

Background

Asthma is characterised by distinct phenotypes, with 50-70% of severe cases involving type-2 (eosinophilic) inflammation, which can be treated with oral or inhaled steroids and biological treatments. Type-2 inflammatory biomarkers can be easily measured in the community using fractional exhaled nitric oxide (FeNO) and blood eosinophil count (BEC). We hypothesise that some individuals may have type-2 inflammation present without symptoms, which may represent a target for earlier identification and treatment.

Objective:

To identify the distribution of type-2 biomarkers (FeNO and BEC) in healthy young adults without asthma

Methods

Inclusion: Aged 18-50 without respiratory disease

Exclusion: Respiratory disease and/or immunosuppressive medication use.

Participants underwent FeNO measurement using the **NIOX VERO®** (see below) and eosinophil measurement using the **HemoCue® WBC DIFF system** from a fingerprick blood sample.



Following FeNO testing and point-of-care BEC measurement, participants were stratified according to the following biomarker categories:

Biomarker stratification category	FeNO and BEC cut-offs
Low	FeNO < 25 ppb or BEC ≤ 0.15 x 10 ⁹ /L
Moderate	FeNO 25 – 49 ppb or BEC 0.15 – 0.29 x 10 ⁹ /L
High	FeNO ≥ 50 ppb or BEC ≥ 0.3 x 10 ⁹ /L

Conclusion

Our study found that 18.4% of healthy young adults have ≥ 1 raised biomarker indicative of eosinophilic airways disease. This was associated with atopy, suggesting pre-symptomatic of subclinical airways disease. Long-term evaluation of these individuals is required to explore their future airways disease risk and whether pre-symptomatic treatment to reduce inflammation could alter their disease trajectory.

Results

Data from 87 participants is reported:

Mean age = 30.3 years (±9.1) Median FeNO = 18 ppb (range: 12 – 32)
 28 (32%) of participants were male Median BEC = 0.1 x 10⁹/L (range: 0.1 – 0.2)
 11 (13%) of participants were current or ex-smokers

Figure 1:

There was a positive correlation between FeNO and BEC (Spearman $p = 0.227$; $p = 0.009$)

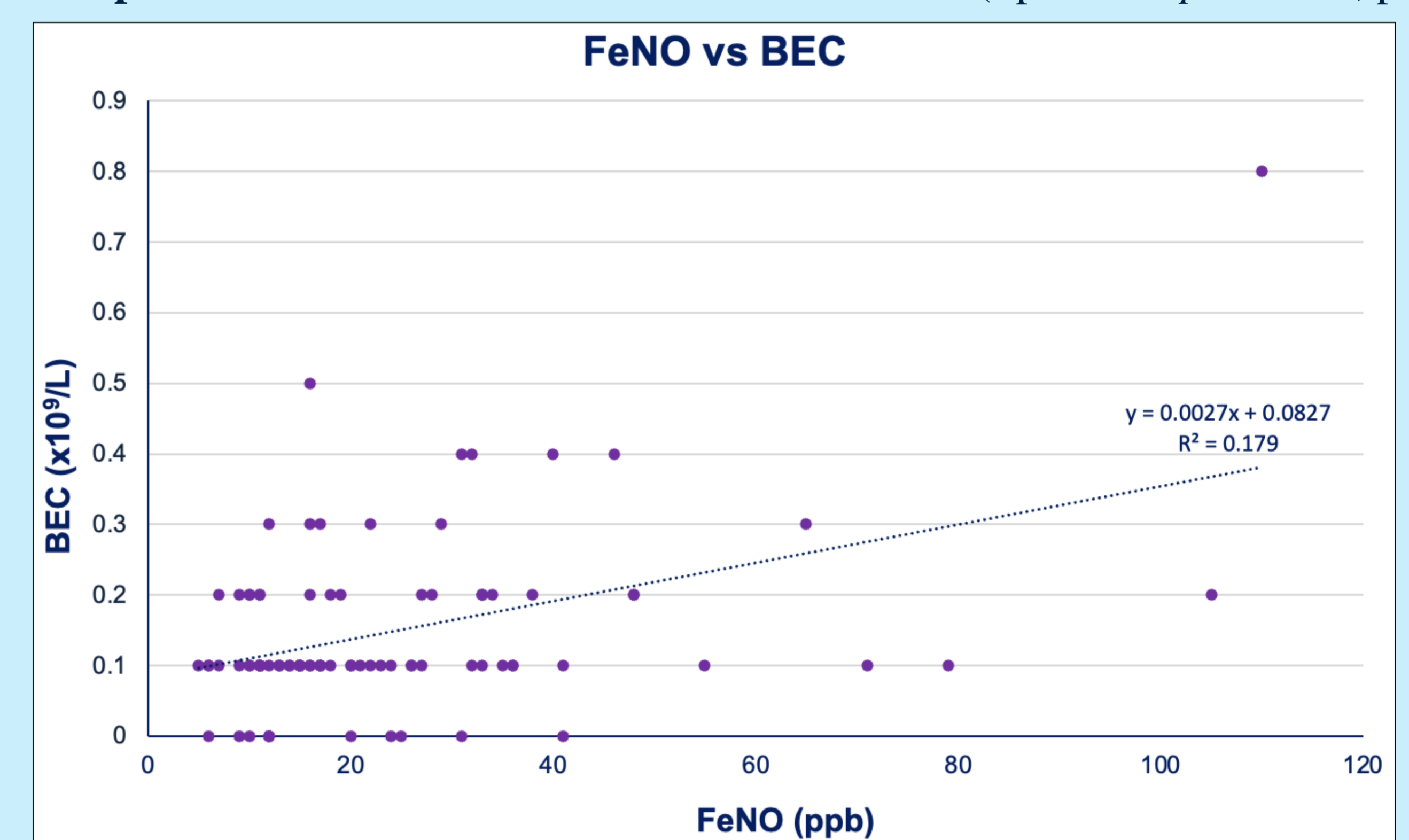


Figure 2:

Males had a higher median FeNO than females (31.5 vs. 16; $p < 0.001$), but there was no significant difference between the sexes for BEC.

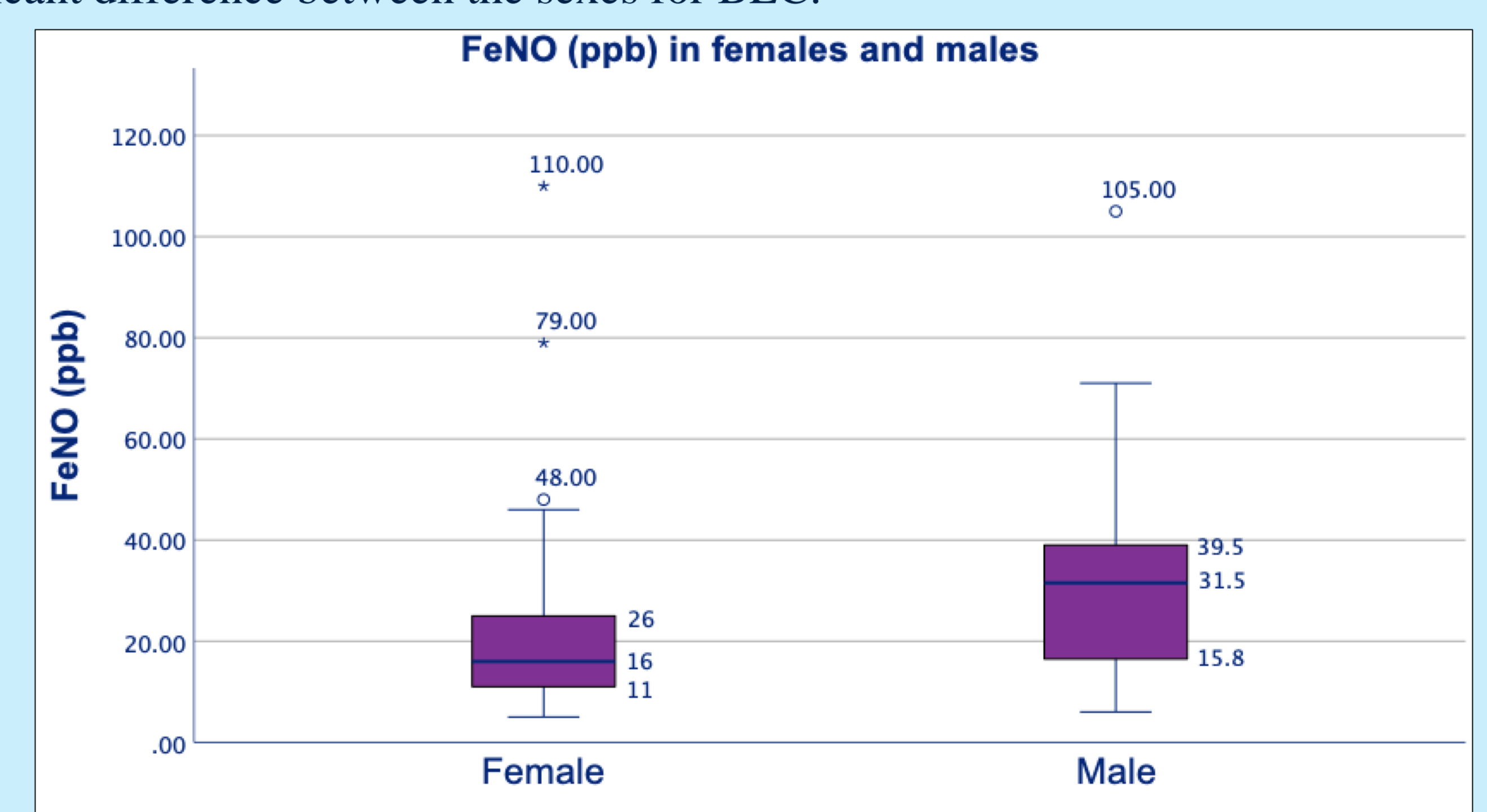


Figure 3:

Higher BEC (≥ 0.3 x 10⁹/L) was associated with increased atopy – 92% of those with BEC ≥ 0.3 x 10⁹/L experienced atopic symptoms compared to 34.7% of those with BEC < 0.3 x 10⁹/L ($p = 0.002$).

