

One way to think about the gas solubilities is to consider alveolar-capillary exchange in the lung. At 37°C, water solubility of CO₂ is 0.574742 ml/ml at 1 atm or 760 mmHg atmospheric pressure as calculated from the following equation:

$$\text{AlphaCO2w37} = 1.052 \cdot \exp(-0.0571 \cdot 37) + 0.6821 \cdot \exp(-0.012 \cdot 37) \quad \text{Eq 1.}$$

where AlphaCO2w37 is the water solubility of CO₂, 37 has units of °C. See equation 4b. of Christmas and Basingthwaighte 2017 for more details about the coefficient values. Figure 1 shows the time course of exchange of CO₂ between a 100 ml solution of CO₂ in water at a pCO₂ of 50 mmHg and CO₂ in 57.4742 ml of alveolar air at a pCO₂ of 10 mmHg. The exchange is driven by the gas tensions on the two sides, and progresses to an equilibrium at pCO₂ of 30 mmHg, at which time the air volume of 55.4742 ml air (dry) contains same amount of CO₂, namely 9.953.10⁻⁵ mol, as are in the 100 ml water.

The time taken for this exchange depends on the membrane permeability-surface area product, P_{s_{alv}}, and the volumes of the spaces. The driving forces for the bidirectional fluxes across the membrane are the partial pressures for CO₂:

$$d pCO2w/dt = - P_{Salv} \cdot (pCO2w - pCO2alv) / (Vw \cdot \text{AlphaCO2w37}); \quad \text{Eq 2.}$$

$$d pCO2alv/dt = P_{Salv} \cdot (pCO2w - pCO2alv) / Valv \quad \text{Eq 3.}$$

where the rates of concentration or pCO₂ changes in the water and alveolar spaces due to the net fluxes of the gas equals the conductance, P_{salv}, times the difference in the partial pressures, divided by the effective volume. To account for the solubility in water, its volume V_w is multiplied by the solubility for the gas. Thus, the solubility, AlphaCO2w37, is like a partition coefficient for V_w, and is less than 1.0 at 37°C.

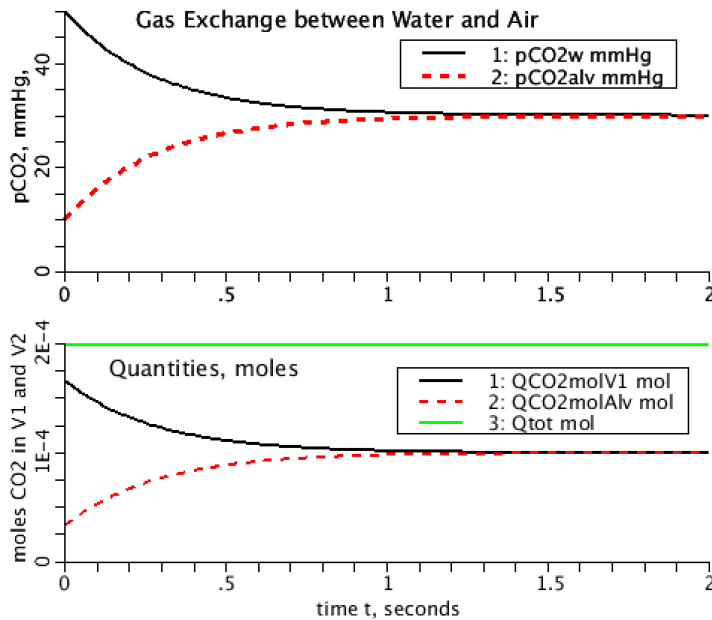


Figure 1. Alveolar-capillary exchange of carbon dioxide between gas phase and solution in water.. *Upper panel:* Partial pressures, mmHg, in gas phase (black) and in water (red dashes), equilibrate. *Lower panel:* Molar amounts in the two phases become identical when the air space volume is

reduced by multiplying by the water solubility for the gas. (Program for figure available at www.physiome.org/jsim/models/webmodel/NSR/AlvCapExch/ .)

REFERENCE:

1. Christmas KM and Bassingthwaighte JB. Equations for O₂ and CO₂ solubilities in saline and plasma:
Combining temperature and density dependencies. J Appl Physiol 122: 1313-1320, 2017.