *Clin. Cancer Res.*, clincanres.0673.2019 (2019).
- Sato, K. et al. Spatially selective depletion of tumor-associated regulatory T cells with near-infrared photoimmunotherapy. *Sci. Transl. Med.* **8**, 352ra110 (2016).
- Ash, C., Dubec, M., Donne, K. & Bashford, T. Effect of wavelength and beam width on penetration in light-tissue interaction using computational methods. *Lasers Med. Sci.* **32**, 1909–1918 (2017).
- Štacko, P. et al. Locked synchronous rotor motion in a molecular motor. *Science* **356**, 964–968 (2017).
- Cosentino, C. et al. Engineering of a light-gated potassium channel. *Science* **348**, 707–710 (2015).
- Gautier, A. et al. How to control proteins with light in living systems. *Nat. Chem. Biol.* **10**, 533 (2014).
- Mickle, A. D. et al. A wireless closed-loop system for optogenetic peripheral neuromodulation. *Nature* **565**, 361–365 (2019).
- Zhang, Z. et al. “Uphill” cation transport: A bioinspired photo-driven ion pump. *Sci. Adv.* **2**, e1600689 (2016).
- Baumann, M. et al. Radiation oncology in the era of precision medicine. *Nat. Rev. Cancer* **16**, 234 (2016).
- Santos Mello, R., Callisen, H., Winter, J., Kagan, A. R. & Norman, A. Radiation dose enhancement in tumors with iodine. *Med. Phys.* **10**, 75–78 (1983).
- Song, G., Cheng, L., Chao, Y., Yang, K. & Liu, Z. Emerging nanotechnology and advanced materials for cancer radiation therapy. *Adv. Mater.* **29**, 1700996 (2017).
- Ni, K. et al. Nanoscale metal-organic frameworks for x-ray activated in situ cancer vaccination. *Sci. Adv.* **6**, eabb5223 (2020).
- Fan, W. et al. Breaking the depth dependence by nanotechnology-enhanced X-ray-excited deep cancer theranostics. *Adv. Mater.* **31**, 1806381 (2019).
- Chen, X., Song, J., Chen, X. & Yang, H. X-ray-activated nanosystems for theranostic applications. *Chem. Soc. Rev.* **48**, 3073–3101 (2019).
- Sun, W., Zhou, Z., Pratz, G., Chen, X. & Chen, H. Nanoscintillator-Mediated X-Ray Induced Photodynamic Therapy for Deep-Seated Tumors: From Concept to Biomedical Applications. *Theranostics* **10**, 1296–1318 (2020).
- Barosi, A. et al. Synthesis and activation of an iron oxide immobilized drug-mimicking reporter under conventional and pulsed X-ray irradiation conditions. *RSC Adv.* **10**, 3366–3370 (2020).
- Wu, S.-Y. et al. Radiation-sensitive dendrimer-based drug delivery system. *Adv. Sci.* **5**, 1700339 (2018).
- Fu, Q. et al. External-radiation-induced local hydroxylation enables remote release of functional molecules in tumors. *Angew. Chem. Int. Ed.* **59**, 21546–21552 (2020).
- Zhang, F. et al. X-ray-triggered NO-released Bi-SNO nanoparticles: all-in-one nano-radiosensitizer with photothermal/gas therapy for enhanced radiotherapy. *Nanoscale* **12**, 19293–19307 (2020).
- Fan, W. et al. Generic synthesis of small-sized hollow mesoporous organosilica nanoparticles for oxygen-independent X-ray-activated synergistic therapy. *Nat. Commun.* **10**, 1241 (2019).
- Su, M., Guggenheim, K. G., Lien, J., Siegel, J. B. & Guo, T. X-ray-mediated release of molecules and engineered proteins from nanostructure surfaces. *ACS Appl. Mater. Interfaces* **10**, 31860–31864 (2018).
- Zhou, Z. et al. Synchronous chemoradiation nanovesicles by X-Ray triggered cascade of drug release. *Angew. Chem. Int. Ed.* **57**, 8463–8467 (2018).
- Deng, W. et al. Controlled gene and drug release from a liposomal delivery platform triggered by X-ray radiation. *Nat. Commun.* **9**, 2713 (2018).
- Kuncic, Z. & Lacombe, S. Nanoparticle radio-enhancement: principles, progress and application to cancer treatment. *Phys. Med. Biol.* **63**, 02TR01 (2018).
- Bort, G. et al. EPR-mediated tumor targeting using ultrasmall-hybrid nanoparticles: From animal to human with theranostic AGuIX nanoparticles. *Theranostics* **10**, 1319–1331 (2020).
- Broichhagen, J., Frank, J. A. & Trauner, D. A roadmap to success in photopharmacology. *Acc. Chem. Res.* **48**, 1947–1960 (2015).
- Dong, M., Babalhavaeji, A., Samanta, S., Beharry, A. A. & Woolley, G. A. Red-shifting azobenzene photoswitches for in vivo use. *Acc. Chem. Res.* **48**, 2662–2670 (2015).
- Xie, J. et al. Emerging strategies of nanomaterial-mediated tumor radiosensitization. *Adv. Mater.* **31**, 1802244 (2019).
- Liu, Y. et al. Metal-based NanoEnhancers for future radiotherapy: Radiosensitizing and synergistic effects on tumor cells. *Theranostics* **8**, 1824–1849 (2018).
- McMahon, S. et al. Biological consequences of nanoscale energy deposition near irradiated heavy atom nanoparticles. *Sci. Rep.* **1**, 18 (2011).
- McMahon, S. J., Paganetti, H. & Prise, K. M. Optimising element choice for nanoparticle radiosensitisers. *Nanoscale* **8**, 581–589 (2016).
- Lux, F. et al. AGuIX® from bench to bedside—Transfer of an ultrasmall theranostic gadolinium-based nanoparticle to clinical medicine. *Brit. J. Radiol.* **91**, 20180365 (2018).
- Bonvalot, S. et al. NBTXR3, a first-in-class radioenhancer hafnium oxide nanoparticle, plus radiotherapy versus radiotherapy alone in patients with locally advanced soft-tissue sarcoma (Act.In.Sarc): A multicentre, phase 2-3, randomised, controlled trial. *Lancet Oncol.* **20**, 1148–1159 (2019).
- Bobrowski, K. *Applications of Ionizing Radiation in Materials Processing* Vol. 1 (eds Yongxia Sun & Andrzej Grzegorz Chmielewski) Ch. 4, p. 81–116 (Institute of Nuclear Chemistry and Technology, Warszawa, 2017).
- Evans, R. D. *The Atomic Nucleus* (ed T.M.H. Edition) Ch. 25, p. 711–745 (Tata McGraw-Hill, Bombay-New Delhi, 1955).
- Francese, S. in *PENELOPE-2014 - A Code System for Monte Carlo Simulation of Electron and Photon Transport*. NEA/NSC/DOC(2015)3 (Workshop Barcelona, Spain, 2015).
- Frelon, S., Douki, T., Favier, A. & Cadet, J. Comparative study of base damage induced by gamma radiation and Fenton reaction in isolated DNA. *J. Chem. Soc., Perkin Trans. 1*, 2866–2870 (2002).
- Miller, C. J., Rose, A. L. & Waite, T. D. Importance of iron complexation for Fenton-mediated hydroxyl radical production at circumneutral pH. *Front. Mar. Sci.* **3**, 4799 (2016).
- Goulet-Hanssens, A. et al. Hole catalysis as a general mechanism for efficient and wavelength-independent Z → E azobenzene isomerization. *Chem* **4**, 1740–1755 (2018).
- Buxton, G. V. *Radiation Chemistry* (ed. Mehran Mostafavi Mélanie Spothheim-Maurizot, Thierry Douki & Jacqueline Belloni) Ch. 1, p. 3–16 (EDP Sciences, France, 2008).
- Buxton, G. V., Greenstock, C. L., Phillips Helman, W. & Ross, A. B. Critical Review of rate constants for reactions of hydrated electrons, hydrogen atoms and hydroxyl radicals ($\cdot\text{OH}/\cdot\text{O}^-$) in Aqueous Solution. *J. Phys. Chem. Ref. Data* **17**, 513–886 (1988).
- Fulford, J., Bonner, P., Goodhead, D. T., Hill, M. A. & O'Neill Experimental determination of the dependence of OH radical yield on photon energy: A comparison with theoretical simulations. *J. Phys. Chem. A* **103**, 11345–11349 (1999).
- Singh, T. S., Madhava Rao, B. S., Mohan, H. & Mittal, J. P. A pulse radiolysis study of coumarin and its derivatives. *J. Photochem. Photobiol.* **153**, 163–171 (2002).
- Louit, G. et al. The reaction of coumarin with the OH radical revisited: hydroxylation product analysis determined by fluorescence and chromatography. *Radiat. Phys. Chem.* **72**, 119–124 (2005).
- Sicard-Roselli, C. et al. A New Mechanism for Hydroxyl Radical Production in Irradiated Nanoparticle Solutions. *Small* **10**, 3338–3346 (2014).
- Brun, E., Girard, H. A., Arnault, J.-C., Mermoux, M. & Sicard-Roselli, C. Hydrogen plasma treated nanodiamonds lead to an overproduction of hydroxyl radicals and solvated electrons in solution under ionizing radiation. *Carbon* **162**, 510–518 (2020).
- Baroncini, M. & Bergamini, G. Azobenzene: A photoactive building block for supramolecular architectures. *Chem. Rec.* **17**, 700–712 (2017).
- Tochitsky, I. et al. Optochemical control of genetically engineered neuronal nicotinic acetylcholine receptors. *Nat. Chem.* **4**, 105–111 (2012).
- Beharry, A. A. & Woolley, G. A. Azobenzene photoswitches for biomolecules. *Chem. Soc. Rev.* **40**, 4422–4437 (2011).
- Iamsaard, S. et al. Conversion of light into macroscopic helical motion. *Nat. Chem.* **6**, 229–235 (2014).

58. Iwaso, K., Takashima, Y. & Harada, A. Fast response dry-type artificial molecular muscles with [c2]daisy chains. *Nat. Chem.* **8**, 625–632 (2016).
59. Kowalik, L. & Chen, J. K. Illuminating developmental biology through photochemistry. *Nat. Chem. Biol.* **13**, 587–598 (2017).
60. Wyart, C. et al. Optogenetic dissection of a behavioural module in the vertebrate spinal cord. *Nature* **461**, 407–410 (2009).
61. Sebai, S. C. et al. Photocontrol of the translocation of molecules, peptides, and quantum dots through cell and lipid membranes doped with azobenzene copolymers. *Angew. Chem. Int. Ed.* **51**, 2132–2136 (2012).
62. Nagle, J. F. & Tristram-Nagle, S. Structure of lipid bilayers. *Biochim. Biophys. Acta* **1469**, 159–195 (2000).
63. Koley, D. & Bard, A. J. Triton X-100 concentration effects on membrane permeability of a single HeLa cell by scanning electrochemical microscopy (SECM). *Proc. Natl Acad. Sci. USA* **107**, 16783–16787 (2010).
64. Mackey, J. R. et al. Functional nucleoside transporters are required for gemcitabine influx and manifestation of toxicity in cancer cell lines. *Cancer Res* **58**, 4349–4357 (1998).
65. Schneider, C. A., Rasband, W. S. & Eliceiri, K. W. NIH Image to ImageJ: 25 years of image analysis. *Nat. Methods* **9**, 671 (2012).
66. Guesdon-Vennerie, A. et al. Breaking photoswitch activation depth limit using ionising radiation stimuli adapted to clinical application, CERN (Zenodo), <https://doi.org/10.5281/zenodo.6379759>, 2022.

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Author contributions

G.Bo. devised, designed and supervised the project. G.Bo., C.L.M., F.A., and A.G.-V. carried out organic syntheses. G.Bo. and S.Mé. conducted the relaxivity measurements.

G.Bo. and A.G.-V. performed the experiments to determine conversion upon IR. G.Bo., A.G.-V. and F.P. carried out and discussed the experiments upon IR in the presence of scavengers. G.Bo. performed the experiments to validate the role of hydroxyl radicals (using scavengers, gas-saturated solutions, Fenton chemistry) and to investigate the mechanism of activation (using different concentrations of GdAzo, quantification of hydroxyl radicals). I.M.-R. performed the Monte Carlo simulation. G.Be. and G.Bo. conducted the DFT calculations. C.B., F.-X.L., and G.Bo. performed the SAXS and DSC experiments. S.Ma., S.T., and G.Bo. carried out the EELS-TEM experiment. G.Bo. carried out the experiments on cancer cells using microscopy and flow cytometry. G.Bo., C.R., and F.P. carried out the cytotoxicity experiments on cancer cells. G.Bo., P.C., and S.Mu. discussed the results. G.Bo. wrote the article. All the authors discussed the results and validated the last version of the article.

Competing interests

A patent associated with this work has been filed by G.Bo., P.C., S.Mu. and F.P.(WO/2020/161308). All other authors declare no competing interests.

Additional information

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