CPAP. We studied 20 (9 male), ambulatory adult patients with EDAC and/or TBM: mean $\pm$ SD age 60 ± 13 years, height 1.67 ± 0.86 m, and BMI 34.1 ± 6.6 kg/m². The NRDI was 356 ± 182 AU while self-ventilating and reduced when CPAP was applied (231 ± 122 AU; $p < 0.001$). The 6MWT while on optimal CPAP was increased comparing to self-ventilation and sham CPAP (296 ± 112 m vs 264 ± 120 m vs 252 ± 125 m, respectively; $p < 0.001$) (figure 1). The treatment effect between sham and optimal CPAP was 31 ± 39 m (95% CI: 13 to 50 m). While on optimal CPAP, 12 patients increased their 6MWT more than 30 m compared to self-ventilation (responders). Comparing responders with non-responders, differences were identified for NRDI (-167 ± 110 AU vs. -63 ± 90 AU, respectively; $p = 0.039$) and 6MWT while self-ventilating (203 ± 94 m vs. 336 ± 133 m, respectively; $p = 0.022$).

In conclusion, CPAP reduces neural respiratory drive and increases exercise capacity in patients with EDAC/TBM. Furthermore, the degree of functional limitation and off-loading of the respiratory muscles on CPAP can identify those likely to have a reduction in neural respiratory drive and enhanced exercise tolerance.

S135 TIMING OF ACIDAEMIA ONSET IN EXACERBATIONS OF COPD REQUIRING ASSISTED VENTILATION AND IN-HOSPITAL MORTALITY

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Introduction Predicting which patients are likely to benefit from assisted ventilation in exacerbations of COPD (ECOPD) is difficult. Existing prognostic scores did not assess the prognostic value of timing of acidaemia onset relative to admission during development. The 2011 National Chronic Obstructive Pulmonary Disease Resources and Outcomes Project (NCROP) report identified this as a potentially important marker of non-invasive ventilation (NIV) success but further study has been limited. We investigated the relation of timing of respiratory acidaemia to NIV outcomes in ECOPD. Of importance, in both participating hospitals few patients are denied NIV on grounds of assumed futility.

Methods A retrospective cohort of consecutive, unique patients, hospitalised with a primary diagnosis of ECOPD were identified from known cohorts, hospital coding records coding and ventilation service records. Other selection criteria included: age 35+ years, smoking history 10+ pack years, air-flow obstruction on spirometry; received assisted ventilation (either NIV or invasive) for acidaemic respiratory failure; and absence of comorbidity expected to limit survival to <12 months (principally metastatic cancer). Population descriptors, ventilation details and outcome data were collected from notes and electronic records.

Results 489 consecutive patients were identified between 30/11/08 and 19/5/13; 124 (25.4%) died in-hospital. Median time to ABG prompting ventilation was 2 hour 42 m (IQR 1 hour 2 m – 15 hour 28 m). Most (94.5%) received NIV alone, 5.5% received invasive ventilation (+/-NIV). In patients requiring assisted ventilation within 12 hours of admission, mortality was 18.3% (65/356), compared to 44.3% (59/133) in all those treated after 12 hours ($p < 0.01$).

Discussion Our study has several strengths including objective confirmation of COPD, capture of consecutive patients and liberal NIV use. Compared to patients with respiratory acidaemia on or shortly after admission, later development was associated with progressively higher mortality. 12 and 48 hours were identified as clinically useful thresholds. Of note, lower pH, FEV1 and prior LTOT prescription do not account for worse outcome. Older age, greater comorbidity, frailty (eMRC5b: requiring help washing and dressing when recently stable), and a strong trend towards increasing pneumonia are associated with later development of acidaemia. Timing of acidaemia should be considered when deciding whether to initiate NIV.

Abstract S135 Table 1 Key population descriptors and inpatient mortality grouped by timing of acidaemia onset

Time	Up to 12 hours	12–48 hours	>48 hours
N	356	69	64
Age	71.9 (9.9)	73.8 (10.0)	76 (9.8) *
FEV1%	37.1 (15.6)	42.8 (18.9) *	38.1 (17.4)
BMI	24.9 (7.5)	24.7 (5.9)	22.8 (7.2) †
LTOT	109 (30.6%)	18 (26.1%)	16 (25.0%)
eMRC5b	5a (4–5a)	5a (4–5a)	5a (4–5b) †
eMRC5a	120 (33.7%)	25 (36.2%)	19 (29.7%)
eMRC5b	63 (17.7%)	13 (18.8%)	24 (37.5%) *
Charlson Index	1 (1–2)	2 (1–3)	2 (1–3) *
Consolidation at NIV	157 (44.1%)	38 (55.1%)	36 (56.3%)
pH at NIV	7.23 (0.09)	7.26 (0.08) †	7.26 (0.08) †
PCO2 at NIV	10.3 (2.5)	10.0 (2.4)	9.3 (1.8) *
Peak Pressure (IPAP)	20 (18–22)	19 (17–20)	18 (16–20) †
Inpatient Mortality	65 (18.3%)	22 (31.9%) †	37 (57.8%) †

t-test, Mann Whitney U or χ^2 test for parametric, non-parametric and categorical data respectively.

* $p < 0.05$; † $p < 0.01$: compared to "Up to 12 hours group"

S136 LUNG PROTECTIVE MECHANICAL VENTILATION FOR ACUTE RESPIRATORY FAILURE IS NOT BEING IMPLEMENTED IN UK CLINICAL PRACTICE

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Introduction The benefits of lung protective ventilation have been replicated in multiple trials.^{1,2} However, we suspected that adherence to this standard of care remained poor. Using the NIHR critical care Health Informatics Collaborative (ccHIC) database, we analysed data from 11 teaching hospital intensive care units (22 524 patient episodes) to investigate real-world clinical practice.

Methods 1248 patient episodes, where ventilation was continued for ≥ 48 hours, with 232,600 hours of concurrent mechanical ventilation and blood gas data were identified as suitable for analysis. Short gaps in ventilation (<6 hours) were imputed based on the median of nearest known values, and