Serum cytokine profiling and enrichment analysis reveal the involvement of immunological and inflammatory pathways in stable patients with chronic obstructive pulmonary disease

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Abstract: Chronic obstructive pulmonary disease (COPD) is a major global health problem. It results from chronic inflammation and causes irreversible airway damage. Levels of different serum cytokines could be surrogate biomarkers for inflammation and lung function in COPD. We aimed to determine the serum levels of different biomarkers in COPD patients, the association between cytokine levels and various prognostic parameters, and the key pathways/networks involved in stable COPD. In this study, serum levels of 48 cytokines were examined by multiplex assays in 30 subjects (control, n=9; COPD, n=21). Relationships between serum biomarkers and forced expiratory volume in 1 second, peak oxygen uptake, body mass index, dyspnea score, and smoking were assessed. Enrichment pathways and network analyses were implemented, using a list of cytokines showing differential expression between healthy controls and patients with COPD by Cytoscape and GeneGo Metacore™ software (Thomson-Reuters Corporation, New York, NY, USA). Concentrations of cutaneous T-cell attracting chemokine, eotaxin, hepatocyte growth factor, interleukin 6 (IL-6), IL-16, and stem cell factor are significantly higher in COPD patients compared with in control patients. Notably, this study identifies stem cell factor as a biomarker for COPD. Multiple regression analysis predicts that cutaneous T-cell-attracting chemokine, eotaxin, IL-6, and stem cell factor are inversely associated with forced expiratory volume in 1 second and peak oxygen uptake change, whereas smoking is related to eotaxin and hepatocyte growth factor changes. Enrichment pathways and network analyses reveal the potential involvement of specific inflammatory and immune process pathways in COPD. Identified network interaction and regulation of different cytokines would pave the way for deeper insight into mechanisms of the disease process.

Keywords: COPD, Bio-Plex assay, biomarkers, pathways, networking

Introduction
Chronic obstructive pulmonary disease (COPD) is a major global health problem. In 2002, it was the fourth worldwide leading cause of death, and it is anticipated to be the third leading cause of death by 2030.1 COPD results from chronic inflammation and eventual irreversible damage to the airways. Several factors, including exposure to environmental pollution and smoking, contribute to the airway damage and obstruction.2 Inflammatory response in COPD is characterized by the increased number of macrophages, neutrophils, and cytotoxic T lymphocytes in the airways and lung parenchyma. Airway inflammation further results in systemic inflammation and other COPD-related manifestations.3,4
Spirometry (forced expiratory volume in 1 second; FEV₁), exercise tests (peak oxygen uptake; VO₂), and dyspnea scores (a scale to determine the degree of difficulty in breathing) are used for diagnosing and determining the prognostic factors of COPD. These tests are also used for evaluating the treatment efficacy in COPD patients. However, the test results do not fully reflect the burden of COPD. Aaron et al have studied 19 serum markers and found that only C-reactive protein, myeloperoxidase, and vascular endothelial growth factor show intra- and interpatient reliability and correlation with disease severity. Franciosi et al performed a meta-analysis by considering the spirometric, demographic, clinical, cytological, and biochemical variables in stable COPD patients. Their meta-analysis indicates that the concentrations of tumor necrosis factor (TNF) and C-reactive protein have suggestive correlation with disease severity. Further, the existence of high variability among biomarker levels in different studies has also been observed. Therefore, additional studies with a well-defined group of patients are needed to identify the sets of potential biomarkers and pathways involved in COPD from different populations.

Cytokines play a critical role in COPD-associated inflammation. Interactions among these cytokines are complex, as some cytokines can have different actions under a specific context. Thus, studying a panel of cytokine biomarkers, their correlation with pulmonary physiological functions and systemic parameters and the involvement of various pathways and networks would provide a better understanding of the mechanisms underlying COPD. This will also help to identify a nonredundant set of biomarkers for COPD. Measuring cytokine panels from the same samples in multiplex assays will be advantageous for reducing technical variability. Only a limited number of studies have used panels of multiple serum cytokines to determine biomarkers in COPD patients simultaneously. Furthermore, no studies with multiple biomarkers and pathway analysis have been reported on COPD patients of Indian descent.

In the present study, we measured the serum levels of 48 cytokines in stable COPD and healthy subjects by immunobead Bio-Plex assays (Bio-Rad Laboratories Inc., Hercules, CA, USA) and analyzed their association with FEV₁ (percentage predicted), maximum oxygen uptake (peak VO₂ predicted), smoking index, dyspnea scores, and body mass index (BMI) by multiple regression analysis. Furthermore, our study identifies that changes in a few key cytokine concentrations explain the changes in lung function-related parameters (FEV₁ and peak VO₂ and smoking index). We also assessed some of the key biomarkers and their common regulatory factors involved in inflammatory and immunological pathways in COPD.

Materials and methods

Ethics statement

This study was approved by the Ethics Committee of All India Institute of Medical Sciences, New Delhi, India. Prior informed written consent was obtained from all participating healthy and COPD volunteers. Blood work and spirometric tests were conducted under the guidelines of the ethics committee.

Patient selection

COPD patients were diagnosed according to the guidelines of Global Initiative for Chronic Obstructive Lung Disease (GOLD) criteria. They were either current smokers with COPD or ex-smokers with COPD. Individuals older than 40 years were recruited for this study and were categorized into two main groups: Group 1 included healthy controls with no evidence of COPD (n=9); four and five participants were smokers and nonsmokers, respectively, and group 2 included patients diagnosed with COPD (n=21); eleven of whom were ex-smokers with COPD and ten of them were current smokers with COPD. All the COPD patients selected for the study were in stable condition (no exacerbation for at least 1 month). Participants were excluded specifically when they had a history of COPD exacerbation within the last 1-month period, asthma, or were taking oral steroids or had any conditions that were associated with inflammation such as infection, cancer, congestive heart failure, end-stage renal disease, or connective tissue disorders that could affect the levels of COPD biomarkers.

Measurement of pulmonary functions

FEV₁ (percentage predicted) and FVC (forced vital capacity) were measured with a calibrated spirometer (Spiro Air, Medisoft; PK Morgan Ltd., Kent, United Kingdom), using standard methodology at the time of enrollment of the subjects. This pulmonary function test value is critical for diagnosing obstructive and restrictive lung diseases.

Calculation of BMI

BMI is an independent prognostic factor for disease severity and survival after the diagnosis of COPD. BMI is a measure of human body shape/obesity based on an individual’s weight and height. Each body mass index score was calculated by the following formula: BMI = weight (kg)/(height [m])².
Symptom-limited incremental exercise test

In patients with COPD, a symptom-limited incremental step test elicited the maximum cardiopulmonary and metabolic responses. A symptom-limited incremental exercise test was performed on a bicycle ergometer, using a 10 W ramp protocol. Exercise was performed while seated on an electronically braked cycle ergometer (Corival; Lode, Groningen, the Netherlands). Heart rate was monitored continuously throughout the protocol using a heart rate monitor (POLAR [Polar Electro Oy, People's Republic of China]). The polar device used in this study had a T31 transmitter and a chest belt (N2965) to transmit heart rate to the system. Breath-by-breath measurements of oxygen consumption (VO₂) were made by custom software (vacumed/vista MX [VacuMed, Ventura, CA, USA]). Standard 12-lead electrocardiograms were obtained at rest, during exercise and in recovery phase. Blood pressure was monitored using a standard cuff sphygmomanometer at rest, every 3 minutes during exercise, and every 2 minutes during recovery. At the end of exercise, the reasons for termination of exercise were obtained from the patient. The VO₂ max was the highest VO₂ observed during exercise. The exercise was also terminated at exhaustion (intolerable dyspnea, as indicated by the patient); severe desaturation (peripheral capillary oxygen saturation <80%); demand by the patient because of leg cramps, chest pain, or discomfort or request by patients for any other reason.

Evaluation of dyspnea

Dyspnea is one of the common symptoms in COPD either at rest or under conditions of exercise, and this applies to all severities. Dyspnea severity was assessed and scored using the standard Modified Medical Research Council questionnaire.  

Measurements of blood markers

A volume of 5 mL of whole blood was collected into a plain tube and allowed to clot for 1 hour. The sera samples were aliquoted after centrifugation at 1,000 × g for 10 minutes and stored at −80°C until further analysis. Concentrations of serum cytokines were analyzed using two complementary Bio-Plex suspension array systems (Bio-Plex Pro Human Cytokine Group 27-Plex Panel and Bio-Plex Pro Human Cytokine Group 21-Plex Panel) to cover the range of all the cytokine biomarkers potentially involved in the pathophysiology of COPD. Forty-eight biomarkers were assessed simultaneously, using the Bio-Plex system. The selection of specific cytokines in the study was based on the previously available reports and the involvement of these cytokine biomarkers in COPD. Assays were performed in duplicate by following the standard operating protocol provided by the Bio-Plex Multiplex cytokine assay.

Briefly, anticytokine antibody-conjugated beads were added to individual wells of a 96-well filter plate and adhered using vacuum filtration. After washing, 50 µL prediluted standards and serum samples were added into respective wells, and the filter plates were shaken at 300 rpm for 30 minutes at room temperature. Thereafter, the filter plates were washed and 25 µL prediluted multiplex biotin-conjugated detection antibody was added and incubated for another 30 minutes. After washing, 50 µL prediluted streptavidin-conjugated PE was added for 10 minutes, followed by an additional wash and the addition of 125 µL Bio-Plex assay buffer to each well. Then, filter plates were analyzed using the Bio-Plex Protein Array System, and concentrations of each cytokine were determined using software (Bio-Plex Manager version 6.0). Standard curves were generated for each biomarker. The line of best fit was determined for standard curves by standard recovery methods and by calculating the concentration of each standard.

Statistical analysis

Cytokines’ levels in sera showed no differences between smokers and nonsmokers within the control group. There were also no statistical differences detected between ex-smokers with COPD and current smokers with COPD. Therefore, we compared the pooled samples (healthy controls versus COPD), using the two-tailed Mann–Whitney test, as outlined in the experimental design (Figure S1). The association of identified serum biomarkers with FEV₁ (percentage predicted), peak VO₂ (predicted), smoking index, dyspnea scores, and BMI were studied in all subjects, using Spearman correlation analysis. In addition, a step-wise multiple regression analysis with weighted regression equations has also been used for calculating coefficients for FEV₁ (percentage predicted), peak VO₂ (predicted), smoking index, dyspnea scores, and BMI with serum biomarkers. Multiple regression analysis determines the relationship between a dependent or criterion variable of interest (Y) and a set of K independent variables or potential predictor variables (X₁, X₂, X₃, …, Xₖ), where the scores on all variables are measured for n number of cases. A multiple regression equation for predicting Y is expressed as \( Y = a + (b_1 X_1) + (b_2 X_2) + \ldots + b_k X_k \). A stepwise multiple regression analysis eliminated the parameters that provided insignificant contribution to the model. The
coefficient for each factor was obtained from the final model with the lowest $P$-value. Calculated and observed values were plotted in a regression line after calculating a common factor for each equation. All the statistical analyses were performed using the Prism version 5 (GraphPad Software, Inc., La Jolla, CA, USA) and SPSS (version 16; SPSS Inc., Chicago, IL, USA) software.

**Enriched pathways and network analysis**

The mean fold changes of the detected biomarkers were calculated between COPD and control subjects. Cytokines showing 1.5-fold changes were used for biological processes, pathways, and network enrichment analyses. The purpose of this analysis was to identify the enriched biological processes and the pathways associated with COPD. Interaction networks among the significantly elevated cytokines in COPD were constructed using Cytoscape and GeneGo Metacore™ software (Thomson Reuters, St Joseph, MI, USA).

**Results**

**Biomarkers CTACK, eotaxin, HGF, IL-6, IL-16, and SCF are elevated in the sera of stable COPD patients**

We assessed 48 candidate biomarkers in the serum samples of healthy control and COPD subjects by Bio-Plex assays. Of these, the concentrations, in picograms per milliliter, of 29 cytokines were measurable. The concentrations of the rest of the 19 cytokines were below the detection limit. Mean comparison procedures show that serum levels of six biomarkers (cutaneous T-cell attracting chemokine [CTACK], eotaxin, hepatocyte growth factor [HGF], interleukin 6 [IL-6], IL-16, and stem cell factor [SCF]; $P<0.05$) are significantly higher in COPD patients compared with control subjects (Figure 1). The data of cytokines not showing significant differences between the groups and the rest of the biomarkers tested in the study are listed in Table S1.

![Figure 1](https://www.dovepress.com/)

*Figure 1* Serum concentrations of cutaneous T-cell attracting chemokine, eotaxin, hepatocyte growth factor, interleukin 6 (IL-6), IL-16, and stem cell factor are elevated in chronic obstructive pulmonary disease patients compared with control subjects. Serum level of (A) cutaneous T-cell attracting chemokine ($P<0.01$), (B) eotaxin ($P<0.05$), (C) hepatocyte growth factor ($P<0.05$), (D) IL-6 ($P<0.05$), (E) IL-16 ($P<0.01$), and (F) stem cell factor ($P<0.05$) are significantly higher in chronic obstructive pulmonary disease patients ($n=21$) compared with control subjects ($n=9$). Significant differences between chronic obstructive pulmonary disease and control subjects are denoted by $^\ast P<0.05$ and $^\ast\ast P<0.01$, as measured by two-tailed Mann–Whitney test. The data are represented as mean ± standard error.
CTACK is a T-cell chemokine, whereas eotaxin is a potent chemoattractant for eosinophils and basophils. HGF is a multifunctional mitogen. IL-6 is a permeability-increasing inflammatory cytokine released mainly by macrophages and epithelial cells, and IL-16 is a proinflammatory cytokine released by bronchial epithelium and dendritic cells, cluster of differentiation (CD)8+ and CD4+ T cells. Hematopoietic factor SCF is secreted by various stroma and inflammatory cells. Collectively, all these cytokines are important for recruiting different types of immune cells, cell proliferation, and tissue repair. The cytokines/biomarkers identified by these analyses are highly relevant to COPD and act as putative candidate biomarkers of COPD patients.

**Eotaxin is negatively correlated with FEV1, whereas CXCL1 is positively correlated with HGF.** To determine the key biomarkers affecting FEV1 (percentage predicted), peak VO2 (predicted), smoking, dyspnea score, and BMI in COPD patients, bivariate correlation analysis was performed. All recruited subjects had complete demographic data, and as expected by the study design, COPD patients have lower FEV1 (percentage predicted) (mean ± standard error: 48.7%±3.9%) compared with age-matched controls (mean ± standard error: 83.6%±1.7%; P<0.001; Table 1). Bivariate analysis of cytokines with each parameter shows that eotaxin concentrations have a strong inverse relationship with FEV1 (percentage predicted; Y=0.106x + 87.08; r=-0.71; P≤0.05; Figure 2A). These data suggest that eotaxin could be one of the important contributors to the decaying lung function. Notably, chemokine (C-X-C motif) ligand 1 (CXCL1) or the growth-related oncogen α (GRO-α) show a strong positive correlation with HGF (Y=8.675x + 124.77; r=0.76; P<0.05; Figure 2B); therefore, CXCL1 and HGF are likely to be dependent variables and/or affected by the same transcription factor. CXCL1 shows statistically significant but low levels of positive correlation with IL-6 (r=0.23; P<0.05) and an inverse relationship with dyspnea (r=-0.53; P<0.05). CXCL1 is a chemoattractant factor for monocytes and neutrophils. This protein is produced by a variety of cells including monocytes, endothelial cells, and fibroblasts. Therefore, CXCL1 may be involved in regulating various components of COPD.

**Table 1 Demographic characteristics of the participants in the study**

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Healthy controls</th>
<th>Patients with COPD</th>
<th>P-values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sample size</td>
<td>9</td>
<td>21</td>
<td>–</td>
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<tr>
<td>Age, years</td>
<td>67.0 (58.5–73.5)</td>
<td>65.5 (53.0–76.0)</td>
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<tr>
<td>Forced expiratory volume in 1 second,</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>% predicted</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Peak oxygen uptake</td>
<td>–</td>
<td>64.0 (44.0–81.0)</td>
<td>–</td>
</tr>
<tr>
<td>Dyspnea score</td>
<td>2.0 (1.0–3.0)</td>
<td>–</td>
<td></td>
</tr>
<tr>
<td>Body mass index</td>
<td>22.0 (19.5–24.5)</td>
<td>22.5 (19.5–23.5)</td>
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<tr>
<td>BODE-index</td>
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<tr>
<td>Smoking history or status†</td>
<td>11.1 (10.3–14.1)</td>
<td>18.0 (14.0–23.5)†</td>
<td>0.08</td>
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</tbody>
</table>

*Notes:* †All the values are given as in median (interquartile range); ‡pack years; §only for smokers; ¶healthy controls have no evidence of COPD (n=9); †four and five subjects were smokers and nonsmokers, respectively. Patients with COPD (n=21); eleven of them are ex-smokers with COPD, whereas ten of them are current smokers with COPD.

*Abbreviations:* COPD, chronic obstructive pulmonary disease; BODE, Body mass index, degree of airflow Obstruction and Dyspnea, and Exercise capacity.
Table 2 Multiple regression analysis between studied physiological parameters and cytokines showing significant differences between controls and patients with COPD

<table>
<thead>
<tr>
<th>Parameters and predictors</th>
<th>Coefficient</th>
<th>SE</th>
<th>P-value*</th>
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<tr>
<td>Forced expiratory volume in 1 second, % predicted</td>
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<td>Eotaxin</td>
<td>−0.20</td>
<td>0.02</td>
<td>0.05</td>
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<td>Interleukin 6</td>
<td>−0.37</td>
<td>0.44</td>
<td>0.04</td>
</tr>
<tr>
<td>Stem cell factor</td>
<td>−0.22</td>
<td>0.03</td>
<td>0.05</td>
</tr>
<tr>
<td>Peak oxygen uptake (predicted)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Eotaxin</td>
<td>−0.46</td>
<td>0.01</td>
<td>0.04</td>
</tr>
<tr>
<td>Cutaneous T-cell attracting chemokine</td>
<td>−0.44</td>
<td>0.02</td>
<td>0.05</td>
</tr>
<tr>
<td>Smoking history</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Eotaxin</td>
<td>+0.45</td>
<td>0.03</td>
<td>0.04</td>
</tr>
<tr>
<td>Hepatocyte growth factor</td>
<td>−0.10</td>
<td>0.01</td>
<td>0.05</td>
</tr>
</tbody>
</table>

Note: *A P-value ≤0.05 is considered to represent a statistically significant factor.

Abbreviations: COPD, chronic obstructive pulmonary disease; SE, standard error.

Increase in CTACK, eotaxin, IL-6, and SCF significantly contributes to the reduction in FEV₁ and peak VO₂

To further determine the contribution of all the cytokines to each of these parameters, we conducted a stepwise multiple regression analysis. This systematic analysis identified that eotaxin, IL-6, and SCF are statistically significantly, contributing to the reduction in lung function measured by FEV₁ (percentage predicted). Contributions of each parameter are described by the following equation: \( Y = (-0.02 \times \text{eotaxin}) + (-0.37 \times \text{IL-6}) + (-0.22 \times \text{SCF}) \) (Table 2; \( P < 0.05 \)). A constant factor of −1.92 converts the Y values equivalent to observed FEV₁ (percentage predicted; Figure 3A; \( r = -0.71; P < 0.01 \)). These analyses suggest that an increase in serum concentrations of CTACK, eotaxin, and SCF reflects the decrease in FEV₁ observed in the present study, and notably, these three cytokines are sufficient to explain most of the changes seen in FEV₁.

Furthermore, eotaxin and CTACK significantly contribute to the observed reduction in peak VO₂ (predicted). Contributions of these two cytokines to peak VO₂ are described by the following equation: \( Y = (-0.44 \times \text{CTACK}) + (-0.46 \times \text{eotaxin}) \) (Table 2; \( P < 0.05 \)). A constant factor of −9.1 converts the Y values equivalent to measured peak VO₂ (predicted; Figure 3B; \( r = -0.70; P < 0.01 \)). These analyses suggest that increase in CTACK and eotaxin concentrations is related to decrease in peak VO₂ observed in these individuals. Once again, these two cytokines are sufficient to explain most of the changes observed in peak VO₂. Collectively, these results show that the four key cytokines (CTACK, eotaxin, IL-6, and SCF) explain the significant

Figure 3 Cutaneous T-cell attracting chemokine, eotaxin, hepatocyte growth factor, interleukin 6 (IL-6), and stem cell factor concentrations estimate the changes in forced expiratory volume in 1 second (FEV₁; percentage predicted), peak oxygen uptake (VO₂; predicted), and smoking status by multiple regression analysis. (A) Stepwise multiple regression analysis identified that the variables eotaxin, IL-6, and stem cell factor significantly contribute to the changes in FEV₁ (percentage predicted). The values calculated on the basis of these three cytokines correlate well with the changes observed in FEV₁ values. (B) Similar analysis shows that cutaneous T-cell attracting chemokine and eotaxin significantly contribute to the changes in the peak VO₂. The values calculated based on these two cytokines correlate well with the changes observed in peak VO₂. (C) The smoking indices are related to eotaxin and hepatocyte growth factor, and the values correlate well with the predicted and observed smoking indices.
changes in lung function-related parameters observed in the study population.

Smoking affects the changes in eotaxin and HGF concentrations

To determine the effect of smoking on the changes in cytokine concentrations, we have conducted a stepwise multiple regression analysis of the biomarkers, as explained earlier. This analysis shows that changes in the sera concentrations of eotaxin and HGF are associated with the smoking levels (pack year): Y=(0.45×eotaxin) + (−0.10×HGF) (Table 2; P<0.05). A constant factor of 12.6 converts the Y values equivalent to the observed smoking index (Figure 3C; r=0.58; P<0.05). These data suggest that smoking partially contributes to the increase in eotaxin and decrease in HGF concentrations observed in the sera of smokers in this study population.

Specific immune and inflammatory response pathways are involved in stable COPD

Complex interactions among cytokines, chemokines, and growth factors regulate various immune and inflammatory responses. To determine the pathways that are potentially regulated by these cytokines in COPD, we have conducted enriched pathway analyses. Of the 29 measurable biomarkers, 16 showed higher concentrations (≥1.5 upregulated), whereas two showed lower concentrations (downregulated, ≤1.5) in COPD patients compared with in healthy controls. Pathway enrichment analysis was conducted using these cytokines by Metacore™ platform, with a priori setting of a cutoff threshold (False Discovery Rate (FDR)p, probability value =0.05). Pathways and biological processes regulated by these cytokines include immune responses (mediated by T-helper cell differentiation, riggering receptor expressed on myeloid cells 1 (TREM1) signaling, prostaglandin E2 signaling, and Th17 [T helper 17 cells]-derived cytokines), inflammation mediated by JAK/STAT (Janus kinase/signal transducers and activators of transcription) pathway, histamine signaling), lymphocyte proliferation and regulation, and leucocyte chemotaxis (Figure 4). Therefore, this analysis suggests the involvement of immune response-related pathways in COPD.

GCR-α, NF-κB and C/EBP-β regulate the major cytokine networks in COPD

Specific transcription factors and regulators determine cytokine response.29 To determine the transcription factors that regulate the key cytokines involved in COPD, we have conducted a network analysis. Thirteen of the 18 cytokines (nodes) generated a large single regulatory network. Three main hubs NF-κB (nuclear factor kappa-light-chain-enhancer of activated B cells), C/EBP-β (CCAAT/enhancer-binding protein beta), and GCR-α (glucocorticoid receptor alpha) regulate these networks (Figure 5). These enriched networks mainly affect mediators involved in immune, inflammatory, and chemotaxis responses. Interconnecting cytokine biomarkers including eotaxin, HGF, IL-6 and SCF are induced either by two or three hubs. Notably, GCR-α directly regulates all four of these cytokines, which are significantly upregulated in stable COPD patients. Therefore, GCR-α is likely a major potential regulator of these cytokine biomarkers in COPD.

Discussion

Serum biomarkers and their effects on COPD-related parameters and pathways that affect COPD have not been fully established. We have assessed the levels of 48 serum biomarkers in COPD patients of Indian descent, using multiplex immunobead-based assays. We have identified some of the key biomarkers involved in pathways associated with stable COPD. In particular, the serum levels of CTACK, eotaxin, HGF, IL-6, IL-16 and SCF are significantly higher in patients with COPD compared with in control subjects (Figure 1). These cytokine levels are expected to reflect long-term effects because our study focuses on stable COPD patients with no exacerbation during the previous 1-month period. To the best of our knowledge, this is the first study that identified an increased serum level of SCF in COPD patients compared with healthy controls. Eotaxin, IL-6, and SCF inversely contribute to the changes in FEV1 (percentage predicted), whereas eotaxin and HGF inversely contribute to the changes in peak VO2 (predicted). Smoking is related to increased eotaxin and decreased HGF concentrations. CXCL1 is positively correlated with IL-6 and HGF, but it is negatively related to dyspnea. Enrichment pathways and network analyses revealed the involvement of inflammatory, chemotaxis, and immune response-related pathways in COPD.

Previously, studies have reported that TNFα can contribute to COPD pathogenesis. Our data show a trend toward increased levels of TNFα in COPD patients (21.1–112.6 pg/mL) compared with in controls (21.1–34.1 pg/mL), but the values are not statistically different (Table S1). This could be a result of a lower level of TNFα in stable COPD patients or because of lower statistical power to detect the change in
Bade et al

this cytokine. Regulatory network analysis showed overlapping connectivity among eotaxin, HGF, IL-6, and SCF via C/EBP-β, NF-κB, and GCR-α transcription factors. In particular, GCR-α is involved in the regulation of all four of these key cytokines in stable COPD patients. The involvement of these cytokines (eotaxin, HGF, IL-6 and SCF) in various pathways and networks indicates their potential regulatory roles in COPD.

CTACK belongs to the C-C motif chemokines family, which plays an important role in the immune-inflammatory processes in many skin diseases. CTACK is constitutively produced by epidermal keratinocytes and participates in

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**Figure 4** Pathway enrichment analysis reveals specific immune and inflammatory response pathways in chronic obstructive pulmonary disease. (A) The top most significant enriched biological processes and pathways including immune responses, inflammation, lymphocyte proliferation and regulation, and leukocyte chemotaxis are represented along with the coenrichment P-value (log scale) in the bar graph. (B) The numbers and names of gene sets overlapping with the significant pathways and processes are shown in the table format.

**Abbreviations:** PGE2, prostaglandin E2; IL-2, interleukin 2; IFN, interferon; GM-CSF, granulocyte-macrophage colony-stimulating factor; CCL2, chemokine ligand 2; SOS1, son of sevenless homologue 1; VEGF, vascular endothelial growth factor; MIP-1, macrophage inflammatory protein 1; HGF, hepatocyte growth factor; TREM1, triggering receptor expressed on myeloid cells 1; MIG, monokine induced by gamma-interferon; IP10, interferon gamma-induced protein 10; MGF, mechano growth factor; STAT, signal transducers and activators of transcription; G-CSF, granulocyte colony-stimulating factor; Th17, T helper 17.
Increased level of CTACK present in COPD participants compared with controls (Figure 1A) suggests the involvement of CTACK in inflammatory processes related to the disease. In a study involving 48 patients, Pinto-Plata et al have also noted that CTACK levels are elevated in COPD subjects by protein microarray analysis. Our study further shows that CTACK negatively affects peak VO\textsubscript{2} (Table 2). Therefore, including CTACK in future studies would be useful to confirm the relevance of this marker in other COPD populations.

Eotaxin is a potent chemoattractant for human eosinophils and basophils. Our study shows that serum level of eotaxin in stable COPD patients is elevated (Figure 1B). The COPD patients with eosinophil phenotypes show good response to corticosteroids. D’Armiento et al have reported that eotaxin levels in plasma and bronchoalveolar lavage can predict rapid deterioration in lung function (FEV\textsubscript{1}) in COPD patients. Similarly, our study shows an inverse relationship between serum eotaxin levels and FEV\textsubscript{1} (percentage predicted), as well as peak VO\textsubscript{2} in the COPD subjects of Indian descent (Figure 3A and Table 2). Furthermore, smoking is positively correlated with eotaxin, suggesting that smoking potentially contributes to the increase of this cytokine. A previous study reports that smoking increases eotaxin levels in asthmatic subjects, but the differences in eotaxin levels are not clearly established in COPD subjects. Pathway enrichment analysis further shows that transcription factors NF-κB and GCR-α upregulate this cytokine production. Therefore, eotaxin is an important cytokine in COPD.
HGF is a multifunctional heterodimeric protein with mitogenic properties. HGF is synthesized by fibroblasts, macrophages, smooth muscle cells, and epithelial cells, and it is an important cytokine for lung development and repair. HGF concentration is higher in COPD patients compared with control subjects (Figure 1C). Smoking is negatively correlated with HGF (Figure 3C and Table 2), which could affect the repair process. Notably, the human HGF gene has an IL-6 response element, and therefore, HGF is expected to be upregulated during inflammation. Collectively increased HGF identified in these patients may reflect the attempt by the host to repair the damaged lung.

IL-6 is a classical marker for inflammation and is associated with many inflammatory disorders. An increased concentration of IL-6 detected in stable COPD patients (Figure 1D) is consistent with previous studies showing the increased level of this cytokine in the plasma of COPD patients. Monocytes of COPD patients are known to overreact and release more IL-6 than cells from normal subjects. IL-6 is negatively related to FEV$_1$ (percentage predicted; Figure 3A and Table 2). Therefore, IL-6 is an important cytokine that could affect systemic inflammation and worsen COPD comorbidity.

IL-16 is a proinflammatory biomarker released by bronchial epithelium and dendritic cells, CD8$^+$ and CD4$^+$ T cells. It acts as a key element in COPD pathobiology. IL-16 secreted by CD8$^+$ cells is involved in regulating the recruitment and activity of CD4$^+$ cells. The present study shows an increase in serum level of IL-16 in the COPD subjects (Figure 1E), reflecting its potential involvement in the inflammatory process of the studied patients. Pinto-Plata et al have studied the serum biomarkers related to inflammation and injury in COPD patients and observed that the levels of the inflammatory markers, including IL-6 and IL-16, were higher in more advanced disease. The present finding is consistent with the involvement of these proinflammatory cytokines in the regulation of the disease process.

SCF is a hematopoietic factor. Various stroma and inflammatory cells express this factor. SCF stimulates mast cells and induces these cells to adhere to extracellular matrix, and lead to the production of proinflammatory cytokines and chemokines in the tissue. For the first time, we have identified that COPD patients have higher levels of SCF than control subjects (Figure 1F). This finding suggests that the SCF could directly remodel airway tissues in these patients. Dolgachev et al have shown that SCF, along with IL-31, is involved in remodeling and fibrosis of airways by promoting recruitment and differentiation of bone marrow-derived fibroblast precursors in mice sensitized and chronically challenged with allergens. In patients with diffused interstitial lung fibrosis, alveolar fibroblasts secrete high levels of SCF. COPD is also characterized by fibrosis and remodeling of airways; hence, SCF is likely to be a key factor involved in permanently altering lung architecture in COPD.

CXCL1, or GRO-$\alpha$ is a chemotactic factor produced by a variety of cells including monocytes, endothelial cells, and fibroblasts. Monocytes and neutrophils from patients with COPD show enhanced chemotactic response to CXCL1. Induced sputum of COPD patients has increased levels of CXCL1. CXCL1 level shows negative correlation with HGF (Figure 3B) and positively correlates with IL-6 and dyspnea scores. The identified key cytokines/biomarkers may be involved in the regulation of the pathophysiology of the disease.

The emerging field of network and pathways analysis can help identify the relationships between systemic inflammatory biomarkers and disease state. The common distinct functional and regulatory pathways identified in the present study are T$_{fi}$ cell differentiation, TREM1 signaling, prostaglandin E2 signaling, Th1 7-derived cytokines, inflammation (mediated by the JAK-STAT pathway, histamine signaling), and leukocyte chemotaxis (Figure 4). One study shows that prostaglandin E2 is important for the mucosal innate immunity of COPD patients. As evident from Figure 5, the network comprises three main seeding nodes (NF-$\kappa$B, C/EBP-$\beta$ and GCR-$\alpha$), which regulate inflammation and immune regulation. Many of the differentially expressed cytokines are regulated by any of these three nodes, and of them, IL-6, HGF, eotaxin and SCF are significantly higher in COPD subjects. A fuller understanding of these processes and pathways will provide a better panel of COPD serum biomarkers and help us better understand the disease.

One of the limitations of the study is the use of a relatively small sample size. Although 29 stable COPD patients who were selected with stringent exclusion criteria helped us identify key cytokines that are sufficient to explain most of the variability in FEV$_1$, and certain other parameters, and to identify pathways and networks, the cohort is not large enough to identify all of the potential biomarkers. This patient cohort is also not large enough to test comorbidities and to fully determine the effect of smoking on COPD biomarkers. Further studies with large sample size are necessary to validate these biomarkers, and molecular/cell biology-based studies are necessary to determine the mechanisms regulated by these cytokines in COPD.
Conclusion
In the present study, six biomarkers (CTACK, eotaxin, HGF, IL-6, IL-16 and SCF) are significantly elevated in the sera of COPD patients compared with healthy controls. Multiple regression analysis shows that eotaxin, IL-6, and SCF are inversely correlated with FEV1. CTACK and eotaxin are negatively related to peak VO2. Smoking may increase eotaxin and decrease HGF. Investigated enriched pathways and networks suggest involvement of inflammatory JAK-STAT and histamine signaling, immune response mediated by TREM1 and prostaglandin E2 signaling and the regulation of lymphocyte proliferation in the progression and pathogenesis of COPD. Notably, the four cytokines IL-6, HGF, eotaxin and SCF that are higher in COPD showed overlapping interactions with the seeding nodes. The involvement of these cytokines in various pathways and network analyses indicates their potential regulatory roles in the disease process.

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Author contributions
AT, GB and RG developed the concept behind the study. GB, AKS, PK, and KKS designed the study and carried out the experiments with the help of MAK. GB, MAK, NP and AT designed the analysis approach and performed the data analysis, pathways analysis, interpretations and wrote the manuscript. All authors helped in manuscript revision and approved the final manuscript.

Disclosure
The authors report no conflicts of interest in this work.

References


Supplementary material

Figure S1 Flow diagram of the experimental design showing overall aim and analysis steps involved in the present study.

Abbreviations: COPD, chronic obstructive pulmonary disease; FEV\textsubscript{1}, forced expiratory volume in 1 second; VO\textsubscript{2}, maximum oxygen uptake; BMI, body mass index; GOLD criteria, Global Initiative for Chronic Obstructive Lung Disease.
### Table S1 Concentrations (pg/mL) of biomarkers detected in the sera samples among groups

<table>
<thead>
<tr>
<th>Biomarkers (pg/mL)</th>
<th>Non-smokers controls, n=5</th>
<th>Smokers controls, n=4</th>
<th>Ex-smokers with COPD, n=11</th>
<th>Smokers with COPD, n=10</th>
</tr>
</thead>
<tbody>
<tr>
<td>MIG</td>
<td>907 (697–1,451)</td>
<td>2,324 (1,874–3,174)</td>
<td>5,201 (1,401–7,699)</td>
<td>3,004 (1,738–5,413)</td>
</tr>
<tr>
<td>IL-4</td>
<td>3.4</td>
<td>3.4</td>
<td>4.0</td>
<td>2.9</td>
</tr>
<tr>
<td>IL-13</td>
<td>(3.4–4.4)</td>
<td>(3.1–4.0)</td>
<td>(3.3–6.2)</td>
<td>(1.3–4.1)</td>
</tr>
<tr>
<td>SCGFβ</td>
<td>17,831 (7.0–16.7)</td>
<td>12,218 (13.8–21.8)</td>
<td>26,283 (12.6–19.1)</td>
<td>18,200 (8.2–15.0)</td>
</tr>
<tr>
<td>IL-2</td>
<td>7.3</td>
<td>6.7</td>
<td>11.8</td>
<td>11.2</td>
</tr>
<tr>
<td>GM-CSF</td>
<td>24.1</td>
<td>9.9</td>
<td>49.8</td>
<td>52.1</td>
</tr>
<tr>
<td>RANTES</td>
<td>(6.3–9.8)</td>
<td>(4.8–8.5)</td>
<td>(6.9–15.1)</td>
<td>(8.4–14.5)</td>
</tr>
<tr>
<td>CXCL1 or GRO-α</td>
<td>47.0</td>
<td>48.7</td>
<td>92.3</td>
<td>116.2</td>
</tr>
<tr>
<td>IL-12</td>
<td>(20.2–111.2)</td>
<td>(46.4–50.2)</td>
<td>(58.5–114.5)</td>
<td>(66.1–149.9)</td>
</tr>
<tr>
<td>VEGF</td>
<td>47.5</td>
<td>80.4</td>
<td>148.6</td>
<td>89.7</td>
</tr>
<tr>
<td>MIP-1α</td>
<td>137.2</td>
<td>122.0</td>
<td>103.8</td>
<td>84.4</td>
</tr>
<tr>
<td>IL-9</td>
<td>30.2</td>
<td>36.7</td>
<td>42.3</td>
<td>70.6</td>
</tr>
<tr>
<td>FGF-β</td>
<td>19.1</td>
<td>16.0</td>
<td>24.9</td>
<td>17.6</td>
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<tr>
<td>MCP-1</td>
<td>22.2</td>
<td>24.6</td>
<td>26.7</td>
<td>33.3</td>
</tr>
<tr>
<td>TNF-α</td>
<td>25.7</td>
<td>23.9</td>
<td>32.7</td>
<td>35.1</td>
</tr>
<tr>
<td>PDGF</td>
<td>10,103 (22.6–34.1)</td>
<td>8,492 (21.1–29.9)</td>
<td>10,862 (22.2–89.2)</td>
<td>8,211</td>
</tr>
<tr>
<td>IL-17</td>
<td>82.2</td>
<td>85.6</td>
<td>96.7</td>
<td>80.7</td>
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<tr>
<td>IL-8</td>
<td>(81.2–86.5)</td>
<td>(79.2–95.6)</td>
<td>(80.9–106.2)</td>
<td>(67.7–93.4)</td>
</tr>
<tr>
<td>IFN-γ</td>
<td>9.4</td>
<td>20.4</td>
<td>14.9</td>
<td>11.7</td>
</tr>
<tr>
<td>IP-10</td>
<td>(7.9–45.0)</td>
<td>(7.3–191.4)</td>
<td>(9.2–180.2)</td>
<td>(9.9–20.3)</td>
</tr>
<tr>
<td>G-CSF</td>
<td>178.1</td>
<td>184.6</td>
<td>206.6</td>
<td>191.2</td>
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<tr>
<td>TRAIL</td>
<td>58.7</td>
<td>722.4</td>
<td>604.8</td>
<td>565.4</td>
</tr>
<tr>
<td>MIP-1β</td>
<td>14.1</td>
<td>15.5</td>
<td>13.7</td>
<td>16.0</td>
</tr>
<tr>
<td>IL-6</td>
<td>(11.9–18.2)</td>
<td>(11.5–17.3)</td>
<td>(10.6–17.2)</td>
<td>(11.3–19.6)</td>
</tr>
<tr>
<td>GRO-β</td>
<td>58.2</td>
<td>42.7</td>
<td>40.9</td>
<td>70.8</td>
</tr>
<tr>
<td>MIP-1β</td>
<td>7.2</td>
<td>7.5</td>
<td>6.4</td>
<td>6.6</td>
</tr>
<tr>
<td>IL-17</td>
<td>(5.2–9.1)</td>
<td>(5.4–13.3)</td>
<td>(5.8–14.5)</td>
<td>(5.7–7.4)</td>
</tr>
</tbody>
</table>

**Notes:** All the values are given as median (interquartile range). *These detected cytokines in the sera samples showed no significant differences in between either of the subgroups including controls smoker and non-smoker and also in between ex-smoker with COPD and smoker with COPD. The following cytokines are undetectable (below the detection limit of the assay) in the sera: NGF-β, IFN-α2, IL-10, IL-15, IL-1β, IL-1α, IL-1β, IL-1Ra, IL-2Ra, IL-3, IL-5, ILF, MCP-3, M-CSF, MIF, MIP-1β, SDF-1α and TNF-β.

**Abbreviations:** COPD, chronic obstructive pulmonary disease; MIG, monokine induced by gamma-interferon; SCGF, stem cell growth factor; GM-CSF, granulocyte macrophage colony-stimulating factor; RANTES, regulated on activation, normal T cell expressed and secreted; GRO, growth-related oncogene; VEGF, vascular endothelial growth factor; FGF, fibroblast growth factor; PDGF, plate-derived growth factor; IP-10, interferon gamma-induced protein 10; G-CSF, granulocyte colony-stimulating factor; NGF, nerve growth factor; IFN, interferon; IL, interleukin; R, receptor; Ra, receptor alpha; LIF, leukemia inhibitory factor; MCP, monocyte chemoattractant protein; M-CSF, macrophage colony-stimulating factor; MIF, migration inhibitory factor; MIP, macrophage inflammatory protein; SDF, stromal cell-derived factor; TNF, tumor necrosis factor.