PULMONARY FUNCTION TESTING INDICATIONS AND INTERPRETATIONS

A Project of the California Thoracic Society

Edited by

Archie F. Wilson, M.D., Ph.D.



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Pulmonary Function Testing: Indications and Interpretations is meant to serve as a sequel to Pulmonary Function Testing: Guidelines and Controversies. Like its predecessor, this handbook is a product of the California Thoracic Society and was compiled prior to, during, and immediately after a postgraduate course that was given by leaders in this field. The chapters were written by individual members of the Pulmonary Physiology Committee after prolonged discussion with the faculty of the postgraduate course and other committee members.

It was our purpose to provide a handbook that would be useful to the pulmonary clinician who uses the laboratory to study disordered respiratory physiology for clinical purposes. We have included not only discussions of all tests likely to be obtained in both community and medical center pulmonary function laboratories but also applications of the testing to a number of nontraditional sites including the work place, intensive care units, and exercise and sleep laboratories. Evaluation of respiration testing for preoperative assessment, occupational disability, and pediatric patients is included. We have also discussed the use of the computer in pulmonary function interpretation.

We hope this handbook will provide a clear guide to diagnosis of pulmonary disorders by function testing. We also hope that we have adequately pointed the way to understanding both the limitations and some of the new areas in which pulmonary function testing is likely to expand.

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Terms and Abbreviations

The variety of abbreviations used in clinical pulmonary function reports (e.g., MEF 50%, FEF 50%, Vmax 50%, and V 50%) often leads to considerable confusion, especially for physicians without specific training in pulmonary medicine. Although not perfect, the terminology and abbreviations suggested by an American College of Chest Physicians/American Thoracic Society (ACCP/ATS) joint committee are the best available and should be used whenever possible. Those most relevant to subsequent chapters are given below. Abbreviations marked with an asterisk were not cited by the ACCP/ATS joint committee, but are used in this book.

	ne committee work sales have been a been as the
A	Alveolar
a	Arterial
an	Anatomic
ATPD	Ambient temperature and pressure, dry
ATPS	Ambient temperature and pressure, saturated with water vapor at these conditions
В	Barometric
BTPS	Body conditions: Body temperature, ambient pressure, and saturated with water vapor at these conditions
he FVC, D	
c of ravile	Capillary and at sec-as TEH bms OVH out
C/V _L	Specific compliance
CD* at angrape	
Cdynap and	flow at the mouth during active breathing. The respiratory frequency should be designated; e.g., C _{dyn} 40
Camulova	Static compliance, compliance determined from measurements

Dead space or wasted ventilation (qualifying symbol, e.g., V_D)

D/VA Diffusion per unit of alveolar volume

Diffusion coefficient or permeability constant as described by Dk

Krogh; it equals D · (P_B - P_{H₂O})/V_A

Diffusing capacity of the alveolar capillary membrane (STPD) D_m

D_x (or D_{LCO}) Diffusing capacity of the lung expressed as volume (STPD) of gas (x) uptake per unit alveolar-capillary pressure difference for the gas used. Unless otherwise stated, carbon monoxide is assumed to be the test gas, i.e., D is Dco. A modifier can be used to designate the technique, e.g., DSB is single breath carbon monoxide diffusing capacity and Dss is steady state CO diffusing capacity. (Editor's note: This recommendation has not widely been accepted. DLCO, DLCOSB, and DLCOSS are still the most commonly used abbreviations.)

Expired E

Expiratory reserve volume; the maximal volume of air exhaled ERV

from the end-expiratory level

est

FEF750%

FEF.

f Respiratory frequency per minute F Fractional concentration of a gas

FEFmax The maximal forced expiratory flow achieved during an FVC Mean forced expiratory flow during the middle half of the FVC FEF25-75% (formerly called the maximum mid-expiratory flow rate)

Instantaneous forced expiratory flow after 75% of the FVC has

been exhaled

FEF₂₀₀₋₁₂₀₀ Mean forced expiratory flow between 200 ml and 1200 ml of the FVC (formerly called the maximum expiratory flow rate)

Forced expiratory flow, related to some portion of the FVC curve.

Modifiers refer to the amount of the FVC already exhaled when

the measurement is made

The forced expiratory time for a specified portion of the FVC; FET_x e.g., FET_{95%} is the time required to deliver the first 95% of the FVC and FET 25-75% is the time required to deliver the FEF 25-75%

Forced expiratory volume (timed) to forced vital capacity ratio, FEV,/FVC% expressed as a percentage

FIF. Forced inspiratory flow. As in the case of the FEF, the appropriate modifiers must be used to designate the volume at which flow is being measured. Unless otherwise specified, the volume qualifiers indicate the volume inspired from RV at the point of the measurement

Functional residual capacity; the sum of RV and ERV (the volume FRC of air remaining in the lungs at the end-expiratory position). The method of measurement should be indicated as with RV

Airway conductance, the reciprocal of Raw Gaw

Gaw/V Specific conductance, expressed per liter of lung volume at which

G is measured (also referred to as SGaw)

Inspired

Inspiratory reserve volume; the maximal volume of air inhaled

from the end-inspiratory level

Inspiratory capacity; the sum of IRV and V_T IC

max Maximal

MIP* Maximal inspiratory pressure MEP* Maximal expiratory pressure

Maximal voluntary ventilation. The volume of air expired in a MVV. specified period during repetitive maximal respiratory effort. The respiratory frequency is indicated by a numerical qualifier; e.g., MVV₆₀ is MVV performed at 60 breaths per minute. If no

qualifier is given, an unrestricted frequency is assumed

p Physiological

P Pressure, blood or gas

PA* Pulmonary artery PD*

Provocative dose; the dose of an agent used in bronchial challenge testing which results in a defined change in a specific physiologic parameter. The parameter tested and the percent change in this parameter is expressed in cumulative dose units over the time following exposure that the positive response occurred. For example, PD₃₅SG_{aw} = x units/y minutes, where x is the cumulative inhalation dose and y the time at which a

35% fall in SGaw was noted

PEF The highest forced expiratory flow measured with a peak flow

meter

Static transpulmonary pressure at a specified lung volume; e.g., Pst P_{st}TLC is static recoil pressure measured at TLC (maximal

recoil pressure)

Capillary blood volume (usually expressed as V_r in the literature, Qc a symbol inconsistent with those recommended for blood volumes). When determined from the following equation, Qc represents the effective pulmonary capillary blood volume, i.e., capillary blood volume in intimate association with alveolar gas:

 $1/D = 1/D_m + 1/(\Theta \cdot Q_c)$

A general symbol for resistance, pressure per unit flow

Raw Airway resistance Rebreathing TO

RQ* Respiratory quotient

Rus	Resistance of the airways on the alveolar side (upstream) of the
	point in the airways where intraluminal pressure equals Ppl, measured under conditions of maximum expiratory flow
RV	Residual volume; that volume of air remaining in the lungs after
inhaled	maximal exhalation. The method of measurement should be
	indicated in the text or, when necessary, by appropriate qualify-
	ing symbols
SBN*	Single breath nitrogen test; a test in which plots of expired N ₂ concentration versus expired volume after inspiration of 100% O ₂ are recorded. The closing volume and slope of Phase III are two parameters measured by this test
STPD	Standard conditions: temperature 0° C, pressure 760 mm Hg, and dry (0 water vapor)
t	Time
T	Tidal
TGV*	Thoracic gas volume; the volume of gas within the thoracic cage as measured by body plethysmography
TLC	Total lung capacity; the sum of all volume compartments or the volume of air in the lungs after maximal inspiration. The method of measurement should be indicated, as with RV
V terostor	Gas volume. The particular gas as well as its pressure, water vapor conditions, and other special conditions must be specified in text or indicated by appropriate qualifying symbols
V	Venous
v	Mixed venous
VA	Alveolar ventilation per minute (BTPS)
Vco2	Carbon dioxide production per minute (STPD)
V_D	Ventilation per minute of the physiologic dead space (wasted
	ventilation), BTPS, defined by the following equation:
V _D	$\dot{V}_D = \dot{V}_E (PaCO_2 - P_ECO_2)/(PaCO_2 - P_1CO_2)$ The physiologic dead-space volume defined as \dot{V}_D/f
V _D an	Volume of the anatomic dead space (BTPS)
VE	Expired volume per minute (BTPS)
VI.	Inspired volume per minute (BTPS)
VisoV*	Volume of isoflow; the volume when the expiratory flow rates
ne, i.e niyeolar	become identical when flow-volume loops performed after breathing room air and helium-oxygen mixtures are compared
V _{O2}	Oxygen consumption per minute (STPD)
V _{max} X	Forced expiratory flow, related to the total lung capacity or the actual volume of the lung at which the measurement is made. Modifiers refer to the amount of lung volume remaining when

the measurement is made. For example: \dot{V}_{max} 75% is instantaneous forced expiratory flow when the lung is at 75% of its

TLC. \dot{V}_{max} 3.0 is instantaneous forced expiratory flow when the lung volume is 3.0 liters. [Editor's note: It is still common to find reports in which modifiers refer to the amount of VC remaining.]

VT XA or Xa Tidal volume; TV is also commonly used

A small capital letter or lowercase letter on the same line following a primary symbol is a qualifier to further define the primary symbol. When small capital letters are not available on typewriters or to printers, large capital letters may be used as subscripts, e.g., $X_A = X_A$

Blood-Gas Measurements

Abbreviations for these values are readily composed by combining the general symbols recommended earlier. The following are examples:

PaCO ₂	Arterial carbon dioxide tension			
C(a-v)O ₂	Arteriovenous oxygen content difference			
CcO ₂	Oxygen content of pulmonary end-capillary blood			
F _E CO*	Fractional concentration of CO in expired gas			
P(A-a)O ₂	Alveolar-arterial oxygen pressure difference; the previously used symbol, A-aDO ₂ is not recommended			
SaO ₂	Arterial oxygen saturation of hemoglobin			
Q _{sp}	Physiologic shunt flow (total venous admixture) defined by following equation when gas and blood data are collected d ing ambient air breathing:			

$$Qsp = \frac{CcO_2 - CaO_2}{CcO_2 - CvO_2} \cdot Q$$

PETO2 TCPO, PO₂ of end tidal expired gas

Transcutaneous PO2

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The Limitations of Pulmonary Function Testing

JOHN F. MURRAY

Pulmonary function tests are widely used in the evaluation and management of patients with known or suspected disorders of respiration. The clinical application of these studies rests on knowledge of pulmonary physiology, which has a sound experimental foundation, although we need more information about the relationship between the structure and function of the lungs in health and disease. Despite having a scientific basis, the results of pulmonary function tests must be interpreted empirically, and the temptation to use certain abnormalities to deduce the presence of specific underlying pathologic changes has proved irresistible, albeit often misleading. In a recent editorial, Butler(1) reviewed the inherent limitations that may involve each step of the testing continuum: the apparatus, the patient, the technician, and the interpreter. The purpose of this discussion is (1) to consider why we perform pulmonary function tests and what we are trying to measure; (2) to remind the reader about certain measurement inaccuracies that are frequently overlooked; and (3) to emphasize the enormous problems that plague interpretation, particularly those concerning the concept of normality and how to isolate a specific abnormality.

WHAT ARE WE MEASURING?

The main reasons for performing pulmonary function studies are listed in Table 1-1. Each of these indications deals with some aspect of the consequences of how particular disorders affect respiratory function. Thus, to examine in

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