Peroxoniobium inhibits leukemia cell growth†

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A new class of polyoxoniobate complex has been synthesized and characterized as a novel anticancer agent for photodynamic therapy. The complex inhibits the growth of chronic myelogenous leukemia cells with an IC₅₀ value of 30 μM, in the dark. However, upon exposure to light (365 nm) there is a fivefold increase in the cytotoxic activity. Light radiation activates the complex with the formation of radical species capable of interacting with DNA according to our experimental and theoretical data.

In this work, we prepared a photosensitive peroxoniobium complex presenting a balance with an active radical phase when illuminated with radiation of 365 nm. A versatile niobium species of amorphous structure was obtained by the reaction of niobium ammonium oxalate with ammonium hydroxide up to pH 7. The material obtained, a niobium oxyhydroxide (NbO₂(OH)) (white solid),[1,4] can be modified with the generation of NbO₂(OH)O₂⁻ peroxo groups (yellow solid).[3] The yellow compound is formed by treatment with H₂O₂. The absorption radiation in the visible region due to the charge transfer transition between the peroxo group and the niobium is shown in Fig. 1.

This complex with the radical as an intermediate is favored in the presence of visible and UV radiation. This property is of interest for photodynamic therapy of cancer (PDT), which involves the exposure of malignant cells containing a photosensitizer molecule to light irradiation, in the presence of oxygen species. The photoactivated drug produces reactive oxygen species that initiate a series of events, resulting in cell death. Selective light activation allows a preferential tumor destruction in comparison to healthy tissues.[4] Several metal complexes exhibit photocytotoxicity under UV or visible light,[5,6] but data about niobium compounds are very scarce in the literature.[7]

The peroxoniobium, generated from niobium oxyhydroxide described here can be very active in the treatment of diseased cells when illuminated with visible or UV radiation due to its light absorption capacity because of the peroxo groups formed. The peroxoniobium complex has some advantages, such as ease of synthesis and in mild conditions, high solubility, low activity under light off, and resistance to inactivation by thiol reagents. Moreover, it is nontoxic and does not employ noble metals like most of the compounds proposed in the literature. Actually, niobium oxide was tested as a bone implant component and showed absence of inflammatory cells or degeneration of the osteoblasts without any sign of damage to the preexisting bone tissue, showing compatibility with the bone tissue.[8–11]

The innovative part in the process of obtaining the polyoxoniobate complex presented in this work consists in the leaching of the complex when treating the niobium oxyhydroxide with H₂O₂. With the treatment, a yellow solid and a leached yellow liquid is obtained, which is the complex containing peroxoniobium in its structure, sensitive to the UV-Vis radiation generating radical species. This species generated with the leaching at neutral pH presents high negative charge and a kinetic volume of 223 nm. The XRD of the lyophilized polyoxoniobate complex presented in this work consists in the leaching of the catalysts.

Fig. 1 UV-Vis profile of the catalysts.

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is clear that the pattern of the decaniobate structure is closer to the experimental spectrum. One should keep in mind that DFT frequencies are normally 10% underestimated with respect to the experimental values. The broader absorption below 600 cm$^{-1}$ can be attributed to the interaction between different decaniobate structures forming the amorphous solid. The calculated peaks at 710, 760 and 860 cm$^{-1}$ are related to the experimental peaks of 800, 870 and 910 cm$^{-1}$ indicating that decaniobate is the local arrangement of the polyoxoniobate complex.

The generation of reactive oxygen under radiation was confirmed by the reaction of the complex with an organic dye, which was monitored by UV-Vis spectroscopy (Fig. 3). The spectrum of the dye solution shows the characteristic peak of the methylene blue (MB) at 663 nm (black trace). It can be clearly seen that in the presence of the peroxoniobium complex and radiation (365 nm) the signal decreased indicating the reaction of the peroxoniobium complex with the dye (blue trace). In the absence of light, there is no decrease in the signal related to the dye, indicating the need of the radiation to activate the oxidation action of the peroxoniobium complex. The equilibrium in which the radical species forms it is not necessary to use a photosensitizer agent, such as porphyrins.$^4$ A further investigation was carried out by $^{31}$P NMR (Fig. S1†) using guanosine as model molecule able to react with the peroxoniobium complex. The $^{31}$P NMR spectrum shown in Fig. S1-a† corresponds to 5-GMP and revealed that the phosphorus atom in the structure exhibits a chemical shift at $\delta$ 5.93. When 5-GMP and the polyoxoniobate are in contact, no significant changes are observed in the $^{31}$P spectrum, only a small displacement of the phosphorus signal to $\delta$ 5.90 (Fig. S1-b†). However, when the 5-GMP and polyoxoniobate mixture is submitted to radiation (Fig. S1-c†), an interaction between the compounds occurs, giving rise to a new species that presents a different chemical shift in the P spectrum ($\delta$ 5.27).

The effect of the peroxoniobium on the growth of K562 cells was evaluated after 4 h of incubation. The compound inhibits K562 cell growth in a concentration-dependent manner, with an IC$_{50}$ of 30.0 ± 1.5 μmol L$^{-1}$. Ammonium niobate(V) oxide was also tested and it has no effect on K562 cells up to 100 μM. The cytotoxic activity of polyoxoniobate increases by 5 times upon 5 min of UV-A light irradiation, with an IC$_{50}$ value of 6.2 ± 0.4 μmol L$^{-1}$ (Fig. 4). The higher activity, when exposed to light, associated to the low toxicity of niobium compounds place the peroxoniobium complex as a candidate for photodynamic therapy.

There are few reports in the literature about the cytotoxic activity of niobium compounds. A peroxoniobium complex with ascorbic acid ([K$_3$[Nb(Asc)(O$_2$)$_3$]]) is moderately active in HL60 human leukemia cells but not in K562 human myelogenous leukemia cells.$^{12}$ A tetrameric Nb$_{28}$-containing cluster inhibits the growth of the human breast cancer MCF-7 cells line with an IC$_{50}$ value of 5.21, after 48 h of incubation.$^{13}$ Methylene blue (MB) is one of the main photosensitizing agents used in PDT due to its good tissue penetration and low
It is active in several types of tumors upon irradiation with red laser light. This fact allied to the ability of the peroxoniobium compound to interact with MB (Fig. 3) prompted us to investigate its effect in the MB photocytotoxicity. We have first checked that exposure to UV-A light did not affect the photocytotoxicity of MB in K562 cells (Table 1). Afterwards, the photocytotoxicity of MB in K562 cells was assessed in the presence of the peroxoniobium complex at its IC₅₀ value. Cells were incubated with increasing concentrations of MB and irradiated for 5 min with UV-A light, in the absence and the presence of 6.5 μM of the peroxoniobium complex. Upon light irradiation, 6.5 μM of peroxoniobium complex doubles the effect of MB. Therefore, the peroxoniobium compound prepared is a promising candidate for PDT therapy, alone or in combination with other sensitizing agents, such as methylene blue dye (Table 1).

The peroxoniobium complex (Fig. S2†) and the DNA/complex systems were thus fully optimized at DFT level, with conjugate gradient and quasi-Newton–Raphson algorithms. The final geometries were obtained with the density functional Becke’s three-parameter exchange functional and the gradient-corrected functional of Lee, Yang and Parr (B3LYP), using LanL2DZ basis set.

The DICKERSON-DREW B-DNA DODECAMER was obtained from the Protein Data Bank (PDB), with code 4C64 and resolution: 1.32 A (ref. 18 and 19) and has shown suitable for our calculations. As previously discussed, the Nb complexes (Fig. S2†) were entirely optimized at the DFT level, to obtain the most stable initial geometries to perform the calculations with the DNA structure. It is important to mention that the more stable complex/DNA system is related to a higher cytotoxicity potential. The following species were considered for this theoretical investigation: Complex a (no radical groups), Complex b (protonated structure) and Complex c (radicals formation). The natural charges of all atoms were elucidated, and according to these data, it is possible to realize more pronounced negative charges referred to the radical species. The protonated substituent (OOH) presented a charge value equals to –0.534 a.u., while the corresponding radical (deprotonated) has shown a charge value significantly lower (–1.170 a.u.). The same is observed for the substituent (OH), with a charge value of –0.379 a.u., and the corresponding radical (–0.739 a.u.). The formation of more negative charges suggests to the highest reactivity of Complex c, in relation to the other complexes. Complex a was put together with DNA in three distinct regions (Fig. S3†), and after performing the optimization, the Nb complex reactivity was analyzed in these zones.

PM6 calculations were performed in order to evaluate the Nb complex (Complex a) affinity in different DNA regions. These results are presented in Table S1† as relative interaction energy values. The Complex a, when put into different regions of DNA, presented quite relevant changes in relation to the intermolecular interaction energy. Thus, Complex a, when docked into the central region of DNA, showed a more favorable energy (Fig. 5). After the optimization structure, the region with the highest interaction between DNA and Nb complexes was considered for other calculations.

According to our theoretical methodology, we have the energy minimizations for the systems: free DNA (E_{DNA}), free complex (E_{complex}) and DNA-complex system (E_{system}). In line with those systems, the affinity energy was calculated using the following equation:

\[ E = E_{DNA/complex} - E_{complex} - E_{DNA} \] (1)

The results from this methodology are described in Table S2,† also as relative interaction energy values. By using the strategy described at the ESI,† it was possible to analyze the efficiency of Nb complexes towards DNA, evaluating which factors contribute most to this reactivity. Our first results indicate that the Nb complex, in general, presents a significant affinity with DNA, with a pronounced increase in the affinity/reactivity in the presence of radical groups (–OO’ radical for example). According to our calculations, Complex a showed an intermediate reactivity towards DNA, presenting a significantly higher energy value than that obtained for Complex c. The energy difference between Complex a and c was 2.88 kcal mol⁻¹.

On the other hand, the energy difference between Complex b and c was remarkable, about 5.01 kcal mol⁻¹. Complex c presented good interaction potential with DNA, undoubtedly due to the radical groups added to the complex structure, coherent with the experimental observations. According to the computational investigation, we can conclude that all Nb complex species (without and with radical groups) presented reactivity and stability when docked into the DNA crystallographic structure. These results corroborate with the experimental data observed in the reaction of guanosine with the niobium complex shown in the ³¹P NMR of Fig. S1.† Furthermore, the addition of radical groups substantially increases the

| Table 1 Effect of the peroxoniobium complex on the photocytotoxicity of methylene blue (MB) in K562 cells |
|---|---|---|
| Compound | IC₅₀⁻⁻ | IC₅₀ irradiated⁻⁻ |
| MB | 7.3 ± 0.4 | 7.0 ± 0.5 |
| MB + NbO₂(OH)−O₂⁻ | 6.3 ± 0.3 | 3.0 ± 0.1 |

⁎ IC₅₀ Methylene blue concentration required to inhibit 50% of cell growth under dark conditions. † IC₅₀ Methylene blue concentration required to inhibit 50% of cell growth after 5 min of UV-A irradiation.

Assays were performed in the presence of 6.5 μM of the peroxoniobium complex.

Fig. 5 Representation of the DNA-complex system.
affinity of the complex towards DNA, supported by the obtaining of more stable structures for the complex/DNA system (lowest energy values), suggesting higher levels of cytotoxicity.

**Conflicts of interest**

The authors declare no competing financial interests.

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**References**


