

(14.8y [12.5y-16.7y]). LCI was significantly higher in those with CF when using both devices (mean difference[95% CI], CF-controls): 2.47[1.4;3.5] for the MS-SF₆ and 2.20[1.2-3.2] for N₂-MBW.

There were no significant group differences between devices for either LCI (mean difference[95% CI] -0.14[-0.45;0.16] or FRC -0.15L[-0.2;-0.08]). Within-subject variability was proportional to mean values (see Figure) and ranged from 0.4-15.7% for LCI and 0.0-19.6% for FRC.

Conclusion Despite some previous reports that N₂ washout results in higher LCI values than MS-SF₆ washout, on average, we found similar values in both healthy school-age children and those with CF. Further work is required to examine causes of within-subject variability and assess validity and sensitivity over a wider age range, including preschool children, before commercial N₂-MBW devices can be confidently used in multi-centre trials.

REFERENCE

1 Aurora et al. *Am J Respir Crit Care Med.* 2011;183:752-8

P100 THE FEASIBILITY OF USING COMMERCIAL MULTIPLE BREATH NITROGEN WASHOUT DEVICES IN SCHOOL-AGED CHILDREN

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10.1136/thoraxjnl-2014-206260.241

Background Multiple breath inert-gas washout (MBW) using sulphur hexafluoride (SF₆) measured by mass spectrometry (MS), is sensitive to early lung disease in children with Cystic Fibrosis (CF)¹ but is not widely available. To increase the accessibility of MBW, commercial devices have been adapted using nitrogen-washout (N₂-MBW). Our aim was to assess the feasibility of two commercial N₂-MBW devices as supplied by the manufacturers compared to a custom-built MS system in school-aged children.

Methods Patients with CF and controls performed MBW on three devices; the Exhalyzer[®]D (ECO MEDICS AG); the EasyOne Pro[®]LAB (nidd Medizintechnik AG) and the MS system (AMIS 2000, Innovision ApS) on the same test occasion (order randomised). Attempts were made to obtain 3 technically acceptable runs/device (maximum 8 attempts on each).

During testing children watched a DVD and were encouraged to breathe normally. Data were analysed using the 'clinical application' setting for both commercial devices, and customised software for the MS. Quality control was in accordance with the ATS/ERS consensus statement¹ and manufacturers' guidelines.

Results 14 control (mean[range]age: 15.0[12.5-16.7]yrs) and 18 children with CF (13.5[7.8-17.4]yrs) were assessed. The median (range) number of runs attempted were: MS 3(3-8), Exhalyzer[®]D 4(3-6), EasyOne Pro[®]LAB 4(3-8). Average calibration time was shorter for EasyOne Pro[®]LAB (5 min) than either MS (11 mins) or Exhalyzer[®]D (12 min). Total test duration was similar between devices and dependent on disease severity.

3 acceptable MBW runs were achieved in all children using the MS, 75% with the EasyOne Pro[®]LAB, and 47% on the Exhalyzer[®]D system (see Table). Reasons for failure with Exhalyzer[®]D were usually due to technical/equipment problems, whereas for the EasyOne Pro[®]LAB these were generally associated with marked changes of breathing pattern at commencement of washout, leading to exclusion of one or more runs.

Discussion Despite use in an experienced MBW centre, our initial attempts to implement commercial MBW devices according

Abstract P100 Table 1 Number (n) of technically satisfactory runs according to MBW device

	MS	EasyOne Pro Lab	Exhalyzer D
n = 3	32	24	15
n = 2	0	7	1
n = 1	0	0	1
None acceptable	0	1	15

to manufacturers' guidelines resulted in a relatively low success rate in schoolchildren when compared to MS. Subsequent feedback to manufacturers has led to further adaptations which should improve feasibility in future, although this has yet to be assessed in very young children.

REFERENCE

1 Robinson et al. *Eur Resp J* 2013

P101 EFFECTS OF USING A MASK VS. MOUTHPIECE ON THE MULTIPLE BREATH INERT GAS WASHOUT TECHNIQUE

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10.1136/thoraxjnl-2014-206260.242

Assessment of ventilation inhomogeneity using the multiple breath washout (MBW) technique has been shown to be more sensitive than spirometry in detecting early cystic fibrosis disease throughout childhood. The current "gold standard" interface for school age children and adults is a mouthpiece. Although masks are better tolerated by infants and younger children, their use increases equipment deadspace-which could influence measured values and hence interpretation of results. The aim of this study was to examine the effect of using a mask vs mouthpiece on values of functional residual capacity (FRC) and the lung clearance index (LCI) derived from MBW.

Method Comparisons were performed in healthy adults. The study design incorporated repeated measures as well as interface comparison. The mask was selected to mimic measurement conditions in infants, the deadspace of 85 mL being approximately 1-2 ml/kg in adults. Mouthpiece (MP) deadspace was ~5 ml. Subjects were randomly allocated to group A (Mask-Mouthpiece-Mouthpiece) or group B (Mouthpiece-Mouthpiece-Mask) protocols. Each subject performed a total of 9 MBW runs, in 3 sets, each consisting of 3 runs, with a 5-minute break between each set. MBW was performed using a mass spectrometer as described previously (Aurora 2005 AJRCCM). Paired t-tests with 95% limits of agreement were used to establish repeatability (MP1 vs. MP2) and any differences between Mask vs. Mouthpiece. This study was approved by the local research ethics committee and written consent obtained from subjects.

Results Technically satisfactory comparative data were obtained on 15 occasions in 14 adults (36% males; age: 22-56 years). Respiratory rate and tidal volume were similar using either approach. Repeatability: Both FRC and LCI were repeatable using the mouthpiece [(Mean (95% CI) diff: FRC: 0.012L (-0.05;0.07); LCI: -0.1(-0.3; 0.1)]; Figure 1A and B. Mask vs. Mouthpiece: FRC and LCI were both significantly higher when assessments were made using a mask compared with a mouthpiece: FRC: 0.101L (0; 0.202); LCI: 0.4 (0.2;0.7); Figure 1C and D.

Conclusion The increase in LCI when using a facemask exceeded normal within test variability in adults and could