Clinical Phenotypes of COPD: Identification, Definition and Implications for Guidelines

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A R T I C L E   I N F O

Article history:
Received 8 August 2011
Accepted 20 October 2011
Available online 1 February 2012

Keywords:
COPD
Phenotype
Asthma
Emphysema
Exacerbation
Treatment

A B S T R A C T

The term phenotype in the field of COPD is defined as "a single or combination of disease attributes that describe differences between individuals with COPD as they relate to clinically meaningful outcomes". Among all phenotypes described, there are three that are associated with prognosis and especially with a different response to currently available therapies. The phenotypes are: the exacerbator, the overlap COPD-asthma and the emphysema-hyperinflation.

The exacerbator is characterized by the presence of, at least, two exacerbations the previous year, and on top of long-acting bronchodilators, may require the use of anti-inflammatory drugs. The overlap phenotype presents symptoms of increased variability of airflow and incompletely reversible airflow obstruction. Due to the underlying inflammatory profile, is used to have a good therapeutic response to inhaled corticosteroids in addition to bronchodilators. Lastly, the emphysema phenotype presents a poor therapeutic response to the existing anti-inflammatory drugs, and long-acting bronchodilators together with rehabilitation are the treatments of choice.

Identifying the peculiarities of the different phenotypes of COPD will allow us to implement a more personalized treatment, in which the characteristics of the patients, together with their severity will be key to choose the best treatment option.

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Fenotipos clínicos de la EPOC. Identificación, definición e implicaciones para las guías de tratamiento

R E S U M E N

El término fenotipo aplicado a la EPOC se define como “aquellos atributos de la enfermedad que solos o combinados describen las diferencias entre individuos con EPOC en relación a parámetros que tienen significado clínico”. De entre todos los descritos, existen tres que se asocian con factores pronósticos y sobretodo con distinta respuesta a los tratamientos disponibles en la actualidad. Estos fenotipos son: el agudizador, el mixto EPOC-asma y el enfisema-hiperinflacion.

El agudizador se caracteriza por la presencia de la menos dos agudizaciones el año previo y además del tratamiento con broncodilatadores de larga duración puede requerir la utilización de fármacos antiinflamatorios. El fenotipo mixto presenta una obstrucción no completamente reversible al flujo aéreo acompañada de una reversibilidad aumentada de la obstrucción. Por su perfil inflamatorio subyacente suele presentar una buena respuesta terapéutica a los corticosteroides inhalados unidos a los broncodilatadores. Por último el fenotipo enfisema presenta una pobre respuesta a los fármacos antiinflamatorios de que disponemos en la actualidad y los broncodilatadores de larga duración, junto a la rehabilitación son la base de su tratamiento.

El reconocimiento de las peculiaridades de los distintos fenotipos de la EPOC nos debe permitir guiar un tratamiento más personalizado, en el que las características del paciente se sumen a su gravedad para dirigir la terapia.

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What Do We Mean by “COPD Phenotype”?  
In recent years, the term “phenotype” has been used to refer to clinical types of patients with chronic obstructive pulmonary disease (COPD). This has been motivated by the boom in studies that propose to identify genetic determinants for developing the disease in its different manifestations. An international group of experts has defined COPD phenotype as “a single or combination of disease attributes that describe differences between individuals with COPD as they relate to clinically meaningful outcomes (symptoms, exacerbations, response to treatment, speed of progression of the disease or death”). Therefore, the phenotype should be able to classify the patients into subgroups with a prognostic value that allow for determining the best therapy in order to achieve better clinical results.

According to the opinion of the majority, the term “COPD phenotype” is reserved for the different clinical types that have therapeutic impact and are identified in COPD patients. In recent years, several researchers have attempted to quantify the different “faces” or phenotypes of COPD in the so-called non-proportional Venn diagram of COPD, that demonstrates the great confusion existing among the different etiopathogenic, clinical and morphologic types of this syndrome that we call COPD, although some have postulated that it should be defined as a group of orphan diseases.

Phenotypes of Clinical Interest in COPD

We should accept that there needs to be a mid-point between the excessive simplification of the term COPD (a definition that encompasses the entire spectrum of patients with non-completely reversible airflow obstruction) and the complexity of considering each patient individually as an orphan disease. This “happy medium” entails the identification and description of some phenotypes that are not only of biological and epidemiological interest but also of prognostic and above all therapeutic interest. Table 1 shows some studies that have identified several clinical phenotypes in COPD. These studies are based on heterogeneous populations, using diverse methodologies to analyze different variables, but they all reach similar conclusions: it is possible to distinguish between different patterns of clinical expression in COPD—the so-called phenotypes. The majority distinguish between 3 and 5 phenotypes based on a series of factors that are enumerated in Table 2.

After analyzing these studies, we can conclude that there is evidence to define at least three different phenotypes with clinical, prognostic and therapeutic repercussions: (1) “overlap” or mixed COPD-asthma; (2) exacerbator; and (3) emphysema-hyperinflation.

Other possible phenotypes have been defined, but these have had little clinical transcendence. Thus, the so-called “fast decliner” would be a patient who suffers a loss of lung function, expressed by FEV₁, that is faster than average. The practical problem is that it is impossible to identify this phenotype without a strict follow-up of the lung function for at least 2 years; on the other hand, no specific treatment has been identified for this type of patients. Another possible phenotype would be chronic bronchitis, defined as cough and expectoration for at least 3 months of the year for 2 consecutive years. This phenotype is usually associated with airway disease, which can be visualized with high-resolution computed tomography (HRCT). Nevertheless, chronic bronchitis can accompany any of the three phenotypes indicated beforehand: “mixed”, exacerbator and emphysema. We therefore prefer to describe it as a modifying factor in any of the 3 main phenotypes. A systemic phenotype has also been defined in patients who present obesity, cardiovascular disease, diabetes or systemic inflammation. It is true that these patients present a different prognosis, but we cannot call “systemic” COPD a phenotype as it does not meet the anterior definition, as the systemic manifestations (or comorbidities) have not been shown to be a manifestation of the COPD itself. The systemic manifestations or comorbidities are very important, but they should be considered apart from the phenotype.

Last of all, one special phenotype is emphysema due to alpha-1-antitrypsin deficiency, which is characterized by predominantly basal emphysema that appears at early stages in life, especially in smokers, and it has a genetic base. Due to its limited prevalence, we prefer to consider it apart from the general classification.

Mixed COPD-Asthma Phenotype

When a patient presents characteristics of more than one obstructive airway disease, we say that he/she has an overlap or mixed syndrome. The guidelines of the American Thoracic Society (ATS) from 1995 defined obstructive disease and identified 11 different syndromes, 6 of which were overlap syndromes. A study that used data from a very extensive population observed that 19% of patients with airflow obstruction had more than one disease present. The most representative and frequent diseases or processes within these subgroups were chronic airflow obstruction and asthma. Therefore, it should not seem strange that there are a good number of patients who share characteristics that are attributed to COPD and asthma. This population is of special interest as it usually sidelines in clinical pharmaceutical trials. Asthma studies tend to exclude smokers, and COPD studies usually exclude individuals with a previous history of asthma. Some even exclude those individuals with a positive bronchodilator test.

Definition of the Mixed Phenotype (COPD-Asthma)

The mixed phenotype in COPD is defined as an airflow obstruction that is not completely reversible, accompanied by symptoms or signs of increased obstruction reversibility.

Justification of the Mixed Phenotype

Pathogenesis and Prevalence

Within the spectrum of chronic airway obstruction, there are individuals with asthma who smoke, asthmatics who develop airflow obstruction that is not completely reversible and non-smokers who develop chronic airflow obstruction. Smokers with asthma have features that mimic COPD, with less response to corticosteroids, a lower frequency of eosinophilic inflammation and a greater probability of neutrophilia in the airways. On the other hand, there are epidemiological studies about the incidence of COPD which demonstrate that young asthmatics who develop COPD have a disease with different characteristics than the non-asthmatics who also develop COPD. In the first case, allergic rhinitis is more frequent, along with nonspecific bronchial hyperreactivity and the presence of wheezing, and plasma IgE concentrations are higher; all of which indicate that it is a mixed asthma-COPD syndrome.

The prevalence of the mixed phenotype is unknown, but there are different estimations of its importance in the context of COPD. One initial study that was small in size estimated that 25% of COPD patients with COPD have significant reversibility and presented clinical response to inhaled corticosteroids (IC). Soriano et al. estimated that approximately 23% of COPD patients between the ages of 50 and 59 could have a mixed phenotype, which increased with age up to 52% between the ages of 70 and 79. Other studies have quantified the prevalence of the mixed phenotype (identified by eosinophilia in sputum) in patients with COPD at 38%, directly associated with the therapeutic response to IC.
bronchodilator test as a reference, 31.5% of the patients identified with COPD in the EPI-SCAN epidemiological study had a positive test. Based on these results, we can conclude that, together, between 20% and 40% of COPD patients can be carriers of a mixed phenotype.

Differential Treatment

The clinical justification for the mixed phenotype lies in its demonstrated sensitivity to the anti-inflammatory action of IC. The basis that explains the response to corticosteroids in COPD patients with greater reversibility lies within the etiopathogeny of the disease. Papi et al. demonstrated that reversible patients, even those who were only partially reversible (increase in FEV₁ >200 ml, but <12%) had greater eosinophilic bronchial inflammation compared with the irreversible ones, in whom neutrophilic inflammation predominated. In fact, several studies have used the greater airflow reversibility, a high concentration of eosinophils in spontaneous or induced sputum or a greater concentration of exhaled NO as markers of the response to IC in COPD, both at the lung function level as well as the improvement of the symptoms. A more recent study classified a small group of patients into 3 different phenotypes, according to the findings from chest computed tomography (CT). The authors demonstrated a relationship between the response to the bronchodilator test, the response to treatment with IC and the concentration of eosinophils in sputum in each of the 3 phenotypes. There is even a randomized clinical assay that compared the treatment with IC in patients with COPD, directed in accordance with the current guidelines.

Table 1

<table>
<thead>
<tr>
<th>Author and Year</th>
<th>Population</th>
<th>Methods</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Casanova, 2005</td>
<td>689 patients with COPD followed for 34 months, on average</td>
<td>Survival analysis according to hyperinflation</td>
<td>Defines the relevance of the emphysema-hyperinflation phenotype</td>
</tr>
<tr>
<td>Wardlaw, 2005</td>
<td>49 patients: 27 with asthma and 22 with COPD</td>
<td>Cluster analysis</td>
<td>Defines 4 phenotypes: COPD, mixed COPD-asthma, asthma with minimal eosinophilia and low IgE</td>
</tr>
<tr>
<td>Kitagushi, 2006 and Fujimoto, 2006</td>
<td>172 patients with stable COPD</td>
<td>Chest CT, marker for inflammation and peripheral cellularity and in sputum</td>
<td>Defines 3 phenotypes: mild emphysema with or without bronchial thickening (BT), emphysema without BT and emphysema with BT classifies the patients into 3 groups according to severity of emphysema on CT</td>
</tr>
<tr>
<td>Makita, 2007</td>
<td>274 COPD patients in stable phase</td>
<td>Chest CT, quality of life and spirometry</td>
<td>Defines the sub-phenotype of chronic bronchitis with greater eosinophilia inflammation</td>
</tr>
<tr>
<td>Marsh, 2008</td>
<td>469 individuals &gt;50 years of age</td>
<td>Questionnaires, chest CT and spirometry</td>
<td>Proportional classification of up to 16 different phenotypes: airway or parenchyma disease</td>
</tr>
<tr>
<td>Pistolesi, 2008</td>
<td>322 COPD patients (development group) and 93 (validation group)</td>
<td>Multidimensional scale and cluster analysis</td>
<td>Nine variables define two main phenotypes: asthma with emphysema and COPD-asthma with emphysema</td>
</tr>
<tr>
<td>Sneeck-Stroband, 2008</td>
<td>114 patients with COPD</td>
<td>Bronchial biopsies and induced sputum</td>
<td>Defines the characteristics of the mixed COPD-asthma mixed phenotype</td>
</tr>
<tr>
<td>Weatherall, 2009</td>
<td>Randomized population sample of 175 individuals aged 25 to 75</td>
<td>Questionnaires, chest CT, spirometry, Fen, blood analysis; cluster analysis</td>
<td>Defines 5 airway disease phenotypes</td>
</tr>
<tr>
<td>García-Río, 2009</td>
<td>110 patients with COPD</td>
<td>Analysis of physical activity related to dynamic hyperinflation</td>
<td>Contributes to characterizing the emphysema-hyperinflation phenotype</td>
</tr>
<tr>
<td>Roy, 2009</td>
<td>127 COPD patients</td>
<td>Spirometry, FeNO, CRO and TNFα in plasma, sputum analysis. Multivariate analysis of linear regression</td>
<td>Identifies 4 major components that explain the different types of COPD</td>
</tr>
<tr>
<td>Gibson and Simpson, 2009</td>
<td>NA</td>
<td>Bibliographic review</td>
<td>Defines the characteristics of the mixed or COPD-asthma mixed phenotype; 3 main components that explain 61% of the variance; 4 phenotypes identified</td>
</tr>
<tr>
<td>Burge, 2010</td>
<td>322 patients with COPD</td>
<td>Clinical symptoms plus spirometry, quality of life and anxiety-depression; analysis of main component and clusters with 31 variables</td>
<td>6 factors identified that explain 75% of the variability and 4 different phenotypes</td>
</tr>
<tr>
<td>Cho, 2010</td>
<td>308 patients with severe emphysema</td>
<td>Survival analysis according to hyperinflation and peripheral cellularity with sputum analysis</td>
<td>4 phenotypes identified according to severity and reversibility</td>
</tr>
<tr>
<td>Hurst, 2010</td>
<td>2,138 COPD patients</td>
<td>Spirometry, FeNO, CRO and TNFα in plasma, sputum analysis. Multivariate analysis of linear regression</td>
<td>4 phenotypes identified according to severity and reversibility</td>
</tr>
<tr>
<td>Jo, 2010</td>
<td>191 patients &gt;60 with obstruction or respiratory symptoms</td>
<td>Spirometry, FeNO, CRO and TNFα in plasma, sputum analysis. Multivariate analysis of linear regression</td>
<td>Defines 3 phenotypes: severe respiratory COPD, moderate respiratory COPD and systemic COPD</td>
</tr>
<tr>
<td>García-Aymerich, 2011</td>
<td>342 COPD after first hospitalization</td>
<td>Symptoms, spirometry, quality of life, exercise capacity, nutritional state, biomarkers and thoracic CT</td>
<td>Identifies 3 phenotypes: severe respiratory COPD, moderate respiratory COPD and systemic COPD</td>
</tr>
<tr>
<td>Márquez-Martín, 2011</td>
<td>64 patients with stable COPD</td>
<td>Thoracic CT, spirometry and exercise tests</td>
<td>Compares patients with and without emphysema</td>
</tr>
</tbody>
</table>

IgE: immunoglobulin E; CT: computed tomography; FeNO: exhaled fraction of nitric acid; CRP: C-reactive protein; TNFα: tumor necrosis factor-alpha.
Eosinophilia in sputum

When the results were analyzed separately, the reversible patients achieved a maximum bronchodilator effect of 319 ml FEV\textsubscript{1}, while the irreversible patients reached 195 ml.\textsuperscript{30}

The limited reduction in mortality observed in the Towards a Revolution in COPD Health (TORCH) study with FSC can be explained, at least in part, by the patient selection. One of the inclusion criteria was to have a negative bronchodilator test, which is reflected in a mean reversibility of the participants of only 3.7%.\textsuperscript{31} Therefore, TORCH explains the long-term effect of the FSC combination in those patients who are less susceptible of being responders to IC. Contrarily, a more recent study has compared FSC with salmeterol in treating patients with severe COPD (FEV\textsubscript{1} <50%).\textsuperscript{32} Its results showed a significant reduction (35%) during one year in the rate of moderate or severe exacerbations with FSC compared with salmeterol alone. This study did not take into account the reversibility of FEV\textsubscript{1} among its inclusion criteria, and in fact the mean reversibility of its patients was 7%, almost the double of TORCH, and the result was a spectacular reduction in the frequency of exacerbations by adding fluticasone to the treatment with salmeterol. In the same direction, in a recent study it was demonstrated that treatment with FSC at a dosage of 250/50 every 12 h produced an increase in the area under the curve 6 h after FEV\textsubscript{1} that was more than double in reversible patients (1.98 l in week 8) than in irreversible ones (0.74 l). This provides more evidence of the different response to IC or combined treatment depending on the response to the bronchodilator test.\textsuperscript{20} It has also been recently shown that there is a direct and significant correlation between the response to the bronchodilator test with salbutamol in COPD and the improvement in lung function after 3 months of treatment with LABA+IC.\textsuperscript{21} In contrast, patients with a defined phenotype such as emphysema-dominant did not present any improvement in lung function with the same treatment.\textsuperscript{21} It should be expected that these findings described in the studies carried out with salmeterol or with SAL/FLU may be extrapolatable to other IC or LABA/IC combinations.

In short, the conclusions that we may draw from the existing studies are: (a) the patients with mixed phenotype, who present certain characteristics (sputum or peripheral eosinophilia, history of asthma and/or atopy, frequent exacerbations, very positive bronchodilator test or wheezing as a guiding sign) are susceptible for presenting a good response to IC, whatever the lung function; (b) COPD patients who do not present the former characteristics will obtain marginal clinical benefits with the use of IC added to long-acting bronchodilators.

### Diagnosis of the Mixed Phenotype (COPD-Asthma)

In order to be able to identify the mixed phenotype, the clinical history will serve as a guide, showing history of asthma and atopy in childhood and youth, less intense smoking exposure, frequency of exacerbations and key symptoms such as wheezing, among others. But in order to characterize a patient with COPD as mixed, it is also necessary to carry out a series of tests. Spirometry, in addition

### Table 2

Factors or Variables Identified as Significant for the Classification of COPD Patients.

<table>
<thead>
<tr>
<th>Author and Year</th>
<th>Factors</th>
</tr>
</thead>
<tbody>
<tr>
<td>Casanova, 2005\textsuperscript{59}</td>
<td>IC/TLC, Eosinophilia in sputum, Reversibility of the obstruction, IgE, Quantiﬁcation of CT for parenchyma and airway, Eosinophilia in sputum and peripheral, Respiratory symptoms, Emphysema by CT, BMI, Quality of life, Sputum volume and appearance, Quantiﬁcation of CT for parenchyma and airway, Pulmonary sounds, FEV\textsubscript{1}/FVC, Air trapping, Eosinophilia in sputum in chronic bronchitis, Age, FEV\textsubscript{1} and FEV\textsubscript{1}/FVC, Reversibility, percentage, Kco, percentage, FRC, percentage, IgE, FeNO, smoking, pack-years, Physical activity related with dynamic hyperinﬂation, Neutrophilia in sputum, IL-8 and TNFα, Eosinophilia in sputum and FeNO, Reversibility to bronchodilator, FEV\textsubscript{1} and IC, CRP, Symptoms, FEV\textsubscript{1}, Bronchial hyperreactivity, Eosinophilia in sputum, FEV\textsubscript{1}, Age, Symptoms, Comorbidity, FEV\textsubscript{1} post-bronchodilator, Bronchodilator response, percentage, Quantitative measurement of emphysema and bronchial wall on CT, FRC, FEV\textsubscript{1}, Frequency of exacerbations in the past, Gastroesophageal reﬂux, Quality of life, Age, Reversibility, percentage, Post-bronchodilator FEV\textsubscript{1}, Severity of respiratory symptoms, Comorbidity and systemic inﬂammation, Emphysema by CT, Peripheral muscle strength, Exercise capacity, BMI</td>
</tr>
<tr>
<td>Wardlaw, 2005\textsuperscript{50}</td>
<td>CT, computed tomography; FEV\textsubscript{1}: forced expiratory volume in one second; BMI: body mass index; IC/TLC: inspiratory capacity/total lung capacity; IgE: immunoglobulin E; VC: vital capacity; FVC: forced vital capacity; Kco: carbon monoxide transfer coefficient; FRC: functional residual capacity; FeNO: exhaled fraction of nitric acid; IL: interleukin; TNFα: tumor necrosis factor alpha; CRP: C-reactive protein.</td>
</tr>
</tbody>
</table>
to diagnosing the disease, provides a measurement of its severity, and the magnitude of the reversibility in the bronchodilator test will guide us towards the possible diagnosis as mixed. The blood work-up reveals if there is eosinophilia, and in an ideal situation the cytologic analysis of the sputum may be able to indicate the intensity of the eosinophilic inflammation, while a high concentration of exhaled nitric oxide could also help to identify patients with mixed COPD-asthma phenotype.

The utility of the bronchodilator test in determining the response to IC has been brought into doubt after the results reported by Calverley et al. In the screening phase of the ISOLDE study, they analyzed a total of 660 patients. By carrying out 3 bronchodilator tests in a 2-month period, they observed that a significant number of patients could be either positive or negative for the different tests; therefore, they concluded that it was not adequate to classify the patients as reversible or irreversible. Nevertheless, it must be kept in mind that they excluded the patients who on the first test had a reversibility of FEV₁ >10%, and that the protocol of the test was different on the 3 different occasions that it was performed. This study does not invalidate the utility of the bronchodilator test, but it reminds us that response is a continuous variable and that it is more reliable to interpret the magnitude of the response in each case than to classify a patient as either reversible or irreversible according to an arbitrary cut-point.

In short, the diagnosis of the mixed phenotype will be established by the presence of a combination of the following factors: history of asthma and/or atopy, reversibility in the bronchodilator test, notable eosinophilia in respiratory and/or peripheral secretions, high IgE, positive prick test to pneumoallergens and high concentrations of exhaled NO.

**Exacerbator Phenotype**

The clinical course of COPD is frequently punctuated by episodes of clinical instability, which we refer to as exacerbations. It is estimated that COPD patients suffer between 1 and 4 exacerbations per year; however, their appearance does not follow a normal distribution. Some patients do not suffer any exacerbations, while others experience them repeatedly. In the Evaluation of COPD Longitudinally to Identify Predictive Surrogate Endpoints (ECLIPSE) study, which is a prospective observational study including 2138 patients with moderate-severe COPD followed for 3 years, 23% of the patients did not have any exacerbations, while 12% of the cases had 2 or more exacerbations per year over the course of the 3-year study period. The exacerbators maintained a notable stability over time to the point that somewhat more than 60% of the patients with 2 or more exacerbations in the first year also had frequent exacerbations in the second year of follow-up, and out of these more than 70% continued having repeated decompensation in the third year. Given this stability over time, it has been suggested that these patients could present individual susceptibility for suffering frequent decompensations. This fact, and also the fact that we are faced with a patient group with a high risk for morbidity and mortality, whose treatment could be differentiated, constitute the rationale for defining the “exacerbator” phenotype. The cut-point of the number of exacerbations for considering a patient an exacerbator has varied over time, but currently exacerbators are considered those patients who present 2 or more exacerbations per year.

**Definition of “Exacerbator”**

“Exacerbators” are defined as those COPD patients who present with 2 or more exacerbations per year. These exacerbations should be separated by at least 4 weeks after the end of treatment of the previous exacerbation or 6 weeks after the onset of the exacerbation in cases that have received no treatment. This is in order to be able to differentiate between the new event and previous therapeutic failure.

**Justification of the Exacerbator Phenotype**

**Individual Susceptibility for Suffering Frequent Exacerbations**

Table 3 compiles the main risk factors linked to the presence of repeated exacerbations. The severity of the airflow limitation is without doubt one of the most well-known factors. However, the relationship between FEV₁ and number of exacerbations is not linear, and in fact close to 40% of the severe or very severe patients do not present exacerbations, while more than 20% of the moderate patients frequently present them. This suggests the existence of other conditioning factors. Of all of them, the history of previous exacerbations is the most frequent factor referenced in the literature, which emphasizes the existence of a certain individual susceptibility that may either be hereditary or acquired.

**Individual Acquired Susceptibility**

**Chronic bronchial-bronchitis hypersecretion.** The presence of cough and chronic expectoration is associated with a greater risk for repeated exacerbations. Foreman et al. found that the odds ratio (OR) for exacerbation was 3.7 for patients with chronic expectoration, much higher than the risk observed related to accumulated tobacco consumption (OR: 1.01, for each pack-year) or post-bronchodilator FEV₁% (OR: 0.98). Similar results have also been reported by Miravitlles et al. who observed a significant association between chronic mucus hypersecretion and the presence of two or more exacerbations in the previous year (OR, 1.54). Likewise, Burgel et al. verified that among the patients with frequent exacerbations, 55% had associated chronic expectoration and cough, compared with 22% of cases without bronchial hypersecretion (P<.001), with a greater risk for hospitalization among the hypersecretors. The association between frequent exacerbations and chronic bronchial hypersecretion was independent of other known risk factors for repeated exacerbations, such as FEV₁, age, cardiovascular comorbidity or active smoking, which confirm chronic expectoration as a notable marker for exacerbation.

**Bronchial hypersecretion has been associated with greater airway inflammation and greater risk for respiratory infection,** which could explain the link with the appearance of repeated exacerbations. The same could be said of bronchiectasis, which is very...
The cause of this greater inflammation—which once again reinforces the inflammatory-infectious hypothesis as an etiopathogenic factor that underlies in the exacerbator phenotype.

Viral infection can also play a relevant role in modulating the inflammatory response of the airway, altering the fragile balance between the presence of bacteria in the airway and the response of the host. In fact, patients with frequent colds (for example, viral infections by rhinovirus) also experience more bacterial exacerbations.68

Recently, it has been suggested that gastroesophageal reflux disease (GERD) may also predispose patients to frequent exacerbations.35,63 The intimate mechanism that links GERD and exacerbation is not clear, but some authors suggest the existence of alterations in the swallowing reflux and the possible existence of microaspiration,70 which once again reinforces the relationship between infection and the inflammation caused by repeated exacerbations.

Although in most exacerbations, especially if they are repeated, there is a potential underlying infectious mechanism, the truth is that the greater inflammation observed in exacerbator patients could have other origins. As mentioned in the previous section, there is a specific group of patients (mixed COPD-asthma phenotype) where certain asthmatic-type characteristics underlay. Recently, an observational, case–control study has shown evidence of a greater risk for suffering frequent exacerbations among patients who had an asthma diagnosis made by their physicians,52 which could suggest an alternative mechanism to the inflammatory-infectious one. However, the study was retrospective and observational; therefore the conclusions should be cautious. In addition, these patients with a reported diagnosis of asthma did not present greater reversibility in the bronchodilator test, nor did they have any objective measurements that confirmed the asthma diagnosis or, if not, the mixed asthma-COPD phenotype instead. Autoimmunity phenomena could also be connected with the persistence of greater airway inflammation. However, to date there is hardly any evidence to establish an association between said autoimmunity and the presence of repeated exacerbations.

Cardiovascular disease and repeated exacerbations. Confronting the inflammatory hypothesis (infectious or non-infectious), a significant association has been described among different cardiovascular pathologies and a greater frequency of exacerbations.36,40,46,54 In
a prospective study with a case–control design in patients with severe COPD, the exacerbators presented a greater number of cardiovascular events than the subjects with COPD of similar severity, but without exacerbations. The direction of this association has not been clearly outlined. While some studies suggest that the exacerbations cause or trigger the cardiovascular manifestations through different mechanisms such as systemic inflammation, hypoxemia or endothelial dysfunction, it is not clear if it is the cardiovascular events themselves, such as some rhythm disorders (auricular fibrillation, flutter, etc.), episodes of myocardial ischemia or ventricular failure, those that could mimic an exacerbation with difficult differential diagnosis, due to, among other reasons, the non-specificity of the clinical symptoms. In fact, almost 30% of severe exacerbations present symptoms suggestive of heart failure, and we frequently observe higher levels of troponin, a marker of myocardial injury, during COPD exacerbations. Be they either a cause or consequence, the truth is that these cardiovascular episodes are especially relevant in severe exacerbations. In a series of patient deaths during hospitalization due to COPD exacerbation, heart failure was identified as the cause of death in more than one-third of the patients. Pulmonary embolism, which is also difficult to diagnose, explained more than 20% of the deaths.

Individual Genetic Susceptibility

Although there is very little information available, the existence of a marked heterogeneity in the defense mechanisms of the host against the pathogen could indicate a certain genetic susceptibility. This fact has been supported by the finding of some polymorphisms in patients with frequent exacerbations. Differences in the genotype-dependent expression of the CCL1 protein, a chemotactic factor for the monocytes and macrophages, could produce alterations in the activation of the innate immune system versus respiratory infections. Likewise, polymorphisms have been described in the MBL2 (mannose binding lectin) connected with a greater frequency of hospitalizations. MBL is a protein of the innate immune system that inactivates a large number of microorganisms by means of the activation of the complement. Its deficiency, due to MBL2 polymorphisms, can potentially increase the susceptibility to infection.

Greater Risk for Morbidity and Mortality

Traditionally, COPD exacerbations have been considered clinical decompensations that are more or less transitory, whose repercussions were limited to the duration of the event itself. However, today we know that an important proportion of patients do not completely recover after an exacerbation, and this can cause later consequences, both pulmonary as well as systemic. In the cases in which repeated exacerbations are produced (exacerbator phenotype), the consequences can accumulate. Numerous studies have demonstrated the existence of a close relationship between the frequency of the exacerbations and the deterioration of the health-related quality of life (HRQL). The same is true with different extrapulmonary manifestations, such as depression, myopathy, myocardial infarction or GERD. Much more frequent among the “exacerbator” patients. An accelerated deterioration in lung function has also been documented, which has been estimated at 8 ml/year more among the patients with frequent exacerbations and even a persistent worsening of the BODE index.

Finally, and perhaps as a consequence of all that has been mentioned, a poorer prognosis has been demonstrated because, as the frequency of exacerbations increases, so does the risk for death, regardless of the baseline severity of the disease. Thus, we believe that “exacerbator” patients form a special group of patients with a high risk for morbidity and mortality, whose therapeutic approach should be different and intensive. These patients are also an enormous burden for the health-care system as it is estimated that they are responsible for 60% of hospital services rendered.

Differential Treatment

Long-acting bronchodilators, which are the first step in treating COPD, have been shown to reduce the frequency of exacerbations. When exacerbations persist despite bronchodilator treatment, the introduction of anti-inflammatories is indicated. In this context, various clinical practice guidelines recognize the usefulness of the use of IC in patients who present frequent exacerbation as their use, especially when associated with bronchodilators, produces a significant reduction in the number of exacerbations and an improvement in HRQL. Traditionally, this effect has been accepted for severe or very severe patients (FEV1 <50%) with frequent exacerbations. However, some studies in patients with less functional severity also back the use of these drugs; therefore, it seems that the main determinant of the benefit is precisely the presence of repeated exacerbations. Roflumilast is a new oral anti-inflammatory drug that acts by selectively inhibiting phosphodiesterase IV which has been approved for preventing exacerbations in patients with severe COPD who present cough and chronic expectoration and also suffer frequent exacerbations; therefore it is indicated for the exacerbator phenotype with chronic bronchitis. Macrolides, administered for a prolonged amount of time, could also have a specific indication for some of these patients as they have anti-inflammatory and immunomodulatory actions in addition to their possible antibacterial action. Some clinical assays suggest that the use of these drugs in stable patients with severe COPD significantly reduces the number of exacerbations, but with a possible increase in the risk of the appearance of bacterial resistances.

Finally, and given the potential role of PPM, it has also been suggested that the use of antibiotics during periods of stability (antibiotic chemoprophylaxis or treatment of the chronic bronchial infection) could be useful for reducing exacerbations. In this direction, the PULSE (Pulsed moxifloxacin Usage and its Long-term impact on the reduction of Subsequent Exacerbation) study is a clinical trial that studied the efficacy of the administration of 5-day cycles of 400 mg of moxifloxacin every 8 weeks in patients with stable COPD. The results indicate that this treatment reduced the risk for exacerbation by 20% in the intention-to-treat (ITT) analysis, 25% in the per-protocol (PP) analysis and 45% in patients who presented purulent or mucopurulent sputum, also by means of a PP analysis, without a significant increase in bacterial resistances. In another study done in patients with severe COPD colonized by Psedomonas aeruginosa, the administration of nebulized tobramycin reduced the number of severe exacerbations by 42%, also reducing bronchial inflammation. More studies are required to help adequately profile candidate patients as well as the duration and the type of antibiotic treatment necessary. It is very likely that the patients who would most benefit from this option would be those who present frequent exacerbations and purulence in sputum during stable phases.

Diagnosis of the Exacerbator Phenotype

The exacerbator phenotype is identified when the following criteria are met: existence of two or more exacerbations per year; the exacerbations should be separated at least 4 weeks after the end of the treatment of the previous exacerbation or 6 weeks from the start of the exacerbation in cases that have not received treatment. In cases in which the exacerbator phenotype is finally established, it is necessary to properly characterize the patient, searching for the existence of chronic bronchial infection and/or the presence...
of bronchiectasis. The use of anti-inflammatory and/or antibiotics can be especially useful among these patients.

**Emphysema-Hyperinflation Phenotype**

In recent years, many studies have demonstrated that variables such as dyspnea, exercise capacity and hyperinflation predict mortality independently from the lung function, and they are even better predictors than FEV\textsubscript{1} itself. This justifies defining and establishing the emphysema-hyperinflation COPD phenotype as a group of patients with a higher risk for mortality which presents certain differences with regards to treatment guidelines.

Pulmonary emphysema is defined, in anatomicopathologic terms, as the permanent destruction of the air spaces beyond the terminal bronchioles. We know that the loss of elastic retraction and the development of expiratory flow limitation make alveolar emptying difficult, originating air trapping and hyperinflation. This phenomenon has been associated with the limitations in the functional capacity of COPD patients and it is more closely related with dyspnea and tolerance to exertion than airflow obstruction. Moreover, it is known that the correlation between the extension and the severity of the macroscopic emphysema and the degree of obstruction (FEV\textsubscript{1}) is low. Nevertheless, the extension of the emphysema measured by HRCT does explain a large part of the variability of the carbon monoxide (CO) diffusing capacity. There are studies that have demonstrated an inverse correlation between body mass index (BMI) and degree of emphysema, evaluated by HRCT.

Hyperinflation in patients with emphysema is usually divided into static and dynamic. Static hyperinflation is the most common, caused by the loss of retraction of the pulmonary parenchyma in patients with emphysema. Although we do not understand its development in the natural evolution of COPD, it appears more frequently and with greater intensity as the FEV\textsubscript{1} diminishes. Dynamic hyperinflation can occur either independently or associated with static hyperinflation, and it appears in patients with any degree of severity. Dynamic hyperinflation is produced when inspiration begins before reaching complete expiration, which determines that with each breath a certain amount of air becomes trapped in the lungs. In patients with COPD, dynamic hyperinflation is produced when there is a limitation of expiratory airflow due to the airway obstruction, secondary to the increase in cholinergic tone, inflammation and mucus plugs. At the same time, it is favored by the increased collapsibility of the airways, which increases their resistance and prolongs the time necessary for completing the expiration. Hyperinflation entails an inspiratory load of the threshold type as in these patients the inspiration begins when there still has not been complete pulmonary emptying and the inspiratory muscles should first surpass the elastic retraction pressure of the lungs that still favors expiration (auto-PEEP or intrinsic PEEP). Hyperinflation is reversible in character; therefore, it is an attractive therapeutic target.

**Definition of the Emphysema-Hyperinflation Phenotype**

The emphysema-hyperinflation phenotype defines COPD patients who present dyspnea and intolerance to exercise as the predominating symptoms, which are frequently accompanied by signs of hyperinflation. Patients with emphysema phenotype present a tendency towards a lower BMI.

This clinical form of COPD is characterized by the presence of functional data of hyperinflation, the existence of emphysema on the HRCT study and/or a diffusion test lower than the reference value, measured with the DLCO/VA ratio adjusted for hemoglobin. The presence of emphysema has not been associated with a greater risk for exacerbations, except if it coexists with chronic bronchitis; in this case, the patient would be classified as exacerbator, and the treatment should prioritize reducing exacerbations.

**Justification of the Emphysema-Hyperinflation Phenotype**

**Genetic Susceptibility**

The different phenotypic expression of the pulmonary disease in smokers is determined in part by genetic factors. Specifically, in family studies using HRCT, we have observed that there is family aggregation, regardless of the emphysema phenotype, which indicates the presence of genetic determinants that define this phenotype. More recent studies have identified single nucleotide polymorphisms (SNP) that are significantly associated with the extension of low-density areas in pulmonary HRCT, and even certain genetic loci have been significantly related with the presence and extension of the emphysema in large COPD patient populations. These studies justify the differentiation of the emphysema phenotype as a characteristic genetic-base process in smokers.

One special case is congenital emphysema due to alpha-1-antitrypsin deficiency, as it is produced by a genetic mutation in the gene that codifies this protein. Patients who are homozygote for the deficiency mutation have an increased risk for developing emphysema that is predominantly basal and early-onset. This type of emphysema has been used as a model for understanding the physiopathology of emphysema in smokers.

**Greater Risk of Morbidity and Mortality**

The clinical importance of identifying the emphysema-hyperinflation phenotype is based on the fact that the degree of dyspnea, intolerance to exercise and hyperinflation are predictors for mortality that are independent of the severity of the obstruction. In a prospective study with a 5-year follow-up, Casanova et al. observed an inverse relationship between the degree of hyperinflation and survival. They demonstrated that patients with COPD and an IC (inspiratory capacity)/TLC (total pulmonary capacity) ratio less than 0.25 were 3.15 more likely to die than those with higher ratios. In this study, the multivariate analysis demonstrated that this ratio (IC/TLC) continued to be a risk factor, regardless of other parameters, such as FEV\textsubscript{1}, age, dyspnea, exercise capacity or comorbidity.

A positive relationship has also been demonstrated between the magnitude of the emphysema measured by HRCT, hyperinflation and the BODE index, although differences have not been observed in pulmonary attenuation among different quartiles of the BODE index, probably due to the participation of extrapulmonary factors in this prognostic score. Nevertheless, the presence of hyperinflation on HRCT in smokers with normal FEV\textsubscript{1} is associated with a faster fall in FEV\textsubscript{1} and, finally, a significant association has been demonstrated between the magnitude of the emphysema evaluated by HRCT with a greater mortality in COPD, regardless of the severity measured by FEV\textsubscript{1}. In this manner, we see the growing evidence of the need for HRCT when evaluating COPD patients for the study of the emphysema as well as to evaluate the possible presence of bronchiectasis.

The impact of emphysema on mortality was also observed in the National Emphysema Treatment Trial (NETT), in which the 3 factors—emphysema, hyperinflation and the BODE index—were independent predictors of mortality. However, we should be reminded that this is a cohort of patients with very severe COPD and, therefore, these results are not extrapolatable to all COPD patients. These data are corroborated by studies such as that by Nishimura et al. and the one by Martinez et al. where the patients with higher residual volume had a greater mortality.
with a similar tendency in cases where the IC/TLC ratio was lower.

Indirectly, dynamic hyperinflation can contribute to a poorer prognosis of COPD by significantly reducing the exercise capacity of patients who are affected by it. Thus, it has been demonstrated that physical activity in patients with moderate-to-severe COPD correlates inversely with degree of dynamic hyperinflation, and the patients with less physical activity are those who present a higher rate of hospitalizations and greater mortality. According to the current guidelines, long-acting bronchodilators are the foundation of the pharmacological treatment of COPD. They improve symptoms and exercise capacity and, consequently, improve the state of health as perceived by the patient, with statistically significant and clinically relevant changes. Nevertheless, occasionally the benefits reached at the clinical level do not translate into an improvement of the degree of obstruction (changes in FEV₁) but of hyperinflation instead by reducing dynamic hyperinflation with improvements in the inspiratory capacity, the degree of dyspnea and exercise tolerance.

The current guidelines recommend the association of bronchodilators in order to try to achieve an additional effect, without increasing the adverse effects in the patients with poorly controlled symptoms in spite of treatment with a bronchodilator. In this direction, the use of double bronchodilator therapy (formoterol and tiotropium) versus bronchodilator monotherapy or versus the fluticasone-salmeterol combination offers an added functional benefit with reduction of the need for rescue medication, improvement in the symptoms and quality-of-life questionnaires. These results can be applied to other LABA/IC combinations.

Anti-inflammatory treatment with inhaled corticosteroids, whose main objective is the prevention of exacerbations, has not been shown to be as effective in the emphysema-hyperinflation phenotype. Nor has the oral anti-inflammatory roflumilast shown good results for the reduction of exacerbations in patients with emphysema, except in those who associated symptoms of chronic bronchitis.

In short, patients with an emphysema-hyperinflation phenotype could benefit more from a double bronchodilator therapy and, of course, from respiratory rehabilitation due to its beneficial effects on dyspnea and exercise tolerance.

**Diagnosis of the Emphysema-Hyperinflation Phenotype**

The lung function parameter that best evaluates the presence of emphysema is the carbon monoxide transference capacity (DLCO), which correlates well with the severity of pulmonary emphysema. Nevertheless, one of its limitations is that it analyzes the entire lung as a whole, unlike HRCT, which is able to detect localized destructive changes, and currently this imaging technique is often used for detecting pulmonary emphysema. In addition, recent studies show that quantifying the magnitude of pulmonary emphysema using densitometry parameters could be a sensitive and specific exploration in the evaluation and the follow-up of pulmonary emphysema. Thus, there are studies that have demonstrated that the analysis of the density of the pulmonary parenchyma in HRCT correlates with the pathological alterations observed in tissue samples and with lung function deficiency (airflow obstruction and diffusion capacity), which would allow for a radiological estimation of the COPD severity.

Hyperinflation is evaluated by means of the determination of static lung volumes. Nevertheless, IC obtained with slow spirometry provides an indirect estimation of the magnitude of the hyperinflation in a simpler, reproducible manner. IC correlates well with dyspnea and with exercise capacity in COPD patients. It has been observed that the reduction in IC correlates with an increase in dyspnea and a decrease in exercise capacity. This fact is justified because the most important physiopathological factor that determines exertion dyspnea in COPD patients is the development of air trapping and dynamic hyperinflation triggered by exercise, including the physical exertion associated with carrying out the activities of daily life.
In this review, we give reasons with prognostic and therapeutic significance. In this way, we may to COPD has resulted in the definition of different types of patients. The development of different options for pharmacological and non-pharmacological treatments has led to the demonstration that the clinical response can be different according to the characteristics of the disease. The concept of phenotype applied to COPD has resulted in the definition of different types of patients with prognostic and therapeutic significance. In this way, we may take on a more personalized treatment according to not only the severity of the airflow obstruction, but also conditioned by the clinical phenotype. In this review, we give reasons for considering three fundamental phenotypes: exacerbator, mixed and emphysema. It is evident that not all the patients will meet the criteria to be able to classify them unequivocally into one of the subgroups, and it will always be the physician’s clinical judgment which will classify the patient into the most relevant phenotype for its prognosis. In this direction, the simple question of, “How many exacerbations did the patient have the previous year?” will classify the patient as an exacerbator if the answer is “two or more”, whatever the clinical or functional characteristics of the patient, as the treatment should prioritize the prevention of exacerbations. If the response is “one or none”, we should confirm whether the patient is either an emphysema or mixed phenotype. The following step will be to recognize these clinical phenotypes in the new COPD treatment guidelines.

Conflict of Interests

Marc Miravitlles has received professional fees for scientific consulting and/or for giving conferences from Almirall, AstraZeneica, Bayer Schering, Boehringer Ingelheim, Grupo Ferrer, GlaxoSmithKline, Laboratorios Esteve, Pfizer, Novartis, Merck Sharp & Dhome and Nycomed. Myriam Calle has received professional fees for giving conferences from Almirall, AstraZeneica, Boehringer Ingelheim, GlaxoSmithKline, Pfizer, Novartis, Merck Sharp & Dhome and Nycomed. Juan José Soler Cataluña has received professional fees for scientific consulting and/or for giving conferences from Almirall, AstraZeneica, Boehringer Ingelheim, Ferrari, GlaxoSmithKline, Laboratorios Esteve, Pfizer, Novartis, Merck Sharp & Dhome and Nycomed.

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