Electronic State of Iron in Hemoglobin, Myoglobin, and Derivatives, as Inferred from X-Ray Fluorescence Spectra

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The Fe Kβ fluorescence emission spectrum was used to study the coordination of iron in some heme proteins. The spectrum was found to be dependent only on the direct environment of the iron. The iron atom in oxyhemoglobin can be regarded as trivalent, with a considerable negative charge on the outer oxygen atom of the O₂ ligand. Carbonmonoxide hemoglobin contains divalent, zero spin iron. Methemoglobin is a low spin compound.

INTRODUCTION

The way in which the iron atom is liganded in the heme proteins and related compounds has been the subject of considerable effort, both experimental and theoretical. The main experimental techniques thus far have been measurement of the magnetic moment, visible and ultraviolet absorption spectrometry, and, in recent years, Mössbauer spectrometry and electron paramagnetic resonance. Each of these techniques sheds a different light on the complex unity which a heme protein is, and gradually the role of the iron atom in the substances has become more clear. However, there still exist inconsistencies. The present study was undertaken to provide another tool for probing the iron atom.

USE OF THE FE Kβ X-RAY SPECTRUM

The Fe Kβ spectrum consists of two main lines, Kβ1,3 which corresponds to a transition between the M_{II,III} level and the K level of the iron atom, and Kβ2, much weaker, a transition between the M_{IV,V} level and the K level. Furthermore, a number of satellites may appear. Information may be drawn from shifts of the main lines (Kβ shifting is more or less proportional to the charge on the iron atom) and the appearance of the other lines. Since the intensities obtainable for the proteins are low, especially of Kβ3, shifts are not likely to be measurable with any precision. The only feature of the spectrum which can be measured rather reliably is the Kβ′ peak. Fortunately, this peak is highly relevant for the present purpose.

Although the existence of Kβ′ (to the low energy side of Kβ,2) has been known since the twenties, its theoretical explanation is still not settled satisfactorily. A review of the theories brought forward has been given by Ekstig et al.; these authors attribute the formation of Kβ′ to the interaction between a hole in the 3p shell and the incomplete 3d shell, causing splitting of the M_{II,III} level. The agreement between calculated and observed spectra is still rather poor, however. Koster and Mendel favor the view that part of the Kβ,3 energy is consumed in promoting an unpaired 3d electron into an excited state (into the conduction band); consequently a low energy satellite of Kβ,3 should appear. This theory requires that the relative intensity of Kβ′ for a given element should be proportional to the number of unpaired 3d electrons present in the compound. Experiments on a large number of compounds of the elements titanium to nickel indicate this to be the case within good approximation. Figure 1 may serve as an illustration. Iron(III) oxide, which contains the 3d⁶ configuration with five parallel spins has a well-pronounced Kβ′ satellite. In K₂Fe(CN)₆, where the six valence electrons of Fe⁶⁺ are engaged in dsp³ hybridization, no Kβ′ peak can be seen; identical Fe Kβ,3 profiles occur in for instance Fe₂ and Fe(CO)₅, where the same situation prevails. Such a profile can be used...
as a reference profile. The Fe Kβ₁,3γ₀ profile of K₃Fe(CN)₆ does show a small Kβ₀ peak, in accordance with the notion that there is one unpaired 3d electron. The relative intensities of Kβ₀ in Fe₂O₃ and K₃Fe(CN)₆ have the ratio 5:1 in good approximation. So we may hope to apply this peak for the heme proteins as well. Hemoglobin and myoglobin were selected, while hemin [iron(III) protoporphyrin chloride] was also studied.

**EXPERIMENTAL**

Two limiting factors are encountered when recording the x-ray spectra. First, heme proteins emit only a weak Fe Kβ fluorescence radiation, since they contain only a small amount of iron. This difficulty can normally be overcome by choosing long counting times, but proteins are rather sensitive to the white x-ray irradiation which is used in x-ray fluorescence equipment, so large numbers of counts cannot be accumulated. Secondly, solid samples must be studied, since aqueous solutions decompose very rapidly when subjected to x-ray excitation.

Commercially available hemoglobins (bovine and horse) were used. They were dissolved in phosphate buffer pH=7.0 to make a 5% solution, which was centrifuged.

To prepare desoxyhemoglobin (Hb), the solution was treated with a 20-fold excess of Na₂S₂O₄ during 20 min and passed through a Sephadex G25 column. The solution was evaporated in vacuum (1 cm Hg) over P₂O₅. The solid samples obtained were completely soluble in buffer pH=7.0, and the optical absorption spectra of their solutions were checked. No deterioration of the proteins could be detected during this drying process, except for HbO₂, which invariably contained 10%-25% of methemoglobin. Only those samples which contained at most 10% metHb were investigated with the x-ray spectrometer.

Freeze drying was also employed to concentrate the protein solutions. The x-ray spectra were similar, but less intense and hence less accurate, because the density of the freeze-dried powders is much less than the density of powders produced as described above.

A commercially available myoglobin preparation (horse) was used in the same way to prepare myoglobin (metMb).

The spectra were recorded on a 2 kW Philips semiautomatic spectrometer PW 1220, using a Mo primary target at 40 kV/40 mA and a LiF analyzing crystal in the second order. A measurement was performed at steps of 0.1° in 2θ, between 121.6°-122.6° 2θ. Care was taken that each sample was subjected to x-ray excitation for at most 20 min, and its optical spectrum was checked after the irradiation. At least three
samples were prepared of each substance, and at least three runs were made of each batch.

RESULTS AND CONCLUSIONS

Representative spectra are shown in Fig. 2. It is apparent that the method essentially probes the iron atom and its immediate neighbors; a sample of a bovine hemoglobin derivative and one of equine origin give the same spectrum, likewise the metMb spectrum is identical with the metHb spectrum. In fact, this rule is found to be obeyed in almost all investigations of x-ray spectra, those bonds only exerting influence which have a considerable energy. The environment of the heme iron can for the present purpose be described as consisting of the four pyrrole groups of the nearly plane heme group, an imidazole nitrogen atom of a proximal histidine somewhere in the globine chain and (except probably in Hb) a sixth ligand on the other side of the heme group.

The Fe Kα,β spectrum of hemin closely resembles that of FeO₃; hemin contains an Fe⁷⁺ ion with five parallel spins. This is to be expected, the Fe⁷⁺ ion not being able to hybridize into a covalent planar square coordination. In fact the Fe lies considerably out of the plane of the four nitrogen atoms (Koenig). The spectrum of Hb shows that the Fe²⁺ ion contains four parallel spins; this was already found by Pauling and Coryell in their classical work on the magnetic moments of hemoglobin derivatives. These magnetic moments were determined in solution; we find the same number of parallel spins in a solid sample, which shows that our way of drying (rather rapidly, and not prolonged after the sample became solid) yields a product whose molecules are comparable to the dissolved molecules as regards the heme environment. If Hb is dried for many hours, a dehydrated molecule is formed, as is witnessed by spectral data of Trautwein, Eicher and Mayer.

The magnetic moment of HbO₂ is zero, and it is assumed that by combining Hb with paramagnetic...
oxygen all spins become paired. The x-ray spectrum shows, however, that the iron atom still possesses one unpaired electron. This contradiction can be explained by recalling that the x-ray fluorescence analysis probes only the iron atom and its immediate vicinity, whereas the magnetic susceptibility is determined by the magnetic moment of the molecule as a whole. The unpaired electron in HbO₂ can only be explained by assuming that the iron is octahedrally coordinated by means of $tsp^3$ hybridization, as in the complex cyanides, and that the iron is actually trivalent. Weiss pointed out already that this might be the case, and since then experimental evidence in favour of this concept has been given by Maling and Weissbluth, using Mössbauer spectroscopy, although Eicher and Trautwein with the same technique came to another viewpoint, and by Wittenberg et al., who combined optical absorption and EPR data. A formal charge of 3+ on the iron means that the O₂⁻ ion is effectively the O₂− ion. The outer oxygen atom of this ion may be supposed to carry most of the negative charge, in accordance with a model of Pauling; in such a way electrostatic bonding occurs with the positively charged imidazole nitrogen atom of the distal histidine. The O₂− ion is known to possess one spin, which combines with the one spin of the iron atom to yield diamagnetic HbO₂. By assuming that the iron atom is essentially trivalent, it is readily understood why this substance is relatively stable to oxidation: it contains already Fe³⁺.

The spectrum of HbCO shows that no unpaired 3d electrons are present; it can be described as an iron(II) compound in which the two bonding electrons of the CO molecule are utilized to form an octahedral complex with $tsp^3$ hybridization. The bonds in HbO₂ and HbCO are thus dissimilar, which seems to follow also from Mössbauer spectra.

The spectra of metHb and metMb indicate that the iron atom is trivalent, with one unpaired electron and six ligands in $tsp^3$ hybridization. They are evidently “low spin” compounds. This is in contradiction to measurements of the magnetic moment, which is consistent with a “high spin” structure of both hydrated and hydroxide of metHb. EPR spectroscopy, however, has shown the hydroxide to be “low spin”; probably a difference in the degree of hydration has brought about these inconsistencies.

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