Biomarkers in Chronic Obstructive Pulmonary Disease: Emerging Roles of Eosinophils and Procalcitonin

Pradeesh Sivapalan, Jens-Ulrik Jensen

Department of Internal Medicine, Respiratory Medicine Section, Herlev and Gentofte Hospital, University of Copenhagen, Hellerup, Denmark; Department of Clinical Medicine, Faculty of Health Sciences, University of Copenhagen, Copenhagen, Denmark

Keywords
Biomarker · Blood eosinophils · Procalcitonin

Abstract
Antibiotics can improve the prognosis in patients with exacerbation of chronic obstructive pulmonary disease. However, the overuse of antibiotics can carry serious adverse effects for patients (gastrointestinal infections) and for society (bacterial resistance). Likewise, systemic corticosteroids may also help these patients, but also carries severe adverse effects like osteoporosis, muscle loss, and diabetes, in many patients. Whenever safe methods exist to reduce these two treatment modalities, they should be implemented. The blood biomarkers procalcitonin and the fraction of leukocytes known as eosinophil granulocytes have been proven in randomized controlled trials (RCTs), to effectively, significantly, and substantially assist in reducing the use of these two potent, yet toxic medication types. In this review, the background and main clinical results are discussed, explaining the rationale for biomarker-guided clinical decisions. Also, the main expected effects, their sizes, and importantly the limitations to such a strategy are described. Clinical evidence is prioritized with main weight on RCTs and meta-analyses of these and regarding outcomes, and focus is set on the safety of such a biomarker-guided strategy, as well as the effects on medicine reduction. In an epoch of increasing demands to physicians from patients and politicians to cure and reduce symptoms, the Hippocratic phrase of "primum non nocere" or "first, do no harm" seems more than ever of contemporary importance.

Introduction

Biomarkers of infection and inflammation have been used for decades in patients with lower respiratory tract infections and flares of inflammatory conditions [1–3]. Exacerbations in chronic obstructive pulmonary disease (COPD) are defined in several ways, although many clinicians and researchers use the classic Anthonisen criteria (increased dyspnea, increased sputum volumes, discolored sputum, and a therapeutic response). Such criteria can lead to the antibiotic and corticosteroid overuse, since they do not necessarily differentiate which patients are in need of such treatments, and in case of doubt, treatment will often be initiated, to avoid undertreating.
For patients with COPD exacerbations, a potentially deadly condition, often requiring hospital admission, the usual biomarkers like the white blood cell count and C-reactive protein seem not to be quite sufficient, when the clinician needs to decide whether to treat with antibiotics and systemic corticosteroids. These common biomarkers are not very sensitive and might be elevated for various reasons, even during stable periods due to systemic inflammation [4], and a low positive predictive value may cause unnecessary use of antibiotics [5]. This is far from satisfactory, since such treatments carry significant risks for the patient, such as serious gastrointestinal infections with Clostridium difficile, superinfections with resistant bacteria, muscle wasting, diabetes mellitus, and osteoporosis.

This review does not attempt to mention all candidate biomarkers for use in COPD patients, rather, the aim is to describe the clinical utility of two biomarkers, blood eosinophils and procalcitonin (PCT), that have been extensively investigated to assist in reducing corticosteroids in patients with severe COPD exacerbations and likewise to reduce unnecessary use of antibiotics. The reader will be able to (i) make a decision of whether there is sufficient evidence to support use of one or both of these biomarkers in his/her own setting, (ii) if decision to use them is made, then propose an evidence-based algorithm for the most effective way to use them.

Much is known about the concrete use of these biomarkers and what should be expected or not expected when using these biomarkers, how to interpret them, and how to implement them. The precise use of them has, however, only been implemented in few hospitals worldwide. Acute exacerbations of COPD may have distinct causes and can be viral, bacterial, or caused by eosinophilic inflammation, each with distinct treatment needs [6]. Due to diagnostic difficulty in distinguishing virally triggered acute exacerbations of COPD from those driven by bacterial infection, there is a potential for unnecessary use of antibiotics in this group of patients. The overuse of antibiotics for respiratory infections is an important driver of multidrug resistance. Therefore, it is imperative to develop effective diagnostic tools to guide both corticosteroid and antibiotic treatment in lung patients and in patients with respiratory infections [6].

**Eosinophilic Inflammation**

COPD exacerbation was initially thought to be mediated primarily by neutrophil-mediated inflammation, while eosinophilic inflammation is thought to be more characteristic in asthma rather than COPD. However, a subset of 20–40% of COPD patients displays eosinophilic airway inflammation, even when carefully excluding patients with possible asthma [7]. Eosinophils are key inflammatory and immune effector cells. Under normal conditions eosinophils are dormant in the blood, but upon exposure to pro-inflammatory cytokines (interleukin [IL]-3, IL-5 IL-6, and granulocyte-macrophage colony-stimulating factor) they activate and migrate to inflamed tissue [8]. Monoclonal antibodies targeting IL-5, the main cytokine involved in the activation of eosinophils, is known to reduce exacerbations of eosinophilic asthma effectively [9]. However, in eosinophil COPD, the effect of anti-IL-5 is still disputed [10], although many researchers have commented on the lack of eosinophilia as an inclusion criterion in these studies and claimed that blocking an eosinophil pathway makes less sense in patients without activation of such a pathway. Corticosteroids are unlike anti-IL drugs, commonly used for regular treatment of COPD, being particularly beneficial in patients with asthmatic or inflammatory features and co-morbidities (such as asthma-COPD overlap syndrome) [11–13]. In healthy persons, eosinophils usually are not found in the lungs, and their presence in the lungs is indicative of an abnormal inflammatory reaction [14]. Eosinophils are typically quantified as a percentage of total leukocyte amounts or as absolute numbers (cells/µL). Both measurements have been shown to be in concordance with each other [7, 15]. Blood eosinophil counts have also been shown to be in concordance with sputum eosinophil counts, and are therefore often used as a proxy for eosinophilic lung inflammation [8]. Eosinophilic inflammation can be present both during exacerbations and stable COPD [7]. Numerous studies have demonstrated associations between blood eosinophil counts and relevant clinical COPD outcomes. Most of the studies did find an association with higher blood eosinophils (eosinophilic COPD phenotype) and worse clinical outcomes in terms of increased risk of readmissions, longer hospital stays, and future exacerbations [16–20]. However, the association is not fully clarified.

**Inhaled Corticosteroid Use and the Blood Eosinophil Count**

According to previous guidelines, patients with moderate to very severe COPD should be treated with inhaled corticosteroids (ICS) [21]. However, the use of ICS in COPD patients has been shown to have a variable effect, which needs to be weighed against the side effects of the treatment such as increased incidence of pneumonia, diabetes [22], osteoporosis, and cataracts [23, 24]. Until re-
Recently, no serum biomarkers had a demonstrated association with COPD treatment response. In 2007, randomized controlled trials (RCTs) showed reduction in COPD exacerbation in patients with a sputum eosinophil count >3% treated with ICS. Subsequently, peripheral blood eosinophils have been extensively studied and seem to be able to better predict the effect of ICS [25]. Several post hoc analyses looking at the benefits of ICS/LABA in comparison to LABA have shown a continuous relation between ICS response and eosinophil counts [15, 26, 27]. Other studies have grouped patients into discrete groups based on blood eosinophils and leading to similar conclusions that higher blood eosinophil levels increase the likelihood of ICS treatment benefit [28–30]. This is reflected in recent GOLD recommendations, that specify that eosinophil counts should be considered when deciding whether to administer ICS. The newest GOLD guidelines recommend ICS usage in patients with blood eosinophils ≥300 cells/µL, and in patients with blood eosinophils >100 cells/µL, and persistent exacerbations while on bronchodilators [31]. Recently, the European Respiratory Society published a guideline for withdrawal of ICS based on the blood eosinophil count. This guideline recommends ICS withdrawal in patients with a blood eosinophil count <300 cells/µL and no frequent exacerbations [32]. All these recommendations are, however, based on post hoc studies and currently, no RCTs have tested this algorithm. At this current time, there is one multicenter RCT seeking to validate these findings (NCT04481555), with planned readouts by 2025.

**Systemic Corticosteroids and the Blood Eosinophil Count**

Unlike stable COPD, acute exacerbations are commonly treated with orally administered systemic corticosteroids such as prednisolone [21]. Systemic corticosteroids are used to improve recovery from symptoms, but they do not improve long-term decline in lung function, future exacerbations after the first month, and the length of stay in intensive care units or mortality [33]. The criteria used to define COPD exacerbations (an acute worsening of respiratory symptoms that result in additional therapy) are broad, and this may lead to the overuse of corticosteroids [31]. The overuse of systemic corticosteroids is associated with increased risk of osteoporotic fractures, cataracts, infections [34], and venous thromboembolism [35, 36]. To reduce the usage of systemic corticosteroids, it has been proposed to use blood eosinophil counts to

---

**Fig. 1.** Treatment algorithm from the CORTICO-COP trial [42].
guide corticosteroid treatment. One RCT used blood eosinophils to categorize patients as having eosinophilic or noneosinophilic exacerbations and found noninferiority when treating noneosinophilic exacerbations with placebo rather than systemic corticosteroids. Notably, there was a reduction of 49% in the total corticosteroid prescription for the eosinophil-guided group ($p < 0.001$) [37]. Similarly, other studies have shown better corticosteroid response in COPD patients with higher eosinophil counts compared to patients with a lower blood eosinophil count [38–40]. The randomized noninferiority CORTICO-COP trial found that daily eosinophil counts could be used to guide treatment to reduce corticosteroid usage in hospitalized patients with COPD exacerbation using the algorithm shown in Figure 1. This led to a reduction in median corticosteroid treatment duration from 5 days to 2 days with approximately two-thirds of the eosinophil-guided patients being off corticosteroids on any given day throughout the study except on day 1 (shown in Fig. 2). This treatment algorithm was shown not to affect days alive and out of hospital within 14 days after recruitment or a 30-day all-cause mortality rate. There were documented fewer side effects in terms of fewer patients with hyperglycemia and fewer patients with worsening of diabetes in patients with preexisting diabetes in the eosinophil-guided corticosteroid group [41]. These analyses indicate that the blood eosinophil count could potentially be used as a biomarker to guide corticosteroid therapy for exacerbations and may help to avoid unnecessary exposure to systemic corticosteroids.

In Table 1, some of the key trials of eosinophil-guided corticosteroid therapy are noted with the main characteristics and results.

**PCT Biology**

PCT is a 114 amino acid, 12.6 kDa polypeptide prohormone of calcitonin [42]. Calcitonin is produced in the neuroendocrine medullary C-cells of the thyroid, which is embryologically derived from the neural crest [43]. PCT produced in the C-cells undergoes posttranslational processing, to form the bioactive calcitonin hormone of 32 amino acids. In animal studies, PCT has been isolated in the tissue from the adrenal gland, spleen, brain, liver, pancreas, colon, lung, fat tissue, spinal cord, testes, and stomach [44]. In bacterial sepsis, the mRNA of the calcitonin (CALC)-1 gene is upregulated more uniformly than the mRNA of other inflammatory cytokines, such as tumor necrosis factor-α, IL-1β, and IL-6 [45]. Mature calcitonin blood levels do not increase significantly during sepsis. The explanation for this has been proposed as a
“shift of pathway,” from a “specific pathway” (where calcitonin precursors are released in secretory granules with enzyme activity to process the precursors to mature calcitonin hormone) to a more “generalized pathway” (where calcitonin precursors are “bulk released” without secretory granules and thus without enzymatic activity to process peptides) [45]. Other calcitonin precursors can be upregulated during bacterial infections and other acute diseases; of these the mid-region pro-adrenomedullin has been claimed to be a valuable severity marker in sepsis [46, 47].

PCT in Diagnosis of COPD Exacerbation

PCT is a biomarker of bacterial infection [48], which is released 2–6 h after IL-1β stimulation; however, it can be almost completely blocked by coadministration of IFNγ [49], this suggesting a potential use in diagnosis and eventual applicability in differentiating between viral and bacterial acute exacerbated COPD (AECOPD). In a diagnostic accuracy study (n = 98) of patients with severe and very severe COPD exacerbation, a PCT of <0.5 ng/mL was observed to have a negative predictive value of 95% for isolation of a human pathogenic bacterium from the airway culture [50]. This has, however, been disputed; and in another small study (n = 72), no difference in PCT was found between patients with viral and bacterial origin of COPD exacerbation. A meta-analysis in this area did though show a clear link between PCT and bacterial infections as the cause of COPD exacerbation, but states low quality of evidence. In conclusion, PCT is not a secure way to discriminate bacterial versus viral cause of COPD exacerbation. Instead, investigators have moved on to find out if PCT can predict which patients need antibiotics during a COPD exacerbation. The rationale for this is that many COPD patients can carry bacteria in the airways, without these causing the current exacerbation (i.e., “innocent bystander”), and further that some bacterial infections are mild and may not need antibiotics, since these also carry side effects. Additionally, it is well-documented that low PCT levels effectively rule out bacteremia [51, 52].

PCT-Guided Antibiotic Treatment of AECOPD

In 2004, Christ-Crain et al. [53] reported on a cluster-RCT, in which an antibiotic-sparing strategy was tested. In general, patients with lower respiratory tract infections who had daily PCT measurements, had antibiotics discontinued, whenever the biomarker was below 0.25 ng/mL. This resulted in a noteworthy reduction in antibiotic use of approximately 50% without any apparent cost on treatment effect. Looking at the subgroup of patients with COPD exacerbation, the effect seemed even higher. The main criticism of the trial was, that it was cluster randomized on institutional level, and thus, that patients were not randomized on an individual patient level. Several individual-level classic RCTs have subsequently confirmed the results of the initial study regarding antibiotic reduction in patients with lower respiratory tract infections, in settings as diverse as primary care [54], departments of respiratory medicine [55], internal medicine, emergency medicine [56], and intensive care [57]. Although results have been very consistent regarding COPD exacerbations (high effect, 50–70% reduction in antibiotics use), when analyzed in subgroups, only few RCTs exist, in which the entire population have been patients with COPD exacerbation. In one RCT (n = 120) [55], the length of antibiotic administration in COPD patients with hospitalization-requiring exacerbation, was reduced from 8.5 days to 3.5 days, \( p < 0.001 \). Another trial included COPD patients with exacerbation, but did not report antibiotic use, rather re-exacerbation within 6 months, which did not seem

| Study type and author | Patient group | N | Primary outcome | Reduction in antibiotic exposure | Strengths/weaknesses | Conclusion |
|-----------------------|--------------|---|----------------|----------------------------------|----------------------|------------|
| RCT, Bafadhel et al. [37] | AECOPD | 109 | Noninferiority in the health status score and treatment failure | Reduced by 49% (no apparent risk) | Treatment based on single exacerbation blood eosinophil count | Successful reduction of corticosteroid exposure |
| RCT, Sivapalan et al. [41] | AECOPD (admitted) | 318 | DAOH14 | Reduced by 60% (no apparent risk) | Open-label | Successful reduction of corticosteroid exposure |

AECOPD, acute exacerbated chronic obstructive pulmonary disease; DAOH14, days alive and out of hospital within 14 days from recruitment; PCT, procalcitonin; RCTs, randomized controlled trials.

| Study type and author | Patient group | N | Primary outcome | Reduction in antibiotic exposure | Strengths/weaknesses | Conclusion |
|-----------------------|--------------|---|----------------|----------------------------------|----------------------|------------|
| RCT, Bafadhel et al. [37] | AECOPD | 109 | Noninferiority in the health status score and treatment failure | Reduced by 49% (no apparent risk) | Treatment based on single exacerbation blood eosinophil count | Successful reduction of corticosteroid exposure |
| RCT, Sivapalan et al. [41] | AECOPD (admitted) | 318 | DAOH14 | Reduced by 60% (no apparent risk) | Open-label | Successful reduction of corticosteroid exposure |

AECOPD, acute exacerbated chronic obstructive pulmonary disease; DAOH14, days alive and out of hospital within 14 days from recruitment; PCT, procalcitonin; RCTs, randomized controlled trials.
different in the randomized arms [58]. In a Chinese RCT, it was explored whether antibiotics improved the prognosis in patients with low (<0.1 ng/mL) PCT; the investigators did not observe any difference, but the use of antibiotics was reduced from 100% (95 of 95 in the antibiotic group) to 17.7% (17 of 96 in the control group) based on the low PCT [59].

The results from PCT-guided antibiotic-sparing regimes in patients with lower respiratory tract infections have been summarized in a meta-analysis of 26 RCTs with 6,708 patients who were individually randomized. The main result was that 30-day all-cause mortality was reduced in PCT-guided patients (286 [9%] of 3,336 PCT-guided patients vs. 336 [10%] in 3,372 controls; adjusted odds ratio 0.83 [95% CI 0.70–0.99], p = 0.037). Additionally, PCT guidance was associated with a 2.4-day reduction in antibiotic exposure (5.7 vs. 8.1 days [95% CI −2.71 to −2.15], p < 0.0001).

In conclusion, PCT guidance of antibiotic use in COPD patients with exacerbations is only examined directly in few and small trials. However, results from these trials are consistent with subgroup analysis from trials recruiting broader, but in which the COPD cohort could be analyzed separately. All these results regarding COPD exacerbations are also consistent with trial results from other patient groups with lower respiratory tract infections, so summarized, they seem solid, and consistently show a reduction in the use of unnecessary antibiotics in a magnitude of 50–70% of all administered antibiotics for these patients.

Such biomarker-guided strategies may prove key in reducing the antibiotic overuse, as summarized in a meta-

| Study type and author | Patient group | N   | Primary outcome | Reduction in antibiotic exposure PCT arm versus control | Strengths/weaknesses | Conclusion                      |
|-----------------------|---------------|-----|----------------|--------------------------------------------------------|----------------------|--------------------------------|
| RCT, Christ-Crain et al. [53] | LRTI (admitted) | 243 | →              | Reduced by 50% (no apparent risk)                      | Cluster randomized, not blinded | Successful reduction of antibiotic exposure in all subgroups including AECOPD |
| RCT, Corti et al. [55] | AECOPD (admitted) | 120 | →              | From 8.5 to 3.5 days (no apparent risk)                | n = 120, open-label | Successful reduction of antibiotics duration |
| Multicenter RCT, Schuetz et al. [56] | LRTI (emergency departments) | 1,359 | →              | From 8.7 to 5.7 days (no apparent risk)               | Heterogeneity in population (pneumonia, AECOPD, and bronchitis) | Successful reduction of antibiotics duration |
| Multicenter RCT, de Jong et al. [58] | Critically ill patients (ICU) | 1,575 | →              | Median consumption from 9.3 to 7.5 daily doses (no apparent risk) | Lower mortality in PCT arm | Successful reduction of antibiotic exposure |
| RCT, Verduri et al. [58] | Severe ECOPD (admitted) | 184 | No change in exacerbation of COPD in PCR-arm | 45 of 88 patients in PCR-arm had 7 days of antibiotics less than control (no apparent risk) | Underpowered (very few patients enrolled) | Successful reduction of antibiotics duration |
| RCT, Wang et al. [59] | AECOPD | 194 | Antibiotics no better than placebo for PCT <0.1 ng/mL | 100% for patients with PCT <0.1 ng/mL (no apparent risk) | Small n, did not look at PCT effect (but could report it) | Successful reduction of antibiotic exposure |
| Meta-analysis of RCTs, Schuetz et al. [60] | Primary care, emergency medicine, respiratory medicine, critically ill | 6,708 | All-cause mortality | 2.7 days reduction, p < 0.0001 | Not powered for all effects in all subgroups | Lower mortality, fewer antibiotic-related side effects, and lower consumption of antibiotics |

LRTI, lower respiratory tract infections; COPD, chronic obstructive pulmonary disease; ECOPD, exacerbated chronic obstructive pulmonary disease; AECOPD, acute exacerbated chronic obstructive pulmonary disease; ICU, intensive care unit; PCT, procalcitonin; RCTs, randomized controlled trials.
Biomarkers to Reduce Side Effects in Treatment of Lung Infections

analysis by Schuetz et al. [60], which is important when facing emerging antibiotic-resistant bacteria. See Table 2, in which some of the most important RCT results regarding PCT-guided antibiotic therapy are noted, including the main characteristics and results. Figure 3 shows a proposed algorithm for use when discontinuing antibiotics in COPD patients with exacerbations, as used in reference 55.

**Conclusion**

In conclusion, there is evidence at level 1B (one RCT) to support the use of blood eosinophils in reducing the use of systemic corticosteroids in patients admitted to hospital with a severe AECOPD; a strategy which is non-inferior regarding death and hospitalization length (days alive and out of hospital). Such a strategy seems applicable to implement in settings where a leukocyte differential count is available for daily analysis. Further, during admission of AECOPD, PCT measurements with approximately every 48 h can lead to a reduction in antibiotics of at least 30%, without any obvious disadvantages; in fact, such a strategy reduces mortality and antibiotic-related side effects in patients with lower respiratory tract infections (level 1A evidence), effects that seem likely also to apply to COPD patients.

**Acknowledgement**

We thank Mr. Valdemar Rømer for careful coordination and helping with the drafting of this manuscript.

**Conflict of Interest Statement**

The authors declare no conflicts of interest.

**Funding Sources**

P.S. was funded by the Danish Regions Medical Fund (grant no. 5894/16), the Danish Council for Independent Research (grant no. 6110-00268B), and Herlev-Gentofte University Hospital.

**Author Contributions**

Both authors contributed equally to the conception, design, and writing of the manuscript.

---

**Fig. 3.** PCT algorithm from [56]. PCT, procalcitonin.
Biomarkers to Reduce Side Effects in Treatment of Lung Infections

44 Muller B, White JC, Nylen ES, Snider RH, Weglohner W, Struck J, Fischer-Schulz C.

42 Becker KL, Nylén ES, White JC, Müller B, Birnbaum RS, Mahoney WC, Burns DM, Bafadhel M, Greening NJ, Harvey-Dunstan TC, Williams JE, Morgan MD, Brightling CE, et al. Blood eosinophils and outcomes in severe hospitalised exacerbations of COPD. Chest. 2016 Aug;150(2):320–8.

40 Serafino-Agrusa L, Scichilone N, Spatafora M, Battaglia S. Blood eosinophils and treatment response in hospitalised exacerbations of chronic obstructive pulmonary disease: a case-control study. Pulm Pharmacol Ther. 2016 Apr;37:89–94.

38 Duman D, Aksoy E, Agca MC, Kocak ND, Ozmen I, Akturk UA, et al. The utility of inflammatory markers to predict readmissions and mortality in COPD cases with or without eosinophilia. Int J Chron Obstruct Pulmon Dis. 2015;10:2469–78.

37 Mohamed N, Masselink R, montage NG, Otto A, Bohuon C, et al. Eosinophilia and mortality in COPD cases with or without eosinophilic inflammation. J Innate Immun. 2022;14:89–97.

DOI: 10.1159/000517161