





DLCO: adjust for lung volume, standardised reporting and interpretation

To the Editor:

The American Thoracic Society (ATS) and European Respiratory Society (ERS) should be congratulated on updating standards for diffusing capacity of the lung for carbon monoxide (DLCO) [1]. I agree that "Besides varying with age, sex, height and possible ethnicity, DLCO also changes with Hb, lung volume, COHb, PIO $_2$..., exercise and body position." and that "adjustments for these factors be made in the predicted rather than the measured DLCO". Reporting transfer coefficient of the lung for carbon monoxide (KCO) rather than DLCO/alveolar volume (VA) will help get away from the mistaken notion that DLCO/VA "corrects" DLCO for lung volume [2]. While the new standards describe how to adjust predicted DLCO for haemoglobin (Hb), COHb and inspired oxygen tension (PIO $_2$), it does not discuss how to adjust predicted DLCO and KCO for lung volume.

The following equations [3] were included in the 2005 ATS/ERS DLCO standards [4], and describe how to adjust DLCO and KCO for lung volume. They were developed studying normal subjects with experimental reductions in inspired volume (VI; and thus VA) and fit the model that DLCO and KCO change in a manner expected from having DLCO reduced proportionate to the surface area for gas exchange with the capillary blood component unchanged. Mathematically, they result in DLCO % predicted for lung volume equaling KCO % predicted for lung volume when using the equation KCO(predicted)=DLCO(predicted)/VA(predicted).

 $DLCO[predicted for lung volume] = DLCO[predicted] \times (0.58 + 0.42 \times (VAm/VAp))$

 $KCO[\text{predicted for lung volume}] = KCO[\text{predicted}] \times (0.42 + 0.58/(V_{Am}/V_{Ap}))$

with VAm/VAp=measured VA/predicted VA.

For example, at VA 50% of predicted, the DLCO predicted for lung volume is 80% and KCO is 160% of that for VA 100% of predicted.

The standards require reporting DLCO and KCO (adjusted, predicted) with specification of the adjustments. Additional reporting requirements should include DLCO (% of adjusted predicted) and VA (% predicted).

Neither the 2005 nor the current standards address how to report DLCO and KCO adjusted for lung volume, or how to interpret DLCO.

In addition to knowing % predicted DLCO and KCO adjusted for all factors except lung volume, it is also very helpful to know % predicted DLCO and KCO when also adjusted for lung volume [2]. Just as adjusting predicted DLCO and KCO for haemoglobin in an anaemic patient yields a better indication of the lung's ability of gas exchange, adjusting DLCO and KCO for lung volume in a patient with low lung volume yields a better indication of the lung's ability of gas exchange.

A shorter nomenclature is needed for DLCO and KCO % predicted also adjusted for lung volume.

I propose DACO and KACO to refer to DLCO and KCO predicted values that have been adjusted for lung volume (the "A" refers to adjusted for lung volume.)

Reporting requirements should include DACO (adjusted, predicted), KACO (adjusted, predicted), as well as DACO (% of adjusted predicted) and KACO (% of adjusted predicted).

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DLCO reports and interpretation should be standardised and include adjusting predicted DLCO and KCO for lung volume http://ow.ly/ywTA30cOh44

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The new standards recommend development of a standardised common report form. I propose the following, one when Hb is not measured (box 1) and a second when Hb is measured (box 2), with both including % of FVC for VI if spirometry was done the same day

BOX 1 Common report form for diffusing capacity when haemoglobin measurements are not taken

Diffusing capacity	Predicted range		Actual	% pred	
	Mean	95%			
D∟co mL·min ⁻¹ ·mmHg ⁻¹	XX.XX	XX.XX	dd.dd	xx	Predicted not adjusted for Hb
VA (BTPS) L	X.XX	X.XX	X.XX	XX	
Kco mL·min ⁻¹ ·mmHg ⁻¹ ·L ⁻¹	XX.XX	XX.XX	X.XX	XX	Predicted not adjusted for Hb
VI (BTPS) L	X.XX	X.XX	XX.XX	XX	xx% of FVC
D _{ACO} mL·min ⁻¹ ·mmHg ⁻¹	XX.XX	XX.XX	dd.dd	уу	Predicted adjusted for lung volume

DLco and Kco are yy% predicted, adjusted for lung volume.

BOX 2 Common report form for diffusing capacity when haemoglobin measurements are taken

Diffusing capacity	Predicted range		Actual	% pred	Hb xx.x from D-MON-YYYY	
	Mean	95%				
<i>D</i> ∟co mL·min ⁻¹ ·mmHg ⁻¹	XX.XX	XX.XX	dd.dd	XX	Predicted not adjusted for Hb	
DLCO mL·min ⁻¹ ·mmHg ⁻¹	XX.XX	XX.XX	dd.dd	XX	Predicted adjusted for Hb	
VA (BTPS) L	X.XX	X.XX	X.XX	XX		
Kco mL·min ⁻¹ ·mmHg ⁻¹ ·L ⁻¹	XX.XX	XX.XX	X.XX	XX	Predicted adjusted for Hb	
VI (BTPS) L	X.XX	X.XX	X.XX	XX	xx% of FVC	
DACO mL·min ⁻¹ ·mmHg ⁻¹	XX.XX	XX.XX	dd.dd	уу	Predicted adjusted for lung volume and Hb	

 \emph{D} LCO and \emph{K} CO are yy% predicted, adjusted for lung volume and Hb.

For both reports, if DLCO and KCO predicted were also adjusted for COHb and/or PIO2, then a line saying "Predicted DLCO and KCO also adjusted for ..." should appear at the end, which includes the data used to make the adjustment, such as "COHb of 2.6% and altitude of 2000m." The 95% values are the lower limit of normal (LLN), with $DACO[LLN] = DACO[adjusted, predicted] \times DLCO[LLN] / DLCO[predicted].$

There is not a clear consensus on interpretation of DLCO. I recommend the following algorithm to interpret DLCO, with DLCO % predicted, adjusted and LLN the lower limit of normal (box 3).

BOX 3 Interpretation of diffusing capacity values					
DLco ≥80% and ≥LLN DLco <80% but ≥LLN DLco ≥60%, <80%, and <lln <40%<="" <60%,="" <lln="" and="" dlco="" th="" ≥40%,=""><th>DLCO is normal DLCO is near lower limit of normal DLCO is mildly reduced DLCO is moderately reduced DLCO is severely reduced</th></lln>	DLCO is normal DLCO is near lower limit of normal DLCO is mildly reduced DLCO is moderately reduced DLCO is severely reduced				

If DLCO is not normal, and DLCO adjusted for lung volume (DACO) is above the LLN as % predicted, then add phrase "due to low lung volume".

If DLCO is not normal, and DLCO adjusted for lung volume is below the LLN as % predicted but more than 10% predicted greater than DLCO, then add phrase "in part due to low lung volume".

As a co-author of the 2005 ERS/ATS DLCO standards, I believe including adjustments of DLCO for lung volume and standardised reports and interpretation would improve the clinical value of DLCO.

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References

- Graham BL, Brusasco V, Burgos F, et al. 2017 ERS/ATS Standards for single-breath carbon monoxide uptake in the lung. Eur Respir J 2017; 49: 1600016.
- Hughes M, Pride N. Examination of the carbon monoxide diffusing capacity (DLCO) in relation to its KCO and VA components. Am J Respir Crit Care Med 2012; 186: 132-139.
- Johnson DC. Importance of adjusting carbon monoxide diffusing capacity (DLCO) and carbon monoxide transfer coefficient (Kco) for alveolar volume. Respir Med 2000; 94: 28-37.
- MacIntyre N, Crapo R, Viegi G, et al. Standardisation of the single-breath determination of carbon monoxide uptake in the lung. Eur Respir J 2005; 26: 720-735.

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From the authors:

D.C. Johnson proposed that an adjustment of the predicted value for the diffusing capacity of the lung for carbon monoxide (DLCO) based on the measured and predicted lung volume be included in the report of DLCO test results. An interpretation algorithm is also proposed.

The rationale is based on the observation that a decreased inspired volume in normal subjects reduces DLCO and that this reduction can be predicted using a linear function of the measured alveolar lung volume (VA) divided by the predicted VA. While this formula may be applicable for normal, healthy lungs and may potentially adjust for effects of reductions in VA due to weakness, chest wall deformities or reduced effort, it is unlikely to be applicable in patients with reduced VA due to pathological processes that also impact the distribution of ventilation, the distribution of perfusion and the development of heterogeneous parenchymal changes in the lung. As Hughes and Pride [1] noted, "any 'correction' of the DLCO for volume must take into account the reason for the volume deficit." Volume adjustment equations for DLCO developed from submaximal inhalation manoeuvres in normal subjects do not consider other mechanisms for a reduced VA.

The 2017 European Respiratory Society (ERS)/American Thoracic Society (ATS) standards for single-breath carbon monoxide uptake in the lung [2] describe technical standards for the measurement of DLCO. The interpretation of DLCO was not included in the mandate of the ERS/ATS task force in the development of these standards. Nevertheless, the reporting of the results must provide all of the necessary information for their interpretation.

The standards address the need for the subject to achieve maximal inhalation of test gas. A failure to reach full inflation to total lung capacity is a reason to repeat the manoeuvre rather than applying a volume correction to the predicted DLCO. Reporting the VA and its z-score together with the DLCO allows the reviewer to interpret the DLCO in the context of whether the VA is reduced, normal or increased. It continues to be the opinion of the ERS/ATS DLCO task force that this information is sufficient for the interpretation of *D*LCO.

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A correction of the predicted value for DLCO based on the measured lung volume is not recommended http://ow.ly/KFn730cPJ9n

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References

- Hughes M, Pride N. Examination of the carbon monoxide diffusing capacity (DLCO) in relation to its KCO and VA components. Am J Respir Crit Care Med 2012; 186: 132–139.
- 2 Graham BL, Brusasco V, Burgos F, et al. 2017 ERS/ATS standards for single-breath carbon monoxide uptake in the lung. Eur Respir J 2017; 49: 1600016.

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