

Introduction Exercise capacity in chronic obstructive pulmonary disease (COPD) is limited both by abnormal pulmonary mechanics, reported as breathlessness, and by leg muscle fatigue. To improve functional capacity it is important to understand the primary physiological constraint. Neural respiratory drive (NRD), measured using the diaphragm electromyogram expressed as a proportion of maximum ($EMG_{di}\%max$), quantifies the load on the respiratory muscles imposed by abnormal pulmonary mechanics, and relates closely to breathlessness. We hypothesised that end-exercise $EMG_{di}\%max$ would be higher in patients stopping because of breathlessness than in those stopping because of leg fatigue.

Methods EMG_{di} , ventilation (V_E), oxygen consumption (VO_2) and ventilatory reserve ($V_E/MVV\%$) were measured in 23 COPD patients (median (IQR) FEV_1 39 (30.0 to 56.8)%predicted) during exhaustive cycle ergometry. Differences in physiological variables between groups of patients stopping because of breathlessness, leg fatigue or both were examined using 1-way ANOVA.

Results $EMG_{di}\%max$ was higher in patients stopping because of breathlessness ($n = 12$, $EMG_{di}\%max$ 75.7 (69.5 to 77.1)%) than in those stopping because of leg fatigue ($n = 8$, $EMG_{di}\%max$ 44.1 (39.4 to 63.3)%, $p < 0.05$). There were no significant differences in end-exercise V_E or VO_2 . $V_E/MVV\%$ tended to higher levels in the breathless group.

Discussion These results suggest that patients limited by breathlessness due to ventilatory constraints can be identified as those reaching near-maximal levels of NRD during exercise. Measurement of $EMG_{di}\%max$ during exercise could prove useful in identifying patients whose functional performance would be best optimised by improving pulmonary mechanics rather than interventions to train peripheral muscle groups.

S54 NEURAL RESPIRATORY DRIVE MEASURED USING PARASTERNAL INTERCOSTAL MUSCLE ELECTROMYOGRAPHY IN PATIENTS WITH INTERSTITIAL LUNG DISEASE

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Introduction Forced vital capacity (FVC) and gas transfer (TLCO) are often used to assess disease severity and monitor progression in patients with interstitial lung disease (ILD). Difficulty in performing the required manoeuvres, particularly in severe disease, and inherent measurement variability makes detection of clinically important changes difficult using these parameters. There is, therefore, a need for new biomarkers in this patient group. Neural respiratory drive (NRD) reflects the load on the respiratory system and the capacity of the respiratory muscles. Parasternal intercostal muscle electromyography (EMGpara) provides a non-invasive measure of NRD which relates to disease severity and breathlessness in obstructive lung diseases. Measurements of EMGpara in ILD could potentially quantify overall disease severity.

Aim The aim of the study was to investigate the relationships between EMGpara, lung function, breathlessness, functional status and quality of life (QoL) in ILD.

Method EMGpara was measured in 45 patients with a range of fibrotic lung diseases using surface electrodes placed in the second intercostal spaces bilaterally. Mean peak root mean square EMGpara per breath was calculated and expressed as a percentage of maximum EMGpara ($EMGpara\%max$). The neural respiratory drive index (NRDI) was derived by multiplying $EMGpara\%max$ by the respiratory rate. Spirometry and lung gas transfer were

performed and the composite physiologic index (CPI) calculated. Six minute walk test (6MWT) and 4 metre gait speed (4MGS) were used to determine functional status. Health-related quality of life was assessed with the King's Brief Interstitial Lung Disease (K-BILD) and the St George's Respiratory Questionnaire (SGRQ). The Baseline Dyspnea Index (BDI) was used to grade breathlessness.

Results NRDI correlated significantly with $VC\%predicted$ ($r = -0.36$, $p = 0.018$) and the CPI ($r = 0.40$, $p = 0.01$). No significant correlations were found between EMGpara or NRDI and breathlessness, QoL or functional status.

Conclusion EMGpara is a feasible measure in ILD. EMGpara correlates with prognostic markers suggesting potential value as a biomarker integrating important pathophysiological changes in lung mechanics in fibrotic ILDs. The lack of association with QoL measures and BDI requires further investigation.

Abstract S54 Table 1 Patient characteristics, $EMGpara\%max$, NRDI, lung function and functional status in forty-five patients with ILD. Correlation coefficients for the relationship between NRDI and individual variables given (* $p < 0.05$)

Parameter	Median	Range	Correlation with NRDI (r=)
Age (years)	65	35–85	-
BMI	27.5	20.6–40.7	-
$FEV_1\%predicted$	82.5	41–148	-0.38*
$FVC\%predicted$	83	43–137	-0.36*
$TLCO\%predicted$	48	21–78	-0.25
$EMGpara\%max$ (%)	8.1	3.4–20.2	-
NRDI a.u.	148	61–514	-
K-BILD	65	21–100	-0.007
SGRQ	37	2.3–80	-0.74
BDI	6	-11	-0.004
CPI	44	9.8–64	0.40*
6MWD (%predicted)	58	18–130	-0.15
4MGS (m/s)	0.88	0.46–1.95	-0.21

S55 NEURAL RESPIRATORY DRIVE USING PARASTERNAL ELECTROMYOGRAPHY IN CLINICALLY STABLE CYSTIC FIBROSIS PATIENTS: A PHYSIOLOGICAL MARKER OF LUNG DISEASE SEVERITY AND EXERCISE CAPACITY

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Introduction Measurement of neural respiratory drive, using parasternal intercostal muscle electromyography (EMGpara), has previously been shown to relate to pulmonary function impairment and exercise-induced breathlessness in advanced cystic fibrosis (CF). This measure reflects the load on the respiratory system and the capacity of the respiratory muscles and therefore may provide a composite measure of overall lung disease severity. In order to utilise EMGpara clinically in CF, its relationship to standard physiological outcome measures requires further investigation across a broad range of disease severities. Aim: To investigate the relationships between EMGpara and standard measures of pulmonary function and exercise performance in patients with CF.

Methods Thirty patients with clinically stable CF were recruited. EMGpara was recorded during five minutes of tidal breathing using electrodes positioned in the second intercostal space directly lateral to the sternum. Peak EMGpara per breath was averaged over the final minute of the recording and expressed as a percentage of EMGpara recorded during a maximal inspiratory manoeuvre ($EMGpara\%max$). Spirometry, lung volumes by body