

## Calculation of O<sub>2</sub> Concentration

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### *Introduction*

The O<sub>2</sub> concentration of the blood includes the greater proportion of oxygen chemically bound to hemoglobin, as well as a smaller amount of physically dissolved oxygen. Whereas the chemically bound O<sub>2</sub> is only found in the erythrocytes, the physically dissolved O<sub>2</sub> is distributed between erythrocytes and plasma.

In the event that the O<sub>2</sub> concentration (O<sub>2</sub> content) of arterial blood must be known but the appropriate measurement technique is not available, an attempt can be made to estimate the O<sub>2</sub> concentration from other available data. For calculating the amount of chemically bound O<sub>2</sub> the Hb concentration (cHb, g/dl) and the O<sub>2</sub> saturation (sO<sub>2</sub>, %) are required; for the physically dissolved O<sub>2</sub> concentration, the O<sub>2</sub> partial pressure (pO<sub>2</sub>, mmHg) and O<sub>2</sub> solubility ( $\alpha$ O<sub>2</sub>, ml/ml/atm) are necessary. If these data are not all available, the O<sub>2</sub> concentration can only be determined approximately. This is illustrated in the two examples given below.

Modern multi-wavelength oxymeters (CO- or Hem-oxymeters) with 4–7 wavelengths are able to calculate the amount of chemically bound O<sub>2</sub> from the cHb and sO<sub>2</sub>; this value, however, neglects the amount of physically dissolved oxygen. This would lead in the case of hyperoxia, for example, to a calculated O<sub>2</sub> concentration that would lie significantly below the true cO<sub>2</sub>.

Blood gas analyzers, some of which are able to measure the Hb concentration, calculate the O<sub>2</sub> saturation from data on the acid-base

status and the measured  $pO_2$ , and thus, on the basis of the cHb, the amount of chemically bound  $O_2$  is derived. From a given value for the  $O_2$  solubility and the measured  $pO_2$ , the amount of physically dissolved  $O_2$  can be estimated, and thus the  $O_2$  concentration can be calculated.

However, this procedure can also only yield an approximate value for the  $O_2$  concentration since several simplifying assumptions must be made (normal  $O_2$  binding curve,  $O_2$  solubility) and only the so-called partial  $O_2$  saturation ( $psO_2$ , %) can be used. This last point ( $psO_2$ ) is considered elsewhere in this book (e.g. in the Appendix).

The data necessary for calculating the exact  $O_2$  concentration of blood will be described here, together with the calculation procedure.

### *Chemically Bound Oxygen*

Since 1 mol of Hb can bind a maximum of 4 mol  $O_2$ , a value is obtained for the so-called theoretical Hüfner number of 1.39 ml  $O_2$ /g Hb, assuming a molecular weight for the Hb molecule of 64,458 and a molar volume for oxygen of 22.394 l/mol. These data are always associated with the name Hüfner, since in 1894 he made the first experimental attempts to confirm them [3]. However, neither Hüfner nor several other authors have succeeded in measuring these parameters. All values measured later have lain between Hüfner's value of 1.34 ml/g and the theoretical value of 1.39 ml/g.

The reasons for this are, on the one hand, that traces of COHb, MetHb and SulfHb cannot be completely eliminated experimentally and, on the other hand, that difficulties have been encountered in the necessary correction of the physically dissolved oxygen. It is therefore recommended that only the theoretical Hüfner number of 1.39 ml/g should be used and that possible traces of COHb, MetHb and SulfHb, which can lead to a decrease in  $O_2$  concentration, be taken into account in assessing the  $O_2$  saturation of the blood. This recommendation has already become normal practice in most modern oxymeters and blood gas analyzers.

The above considerations indicate how the amount of chemically bound oxygen should be calculated. When the Hb concentration (cHb) and the  $O_2$  saturation ( $sO_2$ ) are known, the  $O_2$  concentration is obtained as follows:

$$cO_2 \text{ (ml/dl)} = cHb \text{ (g/dl)} \times sO_2 \times 1.39 \text{ (ml/g)}$$

It should be noted that the true sO<sub>2</sub> (not the partial) must be used and that the saturation is expressed as a fraction rather than as a percentage (e.g. sO<sub>2</sub> = 50% is expressed as 0.5).

At an Hb concentration of 15 g/dl and an O<sub>2</sub> saturation of 97%, for example, a value for the concentration of chemically bound O<sub>2</sub> alone of 20.2 ml/dl is obtained.

### *Physically Dissolved Oxygen*

According to Henry's law, the concentration of physically dissolved O<sub>2</sub> is given by the product of the O<sub>2</sub> partial pressure and the O<sub>2</sub> solubility coefficient,  $\alpha_{O_2}$ . The latter is usually expressed as the so-called Bunsen solubility coefficient, i.e. ml O<sub>2</sub>/ml liquid/atmosphere partial pressure (ml/ml/atm), and is dependent upon temperature.

Although a large number of O<sub>2</sub> solubility coefficients are known for different liquids, also at 37°, determination of the solubility of O<sub>2</sub> in the blood presents a particular difficulty. The comparatively small concentration of physically dissolved O<sub>2</sub> must be determined in the presence of a very high concentration of chemically bound O<sub>2</sub>. With one exception [2], a procedure has always been used in which the hemoglobin is oxidized, with formation of MetHb, so that a reversible binding of O<sub>2</sub> is prevented.

A different procedure has been used with the aim of determining the solubility of O<sub>2</sub> in human blood at 37° under largely physiological conditions [6]. On the one hand, the O<sub>2</sub> concentration was measured in blood samples whose pO<sub>2</sub> values were between 170 and 700 mmHg, i.e., the hemoglobin was always saturated with O<sub>2</sub> and the increased O<sub>2</sub> concentration with increasing pO<sub>2</sub> therefore represented the O<sub>2</sub> solubility. On the other hand, the O<sub>2</sub> concentration was determined in blood samples that had been equilibrated with 30% CO in order to prevent O<sub>2</sub> binding to hemoglobin.

The results of these investigations are summarized in figure 1 and can be outlined as follows: (1) Starting from the O<sub>2</sub> solubility of plasma previously determined [5], the O<sub>2</sub> solubility of blood increased with increasing Hb concentration; (2) the results did not differ between the two methods (exclusion of the influence of Hb by either high pO<sub>2</sub> values or with CO); (3) erythrocyte membranes also had no effect on the O<sub>2</sub> solubility. It is apparent that the values reported here for the O<sub>2</sub> solubility lie clearly

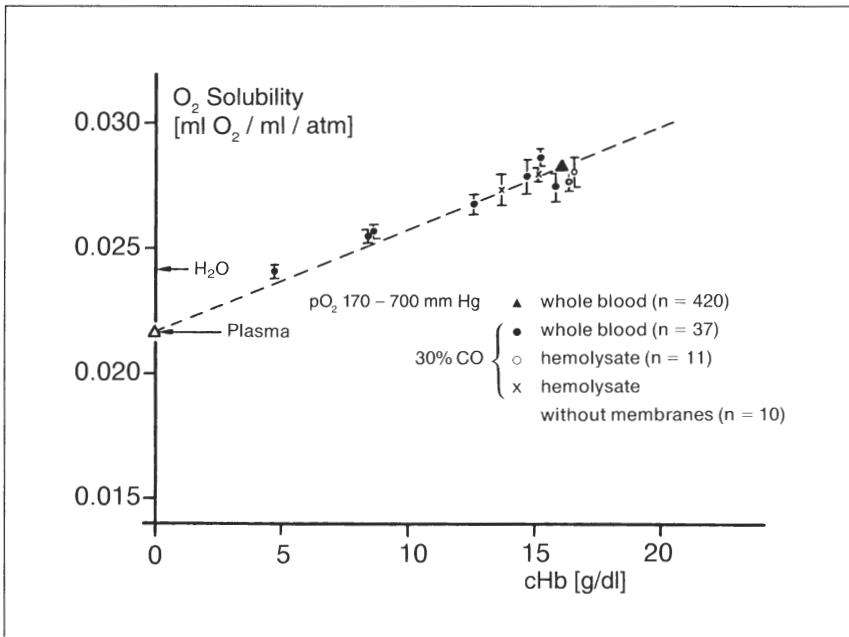


Fig. 1. Measured values for the  $O_2$  solubility of blood in ml/ml/atm at  $37^\circ\text{C}$  as a function of the Hb concentration (cHb) expressed in g/dl (data taken from [6]). Starting from the  $O_2$  solubility of plasma, the  $O_2$  solubility of blood increases linearly with increasing Hb concentration. Erythrocyte membranes have no effect on the  $O_2$  solubility. The experimental procedure for eliminating  $O_2$  binding to Hb ( $pO_2$  170–700 mmHg or inhibition with 30% CO) apparently does not influence the measured values.

above the most cited values from Van Slyke's group [4] (0.0237 ml/ml/atm for normal blood at  $37^\circ\text{C}$ ).

If the Hb concentration (g/dl) or hematocrit (vol.%) are known, the  $O_2$  solubility of the blood in ml/ml/atm can be obtained using the measured  $O_2$  solubility of plasma (0.0217 ml/ml/atm) [5] as follows:

$$\alpha_{O_2} = \text{cHb} \times 0.00041 + 0.0217 \text{ or}$$

$$\alpha_{O_2} = \text{Hct} \times 0.000137 + 0.0217.$$

An Hb concentration of 15 g/dl (Hct = 45%) would thus give an  $O_2$  solubility coefficient of 0.0279 ml/ml/atm.

For practical reasons the  $O_2$  solubility coefficient can be given as

ml/dl/mmHg (instead of ml/ml/atm), from which the following values are obtained:

0.0037 ml/dl/mmHg at cHb = 15 g/dl.

0.0034 ml/dl/mmHg at cHb = 10 g/dl.

0.0031 ml/dl/mmHg at cHb = 5 g/dl.

0.0029 ml/dl/mmHg for plasma.

Calculation of the concentration of physically dissolved O<sub>2</sub> can now be performed from the product of pO<sub>2</sub> and αO<sub>2</sub> if the O<sub>2</sub> partial pressure (pO<sub>2</sub>, mm Hg) is known:

$$cO_2 \text{ (ml/dl)} = pO_2 \text{ (mm Hg)} \times \alpha O_2 \text{ (ml/dl/mm Hg)}.$$

At a pO<sub>2</sub> of 90 mmHg and an αO<sub>2</sub> of 0.0037 ml/dl/mmHg (i.e. normal blood), a concentration of physically dissolved O<sub>2</sub> of 0.33 ml/dl would thus be obtained; at a pO<sub>2</sub> of 600 mmHg (hyperoxia) this value would be 2.2 ml/dl.

### *Calculation of O<sub>2</sub> Concentration*

Since the O<sub>2</sub> concentration (cO<sub>2</sub>; ml/dl) is obtained from the sum of the chemically bound and physically dissolved O<sub>2</sub> concentrations, the relationship is as follows:

$$cO_2 \text{ (ml/dl)} = (cHb \times sO_2 \times 1.39) + (pO_2 \times \alpha O_2).$$

where cHb is expressed in g/dl, sO<sub>2</sub> is expressed as a fraction, pO<sub>2</sub> is expressed in mm Hg and αO<sub>2</sub> in ml/dl/mmHg. In normal blood, for example, with cHb = 15 g/dl, sO<sub>2</sub> = 97% and pO<sub>2</sub> = 90 mmHg, this amounts to an O<sub>2</sub> concentration of 20.6 ml/dl.

For practical purposes it is recommended that the physically dissolved O<sub>2</sub> is taken to be constant at 0.3 ml/dl, as long as the pO<sub>2</sub> lies between 60 and 100 mmHg and the Hb concentration is between 10 and 20 g/dl. Under these conditions the physically dissolved O<sub>2</sub>, and therefore the O<sub>2</sub> concentration of the blood, varies maximally by ±0.1 ml/dl, i.e., always less than ±1% of the O<sub>2</sub> concentration in the blood.

### *Summary*

For calculating the O<sub>2</sub> concentration of blood, knowledge is required of the Hb concentration (cHb, g/dl), the O<sub>2</sub> saturation (sO<sub>2</sub>, %), the theoretical Hüfner number of 1.39 ml/g, the partial pressure (pO<sub>2</sub>, mmHg), and the O<sub>2</sub> solubility (αO<sub>2</sub> ml/dl/mmHg). New data on the O<sub>2</sub> solubility in human blood allow the precise calculation of the O<sub>2</sub> concentration especially in the case of hyperoxia.

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