

# Commencement of fixed dose ICS of varying particle size as a predictor of pneumonia in COPD patients

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**Introduction:** Demographic and clinical characteristics have been used to identify predictive risk factors for pneumonia development in COPD patients.

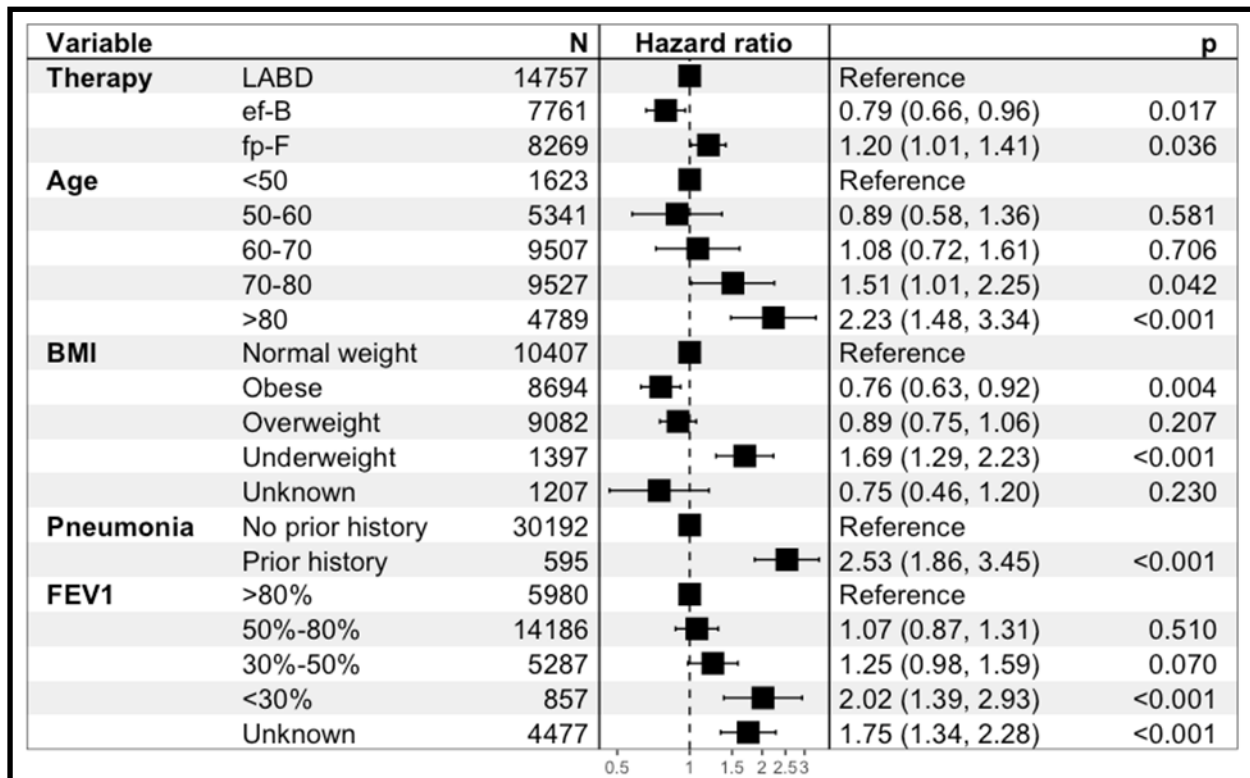
**Aim:** The present investigation extended this modelling to include commencing pharmacological therapies as a variable.

**Methods:** A historical, observational study using patients  $\geq 40$  years, with  $\geq 1$  year of continuous medical data and who were commencing a fixed dose of extrafine beclometasone (ef-B), fine particle fluticasone (fp-F) or long-acting bronchodilators (LABD) was conducted. Cox regression was used to develop the risk prediction model for time to pneumonia. Validation was undertaken using the hold-out method (80 development:20 validation cohort split). A time dependent receiver operating characteristic curve was used to assess performance.

**Results:** 38,484 COPD patients were evaluated. Older age, a low body mass index, low FEV1% predicted and a prior history of pneumonia were associated with an increased risk of pneumonia (Figure 1). Commencing ef-B was associated with a reduced risk of pneumonia when compared to LABD, whereas fp-F was associated with an increased risk. The model

achieved an area under the curve of 0.74 at one year after therapy initiation in the validation cohort.

**Conclusions:** Reduced pneumonia risk in COPD patients commencing ef-B is an important observation, particularly for individuals at high risk due to demographic and clinical factors.



**Figure 1:** Cox regression modelling of risk factors associated with pneumonia development in COPD patients. BMI, body mass index; ef-B, extrafine beclometasone; FEV1, forced expiratory volume in 1 second; fp-F, fine particle fluticasone; LABD, long acting bronchodilator.