

Photoredox Reactions of Lead(IV) and -(II) Hydroxo Complexes

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Ligand-to-metal charge-transfer excitation of $\text{Pb}(\text{OH})_6^{2-}$ in 0.1 M NaOH leads to the formation of $\text{Pb}(\text{OH})_3^-$ with $\phi = 0.07$ at $\lambda_{\text{irr}} = 254$ nm. Hydrogen peroxide was detected as an oxidation product. Upon sp excitation, $\text{Pb}(\text{OH})_3^-$ in 0.1 M NaOH shows a yellow emission at $\lambda_{\text{max}} = 570$ nm with $\phi = 1.5 \times 10^{-3}$ at $\lambda_{\text{irr}} = 254$ nm. In the presence of oxygen, this emission is partially quenched ($\phi = 1.2 \times 10^{-3}$) and $\text{Pb}(\text{OH})_3^-$ is photooxidized to $\text{Pb}(\text{OH})_6^{2-}$ with $\phi = 0.02$. A pseudo-photostationary state with 80% $\text{Pb}(\text{OH})_6^{2-}$ is established. In the absence of oxygen, $\text{Pb}(\text{OH})_3^-$ undergoes a photoreduction with the deposition of elemental lead.

Introduction

Intramolecular photoredox reactions of metal complexes can lead to oxidations of a variety of ligands.^{1–4} Generally, these ligand oxidations are induced by ligand-to-metal charge-transfer (LMCT) excited states. It is remarkable that, despite the importance of photochemical water oxidation, very little is known about such photoredox processes involving water or hydroxide as ligands.^{1–4} This lack of knowledge is certainly related to the high energy which is required to populate the corresponding LMCT states.^{5,6} They may not be accessible at all, or their population interferes with the presence of other excited states. The oxidation of water or hydroxide to stable products such as O_2 or H_2O_2 requires multielectron-transfer processes. Since most metal complexes are not able to participate in photochemical multi-electron-transfer reactions, there is much current interest to identify such systems.^{7,8}

According to these considerations, homoleptic hydroxo complexes of main group metals with an s^0 electron configuration should be good candidates to observe a photochemical oxidation of two hydroxide ligands at one metal since s^0 metals are generally reduced to s^2 complexes as stable products.^{9,10} We explored this possibility and selected the complex $\text{Pb}(\text{OH})_6^{2-}$ for the present study. This choice was based on our experience with halide complexes of s^0 metals which photochemically release halogens upon LMCT excitation.^{9,10}

A further attractive feature of main group metal compounds is the facile interconversion of s^0 and s^2 complexes. As it will be seen, such an interconversion can be achieved photochemically. Applications in photocatalysis are certainly feasible. However, the results of this study are not only of importance with regard to potential applications but are also interesting in their own right since observations on the photochemistry and photophysics of main group metal hydroxo complexes are here reported for the first time.

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Experimental Section

Materials. $\text{Na}_2[\text{Pb}(\text{OH})_6]$ was prepared according to a published procedure.¹¹ Solutions of $\text{Na}[\text{Pb}(\text{OH})_3]$ were obtained^{12,13} by dissolving PbO in aqueous NaOH. The water used in the photochemical experiments was triply distilled.

Photolyses. The photolyses were carried out at room temperature in 1-cm spectrophotometer cells. The light source was a Hanovia Xe/Hg 977 B-1 (1000 W) lamp. Monochromatic light ($\lambda_{\text{irr}} = 254$ nm) was obtained by means of a Schoeffel GM 250-1 high-intensity monochromator. For quantum yield determinations, the complex concentrations were such as to have essentially complete light absorption. The total amount of photolysis was limited to less than 5% to avoid light absorption by the photoproduct. Absorbed light intensities were determined by a Polytec pyroelectric radiometer, which was calibrated and equipped with an RkP-345 detector.

Spectroscopy. Absorption spectra were recorded with a Uvikon 860 double-beam spectrophotometer. Emission spectra were obtained on a Hitachi 850 spectrofluorimeter which was equipped with a Hamamatsu 928 photomultiplier. The luminescence spectra were corrected for monochromator and photomultiplier efficiency. The relative emission intensities were recorded in power units. Absolute emission quantum yields were determined by comparison of integrated emission intensity of $\text{Pb}(\text{OH})_3^-$ with that of quinine sulfate in 0.5 M H_2SO_4 ($\lambda_{\text{max}} = 452$ nm; $\phi = 0.546$)¹⁴ under identical conditions such as exciting wavelength, optical density, and apparatus parameters.

Flash Photolysis. A flash photolysis apparatus of the conventional type was used. The main flash lamp dissipated 300 J at 17.2 kV with a flash half-time of 10 μs . Transient absorption changes were recorded on a OS 250 oscilloscope (Gould Advanced). The argon-saturated solution was taken in a quartz cell with 10-cm path length.

Analyses. Hydrogen peroxide was identified using the peroxide test of Merck (Merckoquant 10011). For the quantitative analysis of H_2O_2 , the enzyme-catalyzed oxidation of scopoletin (7-hydroxy-6-methoxy-2H-1-benzopyran-2-one) and the subsequent decrease of fluorescence intensity was used.¹⁵ The analysis was adjusted to our experimental conditions. After irradiation the solutions were neutralized with 1 M HCl, and the precipitated PbO_2 was removed by centrifugation.

Results

Salts of $\text{Pb}(\text{OH})_6^{2-}$ ¹¹ are stable in aqueous solution only in the presence of an excess of alkali metal hydroxide. The absorption spectrum of $\text{Pb}(\text{OH})_6^{2-}$ (Figure 1) consists of a UV band. Its

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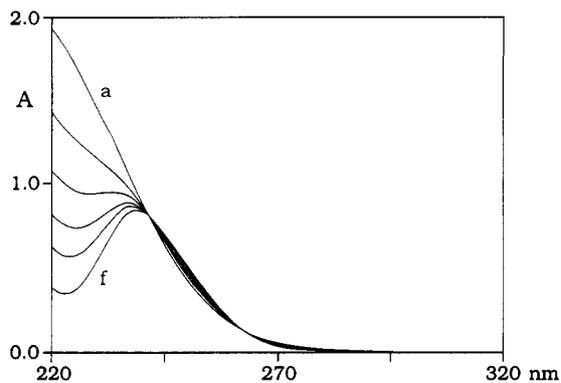
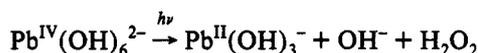


Figure 1. Spectral changes during the photolysis of 1.8×10^{-4} M $\text{Na}_2[\text{Pb}(\text{OH})_6]$ in 0.1 M NaOH at 0 (a), 2, 4, 6, 8, and 12 min (f) of irradiation time ($\lambda_{\text{irr}} = 254$ nm; 1-cm cell).

maximum is assumed to appear below 220 nm. $\text{Pb}(\text{OH})_6^{2-}$ did not show any emission but was light sensitive. The photolysis ($\lambda_{\text{irr}} = 254$ nm) of argon-saturated solutions was assumed to lead to a reductive elimination according to the following equation:



The spectral changes which accompanied the photolysis (Figure 1) include isosbestic points at 242 and 263 nm. This clean photoreaction can be driven to completion. The final spectrum is that of $\text{Pb}(\text{OH})_3^-$ (see below). Hydrogen peroxide was produced in less than stoichiometric amounts, depending on experimental conditions such as concentration, light intensity, and irradiation time. For example, irradiation of 7.2×10^{-4} M $\text{Na}_2[\text{Pb}(\text{OH})_6]$ in 1 M aqueous NaOH yielded 6.3×10^{-5} M $\text{Pb}(\text{OH})_3^-$ after 60-s irradiation at 254 nm and a light intensity of 12.2 mW·cm $^{-2}$. On the basis of the assumed stoichiometry, only 40% of the expected amount of H_2O_2 was recovered. When this irradiation was performed for 120 s, the concentration of $\text{Pb}(\text{OH})_3^-$ was 1.2×10^{-4} M while the relative yield of H_2O_2 dropped to 24%. It is assumed that H_2O_2 underwent a disproportionation to H_2O and O_2 by a secondary photolysis or by other side reactions. The quantum yield for the formation of $\text{Pb}(\text{OH})_3^-$ was $\phi = 0.07 \pm 0.01$ at $\lambda_{\text{irr}} = 254$ nm. In the presence of oxygen, the photolysis of $\text{Pb}(\text{OH})_6^{2-}$ was much slower and a complete conversion to $\text{Pb}(\text{II})$ was not achieved since $\text{Pb}(\text{OH})_3^-$ underwent a photooxidation back to $\text{Pb}(\text{OH})_6^{2-}$.

A low-temperature (77 K) photolysis of $\text{Pb}(\text{OH})_6^{2-}$ in 10 M NaOH glasses was carried out in order to identify radicals by ESR spectroscopy. While $\text{Pb}(\text{III})^{16}$ was not detected, a weak signal with $g = 2.0049$ is compatible with the presence of OH radicals.¹⁷

The flash photolysis of 2.0×10^{-3} M $\text{Pb}(\text{OH})_6^{2-}$ in 1 M NaOH under argon generated an intermediate which shows an absorption at 360 nm. It decayed according to a first-order rate law with a lifetime of $\tau = 5$ μs . Halide or pseudohalide may be used to scavenge OH radicals generated in the flash photolysis. However, this method cannot be used here since these scavengers are known to produce solvated electrons at the irradiating wavelength required for the photolysis of $\text{Pb}(\text{OH})_6^{2-}$.¹⁸ Moreover, halide ions would be incorporated as ligands since s^0 complexes such as $\text{Pb}(\text{OH})_6^{2-}$ are rather labile.^{10,19}

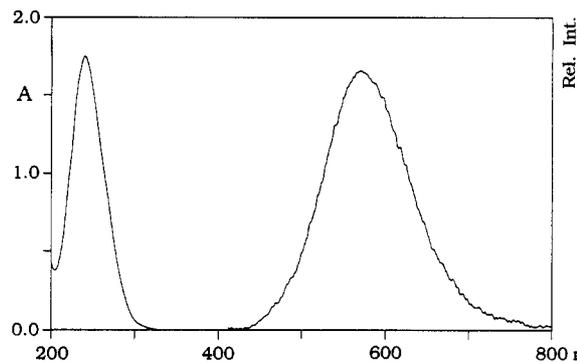
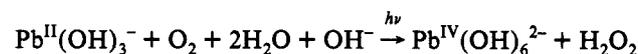


Figure 2. Electronic absorption (left side) and emission (right side) spectra of 3.9×10^{-4} M $\text{Na}[\text{Pb}(\text{OH})_3]$ in 0.1 M NaOH at room temperature (1-cm cell; $\lambda_{\text{exc}} = 254$ nm; intensity in arbitrary units).

Solutions of $\text{Pb}(\text{OH})_3^-$ ^{12,13} were obtained by dissolving PbO in aqueous alkali metal hydroxide. The absorption spectrum of $\text{Pb}(\text{OH})_3^-$ displays a single band (Figure 2) at $\lambda_{\text{max}} = 239$ nm ($\epsilon = 4500$ L·mol $^{-1}$ ·cm $^{-1}$). Argon-saturated solutions of $\text{Pb}(\text{OH})_3^-$ in 0.1 M NaOH show a yellow emission at $\lambda_{\text{max}} = 570$ nm (Figure 2) with $\phi = 1.5 \times 10^{-3}$ at $\lambda_{\text{exc}} = 254$ nm. The excitation spectrum agreed well with the absorption spectrum. In oxygen-saturated solutions, the emission quantum yield dropped to $\phi = 1.2 \times 10^{-3}$.

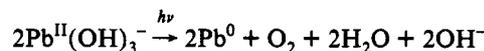
In oxygen-saturated solutions ($\sim 10^{-3}$ M O_2),²⁰ the irradiation of $\text{Pb}(\text{OH})_3^-$ in 0.1 M NaOH with $\lambda_{\text{irr}} = 254$ nm led to the formation of $\text{Pb}(\text{OH})_6^{2-}$. The accompanying spectral changes were simply a reversal of those observed during the photolysis of $\text{Pb}(\text{OH})_6^{2-}$ (Figure 1). Although isosbestic points at 242 and 263 nm indicate a clean photooxidation, a complete conversion could not be achieved since the photoproduct $\text{Pb}(\text{OH})_6^{2-}$ is photoreduced back to $\text{Pb}(\text{OH})_3^-$. A pseudo-photostationary state²¹ with 80% $\text{Pb}(\text{OH})_6^{2-}$ was reached by bubbling a continuous stream of oxygen through the solution during the irradiation. At the beginning of the photolysis of $\text{Pb}(\text{OH})_3^-$, the formation of $\text{Pb}(\text{OH})_6^{2-}$ proceeded with $\phi = 0.020 \pm 0.002$ at $\lambda_{\text{irr}} = 254$ nm.

In analogy to related photooxidations,^{9,10} it was assumed that the stoichiometry corresponds to the equation



However, only traces of H_2O_2 were detected.

In the absence of oxygen, $\text{Pb}(\text{OH})_3^-$ in 1 M NaOH underwent a photoreduction to metallic lead. The irradiation of $\text{Pb}(\text{OH})_3^-$ in argon-saturated solutions was accompanied by the formation of a black colloid which led to an apparent increase of the optical density over the entire UV/visible wavelength region. This increase was larger toward shorter wavelength, in accordance with the optical properties of metal colloids.²² The disappearance of $\text{Pb}(\text{OH})_3^-$ was measured by the decrease of the extinction at 239 nm in the beginning of the photolysis before the metal deposition obscured the spectral measurements. At $\lambda_{\text{irr}} = 254$ nm, $\text{Pb}(\text{OH})_3^-$ disappeared with $\phi = 0.013 \pm 0.003$. Since H_2O_2 as a possible oxidation product was not detected, it is assumed that the photolysis occurs with the overall stoichiometry



However, this stoichiometry does not exclude the primary formation of H_2O_2 , which may undergo a disproportionation to

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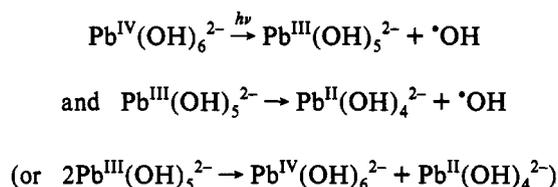
water and oxygen. Metals are well-known to catalyze the decomposition of hydrogen peroxide.²³

Discussion

The complex $\text{Pb}(\text{OH})_6^{2-}$ contains Pb(IV) with an empty valence shell (s^0 electron configuration). Accordingly, only electronic transitions of the LMCT type are possible. In analogy to related complexes such as PbCl_6^{2-} , SnCl_6^{2-} , or SbCl_6^{3-} ,^{9,10} the longest wavelength absorption of $\text{Pb}(\text{OH})_6^{2-}$ is assigned to the $t_{1u} \rightarrow a_{1g}$ transition from filled orbitals of OH^- to the empty antibonding s orbital of Pb(IV). Although Pb(IV) is a strong oxidant, this LMCT transition occurs at very high energies since OH^- is only a very weak reductant.^{5,6}

LMCT excitation of $\text{Pb}(\text{OH})_6^{2-}$ leads to a reductive elimination with the formation of $\text{Pb}(\text{OH})_3^-$ and H_2O_2 , which was found in less than stoichiometric amounts. Since H_2O_2 absorbs in the same wavelength region as $\text{Pb}(\text{OH})_6^{2-}$, it can disappear in a secondary photolysis. The decomposition of hydrogen peroxide is a well-known photoreaction.²⁴ It is also feasible that other side reactions of the photolysis of $\text{Pb}(\text{OH})_6^{2-}$ prevent the stoichiometric production of H_2O_2 (see below).

The photoreaction of $\text{Pb}(\text{OH})_6^{2-}$ may proceed by the release of H_2O_2 in a concerted reaction without the intermediate formation of Pb(III) and/or OH radicals. As an alternative and in analogy to PbCl_4 ,²⁵ the photoreduction of $\text{Pb}(\text{OH})_6^{2-}$ could take place in two consecutive one-electron-transfer steps:



Since $\text{Pb}(\text{OH})_4^{2-}$ is apparently not stable,^{13,26} $\text{Pb}(\text{OH})_3^-$ is formed as final product. The hydroxyl radicals can dimerize to yield H_2O_2 . However, owing to the reactivity of OH radicals, they can also attack other species. For example, hydroxyl radicals are known to react with H_2O_2 in a chain reaction which leads to the disproportionation of H_2O_2 .²⁴

The intermediate formation of OH radicals and Pb^{III} was explored by low-temperature and flash photolysis. Upon irradiation of $\text{Pb}(\text{OH})_6^{2-}$ in glasses at 77 K, an ESR signal developed which is compatible with the presence of OH radicals.¹⁷ This signal was rather weak. However, the room-temperature flash photolysis yielded a distinct transient with an absorption at 360 nm which is assumed to be $\text{Pb}^{\text{III}}(\text{OH})_5^{2-}$ or $\text{Pb}^{\text{III}}(\text{OH})_4^{2-}$.²⁸ It decays with $k = 2 \times 10^5 \text{ s}^{-1}$ according to first-order kinetics. It is suggested that this decay involves the release of a second OH

radical. Unfortunately, for the photolysis of $\text{Pb}(\text{OH})_6^{2-}$, OH radicals cannot be detected by scavenging techniques. Light absorption and photochemistry of suitable traps such as aromatic or olefinic compounds would interfere with the photochemistry of $\text{Pb}(\text{OH})_6^{2-}$.

Although $\text{Pb}(\text{OH})_3^-$ is a well-known species,^{12,13,29} its structure has not yet been determined. However, there is little doubt that it has a trigonal pyramidal geometry (C_{3v}). This structure is generally found for simple three-coordinate s^2 compounds.¹⁹ In analogy to other s^2 complexes,^{9,10,19} including PbCl_3^- ,²⁷ the longest-wavelength absorption of $\text{Pb}(\text{OH})_3^-$ at $\lambda_{\text{max}} = 239 \text{ nm}$ is certainly the A band, which is assigned to the metal-centered sp transition $^1S_0 \rightarrow ^3P_1$. The higher-energy sp transitions (B and C bands) do not occur above 220 nm.

Although many halide complexes of s^2 ions have been shown to emit,^{9,10,19} the photoluminescence of $\text{Pb}(\text{OH})_3^-$ is a remarkable observation. It represents the first example of an emitting hydroxo complex of s^2 metals. The emission of $\text{Pb}(\text{OH})_3^-$ at $\lambda_{\text{max}} = 570 \text{ nm}$ is assigned to the $^3P_1 \rightarrow ^1S_0$ transition. The large Stokes shift $\Delta E = 24\,300 \text{ cm}^{-1}$ indicates a considerable structural rearrangement in the excited state. Other s^2 complexes of the ML_3 type are also characterized by such large Stokes shifts.^{19,27} It is assumed that the trigonal pyramidal structure of $\text{Pb}(\text{OH})_3^-$ rearranges toward a trigonal planar geometry in the sp excited state.

The luminescence quenching by oxygen is associated with the photooxidation of $\text{Pb}(\text{OH})_3^-$ to $\text{Pb}(\text{OH})_6^{2-}$. Similar photooxidations of other s^2 ions such as Tl^+ and Sb^{3+} have been shown to be accompanied by the production of H_2O_2 .^{9,10} However, in the case of $\text{Pb}(\text{OH})_3^-$, only traces of H_2O_2 were detected. Since $\text{Pb}(\text{OH})_3^-$ and H_2O_2 absorb at similar energies, a secondary photolysis of hydrogen peroxide can take place. The photodecomposition of hydrogen peroxide is facilitated by a chain reaction²⁴ (see above).

During the photolysis of $\text{Pb}(\text{OH})_3^-$, the product $\text{Pb}(\text{OH})_6^{2-}$ is accumulating. It starts to absorb light and is photoreduced back to $\text{Pb}(\text{OH})_3^-$. If an excess of oxygen is admitted, a pseudo-photostationary state with about 20% $\text{Pb}(\text{OH})_3^-$ is established. This equilibrium can also be calculated using a simple equation which requires the knowledge of the extinction coefficients at the irradiation wavelength and the quantum yields of both photoreactions.³⁰ The calculated value agrees rather well with the experimental result.

In the absence of oxygen, the photolysis of $\text{Pb}(\text{OH})_3^-$ leads to the formation of elemental lead. Since H_2O_2 was not detected, molecular oxygen can be the only oxidation product of this photolysis. Although the investigation of this photoreaction was not within the scope of the present work, a tentative explanation for these observations is given. The sp excited state of $\text{Pb}(\text{OH})_3^-$ is certainly not only strongly reducing but also oxidizing. The photolysis thus can be induced by an initial intra- and intermolecular electron transfer from hydroxide to Pb^{2+} . Secondary reactions such as a disproportionation of Pb^+ to Pb^0 and Pb^{2+} may finally lead to the deposition of lead.

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(26) PbCl_3^- can add a further Cl^- to yield PbCl_4^{2-} .²⁷ Since the spectrum of $\text{Pb}(\text{OH})_3^-$ does not change upon increasing OH^- concentration, it is concluded that $\text{Pb}(\text{OH})_4^{2-}$ does not exist under these conditions.

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