

CORRESPONDENCE

Pulmonary diffusion impairment following heart transplantation: a prospective study

To the Editor:

Dr EGAN and colleagues [1] have compared changes in transfer factor of the lung for carbon monoxide (TL_{CO}) before and after heart transplantation (HT) and coronary artery bypass grafting (CABG). Both operations involved thoracotomy, so that CABG provided a control for the operative procedure. Thirty days after the operation, the authors observed significant falls in TL_{CO} of 1.0 $\text{mmol}\cdot\text{min}^{-1}\cdot\text{kPa}^{-1}$ (SI unit) after HT and 1.7 SI units after CABG. At this point, they might have deduced that changes in TL_{CO} after HT were due to the operative procedure. Instead, they expressed their results as carbon monoxide transfer coefficient (KCO) (TL_{CO})/alveolar volume (VA); this index showed a decline over the postoperative period for HT, but not for CABG, hence they concluded that "following HT there was an early fall in gas transfer which was independent of surgery and lung bypass, implicating early immunosuppression".

The authors' conclusion presupposes that division by VA standardizes TL_{CO} for variations associated with alveolar volume. The latter index, expressed as total lung capacity (TLC), decreased by 0.5 L following both operations. In fact, KCO is negatively correlated with VA [2], so that use of TL_{CO}/VA overcorrects TL_{CO} for the effect of lung volume. The true relationship of TL_{CO} to VA is not precisely known, but between a litre above functional residual capacity (FRC) and TLC it can be represented as approximately linear. The slope of the relationship is such that, for a decrease in VA of 0.5 L,

the TL_{CO} would have been expected to decrease by approximately 0.5 SI units [3, 4]. Hence, on the assumption that VA was equal to TLC, the postoperative reduction in TL_{CO} at constant lung volume for HT was 0.5 SI units, and for CABG 1.2 SI units. These figures do not support the authors' thesis. However, the corrections should be applied to the actual volumes during breathholding. It would be of interest to know what these were, and if applying a linear correction to them led to a similar conclusion.

References

1. Egan JJ, Lowe L, Yonan N, *et al.* Pulmonary diffusion impairment following heart transplantation: a prospective study. *Eur Respir J* 1996; 9: 663–668.
2. Stam H, Kreuzer FJA, Versprille A. Effect of lung volume and positional changes on pulmonary diffusing capacity and its components. *J Appl Physiol* 1991; 71: 1477–1488.
3. Cotes JE, Hall AM. The transfer factor for the lung: normal values in adults. *In: Normal Values For Respiratory Function in Man.* Arcangeli P, ed. Torino, Panminerva Medica, 1970: pp. 327–343.
4. Cotes JE. Lung Function: Assessment and Application in Medicine. 5th Edn. Oxford, Blackwell Scientific Publications, 1993; pp. 293–294.

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REPLY

From the author:

Cotes and Reed have highlighted the fact that the use of single-breath alveolar volume (VA) versus total lung capacity (TLC) in the estimation of the carbon monoxide transfer coefficient (KCO) is an area of controversy. As they suggest, one cannot absolutely assume that VA is equal to TLC. In our study investigating the influence of heart transplantation on lung physiology, the reported measurements of TLC were completed by plethysmography and not by helium dilution [1]. The potential advantage that estimates of VA have is that it can be considered in physiological terms as the carbon monoxide uptake in the ventilated portion of the lung at "effective" TLC [2]. Controversy does exist as to these interpretations, but it is generally recommended to use

single-breath VA for clinical and epidemiological purposes [2].

Notwithstanding, as suggested by LOVE and SEATON [3], the predicted KCO should be calculated from TLC rather than VA . It is noteworthy that our standardized residual (SR) values for KCO , dependent on the calculation of predicted KCO using TLC, demonstrated a fall in KCO following heart transplantation that was not observed in the patients following coronary artery bypass grafting.

However, the application of KCO standardized residual values ($=$ (observed - predicted)/residual standard deviation (RSD)) also has potential limitations, as RSD values for KCO have not been provided by the European Working Party [4]. Therefore, RSD values for KCO can only be estimated correctly from a sample of the individual population KCO values. We calculated SR values for KCO in this fashion. Accordingly, the fall in standardized residual value of KCO and the observation

by others of a fall in gas transfer following heart transplantation suggests that our conclusions are reasonable [5, 6].

References

1. Egan JJ, Lowe L, Yonan N, *et al.* Pulmonary diffusion impairment, following heart transplantation: a prospective study. *Eur Respir J* 1996; 9: 663–668.
2. Laszlo G. European standards for lung function testing: 1993 update. *Thorax* 1993; 48: 873–876.
3. Love RG, Seaton A. About the ECCS summary equations. *Eur Respir J* 1990; 3: 489.
4. Coates JE, Chinn DJ, Quanjer PH, Roca J, Yernault JC. Standardization of the measurement of transfer factor (diffusing capacity). *Eur Respir J* 1993; 6 (Suppl. 16): 41–52.
5. Ravenscraft SA, Gross CR, Kubo SH *et al.* Pulmonary function after successful heart transplantation: one year follow up. *Chest* 1993; 103: 54–58.
6. Groen HJ, Bogaard JM, Balk AH, Kho SG, Hop WC, Hilvering C. Diffusion capacity in heart transplant recipients. *Chest* 1992; 102: 456–460.

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