

REVIEW

Open Access

Chronic obstructive pulmonary disease and glucose metabolism: a bitter sweet symphony

Aibek E Mirрахимов

Abstract

Chronic obstructive pulmonary disease, metabolic syndrome and diabetes mellitus are common and underdiagnosed medical conditions. It was predicted that chronic obstructive pulmonary disease will be the third leading cause of death worldwide by 2020. The healthcare burden of this disease is even greater if we consider the significant impact of chronic obstructive pulmonary disease on the cardiovascular morbidity and mortality. Chronic obstructive pulmonary disease may be considered as a novel risk factor for new onset type 2 diabetes mellitus via multiple pathophysiological alterations such as: inflammation and oxidative stress, insulin resistance, weight gain and alterations in metabolism of adipokines.

On the other hand, diabetes may act as an independent factor, negatively affecting pulmonary structure and function. Diabetes is associated with an increased risk of pulmonary infections, disease exacerbations and worsened COPD outcomes. On the top of that, coexistent OSA may increase the risk for type 2 DM in some individuals. The current scientific data necessitate a greater outlook on chronic obstructive pulmonary disease and chronic obstructive pulmonary disease may be viewed as a risk factor for the new onset type 2 diabetes mellitus. Conversely, both types of diabetes mellitus should be viewed as strong contributing factors for the development of obstructive lung disease. Such approach can potentially improve the outcomes and medical control for both conditions, and, thus, decrease the healthcare burden of these major medical problems.

Keywords: COPD, Dysglycemia, Insulin resistance, Obesity, Metabolic syndrome, Diabetes mellitus endothelial dysfunction, Vasculopathy

Introduction

Chronic obstructive pulmonary disease (COPD) defines a group of chronic inflammatory pulmonary disorders characterized by partially reversible airflow limitation [1]. The notion that COPD is primarily a lung disease has been challenged, and the broader definition of COPD as a systemic inflammatory syndrome has been proposed [2]. Evidence for this approach has been provided by strong associations with increased rates of cardiovascular diseases [3], anemia [4], musculoskeletal disease [5,6] and malignant conditions seen in COPD patients [7,8].

Moreover, it is predicted that COPD will be the third leading cause of death worldwide by 2020, which will impose an even greater burden on the healthcare system globally [9]. If the mortality from comorbid conditions associated with COPD is taken into count, then this

disease poses an even greater impact on health outcomes. The common risk factors for COPD are air pollution, aging, respiratory infections, bronchial asthma (BA), low socioeconomic status [1]. The COPD grading is presented in Table 1.

Type 2 diabetes mellitus (DM) and metabolic syndrome (MetS) are particularly common medical disorders and are leading causes of morbidity and mortality worldwide. Disturbances in glucose metabolism are more common in COPD patients than in COPD free individuals. COPD, MetS and type 2 DM are associated with advanced age, which may in part explain this observation. It is well known that offspring of affected parents are more likely to develop both COPD and type 2 DM. Smoking during pregnancy can in part explain the association between COPD and type 2 DM, due to delivery of low birth weight infants, which is a known risk factor for both diseases [10]. In addition, smoking can contribute to the occurrence of these diseases via its effects on

Correspondence: amirрахимов1@gmail.com
Kyrgyz State Medical Academy named by I.K. Akhunbaev, Akhunbaev street 92, Bishkek 720020, Kyrgyzstan

Table 1 Global Initiative for Chronic Obstructive Lung Disease (GOLD) Grading system for COPD (Adapted from reference #) [1]

Grade	Severity	Spirometry findings
GOLD 1	Mild	FEV ₁ /FVC<0.7 and FEV ₁ ≥80% predicted
GOLD 2	Moderate	FEV ₁ /FVC<0.7 and 50%≤FEV ₁ ≥80% predicted
GOLD 3	Severe	FEV ₁ /FVC<0.7 and 30%≤FEV ₁ ≥50% predicted
GOLD 4	Very Severe	FEV ₁ /FVC<0.7 and FEV ₁ >30% predicted

Abbreviations: **GOLD:** Global Initiative for Chronic Obstructive Lung Disease; **FEV₁:** Forced Expiratory Flow in 1 second; **FVC:** Forced Vital Capacity.

systemic inflammation and oxidative stress [11,12]. However, the pathogenesis of glucose metabolism dysregulation is likely to be much more complex, whereby myriads of pathways are likely to be implicated, and much is still to be discovered and clarified.

On the other hand, type 1 DM and other forms of DM are much less common than type 2 DM [13], in part due to the growing burden of obesity and its associations with type 2 DM [14]. The risk factors for type 1 DM are not well understood and are of minor relevance to this manuscript. The major risk factors for type 2 DM are obesity, sedentary lifestyle, family history/genetic predisposition, aging, MetS, gestational diabetes and smoking.

Epidemiological data suggest that certain comorbid diseases, including DM are much more common in patients with COPD than in controls [3,15], and are associated with a worse COPD outcome [16]. As shown by Cazzola et al., patients with COPD have a higher burden of type 2 DM [17]. Sode et al. showed that patients with COPD have a greater burden of not only DM, but also of myocardial infarction, lung cancer, depression and hip fracture [18]. In contrast to these findings, Korean researchers did not find any association between COPD and greater DM prevalence, which may be related to high percentage of underweight subjects in the studied population [19].

The goal of this article is to summarize the published data on COPD and dysglycemic states. Firstly, the background of the problem and possible pathophysiological mechanisms will be discussed. Secondly, the article will focus on the clinical data and epidemiology of type 2 DM in COPD and the impact of both conditions on the clinical course of each other. Thirdly, we will briefly discuss the data on overlap syndrome, coexistence of obstructive sleep apnea (OSA) and COPD on the risk of type 2 DM.

Pathophysiology of glucose metabolism disturbances in COPD

Weight as a risk factor for COPD

Obesity is a growing health concern worldwide, primarily due to modern lifestyle changes and sedentarism. Obesity

is one of the major criteria of MetS and a well established risk factor for new onset type 2 DM [20].

Obesity is known to affect pulmonary function and lung volumes. Obesity is associated with a decrease in expiratory reserve volume (ERV) and functional residual capacity (FRC), due to its extrapulmonary restrictive component [21]. Aside from this, obesity can perpetuate both systemic and pulmonary inflammation, since excessive adipose tissue is able to produce various proinflammatory cytokines including interleukin-6 (IL-6) and tumor necrosis factor alpha (TNF-α). On the other hand, excessive central adiposity is associated with a decrease in adiponectin levels, which is known for its anti-inflammatory properties, and this can contribute to the pulmonary and vascular damage [22].

Leone et al. studied the association between the components of MetS and airflow obstruction in 129,965 men [23]. MetS was associated with low forced expiratory volume in one second (FEV₁) (odds ratio (OR) 1.28; 95% confidence interval (CI) 1.20-1.37) and forced vital capacity (FVC) (OR 1.41; 95% CI 1.31-1.51), independently from possible confounding factors. Abdominal obesity was strongly associated with low FEV₁ (OR 1.94; 95% CI 1.80-2.09) and FVC (OR 2.11; 95% CI 1.95-2.29). It is particularly pertinent to note that hyperglycemia, high blood pressure and dyslipidemia were related to airflow obstruction. This finding indirectly supports a potential impact of MetS and type 2 DM on the pathogenesis and clinical course of COPD.

Similar results were obtained in a study by Lam et al., who enrolled 7,358 adults [24]. These researchers showed that abdominal obesity was associated with airflow obstruction independently from smoking (OR 1.43; 95% CI 1.09-1.88). However, a study performed by Paek et al., who enrolled 4,001 individuals failed to document any relationship between MetS and airflow limitation [25].

Marquis et al. enrolled 38 subjects with COPD and 34 controls to test whether MetS and its components were more common in COPD patients [26]. Overall 47% of COPD patients had three or more MetS components in comparison with 21% of control subjects, and the prevalence of obesity was approximately two times higher in the COPD group.

Steuten et al. recruited 317 subjects with COPD to study the prevalence of weight distribution in COPD patients in a Dutch primary care setting [27]. Obesity was much more prevalent in mild to moderate COPD (16-24%) disease than in severe disease (6%) in this study.

A research group from the University of California in San Francisco enrolled 355 patients with COPD to study the impact of body composition on functional status in patients with various stages of COPD [28]. Obesity (BMI≥30 kg/m²) was diagnosed among 54% of

study participants, which is much higher than in the general US population.

Guerra et al. in a nested case control study tested whether BMI was associated with emphysema and bronchitis phenotypes of COPD [29]. In subjects with bronchitis, 25% of COPD cases had a BMI ≥ 28 kg/m² compared with 16% of controls. Moreover, a BMI ≥ 28 kg/m² was associated with an increased risk of being diagnosed with bronchitis (OR 1.80; 95% CI 1.32-2.46), whereas, the presence of emphysema was related to decreased weight.

However, some previously published data suggests that low BMI is a risk factor for COPD [30] and others have reported that obese men have a lower annual FEV₁ decline [31].

It is well known, that a low BMI in patients with COPD is considered to be a risk factor for all cause related and pulmonary related mortality [32]. This was first demonstrated by Celli et al., who validated a globally used prognostic tool, more known as the BODE Index. Others have validated three item questionnaire for a poor COPD related quality of life, with a low BMI being one of the criteria [33]. However, it is worth emphasizing that a decrease in BMI seen in advanced COPD is likely to be explained by a loss of fat free mass, such as skeletal muscle [34]. Skeletal muscle loss in advanced COPD may further contribute to the deterioration in the clinical course of the disease, which is pathophysiologically similar to cancer cachexia.

Landbo et al. recruited 1,218 men and 914 women with FEV₁/FVC < 0.7 and prospectively studied the impact of nutritional status on mortality in COPD from all causes [35]. Importantly, smoking status, spirometry data, BMI and respiratory symptoms were assessed at baseline and after 17 years. After adjustment for possible confounders, a lower BMI was associated with greater mortality compared with a normal or high BMI (relative risk (RR) 1.64; 95% CI 1.20-2.23); this association was strongest in patients with COPD.

The same research group tested whether a change in BMI was associated with mortality in a cohort of patients with COPD and controls [36]. Interestingly, a loss of > three BMI units was associated with increased all-cause mortality in controls (RR 1.63; 95% CI 1.38-1.92) as well in COPD patients (RR 1.71; 95% CI 1.32-2.23). Weight loss was related to an increased risk of COPD related mortality (RR 2.14; 95% CI 1.18-3.89), whereas weight gain was associated with increased mortality only in controls.

Jee et al. studied 1,213,829 Koreans to prospectively study the impact of weight on mortality in patients with COPD [37]. These researchers found that a lower BMI was associated with increased mortality from respiratory causes, whereas a relationship was observed between

high BMI and increased mortality from cardiovascular diseases or malignancies.

These data highlight the unresolved issue of the "obesity paradox" seen in many chronic diseases, such as COPD and heart failure (HF), where a higher BMI is associated with a decrease in overall mortality. The discrepancies among COPD studies are likely because of different phenotypes of disease being enrolled, such as emphysema predominant and bronchitis predominant. Indeed, emphysema predominant phenotype is associated with a loss of skeletal muscle, rather than a loss of body fat, which may explain differences in mortality in patients with normal and low BMI. Thus, future studies are needed to address the fat free mass rather than BMI in studying the impact body weight on morbidity and mortality in patients with COPD. Indeed, loss of a skeletal muscle may explain a greater pulmonary function decline in patients with low BMI compared to patients with normal BMI as was shown by Watson et al. [31].

Adipokines and COPD

Adipose tissue is an active endocrine organ producing various substances, which regulate biological processes including glucose and insulin metabolism. Leptin and adiponectin are the most studied adipokines to date. Leptin may decrease the expression of insulin as well as decrease glucose uptake by pancreatic β cells. Adiponectin, on the other hand, may upregulate insulin secretion when it is physiologically required. The reader is referred to some comprehensive review articles on the topic of adipokines and glucose metabolism [38,39]. A summary of the adipokines and their relationship to DM and COPD is presented in Table 2.

Overall there is a paucity of published data on COPD and adipokines. Some of the published reports suffer from flaws in assessing the effects of adipokines on lung function and on the course of COPD. Nevertheless, the current data on this topic will be briefly reviewed.

Broekhuizen et al. enrolled 14 patients with moderate COPD to study the relationship between sputum leptin and other inflammatory markers [40]. Sputum leptin levels were related to sputum C-reactive protein (CRP) ($p < 0.001$) and TNF- α ($p < 0.01$).

Bruno et al. recruited 15 smokers without COPD and 27 smokers with mild to moderate COPD to study the pulmonary expression of leptin and its receptor [41]. They found that patients with COPD had increased pulmonary leptin expression, which is associated with air-flow inflammation and obstruction.

A research group from Maastricht University, in the Netherlands enrolled 15 male subjects with bronchitis predominant COPD and 27 male subjects with emphysema predominant disease to study the association between systemic inflammation, leptin and energy balance

Table 2 Key proinflammatory cytokines and adipokines and their potential role in COPD and DM (Adapted from references #) [38-77]

Name	Major origin	Physiological function	Data on COPD	Data on DM
CRP	Liver/Hepatocyte	Proinflammatory.	COPD is independently associated with increased levels of CRP. Moreover, CRP may predict the future onset of COPD.	Elevated CRP levels may predict the development of onset of type 2 DM.
TNF- α	Macrophages, other leukocytes and adipocytes	Proinflammatory and proapoptotic. Possible insulin antagonism.	COPD is independently associated with increased levels of TNF- α	May be a risk factor for the development of new onset type 2 DM.
IL-1	Macrophages, other leukocytes, dendritic cells etc.	Proinflammatory. Lymphocyte activation	IL-1 is implicated in the pathogenesis of COPD related inflammation.	An increase in IL-1 β may predict the development of new onset type 2 DM.
IL-6	Liver/Hepatocyte, macrophages, other leukocytes, adipocytes etc.	Proinflammatory. Upregulation of the synthesis of CRP and other proinflammatory cytokines in the liver.	COPD is independently associated with increased levels of IL-6.	IL-6 was shown to increase the risk for the new onset type 2 DM.
Fibrinogen	Liver/Hepatocyte	Proinflammatory. Active participation in coagulation.	COPD is independently associated with increased fibrinogen levels.	No data on fibrinogen and the risk of new onset type 2 DM.
Leptin	Adipocyte	Appetite regulation. Possible proinflammatory actions.	Leptin levels are increased in patients with COPD. May contribute to COPD related weight loss and pulmonary function decline.	Leptin may increase the risk of type 2 DM. Leptin may participate in the development of DM related complications via its proinflammatory actions.
Adiponectin	Adipocyte	Antiinflammatory. Increase in insulin synthesis and increase in insulin sensitivity.	Adiponectin levels are increased in patients with COPD and low BMI, which may explain decreased mortality from cardiovascular causes in advanced COPD.	Adiponectin may prevent the development of type 2 DM via its anti-inflammatory and proinsulin actions.
Resistin	Leukocytes	Proinflammatory and insulin antagonizing actions.	Resistin levels may be increased in COPD and mediate IR.	Resistin may directly participate in the development of IR.

Abbreviations: *CRP*: C-Reactive Protein; *DM*: Diabetes Mellitus; *IL*: Interleukin; *TNF- α* : Tumor Necrosis Factor alpha.

in COPD [42]. Emphysematous patients had lower leptin concentration, most likely due to lower body weight ($p= 0.02$). A positive correlation between serum leptin and soluble TNF- α receptor levels was observed in patients with emphysema.

Opposing results were obtained by Takabatake et al., who recruited 31 people with COPD and 15 controls [43]. The mean BMI in COPD patients was significantly lower than in controls (BMI = 18.1 \pm 2.7 kg/m² versus 22.8 \pm 2.2 kg/m² respectively; $p < 0.0001$), which may explain the lower leptin levels seen in COPD patients ($p < 0.05$). These investigators failed to find any association between leptin levels and inflammatory biomarkers.

Karakas et al. enrolled 30 subjects with COPD and 20 healthy controls to assess the possible association between serum leptin levels and body composition [44]. Patients were grouped based on their BMI and serum leptin concentration. These researchers found an inverse relationship between serum leptin levels and BMI.

Wang et al. enrolled 57 patients with stable COPD and 31 controls to study the association between nutritional status and both leptin and resistin levels [45].

Leptin levels were positively correlated with resistin levels and both were associated with lower BMI. In support of this notion, a study performed by Al Mutairi et al., who studied the association between circulating resistin and inflammation showed a direct association between resistin levels and inflammatory obstructive airway disease [46]. These researchers speculated that resistin may be a novel disease marker and play a role in the development of insulin resistance (IR) in COPD.

Hansel et al. enrolled 429 subjects to study the relationship between leptin receptors and the rate of pulmonary function decline in patients with COPD [47]. They identified 21 single nucleotide polymorphisms (SNP) of the leptin receptor gene which were significantly associated with accelerated loss of lung function. This study provided information that certain leptin receptor gene polymorphisms may act as predisposing factors for COPD incidence among smokers, which could explain why not all smokers develop obstructive airway disease.

Calikoglu et al. recruited 26 patients with stable COPD, 16 patients with COPD exacerbation and 15

controls to study the association between leptin, TNF- α and nutritional parameters [48]. These investigators found that leptin levels were lower in patients with stable COPD in comparison with controls; however, TNF- α was higher in stable COPD than in controls and COPD exacerbation was associated with an increase in serum leptin and TNF- α . Dutch researchers found similar results with higher leptin levels during COPD exacerbation. It was speculated that this phenomenon could be explained by systemic inflammation and/or through high dose systemic corticosteroid treatment [49]. It was proposed that leptin may participate in the pathogenesis of pulmonary cachexia through its anorexigenic and metabolic effects [42].

Pan et al. enrolled 56 patients experiencing COPD exacerbation to study the association between dysglycemia, IR and in hospital clinical course [50]. The authors concluded that leptin induced IR and hyperglycemia prolonged hospital stay and adversely affected pulmonary function. Kythreotis et al. found that leptin levels were elevated during exacerbation and 2 weeks after the discharge [51].

However, an association between leptin and TNF- α was not found in a study performed by Yang et al. [52]. Small sample size and the population studied may underlie the lack of association in this work compared to other studies.

Tomoda et al. recruited 31 patients with COPD and 12 controls to study the role of adiponectin in patients with COPD [53]. Plasma adiponectin was higher among patients with COPD and correlated with residual volume (RV) ($p < 0.05$), TNF- α ($p < 0.05$) and was inversely correlated with BMI ($p < 0.01$). These researchers speculated that high adiponectin levels may explain why COPD patients with a low BMI have decreased mortality due to cardiovascular causes.

Kirdar et al. enrolled 36 patients with COPD exacerbation and 17 controls to assess the role of adiponectin as a potential COPD biomarker [54]. Serum levels of adiponectin were higher among COPD patients ($p < 0.001$); however, leptin metabolism was not altered in COPD exacerbation.

Miller et al. reported that patients with emphysema had a higher expression of pulmonary adiponectin [55], whereas Summer et al. showed that mice lacking adiponectin were prone to emphysema development [56].

Yoon et al. measured adiponectin levels in patients enrolled in the Lung Health Study to investigate the relationship between this adipokine and COPD [57]. These researchers found that increased adiponectin was related to a decrease in cardiovascular mortality, but was associated with an increase in mortality due to respiratory causes.

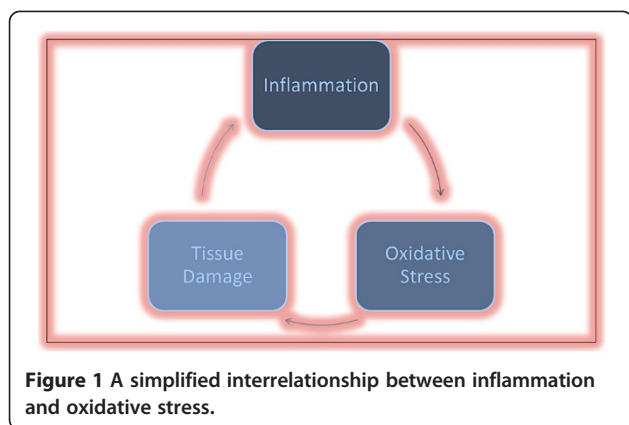
Breyer et al. studied the various adipokines and their expression in patients with COPD [58]. These researchers showed that patients with COPD had higher levels of CRP, IL-6, fibrinogen and adiponectin. Gender specific analysis demonstrated, that males with COPD had higher CRP, IL6 and fibrinogen levels, whereas, females with COPD had higher levels of CRP and fibrinogen only. Moreover, in female patients with COPD, leptin was correlated with CRP and fibrinogen, but this correlation was not observed in males with COPD. A negative correlation was shown between adiponectin and CRP in patients with COPD.

Based on the current data, leptin may propagate both pulmonary and systemic inflammation and together with resistin contribute to the pathogenesis of related dysglycemia. Data on the adiponectin and mortality is quite intriguing and conflicting, which merits further research to explore potential mechanisms. In a mouse model of emphysema, it was shown that adiponectin may protect against the emphysema development. Thus, more studies are needed on the role of adipokines in the development and course of COPD, and whether they are not simply the markers of COPD progression.

COPD, systemic inflammation and oxidative stress

As in many other chronic medical conditions, COPD is associated with low grade systemic inflammation. Various factors can contribute to this finding such as the spill of inflammatory mediators from the pulmonary system into the circulation, hypoxia, effects of obesity and hormonal disturbances. Current evidence suggests that excessive oxidative stress can be a risk factor for new onset type 2 DM and conversely, oxidative stress may be a consequence of new onset type 2 DM [59]. COPD as well as other pathologies, in which hypoxia is a feature is associated with an excessive oxidative state [60]. The reader is referred to a comprehensive review of hypoxia and metabolism for a more detailed discussion of this topic [61]. A summary of some of the key proinflammatory mediators and their relationship with DM and COPD is presented in Table 2. A simplified interrelationship between inflammation and oxidative stress is presented in Figure 1.

It is believed that inflammation increases the risk of future type 2 DM. Pradhan et al. studied 27,628 females to assess the impact of inflammatory biomarkers on the new onset type 2 DM [62]. They group found that the highest quartile of IL-6 and CRP gave a RR of 2.3 (95% CI 0.9-5.6) and 4.4 (95% CI 1.5-12.0) respectively. Spranger et al., studied 27,548 individuals to assess the relationship between inflammation and new onset type 2 DM [63]. These researchers found that interleukin-1 β (IL-1 β) and IL-6 may be useful predictors of the development of new onset type 2 DM.



Landmark studies performed by Zinman et al. [64] and Hu et al. [65] showed a positive association between TNF- α and the development of IR and overt type 2 DM. Similar data came from the Atherosclerosis in Communities Study, which showed that low grade systemic inflammation predicts the incident type 2 DM [66].

Botelho et al. demonstrated that interleukin-1 α (IL-1 α) plays a major role in the development of neutrophilia in smoke induced inflammation in a mouse model [67].

Altaran et al. enrolled 50 patients with COPD to study the association between IL-6 and COPD severity [68]. These researchers showed an increase in IL-6 among patients with COPD and a possible independent contribution of IL-6 to airflow limitation. However, the results of the study cannot be generalized, since the majority of study participants had a history of sulfa mustard exposure. He et al. showed that a SNP in the IL-6 gene 174G/C was associated with a rapid decline in FEV₁ and COPD susceptibility among smokers [69]. Van Durme et al. showed that an increase in IL-6 was associated with an increased risk of COPD development; however, no association was found between polymorphisms in the IL-6 gene and such risk [70]. Data from further study, performed by Yanbaeva et al. supported the hypothesis of an increased risk of future COPD development associated with high IL-6, particularly for the IL-6 H2 haplotype [71]. Tkacova et al. studied the adipose tissue expression of CD40, a key receptor for the TNF in 20 patients with COPD [72]. These researchers showed that patients with severe COPD had a higher expression of CD40, which was inversely related to a lower partial pressure of oxygen. Thus, COPD related adipose tissue dysfunction may further enhance systemic inflammation and IR.

According to a meta-analysis performed by Gan et al., patients with COPD have higher levels of circulating neutrophils, CRP, fibrinogen and TNF- α . [73]. Vozarova et al., showed that a higher white blood cell count was

associated with an increased risk of type 2 DM development [74]. Aside from this, elevated CRP has been shown to predict hospitalization and death in COPD [75], and CRP levels in COPD have a strong genetic background [76]. Moreover, a high CRP levels may act as a risk factor for future development of COPD among smokers [71,77]. The fact that only a small number of smokers develop COPD may be based on genetic alterations of common proinflammatory cytokines such as CRP and IL-6.

Indeed, the genetically mediated proinflammatory state in some individuals with COPD may underlie the clinical phenotype of patients with frequent COPD exacerbations. All of the findings may help in our understanding of the increase in cardiovascular morbidity and mortality, as well as the increased metabolic risk seen in COPD.

Both DM and COPD are associated with systemic inflammation and, on the other hand, systemic inflammation is associated with the disease risk, leading to a vicious cycle. Indeed, it was shown that certain genetic polymorphisms of common pro-inflammatory molecules may increase the risk for COPD in smokers.

Hypoxia and glucose metabolism

Pallayova et al. hypothesized that pancreatic β cells may be damaged by hypoxia, but the focus of their study was primarily on obstructive sleep apnea, which is associated with intermittent hypoxia, rather than chronic hypoxia seen in COPD [78]. However, both of these diseases may share some pathophysiological pathways, which may be mediated by the hypoxia inducible factor family (HIF) [61].

Based on the available data it is likely that different tissues have various metabolic responses to hypoxia. Interestingly, skeletal muscles under hypoxia have a higher glucose uptake [79], and increased sensitivity of skeletal muscles to insulin during exercise [80]. However, patients with COPD receiving oxygen therapy have less muscular IR [81]. This may be explained by the fact that long standing hypoxia may detrimentally affect skeletal muscle insulin sensitivity compared with acute muscular hypoxia seen in exercise.

HIF-1 α expression is increased in hypoxic pancreatic β cell regions, undergoing programmed cell death, which may simply indicate an effect of hypoxia on the expression of HIF-1 α , with the latter having no role in the pathogenesis of apoptosis. Alternatively HIF-1 α may exert unique effects on the β cells [82]. Cheng et al. showed that HIF-1 α has significant and beneficial effects on pancreatic β cell function, and hypothesized that HIF-1 α may be a novel target to improve the function of insulin producing cells [83].

Important results have arisen from a study performed by Oltmanns et al., which exposed healthy volunteers to hypoxia under euglycemic clamp [84]. These researchers found that hypoxia caused glucose intolerance, and this phenomenon could be partly explained by an increase in epinephrine release.

The effects of hypoxia on the function of adipocytes seem to be straightforward. Chen et al. showed that hypoxia decreased the production of adiponectin and increased the production of plasminogen activator inhibitor-1 which may contribute to a greater cardio-metabolic risk [85].

Aside from this, a hypoxia mediated increase in HIF-1 α can induce adipose tissue fibrosis and resistance to insulin [86,87].

On the other hand, hyperglycemia is able to down-regulate HIF-1 α , and via this mechanism contributes to the pathophysiology of angiopathic conditions commonly seen in patients with DM [88].

Based on above findings, it is likely that hypoxia can mediate its detrimental effects on glucose metabolism and IR at least through the effects on insulin sensitivity at the level of adipose tissue and skeletal muscle.

Muscle metabolic pathways in COPD

IR at the level of skeletal muscle plays a role in the pathogenesis of type 2 DM. Earlier reports linked this phenomenon to a decreased amount of mitochondria in diabetic skeletal muscle, but new data suggest a decreased functional status of this cellular organelle [89]. From a theoretical point of view, it is plausible that COPD and associated hypoxia, can detrimentally affect mitochondria. However, it should be noted, that the evidence regarding the role of hypoxia in decreased glucose transport in skeletal muscle is controversial [90]. The information on this topic is relatively scant and will be briefly discussed below.

Wuyam et al. showed that patients with COPD had decreased aerobic capacity compared to healthy controls [91]. However, in contrast to their findings, others explained this association by a decreased level of physical activity at baseline, and a shift in type of muscle fiber, and, they, therefore, doubted the existence of an intrinsic mitochondrial problem in COPD [92].

Jakobsson et al. group reported that patients with advanced COPD had upregulated glycolysis in quadriceps femoris muscles, but lower muscular aerobic capacity [93]. This finding points toward the existence of an adaptation of skeletal muscle to hypoxia, by upregulating glucose carriers, such as GLUT-4.

COPD patients experience a shift in the skeletal muscle fiber [94], which may explain a decrease in oxidative capacity [91,95]; such a fiber shift has been shown to be associated with a greater COPD severity [96]. Patients

with COPD have impaired β oxidation of fatty acids, electron transport chain and citric acid cycle, all of which are known to take place in mitochondria [97].

A key study performed by Green et al. showed that muscles from patients with COPD had lower levels of glucose transporter-4 (GLUT-4) compared with controls, which is known to be closely linked to insulin action [98]. However, as mentioned previously, others have failed to show any detrimental effects of hypoxia on glucose uptake by skeletal muscle cells.

Moreover, IR may synergistically impair muscle function, since it is associated with decreased muscle strength, even in patients without overt DM [99].

It is theoretically plausible, that longstanding hypoxia have different molecular effects on skeletal muscle metabolism compared with muscle hypoxia induced by physical exercise in subjects without pulmonary disease. However, this conclusion is not based on firm evidence and further research is needed to clarify the effects of hypoxia and COPD on glucose metabolism in skeletal muscle.

COPD and hormonal dysregulation

From a theoretical viewpoint the alterations in insulin opposing hormones can explain the association between the development of COPD and dysglycemia. However, it should be noted that the evidence is scant and controversial. Nevertheless, we will briefly discuss the data on COPD and hormones, which are associated with MetS and type 2 DM.

There are some reports linking abnormalities in androgen metabolism with the development of type 2 DM [100]. The exact mechanisms of these associations are not clear, but they may be related to the beneficial effects of testosterone on weight, insulin sensitivity and modulation of inflammation. Svartberg et al. showed that men with lower levels of free and total testosterone had lower numbers of FEV₁ and FVC, independently from potential confounders [101].

Laghi et al. enrolled 101 patients with COPD and measured both free and total testosterone [102]. These researchers found that approximately 40% of patients with COPD were hypogonadal; however, they did not find any association between testosterone levels and pulmonary function. Notably, the prevalence of hypogonadism in patients with COPD has been shown to be greater than in an age-matched population without this disease [103], and the lack of association between androgens and pulmonary function could be explained by the small sample size and lack of a control group in this study.

In contrast to the study performed by Laghi et al. [102], Van Vliet et al. showed that hypogonadism was associated with decreased physical endurance, muscle

weakness and proinflammatory state among patients with COPD [104]. It is difficult to prove causality between low testosterone and COPD, since multiple mechanisms are likely to act simultaneously. However, it is possible that hypoxia may underlie the development of hypogonadism in patients with pulmonary disease [105].

Catecholamine hormones are known to be insulin antagonists and contribute to the occurrence of hyperglycemia [106]. Kanstrup et al. showed an association between prolonged hypoxia and the development of enhanced catecholamine output in eight healthy men [107]. In another study, Scavini et al. showed that patients with COPD had higher catecholamine levels which were independently related to a decrease in FEV₁ [108]. Bratel et al. showed that patients with severe nocturnal hypoxemia treated with long term oxygen therapy experienced a reduction in nocturnal norepinephrine, suggesting a potential causal role of hypoxia in the overactivation of the sympathetic nervous system in patients with COPD [109].

Abnormalities in the renin angiotensin aldosterone system (RAAS) are implicated in the development and pathogenesis of cardiovascular diseases, MetS and type 2 DM [110]. Farber et al., in two published studies, showed that patients with COPD had higher levels of aldosterone and an increase in plasma renin activity [111,112]. Stewart et al. found that patients with COPD had increased levels of vasopressin and aldosterone, which could be explained by COPD, related autonomic neuropathy [113]. Alternatively, the RAAS activity in patients with COPD can be upregulated by increased sympathetic tone, hypoxia and underlying comorbid states such as cardiac and renal pathologies. The importance of RAAS overactivation was highlighted by Mancini et al., who showed that drugs targeting the RAAS may improve survival in patients with COPD [114].

Vitamin D is implicated into the pathogenesis of myriad of extraskeletal diseases including both major types of DM [115,116]. This relationship can be explained by the anti-inflammatory and antioxidant effects of vitamin D as well as through its modulation of the RAAS activity. Moreover, patients with COPD exhibit a trend towards lower levels of vitamin D [117]. However, it is difficult to conclude causality, since patients with COPD may have a lower level of activity and, thus, have less sun exposure.

Evidence for an association between COPD and thyroid function is scant and predominantly negative [118].

To summarize, COPD is associated with abnormalities in metabolism of androgen hormones, vitamin D, catecholamines and RAAS. However, more studies are needed to clarify the role of these hormonal systems in the pathogenesis of COPD related IR.

Clinical data on COPD and the risk of DM/MetS COPD, reduced lung function and the risk of new onset type 2 DM

Reduced lung function and the risk of type 2 DM
Lazarus et al. in a prospective observational study showed that baseline values of FVC, FEV₁ and maximal mid-expiratory flow rate (MMEF) are negatively associated with IR even after adjustment for potential confounders [119].

Ford et al. showed that FEV₁ and FVC values at baseline were inversely associated with the incidence of type 2 DM [120]. It is important to note that restrictive, but not obstructive lung disease was associated with the incidence of new onset type 2 DM. Similar findings were shown by Paek et al. and Wannamethee et al. [25,121], who demonstrated that restrictive lung diseases were strongly associated with new type 2 DM. These results were statistically significant even after adjustment for confounding factors such as age, gender and weight.

Researchers from the Johns Hopkins University studied 11,479 subjects free of type 2 DM and followed them for nine years [122]. They found that lower FVC was independently associated with new onset type 2 DM in both men and women, independently from potential confounders such as age, weight and race. Hsiao et al. showed that reduced baseline FVC and FEV₁ were independently related to a greater risk of future development of MetS [123]. A shared pathophysiology may underlie this association or alternatively, reduced lung volumes may simply be the markers of lower physical endurance in patients at risk for the development of MetS.

Similar data have been reported by Engstrom et al., who showed that reduced pulmonary function was inversely associated with the development of type 2 DM [124]. In a later study, Engstrom et al. showed that low FVC was associated with an increased risk of IR and the development of new onset type 2 DM [125].

In a recent cross-sectional study, Kwon et al. showed that reduced FEV₁ and FVC are directly related to the new onset type 2 DM [126]. The results stayed significant even after adjustment for age, weight, health literacy and exercise.

Cross-sectional studies performed in Japanese [127] and Korean men [128,129] showed an association between reduced FVC and an increased risk of both MetS and type 2 DM.

However, the association between type 2 DM and reduced lung function should be interpreted with caution. It is well known that most of the participants enrolled in these studies were obese, and obesity is associated with reduced respiratory muscle performance, which could underlie this relationship [130,131].

Heianza et al. observed 5,346 men without DM or pulmonary disease to study the impact of decreased lung function on the incidence of type 2 DM [132]. They showed that reduced pulmonary function was associated with increased incidence of type 2 DM independently from BMI, smoking status and baseline glycated hemoglobin (hazard ratio (HR) 1.3; 95% CI, 1.01-1.26).

Thus, the observed association could be explained by three possibilities: reduced lung performance and incidence of type 2 DM share similar pathophysiological pathways (such as low grade inflammation); secondly, that reduced lung performance could be just a marker of reduced physical endurance; thirdly, that reduced lung function may simply portray the overall health status of individuals prone to develop type 2 DM, with obesity being the most important confounding factor.

A summary of the key studies on the topic of reduced pulmonary function and a risk of type 2 DM is presented in Table 3.

COPD and the risk of type 2 DM Several studies assessed the impact of COPD on the risk of new onset type 2 DM. Rana et al. analyzed data from the Nurses' Health Study from 1988–1996 which enrolled 103,614 females [133]. Study participants were followed for eight years, and COPD was found to have a multivariate RR of 1.8 (95% CI 1.1-2.8) for new onset type 2 DM. It is vital to note that the diagnosis of BA was not associated with an increased risk of type 2 DM. However, these results should be interpreted with caution, since spirometry data were not available and the diagnosis of pulmonary disease was based on the patients' questionnaires.

Table 3 Key studies on pulmonary volumes and the risk of new onset type 2 DM

First author and year	Country	Study design	Population studied	Findings
Lazarus et al. [119]; 1998	USA	Prospective cohort study with a mean follow up of 20.9 years	n:1,050 men (with no self-reported DM) included in the final analysis mean age: 41.4 years mean BMI: 25.6 kg/m ²	Reduced FVC, FEV ₁ and MMEF were associated with greater fasting insulin and fasting insulin resistance after logistic regression analysis.
Engström et al. [124]; 2002	Sweden	Prospective cohort study with a mean follow up of 13 years	n: 382 non-diabetic men age at the beginning of the study:55 years mean BMI: 24.4-24.7 years (depends on the pulmonary VC subgroup)	15 new cases of type 2 DM were diagnosed during the follow up. DM and log glucose were inversely associated with baseline VC.
Engström et al. [125]; 2003	Sweden	Population based cohort study with a mean follow up of 13.9 years for men and 9.4 years for women	n: 2,332 non-diabetic subjects (men-1,436, 61.57%) mean age for men: 44.0-44.9 years (depends on the pulmonary FVC subgroup) mean age for women: 49.4-50.4 years (depends on the pulmonary FVC subgroup) mean BMI for men: 24.2-24.9 kg/m ² (depends on the pulmonary FVC subgroup) mean BMI for women: 23.2-23.9 kg/m ² (depends on the pulmonary FVC subgroup)	The association between baseline FVC and IR at follow up remained statistically significant after adjustment for gender, age at screening, follow up time, smoking/ tobacco consumption, BMI and waist to hip ratio, physical activity and baseline log glucose
Kwon et al. [126]; 2012	Korea	Prospective cohort study with a follow up of 5 years	n: 9,220 men non-diabetic at baseline mean age: 41.4 years mean baseline BMI: 24.4 kg/m ² for patients without type 2 DM at follow up and 26.7 kg/m ² for patients with type 2 DM at follow up	207 patients developed type 2 DM with the incidence of 2.2%. FEV ₁ and FVC were negatively associated with type 2 DM. In patients with BMI<25 kg/m ² the lowest quartile of FVC and FEV ₁ had OR of 2.15 (95% CI 1.02-4.57) and 2.19 (95% CI 1.09-4.42) for incident type 2 DM.
Heianza et al. [132]; 2012	Japan	Observational study with a 4 year follow up	n: 5,346 (100% men) mean age: 48.6 years mean BMI: 23.3 kg/m ²	Compared with the highest FEV ₁ quartile, lower quartiles were associated with 1.79-1.86 greater risk for the new onset type 2 DM (95% CI, 1.10-2.91).

Abbreviations: **BMI:** Body Mass Index; **CI:** Confidence Interval; **FEV₁:** Forced Expiratory Volume in 1 second; **FVC:** Forced Vital Capacity; **HR:** Hazard Ratio; **MMEF:** Maximal Mid-Expiratory Flow; **OR:** Odds Ratio; **VC:** Vital Capacity.

Song et al. analyzed data of the Women's Health Study in which 38,570 women without DM were enrolled [134]. As in the previous study, the diagnosis of pulmonary disease was assessed by the presence of self-reported disease. The study participants were followed for a median of 12.2 years. Women who had reported physician diagnosed COPD had a multivariate RR of 1.38 (95% CI 1.14-1.67) for new onset type 2 DM. In contrast to the previous study, these investigators found that BA was associated with an increased risk of type 2 DM with a RR of 1.37 (95% CI 1.2-1.57). Adapted data from the references 132 and 133 are presented in Figure 2.

Feary et al. analyzed the primary care records of 1,204,100 individuals aged over 35 years from 311 primary care practices in the United Kingdom [3]. They showed an OR of 2.04 (95% CI 1.97-2.12) for the development of new onset type 2 DM associated with physician diagnosed COPD.

Bolton et al. recruited 56 non-hypoxemic COPD patients and 29 healthy controls to study the potential association between inflammation and IR [135]. Patients with COPD had higher insulin levels, which were related to inflammatory markers such as CRP, IL-6 and soluble receptors for TNF- α . Indeed, the COPD related inflammatory state may underlie the pathogenesis of an increased risk of type 2 DM in these patients.

In contrast, Bayliss et al. failed to observe any effect of COPD exacerbation on worsened long term glycemic control in patients with type 2 DM [136]. However, as was acknowledged by these researchers, the studies using community electronic records are subject to some limitations such as recall and selection bias; thus, the results should be interpreted with caution.

Based on the current evidence, COPD should be approached as a risk factor for the new onset type 2 DM. This risk is based on myriad of pathological pathways discussed in this manuscript. A summary of the

key studies on COPD and a risk of type 2 DM is presented in Table 4.

Treatment of COPD and the risk of hyperglycemia

Patients with COPD are often treated with corticosteroids, either inhaled (typically stable COPD) or systemic (typically COPD exacerbation) [1]. Corticosteroid therapy is associated with a decreased rate of FEV₁ and pulmonary function decline and fewer disease exacerbations.

However, it is well known that systemic corticosteroid therapy is associated with multiple undesired effects, including the development of dysglycemia and overt DM [137]. Niewoehner et al. showed that systemic glucocorticosteroid therapy was significantly associated with the development of medically relevant hyperglycemia in 271 patients with COPD exacerbation [138]. Burt et al. showed that patients with COPD treated with systemic corticosteroids had a circadian rise of glucose predominantly in the afternoon and evening [139].

Despite being useful in the management of COPD exacerbation, long term systemic corticosteroid therapy is associated with the development of respiratory muscle damage with resultant decreases in both inspiratory and expiratory muscle strengths [140,141]. Detrimental effects of systemic corticosteroids on respiratory performance have been confirmed in patients with BA [142] and in patients with normal baseline pulmonary function [143].

In contrast to systemic corticosteroids, inhaled corticosteroids have much more favorable side effect profile, which is explained by the route of administration and a lower corticosteroid dose being administered [144]. A recently published pooled analysis of 34 studies using inhaled steroids in patients with BA or COPD failed to show any association with new onset type 2 DM or hyperglycemia [145]. However, higher doses of inhaled corticosteroids were associated with higher glucose levels in treated patients in one published study [146].

Thus, it is critically relevant to address the risk of new onset type 2 DM in patients with COPD being treated with high doses of inhaled corticosteroids due to a greater prevalence of other risk factors for DM in this population.

The impact of DM/Mets on the lung

Effects of DM on the pulmonary vasculature and diffusing capacity

DM is directly implicated in the development of vascular damage to both large and small arteries or macroangiopathy and microangiopathy respectively [147,148]. Therefore, it is plausible that long standing DM detrimentally affects the alveolar capillary bed. Key studies in this area will be briefly reviewed.

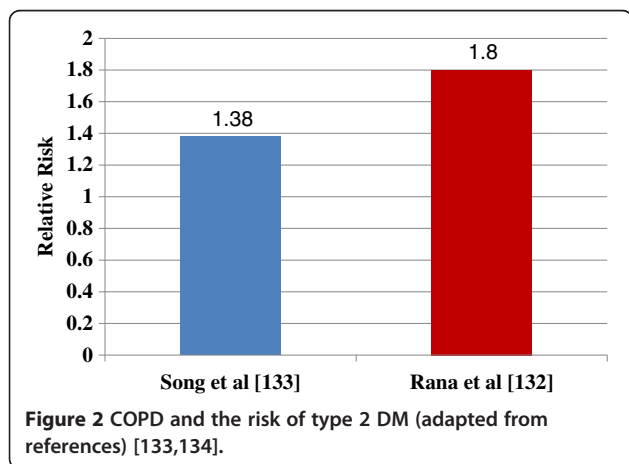


Table 4 Key studies assessing the risk of IR/new onset type 2 DM and control of DM in patients with COPD

First author and year	Country	Study design	Population studied	Findings
Feary et al. [3]; 2010	UK	Cross-sectional study using computerized records from 311 primary care practices	n: 1,204,100 (Men-618,090 subjects-51.3%) age: 35–44 years-327,800 subjects (27.2%), 45–54 years-274,375 subjects (22.8%), 55–64 years-254,472 subjects (21.1%), 65–74 years-177,102 subjects (14.7%) and ≥75 years- 170,361 subjects (14.2%)	COPD had OR of 2.04 (95% CI 1.97-2.12) for the development of new onset type 2 DM.
Song et al. [134]; 2010	USA	Prospective cohort study with a median follow up of 12.2 years	n:38,570 non-diabetic women mean age:-57.8 years(for patients with physician diagnosed COPD) mean BMI: 26.1 kg/m ² (for patients with physician diagnosed COPD) Data on BMI not provided	COPD had a multivariate adjuster RR of 1.38 (95% CI 1.14–1.67) for a new onset type 2 DM.
Bolton et al. [135]; 2007	UK	Cross-sectional study	n: 56 patients (30 men) with non-hypoxemic COPD and 29 healthy subjects (15 men) mean age: 66.7 years for COPD patients and 62.9 years for healthy controls mean BMI: 25.7 kg/m ² for COPD patients and 26.5 kg/m ² for healthy subjects	Patients with COPD had greater levels of CRP, IL-6 and TNF-α sr I and sr II. IR assessed by HOMA was 1.68 in COPD patients vs. 1.13 in healthy subjects. Log ₁₀ IL-6 and BMI were found to be predictive for IR in stepwise regression analysis
Bayliss et al. [136]; 2011	USA	Prospective cohort study in patients with established type 2 DM	2,332 develop chronic pulmonary disease exacerbation (women-1,244, 53.34%), 582 (women-318, 54.64%) developed cancer and 2,959 developed depression (women-1,670, 56.44%) mean age at index date: 62.04 years for depression, 63.2 years for chronic pulmonary disease exacerbation and 65.53 years for cancer mean BMI: 31.09 kg/m ² for cancer, 32.01 kg/m ² for depression and 32.45 kg/m ² for chronic pulmonary disease exacerbation	The researchers did not find any effect of chronic pulmonary disease exacerbation, depression and cancer on the glycated hemoglobin.

Abbreviations: **BMI:** Body Mass Index; **CI:** Confidence Interval; **COPD:** Chronic Obstructive Pulmonary Disease; **CRP:** C-Reactive Protein; **DM:** Diabetes Mellitus; **HOMA:** Homeostatic Model Assessment; **IL-6:** Interleukin-6; **OR:** Odds Ratio; **RR:** Relative Risk; **sr:** Soluble Receptor.

Guazzi et al. investigated the association between type 2 DM with or without HF and the worsening pulmonary diffusion capacity to carbon monoxide (DLCO) [149]. These researchers recruited 15 healthy controls and three groups 15 patients with type 2 DM, type 2 DM+HF and HF alone. They found that patients with both HF and type 2 DM had a lower level of DLCO, which is a known surrogate marker for the alveolar capillary membrane morphological and functional status. Subsequently, the researchers tested the effects of regular insulin on DLCO. They found that the administration of insulin improved DLCO in patients with type 2 DM and type 2 DM+HF, but not in patients with HF alone. The same group tested the effects of administered regular insulin on the alveolar gas conductance in 19 never smokers with type 2 DM [150]. They found an association between type 2 DM and decreased alveolar gas conductance. Administered insulin improved gas conductance, which was believed to occur through the decrease in alveolar capillary membrane impedance.

In fact, the observed beneficial effects of insulin on the pulmonary variables were attractive and further studies were carried out to explore an association with the use of inhaled insulin [151]. However, unfortunately, some studies pointed towards the potential adverse effects of inhaled insulin, in terms of the occurrence of a cough, and a potential reduction in DLCO and FEV₁ [152]. Some experts doubted the usefulness of inhaled insulin, due to an age associated decline in pulmonary function and reserve [153] and the fact that smoking can significantly alter the absorption of inhaled insulin [154]. However, in contrast a recent trial did not show any long term detrimental effects of inhaled insulin on pulmonary variables in patients with BA [155] and a second study did not find any harmful effects of inhaled insulin on pulmonary function in patients with type 2 DM [156]. Despite, these optimistic data, more research is needed before inhaled insulin can be recommended in diabetic patients with or without pulmonary disease.

Guvener et al. enrolled 25 patients with type 2 DM and 12 healthy controls to assess the impact of type 2 DM on alveolar capillary permeability, measured using the ratio of DLCO to alveolar ventilation (VA) [157]. The study demonstrated a statistically significant decrease in alveolar capillary capacity in patients with type 2 DM compared with healthy controls. Age, duration of type 2 DM and the presence of microalbuminuria were associated with the presence of alveolar capillary exchange abnormality. It is pertinent to note that microalbuminuria was predictive for the presence of pulmonary capillaropathy, and it is theoretically plausible that both diabetic related complications share a similar pathophysiology.

Researchers from the Southwestern University, USA enrolled 65 never smokers with type 2 DM to assess the status of pulmonary microvascular reserve [158]. Both non-obese ($BMI < 30 \text{ kg/m}^2$) and obese ($BMI \geq 30 \text{ kg/m}^2$) patients with type 2 DM had reduced levels of the measured pulmonary function variables during exercise. However, the observed changes in obese patients were not fully explained by the baseline lung volumes. Moreover, the presence of retinopathy, neuropathy, microalbuminuria and control of diabetes were associated with a reduced pulmonary microvascular reserve.

It is important to mention the findings of Minette et al., who demonstrated that never smokers with type 1 DM had a thicker alveolar capillary membrane [159]. However, these researchers failed to find any association between type 1 DM and worsened alveolar gas conductance. Scherthaner et al. were unable to find any adverse impact of type 1 DM on lung function [160]. However, it should be noted, that the cross-sectional design of the above studies pose limitations on the study conclusion, since the study participants could be tested at the time, when disease was at a subclinical stage.

In contrast to these negative studies of type 1 DM and lung diffusing capacity, the researchers from Russia demonstrated that a long standing type 1 DM was associated with reduced lung volumes and decreased alveolar capillary gas exchange [161]. However, it is important to keep in mind that patients with type 1 DM are generally younger and may have less pulmonary damage resulting from clinically detectable pulmonary disease compared with type 2 DM at the time of the actual study, due to a greater lifetime exposure to noxious stimuli and an age related physiological decline in lung function.

Asanuma et al. compared FVC and DLCO in patients with DM without pulmonary disease to healthy age-matched controls [162]. They found that patients with DM have lower DLCO and FVC in comparison with controls. This reduction was significantly related to the duration of DM and the presence of diabetic retinopathy.

Sandler et al. showed that patients with type 1 DM had a lower pulmonary capillary flow with a resultant decrease in DCLO, which was associated with the duration of DM [163]. Cooper et al. showed that older patients (>35 years) with type 1 DM had lower DLCO compared with controls, which might be explained by an underlying pulmonary microvascular disease [164].

Ramirez et al. enrolled 18 patients with type 1 DM and assigned them either to a standard insulin regimen or to intensive treatment with measurement of glycated hemoglobin at quarterly intervals [165]. These researchers followed study participants for six years with measurement of DLCO and FVC. Participants from the standard treatment group had higher levels of glycated hemoglobin and worse values of DLCO and FVC. The researchers speculated that the treatment of DM itself may prevent the development of pulmonary dysfunction pointing toward a causative role. In a later study, the same group supported these findings, by demonstrating diminished DLCO and greater breathing effort in patients with type 1 DM [166].

Strojek et al. showed that patients with type 1 DM related complications had lower levels of DLCO compared with patients with type 1 DM and no complications [167]. Similar results were found by Innocenti et al., who showed an association between urinary albumin excretion and diminished DLCO [168]. Schnack et al. showed that patients with type 1 DM and microalbuminuria had lower values of FEV_1 , vital capacity (VC) and DLCO compared with patients without microalbuminuria and to controls [169].

Saler et al. compared DLCO between 80 healthy subjects 44 patients with type 1 DM and 68 patients with type 2 DM [170]. DLCO and the ratio of DLCO to VA were significantly decreased in patients with both types of DM, but not in the control group. As shown in the above mentioned studies the urinary albumin excretion was inversely correlated with a decline in pulmonary diffusing capacity. It is essential to note that even children with type 1 DM have lower DLCO, which can be viewed as an early marker of DM related microangiopathy [171].

Sinha et al. showed that patients with type 2 DM and microangiopathy had lower DLCO in comparison with patients with type 2 DM and no microangiopathy [172].

Boulbou et al. enrolled 16 patients with type 1 DM, 33 patients with type 2 DM and 22 healthy controls to study the expression of adhesion molecules in all groups [173]. Patients with both types of DM had lower levels of DLCO corrected for alveolar ventilation and this was inversely related to increased expression of E-Selectin. Indeed, based on the current data DM can be considered as a risk factor for pulmonary capillaropathy with subsequent alteration in DLCO.

In contrast, a research group from Israel failed to find any association between the presence of DM and decreased DLCO or other pulmonary function parameters [174]. However, some study limitations, such as a small sample size and a relatively young age of the subjects may explain these negative results.

Robust data indicate that DM leads to pulmonary microangiopathy through the mechanisms similar to the DM related nephropathy. This is based on the fact the presence of DM related renal disease was significantly associated with the presence of pulmonary capillary dysfunction. A summary of the key studies on the impact of DM on pulmonary diffusing capacity is presented in Table 5.

Effects of DM on the respiratory muscles and physical endurance

DM is associated with the development of muscle dysfunction, which may not be clinically detectable, and yet may contribute to the occurrence of diabetic complications [99,175]. Moreover, DM is independently associated with decreased physical performance and endurance [176]. This association can be explained by the impact of diabetic neuropathy and the, directly detrimental, effects of DM on the muscle.

The diaphragm, which is a major respiratory muscle, has the morphological structure of skeletal musculature and can be targeted by DM related hyperglycemia [177].

Fierro et al. showed that patients with DM had decreased phrenic nerve conduction velocity, which can underlie a diminished respiratory performance [178]. Several other reports have highlighted the detrimental effects of DM on the diaphragm, which are probably mediated via phrenic neuropathy [179-181].

Polotsky et al. showed in a mouse model that the diaphragm glycosylation is associated with decreased hypercapnic ventilator response [182]. In fact, the muscular proteins are prone to glycation, with resultant impaired functionality, which likely explains the pathophysiology of DM related diaphragm weakness [183]. Wanke et al. supported the notion of impaired respiratory muscle performance, by showing a decrease in inspiratory VC in patients with type 1 DM [184].

Tang et al. reported a case of patient with type 2 DM and associated phrenic neuropathy complicated by the development of acute respiratory failure [185]. A notable strength of their case report was that they performed a diaphragm biopsy. In a later case report, Rison et al. showed beneficial effects of intravenous immunoglobulin in a patient with DM and bilateral phrenic neuropathy [186].

Brannagan et al. reported a case of a patient with type 1 DM and proximal diabetic neuropathy with decreased respiratory performance [187]. Thus, it is

possible that DM may affect multiple respiratory muscles simultaneously and contribute to impaired respiratory performance.

Dharwadkar et al. showed that patients with type 2 DM had lower values of FEV₁, FVC and maximal expiratory pressure which were independently related to hyperglycemia [188]. These investigators speculated that respiratory muscle dysfunction could underlie the observed effects.

Several studies have pointed towards increased peripheral airway resistance in patients with type 1 DM, which may explain why these patients have a greater breathing effort during exercise [189-191].

In a recently published study, Fuso et al. showed that respiratory muscle strength in patients with type 2 DM was significantly related to metabolic control [192].

Hyperglycemia is implicated in the dysfunction of the diaphragm and other respiratory muscles, which lead to decreased physical performance even in patients without overt cardiopulmonary disease. A summary of the key studies on the impact of DM on respiratory muscle performance is presented in Table 5.

DM as a potential risk factor for accelerated decline in lung function

As mentioned above, patients with DM have abnormal DLCO and decreased strength of respiratory muscles. Researchers from Japan and USA showed that patients with DM and no pulmonary disease had lower levels of FVC, which were related to the presence of DM related complications [162,165]. A group from the United Kingdom demonstrated a lower total lung capacity (TLC) in patients with type 1 DM [164]; researchers from Austria showed decreased levels of FEV₁ and VC, which were associated with the presence of diabetic nephropathy [168].

Lange et al. analyzed data from the Copenhagen City Heart Health Study which included 17,506 adults, 266 of whom were diabetic [193]. They found that patients with DM irrespective of gender had lower values of FEV₁ and FVC; however, no accelerated decline in pulmonary function was shown during follow up compared with subjects without DM. As discussed above, patients with DM have impaired respiratory muscle performance and decreased physical endurance which may underlie and confound, the decreased pulmonary variables demonstrated in their work. In a similar study, Walter et al. reviewed the data from the Framingham Heart Study [194]. They found that patients with DM had lower levels of FEV₁ and FVC even after adjustment for smoking, but the diagnosis of DM was not related to COPD independently from gender and smoking.

Lawlor et al. analyzed the data of 3,911 women enrolled in the British Women Heart and Health Study [195]. These researchers showed that the presence of IR

Table 5 Some of the key studies assessing the impact of type 2 DM/MetS on the pulmonary function and COPD

First author and year	Country	Study design	Population studied	Findings
Guazzi et al. [149]; 2002	Italy	In hospital study assessing the effects of regular insulin on the alveolar-capillary conductance. DLCO and its subcomponents were measured.	n: 19 patients (11 men) with type 2 DM and normal cardiac function. mean age: 59.9 years mean weight: 75.8 kg.	DLCO and alveolar capillary membrane conductance were increased by 12% and 14% respectively by insulin therapy.
Guazzi et al. [150]; 2002	Italy	Cross-sectional study.	n: 30 patients with HF (19 men), 15 patients with type 2 DM (8 men) and 15 controls (8 men). mean age: 61.1 years for controls, 62.3 years for patients with type 2 DM, 62.8 years for patients with type 2 DM and HF and 64.1 years for patients with HF alone. mean weight: 75.2 kg for patients with HF and type 2 DM, 76.1 kg for patients with type 2 DM, 76.7 kg for patients with HF and 77.8 kg for controls.	Patients with type 2 DM had a lower DLCO than controls. Regular insulin improved DLCO in patients with type 2 DM and type DM and HF, with no improvement in patients with HF alone. The improvement in patients with type 2 DM and HF was greater than in patients with type 2 DM alone.
Guvener et al. [157]; 2003	Turkey	Cross-sectional study.	n: 25 patients with type 2 DM (9 men) and 12 healthy controls (4 men). mean age: 56.3 years for patients with type 2 DM and 50.1 years for controls mean BMI: 29.9 kg/m ² for patients with type 2 DM and 29.5 kg/m ² for controls.	Patients with type 2 DM had lower ratio of DLCO to VA. In a stepwise regression model with inclusion of age, duration of type 2 DM and microalbuminuria, only microalbuminuria was found to be independent predictor of DLCO/VA.
Chance et al. [158]; 2008	USA	Cross-sectional study assessing alveolar capillary bed in patients with type 2 DM.	n: 69 never smokers with type 2 DM and no overt cardiopulmonary disease (46% women) vs. 45 controls (45% women). mean age: 45 years for controls, 49 years for patients with type 2 DM and BMI > 30 kg/m ² and 45 years for patients with type 2 DM and BMI < 30 kg/m ² . mean BMI: 28.8 kg/m ² for controls, 27.4 kg/m ² and 34.4 kg/m ² for aforementioned type 2 DM groups respectively.	Both non-obese (BMI < 30 kg/m ²) and obese (BMI ≥ 30 kg/m ²) patients with type 2 DM had reduced levels of the measured pulmonary function variables during exercise. However, the observed changes in obese patients were not fully explained by the baseline lung volumes. Moreover, the presence of retinopathy, neuropathy, microalbuminuria and control of diabetes were associated with the reduced pulmonary microvascular reserve
Niranjan et al. [166]; 1997	USA	Prospective observational study of cardio-pulmonary function at 7 years of follow up. Aerobic exercise capacity was measured with cycle ergometry. Lung volume and diffusing capacity were measured with rebreathing technique and ventialatory power was measured by esophageal balloon technique.	n: 18 subjects with type 1 DM (11 men) and 14 controls (10 men). mean age: 31 years for controls and 39 for patients with type 1 DM. mean BMI: 22 kg/m ² for controls and 24.91-25.49 kg/m ² among patients with type 1 DM.	Patients with poor glycemic control had worse restriction of lung volume, pulmonary diffusing capacity and membrane diffusing capacity. Cardiac stroke index was reduced among subjects with poor glycemic control. In the long-term analysis the rate of FEV ₁ and FVC decline was similar to those without DM.

Table 5 Some of the key studies assessing the impact of type 2 DM/MetS on the pulmonary function and COPD (Continued)

Saler et al. [170]; 2009	Turkey	Cross-sectional study assessing alveolar capillary bed in patients with type 1 and type 2 DM.	n: 44 subjects with type 1 DM (29 women), 68 subjects with type 2 DM (49 women) and 80 controls (58 women). mean age: 32.52 years for patients with type 1 DM, 52.4 years for patients with type 2 DM and 40.08 years for controls. mean BMI: 24.4 kg/m ² for patients with type 1 DM, 27.0 kg/m ² for patients with type 2 DM and 25.6 kg/m ² for controls.	DLCO and the ratio of DLCO to VA were significantly decreased in patients with both types of DM, but not in control group.
Wanke et al. [184]; 1991	Austria	Cross-sectional study.	n: 36 patients with type 1 DM (31 men) and 40 controls (33 men). mean age: 33 years for patients with DM and 27 years for controls. mean BMI: 24.3 kg/m ² for patients with DM and 22.3 kg/m ² for controls.	Patients with type 1 DM had significantly lower inspiratory VC, which was in part explained by reduced maximal sniff transesophageal and transdiaphragmatic pressures in patients with type 1 DM.
Walter et al. [194]; 2003	USA	Prospective observational study with a follow up >15 years using the data of the Framingham Heart Study.	n: 3,254 (1,547 men) with 280 subjects having type 2 DM mean age: 53.9 years for patients without type 2 DM and 59.6 years for patients with type 2 DM. mean BMI: 27.3 kg/m ² for patients without type 2 DM and 30.67 kg/m ² for patients with type 2 DM.	Patients with type 2 DM had a lower mean FEV ₁ , FVC. The FVC/FEV ₁ ratio was slightly higher in patients with type 2 DM. Higher fasting glucose levels had association with decreased FEV ₁ , FVC and FVC/FEV ₁ ratio (only in current smokers).
Lawlor et al. [195]; 2004	UK	Cross-sectional study.	N: 3,911 women (9.8% had type 2 DM). Age: post-menopausal women aged 60–79 years Mean BMI: not provided.	FEV ₁ and FVC were inversely related to the HOMA score. After adjustment for age, anthropometric variables, smoking, physical activity, childhood and adult social class and respiratory medications the higher FEV ₁ and FVC were associated with decrease in HOMA score of 3% and 5% respectively. Patients with higher values of FEV ₁ and FVC had lower prevalence of DM.
Davis et al. [196]; 2004	Australia	Prospective study using the data from the Fremantle Diabetes Study with a mean follow up of 7 years. FEV ₁ , FVC, VC and PEF were measured at baseline and during follow up.	n: 125 subjects with type 2 DM. mean age: 61.5 years mean BMI: 29.9 kg/m ²	Higher follow up fasting glucose, greater levels of follow up HbA1c and mean updated HbA1c were associated with a decrease in measured pulmonary parameters. Decreased FEV ₁ was found to be independent predictor of all-cause mortality.
McKeever et al. [197]; 2005	UK	Study analyzing the data of the NHANES III.	n: 4,257 (1,943 men) mean age: 37 years mean BMI: 27.0 kg/m ²	Patients with higher 2 hour 75 g glucose tolerance test had lower levels of FEV ₁ and FVC. Patients with a history of DM had lower levels of FEV ₁ and patients with poor control of DM had lower FEV ₁ than patients with controlled DM.

Table 5 Some of the key studies assessing the impact of type 2 DM/MetS on the pulmonary function and COPD (Continued)

Litonjua et al. [198]; 2005	USA	A nested case-control study using the data of Normative Aging Study.	n: 352 men who developed type 2 DM and 352 controls (all men) mean age: 43.1 years for patients with type 2 DM and 43.2 years for controls. mean BMI: 26.81 kg/m ² for patients with type 2 DM and 25.31 kg/m ² for controls.	Patients with type 2 DM had lower FEV ₁ and FVC values (but not FEV ₁ /FVC) many years prior to the diagnosis of type 2 DM. However, there was no difference in the rate of annual decline of FEV ₁ and FVC in patients with type 2 DM and controls.
Baker et al. [207]; 2006	UK	Cross-sectional study in patients admitted with acute exacerbation of COPD. The patients were divided into 3 groups based on the glucose quartile (<6.0mmol/l, 6.0-6.9 mmol/l 7.0-8.9 mmol/l and >9 mmol/l)	n: 284 subjects (167 men) mean age: from 72.9 to 76.7 years (depends on the glucose quartile, with higher glucose associated with older age) mean BMI: not provided	The RR of death and long hospital stay was greater in patients with glucose from 7.0-8.9 mmol/l and >9.0 mmol/l (RR 1.46 and 1.97 respectively), which was independent from age, gender, COPD severity and prior diagnosis of DM. Each mole increase in glucose was associated with 15% greater risk for adverse clinical outcome. Higher glucose level was associated with the isolation of <i>Staphylococcus Aureus</i> from the sputum.
Chakrabarti et al. [208]; 2009	UK	An observation study on the effects of hyperglycemia on the outcome of non-invasive ventilation (NIV) during COPD exacerbation.	n: 88 in hospital patients (39 men). mean age: 70 years mean BMI: data not provided	Random blood glucose ≥7.0 mmol/l was independently associated with an adverse NIV outcome such as NIV failure and greater risk for pneumonia.
Küpeli et al. [209]; 2010	Turkey	An observational study assessing the correlation between the presence of MetS and COPD exacerbation rate.	n: 106:29 patients with MetS (24 men) and 77 patients without MetS (67 men). mean age: 64.9 years for patients with MetS and 67.3 years for patients without MetS. mean BMI: 30.3 kg/m ² for patients with MetS And 27.2 kg/m ² for patients without MetS.	The mean COPD exacerbation rate was 2.4 in MetS group compared to 0.68 in the control group. Mean length of each exacerbation was 7.5 days in patients with MetS compared to 5 days in patients without MetS. Serum C-reactive protein, fasting blood glucose, and triglycerides were positively correlated with COPD exacerbation rate.

Abbreviations: **BMI:** Body Mass Index; **COPD:** Chronic Obstructive Pulmonary Disease; **DLCO:** Diffusion Capacity for Carbon Monoxide; **DM:** Diabetes Mellitus; **FEV₁:** Forced Expiratory Volume in 1 second; **FVC:** Forced Vital Capacity; **HF:** Heart Failure; **HOMA:** Homeostasis Model Assessment; **NHANES:** National Health and Nutrition Examination Survey; **MetS:** Metabolic Syndrome; **NIV:** Non-Invasive Ventilation; **PEF:** Peak Expiratory Flow; **RR:** Relative Risk; **VA:** Alveolar Ventilation; **VC:** Vital Capacity.

and DM were associated with reduced values of FEV₁. Researchers from Australia studied the data from the Fremantle Diabetes Study and showed that DM was associated with lower values of FEV₁, VC, FVC and peak expiratory flow (PEF) [196]. More importantly, they found that patients with DM had a greater rate of annual decline in pulmonary function and, in addition, that, DM related airflow limitation was associated with increased mortality.

McKeever et al. analyzed data from the Third National Health and Nutrition Examination Survey to

study the association between glucose control and lung function [197]. They showed that patients with DM had lower values of FEV₁ and FVC, but not a decrease in the ratio of FEV₁ to FVC. It is pertinent to note that poor control of DM was associated with worse pulmonary function. Litonjua et al. analyzed data from the Normative Aging Study [198]. They demonstrated that patients with DM had lower FEV₁ and FVC values, even after adjustment for age, gender, smoking, height and weight. However, the presence of DM was not associated with an accelerated decline of

pulmonary function in comparison with patients without DM.

Thus, based on the above data, two major possibilities exist: either that, DM independently contributes to the pathophysiology of accelerated decline in pulmonary function or that DM is just a marker for decreased performance of respiratory muscles and physical endurance. A summary of the key studies on the impact of DM on pulmonary function is presented in Table 5.

Impact of DM on the COPD outcomes

As mentioned above, the presence of COPD is associated with an increased risk of comorbid diseases and DM in particular [17]. Furthermore, comorbid diseases including DM increase the risk of COPD exacerbation and mortality [16].

As shown by Gan et al. and others, the risk of DM development is associated with elevated fibrinogen and other markers of inflammation [73,199]. In fact, this proinflammatory state may act as an independent risk factor and predictor for COPD exacerbations [200]. Interestingly, a lower DLCO is associated with an increased risk of COPD exacerbation, and as discussed above, DM can be considered as a risk factor for the development of alveolar capillaropathy.

Dahl et al. showed that elevated CRP levels increased the risk of COPD flares [75]. Thus, DM may aggravate the disease course via its proinflammatory profile. It should be emphasized that patients with frequent COPD exacerbation have a much more rapid disease progression and related mortality, which merits particular attention to this clinical group [201].

On the other hand, all types of DM are associated with a significantly increased risk of infections, such as pneumonia and bronchitis [202]. DM associated hyperglycemia may increase the risk of pulmonary infections by making glucose present in the respiratory tree, which in turn predisposes to an infectious complication [203]. On the other hand, airway inflammation can increase local

glucose availability, thus causing a vicious cycle. As shown by Phillips et al., glucose in the bronchial tree significantly increased the risk of methicillin resistant staphylococcus aureus (MRSA) infections in mechanically ventilated patients [204]. MRSA infections are well known for difficulties in management and high rates of related morbidity and mortality [205].

McAlister et al. studied the data from 2,471 patients admitted to hospital with community acquired pneumonia (CAP) and analyzed the impact of hyperglycemia on admission to CAP outcomes [206]. These investigators showed that admission glucose levels >11.0 mmol/l were associated with a greater level of in hospital mortality and morbidity. As mentioned previously, researchers from China demonstrated a positive association between IR, hyperglycemia and worsened outcomes during COPD exacerbation [50].

Baker et al. analyzed the data from 284 patients admitted to hospital with COPD exacerbation [207]. Enrolled patients were subdivided into four groups according to their glucose levels: <6.0 mmol/l, 6.0-6.9 mmol/l, 7.0-8.9 mmol/l and >9.0 mmol/l. This group demonstrated that all levels of increase in glucose were associated with increased COPD related morbidity and mortality. Adapted data from their study is presented in Figure 3.

Chakrabarti et al. studied 88 patients with COPD exacerbation, which required the initiation of non-invasive lung ventilation to assess the impact of glucose control on disease outcomes [208]. These researchers showed that hyperglycemia may be a clinically useful predictor of poor outcomes among patients with severe COPD flare up requiring non-invasive ventilation.

Interesting findings were found by Küpeli et al., who studied 106 patients with COPD, including 29 with MetS to test the impact of MetS on COPD exacerbation [209]. It was shown that patients with MetS had a higher rate of COPD flares, and this was related to an increase in fasting glucose, triglyceride level and CRP. It is plausible that the low grade systemic

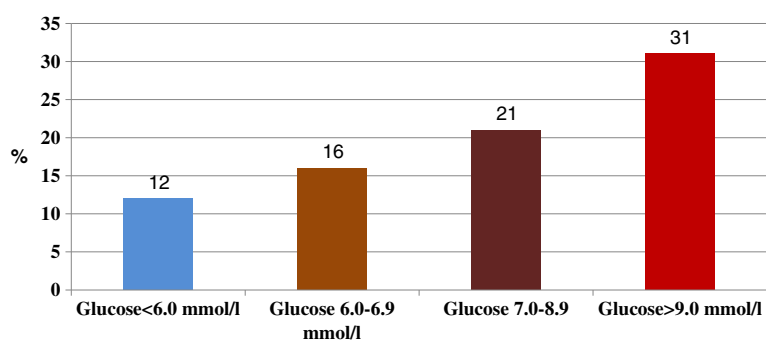


Figure 3 Serum Glucose, COPD exacerbation and in hospital mortality (adapted from reference #) [203].

inflammation seen in MetS contributes to an accelerated progression of COPD. A summary of the key studies on the impact of DM on COPD outcomes is presented in Table 5.

DM therapies other than insulin and pulmonary function

As discussed above insulin therapy may improve DLCO, which acts as a surrogate marker of alveolar capillary function. However, inhaled insulin is not generally recommended for widespread use, because of concerns about poor absorption and theoretically plausible side effects.

Kim et al. retrospectively analyzed the data from 61 patients with type 2 DM with a concomitant diagnosis of COPD [210]. These investigators showed that treatment with an oral insulin sensitizer, such as metformin or thiazolidinedione improved FVC in the recruited individuals; this observation was thought to be due to improved respiratory muscle function. However, this study was critically reviewed and potential limitations were highlighted [211]. From a theoretical point of view, antidiabetic medications may improve endothelial function [212], and may also via this mechanism improve the functionality of pulmonary vasculature and gas diffusion.

Moreover, antidiabetic agents have been shown to be associated with a decreased risk of lung cancer in Taiwanese adults [213]. This is particularly relevant, since COPD is considered to be a risk factor for lung cancer independently from smoking [214]. Indeed, metformin can prevent tobacco induced lung carcinogenesis [215] and can activate apoptosis of lung cancer cells [216,217]. As was shown by Tan et al., metformin may improve the efficacy of chemotherapy targeted against non-small lung cancer in patients with type 2 DM [218]. Metformin may be beneficial in the prevention of cancer in general, in patients with type 2 DM and may in fact decrease mortality in patients with cancer and concomitant type 2 DM [219]. However, Bodmer et al. failed to find any effect of metformin administration on the risk of lung cancer [220].

Furthermore, metformin has been shown to ameliorate ventilator induced lung injury, recently demonstrated in a rabbit model [221]. A simplified interrelationship between COPD and DM is presented in Figure 4.

OSA, overlap syndrome and the risk of type 2 DM

OSA is a common medical disorder affecting up to 24% of the general adult population [222]. OSA is characterized by repetitive full and/or partial collapses of the upper airway (UA) leading to a state of chronic intermittent hypoxia. OSA is associated with low grade systemic inflammation, oxidative stress and fundamental metabolic abnormalities [223,224], including postprandial dyslipidemia [225], an emerging risk factor for vascular

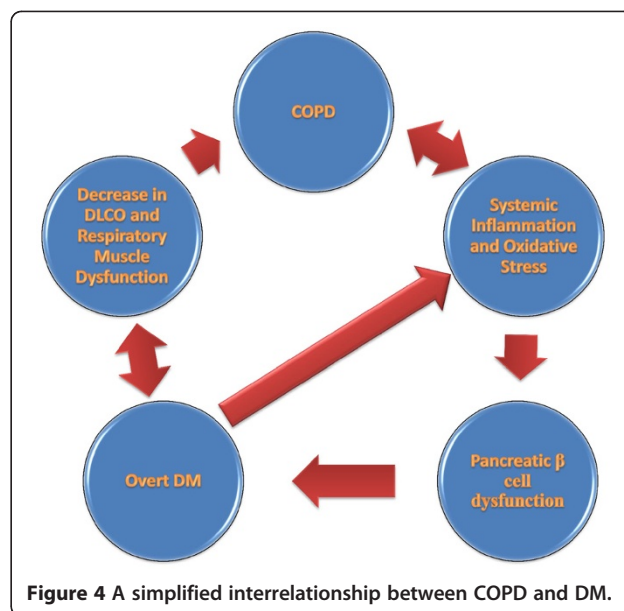


Figure 4 A simplified interrelationship between COPD and DM.

disease [226,227]. These abnormalities are believed to underlie the increased risk for vascular [228-230] and renal disease [231,232] seen in OSA. The disorder is classified based on the number of apnea and/or hypopnea recordings per hour of sleep, which is known as apnea hypopnea index (AHI).

The OSA staging is presented in Table 6. For a more detailed discussion on the topic of general OSA, the reader is referred to a well-written review article [233].

The coexistence of OSA and COPD in an individual is called overlap syndrome. Patients with overlap syndrome often have much more advanced cardiopulmonary disease, including pulmonary hypertension, than either disease alone. This is likely explained by far a greater hypoxia, systemic inflammation and oxidative stress from both diseases simultaneously. It is generally thought that it affects approximately 1% of the adult population [234]. However, the epidemiological data are scant.

From a theoretical point, UA collapses in patients with OSA may lead to bronchial obstruction via a reflex mediated pathway [235]. More importantly, patients with overlap syndrome have greater morbidity and mortality. It was shown that continuous positive airway pressure (CPAP) therapy has been shown to reduce the rates of COPD exacerbations [236,237]. For a detailed discussion on the topic of overlap syndrome, the reader is referred to a well-written review article [234].

Table 6 OSA staging

OSE Severity	Apnea Hypopnea Index
Mild	5-14
Moderate	15-29
Severe	≥30

Unfortunately, the scientific literature on the impact of overlap syndrome on glucose metabolism is scant. However, from a theoretical point of view it is highly likely that it should lead to even greater alteration in glucose metabolism and subsequently to type 2 DM. We will briefly review the current evidence of OSA mediated risk for the new onset type 2 DM.

Several mechanisms are believed to contribute to the pathogenesis of OSA related IR: sleep fragmentation and intermittent hypoxia [238], inflammation and oxidative stress [239,240] and enhanced sympathetic output [224]. It is pertinent to mention that the understanding of how OSA might lead to IR and overt type 2 DM are far from complete. This chapter is not intended to be extensive and the interested reader is referred to some of the well-written review articles on this topic [78,241].

Fredheim et al. showed that OSA is much more common in patients with pre-diabetes and type 2 DM [242]. Indeed, these researchers found that OSA was present in 78% of patients with type 2 DM, compared with 67% of patients with pre-diabetes and 38% of patients without evidence of IR or overt type 2 DM. This was true even after adjustment for age, gender, BMI, insulin sensitivity and high sensitivity for CRP.

Pallayova et al. showed that patients with OSA have lower tissue sensitivity to insulin independent of gender and adiposity [243]. The observed association was believed to be due to an increase in TNF- α and IL-6.

Toqueiro et al. showed that moderate and severe OSA was independently associated with the presence of impaired fasting glucose and markers of IR [244]. Furthermore, several studies showed that OSA is independently related with an increase in glycated hemoglobin [245,246], which is a marker of long term glucose control. Interestingly, it was recently shown that even lean subjects with OSA have greater insulin levels as a result of IR [247], which supports an independent role of OSA in the development of type 2 DM.

Several prospective studies showed that OSA is an independent risk factor for the development for type 2 DM [248,249] and a recently published meta-analysis of prospective studies confirmed this relationship [250].

Sharma et al. studied 86 patients with OSA to assess the impact of CPAP treatment on the metabolic profile, including the IR measurement and glycated hemoglobin in a double-blind placebo (sham CPAP)-controlled trial [251]. These researchers showed that CPAP therapy was significantly associated with a statistically significant reduction in glycated hemoglobin levels. Weinstock et al. showed that CPAP treatment of severe OSA is associated with an improvement in insulin sensitivity [252]. It also has been recently shown that CPAP treatment is independently related to an increase in peripheral insulin sensitivity in patients with concomitant acromegaly

[253]. This is particularly important since acromegaly is associated with both type 2 DM and OS [254]. A recent meta-analysis performed by Yang et al. supported the beneficial effects of CPAP on the glucose metabolism [255].

In concluding this section it is necessary to note that emerging evidence suggests that OSA might be an independent risk factor for the development of non-alcoholic fatty liver disease [256], which is associated with IR and overt type 2 DM [257], as well as vascular [258] and renal diseases [259].

Based on the current evidence, OSA should be considered as an independent risk factor for the development of type 2 DM, and, when coexisting with COPD, the risks are likely to be higher. However, it is essential to note that scientific literature is scant on the topic of overlap syndrome and the risks of type 2 DM. A simplified sketch on how OSA may increase the risk of type 2 DM is shown in Figure 5.

Conclusion

COPD, MetS and DM are common and underdiagnosed medical conditions. It is predicted that COPD will be the third leading cause of death worldwide by 2020. The burden of this disease is even greater if we consider the significant impact of COPD on cardiovascular mortality.

COPD may be considered as a novel risk factor for new onset type 2 DM. The pathophysiology of this is likely to be very complex with several factors being involved, including: inflammation and oxidative stress, administration of glucocorticosteroids, COPD related skeletal muscle dysfunction and abnormalities in adipokine metabolism etc. However, COPD should not be considered as a risk factor for type 1 DM, because of the unique pathophysiology of type 1 DM and different ages at disease presentation.

On the other hand, diabetes may act as an independent factor negatively affecting lung structure and function. Diabetes can cause muscle and neuronal damage,

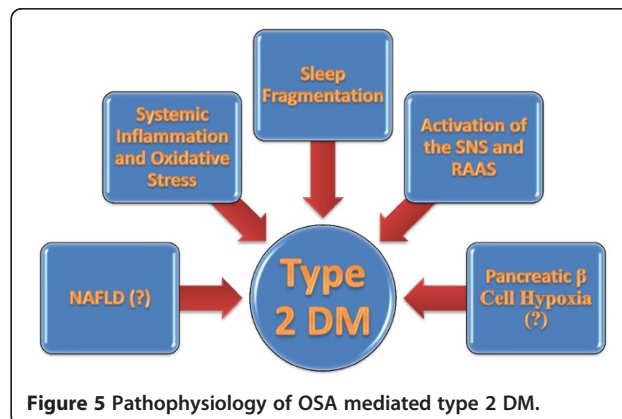


Figure 5 Pathophysiology of OSA mediated type 2 DM.

which is relevant to deficient function of respiratory muscles. Moreover, diabetes is independently associated with lower physical performance, which can be disabling for patients with COPD, who already have some limitation in physical performance. DM is able to detrimentally affect alveolar capillary membrane and decrease DLCO, similarly to other microangiopathic complications, such as diabetic nephropathy. Furthermore, DM is associated with the presence of glucose in airway secretions, and this may contribute to the increased risk of pulmonary infections seen in diabetics. MetS can increase the risk of COPD exacerbation, and diabetes is associated with worsened outcomes of COPD flares. On the top of that, co-existent OSA may increase the risk for type 2 DM in some individuals.

Antihyperglycemic medications may in fact improve DLCO, as has been shown with insulin in patients with DM. However, concerns about safety and pharmacokinetics preclude the recommendation for inhaled insulin to be used at this time. On the other hand, oral antihyperglycemic medications such as metformin and thiazolidinedione may improve FVC in patients with DM. Moreover, metformin has been shown to have antitumor effects and may increase survival in patients with lung cancer.

Thus, it is essential to look at COPD as a potential independent risk factor for the incidence of MetS and type 2 DM and for a complicated course of DM. Conversely, both types of DM and MetS are associated with a worsened clinical course of COPD and a greater degree of morbidity and mortality.

Abbreviations

AHI: Apnea hypopnea index; BA: Bronchial asthma; BMI: Body mass index; CAP: Community acquired pneumonia; CI: Confidence interval; COPD: Chronic obstructive pulmonary disease; CPAP: Continuous positive pressure therapy; CRP: C-Reactive Protein; DLCO: Diffusing capacity for CO; DM: Diabetes Mellitus; ERV: Expiratory reserve volume; FEV₁: Forced expiratory flow in 1 second; FRC: Functional residual capacity; FVC: Forced vital capacity; GLUT: Glucose transporter; GOLD: Global initiative for chronic Obstructive lung disease; HF: Heart failure; HIF: Hypoxia inducible factor; HOMA: Homeostatic model assessment; HR: Hazard ratio; IL: Interleukin; IR: Insulin resistance; MetS: Metabolic syndrome; MMEF: Maximum midexpiratory flow; MRSA: Methicillin resistant staphylococcus aureus; NAFLD: Non alcoholic fatty liver disease; NHANES: National health and nutrition examination survey; NIV: Non-invasive ventilation; OSA: Obstructive sleep apnea; OR: Odds ratio; PEF: Peak expiratory flow; RAAS: Renin angiotensin aldosterone system; RR: Relative risk; RV: Residual volume; SNP: Single nucleotide polymorphism; SNS: Sympathetic nervous system; TLC: Total lung capacity; TNF- α : Tumor necrosis factor alpha; UA: Upper airway; VA: Alveolar ventilation; VC: Vital capacity.

Competing interest

The author has no competing interests.

Received: 31 August 2012 Accepted: 4 October 2012
Published: 27 October 2012

References

1. Vestbo J, Hurd SS, Agusti AG, Jones PW, Vogelmeier C, Anzueto A, Barnes PJ, Fabbri LM, Martinez FJ, Nishimura M, Stockley RA, Sin DD, Rodriguez-Roisin R: **Global Strategy for the Diagnosis, Management and Prevention**

- of Chronic Obstructive Pulmonary Disease, GOLD Executive Summary. *Am J Respir Crit Care Med* 2012, Aug 9. [Epub ahead of print].
2. Fabbri LM, Rabe KF: **From COPD to chronic systemic inflammatory syndrome?** *Lancet* 2007, **370**:797-799.
3. Feary JR, Rodrigues LC, Smith CJ, Hubbard RB, Gibson JE: **Prevalence of major comorbidities in subjects with COPD and incidence of myocardial infarction and stroke: a comprehensive analysis using data from primary care.** *Thorax* 2010, **65**:956-962.
4. Similowski T, Agustí A, MacNee W, Schönhofer B: **The potential impact of anaemia of chronic disease in COPD.** *Eur Respir J* 2006, **27**:390-396.
5. Couillard A, Prefaut C: **From muscle disuse to myopathy in COPD: potential contribution of oxidative stress.** *Eur Respir J* 2005, **26**:703-719.
6. Lehouck A, Boonen S, Decramer M, Janssens W: **COPD, bone metabolism, and osteoporosis.** *Chest* 2011, **139**:648-657.
7. Young RP, Hopkins RJ, Christmas T, Black PN, Metcalf P, Gamble GD: **COPD prevalence is increased in lung cancer, independent of age, sex and smoking history.** *Eur Respir J* 2009, **34**:380-386.
8. McGarvey LP, John M, Anderson JA, Zvarich M, Wise RA: **Ascertainment of cause-specific mortality in COPD: operations of the TORCH Clinical Endpoint Committee.** *Thorax* 2007, **62**:411-415.
9. Murray CJ, Lopez AD: **Alternative projections of mortality and disability by cause 1990-2020: Global Burden of Disease Study.** *Lancet* 1997, **349**:1498-1504.
10. Bakker H, Jaddoe VW: **Cardiovascular and metabolic influences of fetal smoke exposure.** *Eur J Epidemiol* 2011, **26**:763-770.
11. Rimm EB, Manson JE, Stampfer MJ, Colditz GA, Willett WC, Rosner B, et al: **Cigarette smoking and the risk of diabetes in women.** *Am J Public Health* 1993, **83**:211-214.
12. Yanbaeva DG, Dentener MA, Creutzberg EC, Wesseling G, Wouters EF: **Systemic effects of smoking.** *Chest* 2007, **131**:1557-1566.
13. van Belle TL, Coppieters KT, von Herrath MG: **Type 1 diabetes: etiology, immunology, and therapeutic strategies.** *Physiol Rev* 2011, **91**:79-118.
14. Matthews DR, Matthews PC: **Banting Memorial Lecture 2010^A. Type 2 diabetes as an 'infectious' disease: is this the Black Death of the 21st century?** *Diabet Med* 2011, **28**:2-9.
15. Kobylanskiĭ VI, Babadzhanova GI, Suntsov II: **Association between chronic obstructive pulmonary diseases and diabetes mellitus, type I and II.** *Ter Arkh* 2010, **82**:13-18.
16. Mannino DM, Thorn D, Swensen A, Holguin F: **Prevalence and outcomes of diabetes, hypertension and cardiovascular disease in COPD.** *Eur Respir J* 2008, **32**:962-969.
17. Cazzola M, Bettoncelli G, Sessa E, Cricelli C, Biscione G: **Prevalence of comorbidities in patients with chronic obstructive pulmonary disease.** *Respiration* 2010, **80**:112-119.
18. Sode BF, Dahl M, Nordestgaard BG: **Myocardial infarction and other co-morbidities in patients with chronic obstructive pulmonary disease: a Danish nationwide study of 7.4 million individuals.** *Eur Heart J* 2011, **32**:2365-2375.
19. Joo H, Park J, Lee SD, Oh YM: **Comorbidities of chronic obstructive pulmonary disease in Koreans: a population-based study.** *J Korean Med Sci* 2012, **27**:901-906.
20. Alberti KG, Eckel RH, Grundy SM, Zimmet PZ, Cleeman JI, Donato KA, Fruchart JC, James WP, Loria CM, Smith SC Jr: **Harmonizing the Metabolic Syndrome: a Joint Interim Statement of the Association for the Study of Obesity Heart Federation; International Atherosclerosis Society; and International National Heart, Lung, and Blood Institute; American Heart Association; World International Diabetes Federation Task Force on Epidemiology and Prevention.** *Circulation* 2009, **120**:1640-1645.
21. Gifford AH, Leiter JC, Manning HL: **Respiratory function in an obese patient with sleep-disordered breathing.** *Chest* 2010, **138**:704-715.
22. Mancuso P: **Obesity and lung inflammation.** *J Appl Physiol* 2010, **108**:722-728.
23. Leone N, Courbon D, Thomas F, Bean K, Jégo B, Leynaert B, Guize L, Zureik M: **Lung function impairment and metabolic syndrome: the critical role of abdominal obesity.** *Am J Respir Crit Care Med* 2009, **179**:509-516.
24. Lam KB, Jordan RE, Jiang CQ, Thomas GN, Miller MR, Zhang WS, Lam TH, Cheng KK, Adab P: **Airflow obstruction and metabolic syndrome: the Guangzhou Biobank Cohort Study.** *Eur Respir J* 2010, **35**:317-323.

25. Paek YJ, Jung KS, Hwang YI, Lee KS, Lee DR, Lee JU: **Association between low pulmonary function and metabolic risk factors in Korean adults: the Korean National Health and Nutrition Survey.** *Metabolism* 2010, **59**:1300–1306.
26. Marquis K, Maltais F, Duguay V, Bezeau AM, LeBlanc P, Jobin J, Poirier P: **The metabolic syndrome in patients with chronic obstructive pulmonary disease.** *J Cardiopulm Rehabil* 2005, **25**:226–232.
27. Steuten LM, Creutzberg EC, Vrijhoef HJ, Wouters EF: **COPD as a multicomponent disease: inventory of dyspnoea, underweight, obesity and fat free mass depletion in primary care.** *Prim Care Respir J* 2006, **15**:84–91.
28. Eisner MD, Blanc PD, Sidney S, Yelin EH, Lathon PV, Katz PP, Tolstykh I, Ackerson L, Iribarren C: **Body composition and functional limitation in COPD.** *Respir Res* 2007, **8**:7.
29. Guerra S, Sherrill DL, Bobadilla A, Martinez FD, Barbee RA: **The relation of body mass index to asthma, chronic bronchitis, and emphysema.** *Chest* 2002, **122**:1256–1263.
30. Harik-Khan RI, Fleg JL, Wise RA: **Body mass index and the risk of COPD.** *Chest* 2002, **121**:370–376.
31. Watson L, Vonk JM, Löfdahl CG, Pride NB, Pauwels RA, Laitinen LA, et al: **Predictors of lung function and its decline in mild to moderate COPD in association with gender: results from the Euroscop study.** *Respir Med* 2006, **100**:746–753.
32. Celli BR, Cote CG, Marin JM, Casanova C, de Oca Montes M, Mendez RA, Pinto Plata V, Cabral HJ: **The body-mass index, airflow obstruction, dyspnea, and exercise capacity index in chronic obstructive pulmonary disease.** *N Engl J Med* 2004, **350**:1005–1012.
33. Sundh J, Ställberg B, Lisspers K, Montgomery SM, Janson C: **Co-morbidity, body mass index and quality of life in COPD using the Clinical COPD Questionnaire.** *COPD* 2011, **8**:173–181.
34. Wouters EF: **Muscle wasting in chronic obstructive pulmonary disease: to bother and to measure!** *Am J Respir Crit Care Med* 2006, **173**:4–5.
35. Landbo C, Prescott E, Lange P, Vestbo J, Almdal TP: **Prognostic value of nutritional status in chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 1999, **160**:1856–1861.
36. Prescott E, Almdal T, Mikkelsen KL, Tofteng CL, Vestbo J, Lange P: **Prognostic value of weight change in chronic obstructive pulmonary disease: results from the Copenhagen City Heart Study.** *Eur Respir J* 2002, **20**:539–544.
37. Jee SH, Sull JW, Park J, Lee SY, Ohrr H, Guallar E, Samet JM: **Body-mass index and mortality in Korean men and women.** *N Engl J Med* 2006, **355**:779–787.
38. Marino JS, Xu Y, Hill JW: **Central insulin and leptin-mediated autonomic control of glucose homeostasis.** *Trends Endocrinol Metab* 2011, **22**:275–285.
39. Lee YH, Magkos F, Mantzoros CS, Kang ES: **Effects of leptin and adiponectin on pancreatic β -cell function.** *Metabolism* 2011, **60**:1664–1672.
40. Broekhuizen R, Vernooij JH, Schols AM, Dentener MA, Wouters EF: **Leptin as local inflammatory marker in COPD.** *Respir Med* 2005, **99**:70–74.
41. Bruno A, Chanez P, Chiappara G, Siena L, Giammanco S, Gjomarkaj M, Bonsignore G, Bousquet J, Vignola AM: **Does leptin play a cytokine-like role within the airways of COPD patients?** *Eur Respir J* 2005, **26**:398–405.
42. Schols AM, Creutzberg EC, Buurman WA, Campfield LA, Saris WH, Wouters EF: **Plasma leptin is related to proinflammatory status and dietary intake in patients with chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 1999, **160**:1220–1226.
43. Takabatake N, Nakamura H, Abe S, Hino T, Saito H, Yuki H, Kato S, Tomoike H: **Circulating leptin in patients with chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 1999, **159**:1215–1219.
44. Karakas S, Karadag F, Karul AB, Gurgey O, Gurel S, Guney E, Cildag O: **Circulating leptin and body composition in chronic obstructive pulmonary disease.** *Int J Clin Pract* 2005, **59**:1167–1170.
45. Wang QY, Zhang H, Yan X, Kang J, Yu RJ: **Serum resistin and leptin in patients with chronic obstructive pulmonary disease and their relationship to nutritional state.** *Zhonghua Jie He He Hu Xi Za Zhi* 2005, **28**:445–447.
46. Al Mutairi SS, Mojiminiyi OA, Shihab-Eldeen A, Al Rammah T, Abdella N: **Putative roles of circulating resistin in patients with asthma, COPD and cigarette smokers.** *Dis Markers* 2011, **31**:1–7.
47. Hansel NN, Gao L, Rafaels NM, Mathias RA, Neptune ER, Tankersley C, Grant AV, Connett J, Beaty TH, Wise RA, Barnes KC: **Leptin receptor polymorphisms and lung function decline in COPD.** *Eur Respir J* 2009, **34**:103–110.
48. Calikoglu M, Sahin G, Unlu A, Ozturk C, Tamer L, Ercan B, Kanik A, Atik U: **Leptin and TNF-alpha levels in patients with chronic obstructive pulmonary disease and their relationship to nutritional parameters.** *Respiration* 2004, **71**:45–50.
49. Creutzberg EC, Wouters EF, Vanderhoven-Augustin IM, Dentener MA, Schols AM: **Disturbances in leptin metabolism are related to energy imbalance during acute exacerbations of chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 2000, **162**:1239–1245.
50. Pan HY, Lu XZ, Wang DX, Zeng Y, Zhong HB: **The investigation of the relationship between Leptin-insulin resistance and pulmonary function in patients with chronic obstructive pulmonary disease with acute exacerbation.** *Zhongguo Wei Zhong Bing Ji Jiu Yi Xue* 2007, **19**:519–521.
51. Kythreotis P, Kokkini A, Avgeropoulou S, Hadjiannou A, Anastasakou E, Rasidakis A, Bakakos P: **Plasma leptin and insulin-like growth factor I levels during acute exacerbations of chronic obstructive pulmonary disease.** *BMC Pulm Med* 2009, **9**:11.
52. Yang YM, Sun TY, Liu XM: **The role of serum leptin and tumor necrosis factor-alpha in malnutrition of male chronic obstructive pulmonary disease patients.** *Chin Med J (Engl)* 2006, **119**:628–633.
53. Tomoda K, Yoshikawa M, Itoh T, Tamaki S, Fukuoka A, Komeda K, Kimura H: **Elevated circulating plasma adiponectin in underweight patients with COPD.** *Chest* 2007, **132**:135–140.
54. Kirdar S, Serter M, Ceylan E, Sener AG, Kavak T, Karadağ F: **Adiponectin as a biomarker of systemic inflammatory response in smoker patients with stable and exacerbation phases of chronic obstructive pulmonary disease.** *Scand J Clin Lab Invest* 2009, **69**:219–224.
55. Miller M, Cho JY, Pham A, Ramsdell J, Broide DH: **Adiponectin and functional adiponectin receptor 1 are expressed by airway epithelial cells in chronic obstructive pulmonary disease.** *J Immunol* 2009, **182**:684–691.
56. Summer R, Little FF, Ouchi N, Takemura Y, Aprahamian T, Dwyer D, Fitzsimmons K, Suki B, Parameswaran H, Fine A, Walsh K: **Alveolar macrophage activation and an emphysema-like phenotype in adiponectin-deficient mice.** *Am J Physiol Lung Cell Mol Physiol* 2008, **294**:L1035–L1042.
57. Yoon H, Li Y, Man SF, Tashkin D, Wise RA, Connett JE, Anthonisen NA, Chung A, Wright JL, Sin DD: **The complex relationship of serum adiponectin to COPD outcomes.** *Chest* 2012, **142**:885–892.
58. Breyer MK, Rutten EP, Locantore NW, Watkins ML, Miller BE, Wouters EF: **Dysregulated adipokine metabolism in chronic obstructive pulmonary disease.** *Eur J Clin Invest* 2012, **42**:983–991.
59. Evans JL, Goldfine ID, Maddux BA, Grodsky GM: **Oxidative stress and stress-activated signaling pathways: a unifying hypothesis of type 2 diabetes.** *Endocr Rev* 2002, **23**:599–622.
60. MacNee W: **Pulmonary and systemic oxidant/antioxidant imbalance in chronic obstructive pulmonary disease.** *Proc Am Thorac Soc* 2005, **2**:50–60.
61. Semenza GL: **Regulation of oxygen homeostasis by hypoxia-inducible factor 1.** *Physiology* 2009, **24**:97–106.
62. Pradhan AD, Manson JE, Rifai N, Buring JE, Ridker PM: **C-reactive protein, interleukin 6, and risk of developing type 2 diabetes mellitus.** *JAMA* 2001, **286**:327–334.
63. Spranger J, Kroke A, Möhlig M, Hoffmann K, Bergmann MM, Ristow M, Boeing H, Pfeiffer AF: **Inflammatory cytokines and the risk to develop type 2 diabetes: results of the prospective population-based European Prospective Investigation into Cancer and Nutrition (EPIC)-Potsdam Study.** *Diabetes* 2003, **52**:812–817.
64. Zinman B, Hanley AJ, Harris SB, Kwan J, Fantus IG: **Circulating tumor necrosis factor-alpha concentrations in a native Canadian population with high rates of type 2 diabetes mellitus.** *J Clin Endocrinol Metab* 1999, **84**:272–278.
65. Hu FB, Meigs JB, Li TY, Rifai N, Manson JE: **Inflammatory markers and risk of developing type 2 diabetes in women.** *Diabetes* 2004, **53**:693–700.
66. Duncan BB, Schmidt MI, Pankow JS, Ballantyne CM, Couper D, Vigo A, Hoogeveen R, Folsom AR, Heiss G: **Low-grade systemic inflammation and the development of type 2 diabetes: the atherosclerosis risk in communities study.** *Diabetes* 2003, **52**:1799–1805.
67. Botelho FM, Bauer CM, Finch D, Nikota JK, Zavitz CC, Kelly A, Lambert KN, Piper S, Foster ML, Goldring JJ, Wedzicha JA, Bassett J, Bramson J, Iwakura Y, Sleeman M, Kolbeck R, Coyle AJ, Humbles AA, Stämpfli MR: **IL-1 α /IL-1R1**

- expression in chronic obstructive pulmonary disease and mechanistic relevance to smoke-induced neutrophilia in mice. *PLoS One* 2011, **6**:e28457.
68. Attaran D, Lari SM, Towhidi M, Marallu HG, Ayatollahi H, Khajehdaluae M, Ghanei M, Basiri R: **Interleukin-6 and airflow limitation in chemical warfare patients with chronic obstructive pulmonary disease.** *Int J Chron Obstruct Pulmon Dis* 2010, **5**:335–340.
69. He JQ, Foreman MG, Shumansky K, Zhang X, Akhbar L, Sin DD, Man SF, DeMeo DL, Litonjua AA, Silverman EK, Connett JE, Anthonisen NR, Wise RA, Paré PD, Sandford AJ: **Associations of IL6 polymorphisms with lung function decline and COPD.** *Thorax* 2009, **64**:698–704.
70. van Durme YM, Lahousse L, Verhamme KM, Stolk L, Eijgelsheim M, Loth DW, Uitterlinden AG, Breteler MM, Joos GF, Hofman A, Stricker BH, Brusselle GG: **Mendelian randomization study of interleukin-6 in chronic obstructive pulmonary disease.** *Respiration* 2011, **82**:530–538.
71. Yanbaeva DG, Dentener MA, Spruit MA, Houwing-Duistermaat JJ, Kotz D, Passos VL, Wouters EF: **IL6 and CRP haplotypes are associated with COPD risk and systemic inflammation: a case-control study.** *BMC Med Genet* 2009, **10**:23.
72. Tkacova R, Ukropec J, Skyba P, Ukropcova B, Pobeha P, Kurdivova T, Joppa P, Klimes I, Tkac I, Gasperikova D: **Increased adipose tissue expression of proinflammatory CD40, MKK4 and JNK in patients with very severe chronic obstructive pulmonary disease.** *Respiration* 2011, **81**:386–393.
73. Gan WQ, Man SF, Senthilselvan A, Sin DD: **Association between chronic obstructive pulmonary disease and systemic inflammation: a systematic review and a meta-analysis.** *Thorax* 2004, **59**:574–580.
74. Vozarova B, Weyer C, Lindsay RS, Pratley RE, Bogardus C, Tataranni PA: **High white blood cell count is associated with a worsening of insulin sensitivity and predicts the development of type 2 diabetes.** *Diabetes* 2002, **51**:455–461.
75. Dahl M, Vestbo J, Lange P, Bojesen SE, Tybjaerg-Hansen A, Nordestgaard BG: **C-reactive protein as a predictor of prognosis in chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 2007, **175**:250–255.
76. Hersh CP, Miller DT, Kwiatkowski DJ, Silverman EK: **Genetic determinants of C-reactive protein in COPD.** *Eur Respir J* 2006, **28**:1156–1162.
77. van Durme YM, Verhamme KM, Aarnoudse AJ, Van Pottelberge GR, Hofman A, Witterman JC, Joos GF, Brusselle GG, Stricker BH: **C-reactive protein levels, haplotypes, and the risk of incident chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 2009, **179**:375–382.
78. Pallayova M, Lazurova I, Donic V: **Hypoxic damage to pancreatic beta cells—the hidden link between sleep apnea and diabetes.** *Med Hypotheses* 2011, **77**:930–934.
79. Heinonen I, Kempainen J, Kaskinoro K, Peltonen JE, Sipilä HT, Nuutila P, Knuuti J, Boushel R, Kalliokoski KK: **The effects of adenosine, exercise, and acute moderate hypoxia on energy substrate utilization of human skeletal muscle.** *Am J Physiol Regul Integr Comp Physiol* 2012, **302**:R385–R390.
80. Mackenzie R, Elliott B, Maxwell N, Brickley G, Watt P: **The effect of hypoxia and work intensity on insulin resistance in type 2 diabetes.** *J Clin Endocrinol Metab* 2012, **97**:155–162.
81. Jakobsson P, Jorfeldt L: **Oxygen supplementation increases glucose tolerance during euglycaemic hyperinsulinaemic glucose clamp procedure in patients with severe COPD and chronic hypoxaemia.** *Clin Physiol Funct Imaging* 2006, **26**:271–274.
82. Moritz W, Meier F, Stroka DM, Giuliani M, Kugelmeier P, Nett PC, Lehmann R, Candinas D, Gassmann M, Weber M: **Apoptosis in hypoxic human pancreatic islets correlates with HIF-1 α expression.** *FASEB J* 2002, **16**:745–747.
83. Cheng K, Ho K, Stokes R, Scott C, Lau SM, Hawthorne WJ, O'Connell PJ, Loudovaris T, Kay TW, Kulkarni RN, Okada T, Wang XL, Yim SH, Shah Y, Grey ST, Biankin AV, Kench JG, Laybutt DR, Gonzalez FJ, Kahn CR, Gunton JE: **Hypoxia-inducible factor-1 α regulates beta cell function in mouse and human islets.** *J Clin Invest* 2010, **120**:2171–2183.
84. Oltmanns KM, Gehring H, Rudolf S, Schultes B, Rook S, Schweiger U, Born J, Fehm HL, Peters A: **Hypoxia causes glucose intolerance in humans.** *Am J Respir Crit Care Med* 2004, **169**:1231–1237.
85. Chen B, Lam KS, Wang Y, Wu D, Lam MC, Shen J, Wong L, Hoo RL, Zhang J, Xu A: **Hypoxia dysregulates the production of adiponectin and plasminogen activator inhibitor-1 independent of reactive oxygen species in adipocytes.** *Biochem Biophys Res Commun* 2006, **341**:549–556.
86. Regazzetti C, Peraldi P, Grémeaux T, Najem-Lendom R, Ben-Sahra I, Cormont M, Bost F, Le Marchand-Brustel Y, Tanti JF, Giorgetti-Peraldi S: **Hypoxia decreases insulin signaling pathways in adipocytes.** *Diabetes* 2009, **58**:95–103.
87. Halberg N, Khan T, Trujillo ME, Wernstedt-Asterholm I, Attie AD, Sherwani S, Wang ZV, Landskroner-Eiger S, Dineen S, Magalang UJ, Brekken RA, Scherer PE: **Hypoxia-inducible factor 1 α induces fibrosis and insulin resistance in white adipose tissue.** *Mol Cell Biol* 2009, **29**:4467–4483.
88. Catrina SB, Okamoto K, Pereira T, Brismar K, Poellinger L: **Hyperglycemia regulates hypoxia-inducible factor-1 α protein stability and function.** *Diabetes* 2004, **53**:3226–3232.
89. Holloszy JO: **Skeletal muscle "mitochondrial deficiency" does not mediate insulin resistance.** *Am J Clin Nutr* 2009, **89**:463S–466S.
90. Cartee GD, Douen AG, Ramlal T, Klip A, Holloszy JO: **Stimulation of glucose transport in skeletal muscle by hypoxia.** *J Appl Physiol* 1991, **70**:1593–1600.
91. Wuyam B, Payen JF, Levy P, Bensaidane H, Reutenauer H, Le Bas JF, Benabid AL: **Metabolism and aerobic capacity of skeletal muscle in chronic respiratory failure related to chronic obstructive pulmonary disease.** *Eur Respir J* 1992, **5**:157–162.
92. Thompson CH, Davies RJ, Kemp GJ, Taylor DJ, Radda GK, Rajagopalan B: **Skeletal muscle metabolism during exercise and recovery in patients with respiratory failure.** *Thorax* 1993, **48**:486–490.
93. Jakobsson P, Jorfeldt L, Henriksson J: **Metabolic enzyme activity in the quadriceps femoris muscle in patients with severe chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 1995, **151**:374–377.
94. Whittom F, Jobin J, Simard PM, Leblanc P, Simard C, Bernard S, Belleau R, Maltais F: **Histochemical and morphological characteristics of the vastus lateralis muscle in patients with chronic obstructive pulmonary disease.** *Med Sci Sports Exerc* 1998, **30**:1467–1474.
95. Gosker HR, van Mameren H, van Dijk PJ, Engelen MP, van der Vusse GJ, Wouters EF, Schols AM: **Skeletal muscle fibre-type shifting and metabolic profile in patients with chronic obstructive pulmonary disease.** *Eur Respir J* 2002, **19**:617–625.
96. Gosker HR, Zeegers MP, Wouters EF, Schols AM: **Muscle fibre type shifting in the vastus lateralis of patients with COPD is associated with disease severity: a systematic review and meta-analysis.** *Thorax* 2007, **62**:944–949.
97. Green HJ, Bombardier E, Burnett M, Iqbal S, D'Arsigny CL, O'Donnell DE, Ouyang J, Webb KA: **Organization of metabolic pathways in vastus lateralis of patients with chronic obstructive pulmonary disease.** *Am J Physiol Regul Integr Comp Physiol* 2008, **295**:R935–R941.
98. Green HJ, Burnett ME, D'Arsigny CL, O'Donnell DE, Ouyang J, Webb KA: **Altered metabolic and transporter characteristics of vastus lateralis in chronic obstructive pulmonary disease.** *J Appl Physiol* 2008, **105**:879–886.
99. Barzilay JI, Cotsonis GA, Walston J, Schwartz AV, Satterfield S, Miljkovic I, Harris TB: **Insulin resistance is associated with decreased quadriceps muscle strength in nondiabetic adults aged ≥ 70 years.** *Diabetes Care* 2009, **32**:736–738.
100. Jones TH: **Effects of testosterone on Type 2 diabetes and components of the metabolic syndrome.** *J Diabetes* 2010, **2**:146–156.
101. Svartberg J, Schirmer H, Medbø A, Melbye H, Aasebø U: **Reduced pulmonary function is associated with lower levels of endogenous total and free testosterone. The Tromsø study.** *Eur J Epidemiol* 2007, **22**:107–112.
102. Laghi F, Antonescu-Turcu A, Collins E, Segal J, Tobin DE, Jubran A, Tobin MJ: **Hypogonadism in men with chronic obstructive pulmonary disease: prevalence and quality of life.** *Am J Respir Crit Care Med* 2005, **171**:728–733.
103. Traish AM, Miner MM, Morgentaler A, Zitzmann M: **Testosterone deficiency.** *Am J Med* 2011, **124**:578–587.
104. Van Vliet M, Spruit MA, Verleden G, Kasran A, Van Herck E, Pitta F, Bouillon R, Decramer M: **Hypogonadism, quadriceps weakness, and exercise intolerance in chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 2005, **172**:1105–1111.
105. Sempke PD, Beastall GH, Watson WS, Hume R: **Hypothalamic-pituitary dysfunction in respiratory hypoxia.** *Thorax* 1981, **36**:605–609.
106. Barth E, Albuszies G, Baumgart K, Matejovic M, Wachter U, Vogt J, Radermacher P, Calzia E: **Glucose metabolism and catecholamines.** *Crit Care Med* 2007, **35**:S508–S518.
107. Kanstrup IL, Poulsen TD, Hansen JM, Andersen LJ, Bestle MH, Christensen NJ, Olsen NV: **Blood pressure and plasma catecholamines in acute and**

- prolonged hypoxia: effects of local hypothermia. *J Appl Physiol* 1999, **87**:2053–2058.
108. Scalvini S, Volterrani M, Vitacca M, Clark AL, Solfrini R, Panzali AM, Ferrari R, Levi GF: **Plasma hormone levels and haemodynamics in patients with chronic obstructive lung disease.** *Monaldi Arch Chest Dis* 1996, **51**:380–386.
109. Bratel T, Wennlund A, Carlström K: **Impact of hypoxaemia on neuroendocrine function and catecholamine secretion in chronic obstructive pulmonary disease (COPD). Effects of long-term oxygen treatment.** *Respir Med*. 2000, **94**:1221–1228.
110. Kalupahana NS, Moustaid-Moussa N: **The renin-angiotensin system: a link between obesity, inflammation and insulin resistance.** *Obes Rev* 2012, **13**:136–149.
111. Farber MO, Roberts LR, Weinberger MH, Robertson GL, Fineberg NS, Manfredi F: **Abnormalities of sodium and H₂O handling in chronic obstructive lung disease.** *Arch Intern Med* 1982, **142**:1326–1330.
112. Farber MO, Weinberger MH, Robertson GL, Fineberg NS, Manfredi F: **Hormonal abnormalities affecting sodium and water balance in acute respiratory failure due to chronic obstructive lung disease.** *Chest* 1984, **85**:49–54.
113. Stewart AG, Waterhouse JC, Billings CG, Baylis PH, Howard P: **Hormonal, renal, and autonomic nerve factors involved in the excretion of sodium and water during dynamic salt and water loading in hypoxaemic chronic obstructive pulmonary disease.** *Thorax* 1995, **50**:838–845.
114. Mancini GB, Etminan M, Zhang B, Levesque LE, FitzGerald JM, Brophy JM: **Reduction of morbidity and mortality by statins, angiotensin-converting enzyme inhibitors, and angiotensin receptor blockers in patients with chronic obstructive pulmonary disease.** *J Am Coll Cardiol* 2006, **47**:2554–2560.
115. Hyppönen E: **Vitamin D, and increasing incidence of type 1 diabetes-evidence for an association?** *Diabetes Obes Metab* 2010, **12**:737–743.
116. Mitri J, Muraru MD, Pittas AG: **Vitamin D and type 2 diabetes: a systematic review.** *Eur J Clin Nutr* 2011, **65**:1005–1015.
117. Herr C, Greulich T, Koczulla RA, Meyer S, Zakharkina T, Branscheidt M, Eschmann R, Bals R: **The role of vitamin D in pulmonary disease: COPD, asthma, infection, and cancer.** *Respir Res* 2011, **12**:31.
118. Gow SM, Seth J, Beckett GJ, Douglas G: **Thyroid function and endocrine abnormalities in elderly patients with severe chronic obstructive lung disease.** *Thorax* 1987, **42**:520–525.
119. Lazarus R, Sparrow D, Weiss ST: **Baseline ventilatory function predicts the development of higher levels of fasting insulin and fasting insulin resistance index: the Normative Aging Study.** *Eur Respir J* 1998, **12**:641–645.
120. Ford ES, Mannino DM: **Prospective association between lung function and the incidence of diabetes: findings from the National Health and Nutrition Examination Survey Epidemiologic Follow-up Study.** *Diabetes Care* 2004, **27**:2966–2970.
121. Wannamethee SG, Shaper AG, Rumley A, Sattar N, Whincup PH, Thomas MC, Lowe GD: **Lung function and risk of type 2 diabetes and fatal and nonfatal major coronary heart disease events: possible associations with inflammation.** *Diabetes Care* 2010, **33**:1990–1996.
122. Yeh HC, Punjabi NM, Wang NY, Pankow JS, Duncan BB, Brancati FL: **Vital capacity as a predictor of incident type 2 diabetes: the Atherosclerosis Risk in Communities study.** *Diabetes Care* 2005, **28**:1472–1429.
123. Hsiao FC, Wu CZ, Su SC, Sun MT, Hsieh CH, Hung YJ, He CT, Pei D: **Baseline forced expiratory volume in the first second as an independent predictor of development of the metabolic syndrome.** *Metabolism* 2010, **59**:848–853.
124. Engström G, Janzon L: **Risk of developing diabetes is inversely related to lung function: a population-based cohort study.** *Diabet Med* 2002, **19**:167–170.
125. Engström G, Hedblad B, Nilsson P, Wollmer P, Berglund G, Janzon L: **Lung function, insulin resistance and incidence of cardiovascular disease: a longitudinal cohort study.** *J Intern Med* 2003, **253**:574–581.
126. Kwon CH, Rhee EJ, Song JU, Kim JT, Sung KC: **Reduced lung function is independently associated with increased risk of type 2 diabetes in Korean men.** *Cardiovasc Diabetol* 2012, **11**(1):38. Epub ahead of print.
127. Oda E, Kawai R: **A cross-sectional relationship between vital capacity and metabolic syndrome and between vital capacity and diabetes in a sample Japanese population.** *Environ Health Prev Med*. 2009, **14**:284–291.
128. Kim SK, Hur KY, Choi YH, Kim SW, Chung JH, Kim HK, Lee MK, Min YK, Kim KW, Kim JH: **The Relationship between Lung Function and Metabolic Syndrome in Obese and Non-Obese Korean Adult Males.** *Korean Diabetes J* 2010, **34**:253–260.
129. Lim SY, Rhee EJ, Sung KC: **Metabolic syndrome, insulin resistance and systemic inflammation as risk factors for reduced lung function in Korean nonsmoking males.** *J Korean Med Sci* 2010, **25**:1480–1486.
130. Chlif M, Keochkerian D, Mourlhon C, Choquet D, Ahmadi S: **Noninvasive assessment of the tension-time index of inspiratory muscles at rest in obese male subjects.** *Int J Obes (Lond)* 2005, **29**:1478–1483.
131. Yeh F, Dixon AE, Marion S, Schaefer C, Zhang Y, Best LG, Calhoun D, Rhoades ER, Lee ET: **Obesity in adults is associated with reduced lung function in metabolic syndrome and diabetes: the Strong Heart Study.** *Diabetes Care* 2011, **34**:2306–2313.
132. Heianza Y, Arase Y, Tsuji H, Saito K, Amakawa K, Hsieh SD, Kodama S, Shimano H, Yamada N, Hara S, Sone H: **Low Lung Function and Risk of Type 2 Diabetes in Japanese Men: The Toranomon Hospital Health Management Center Study 9 (TOPICS 9).** *Mayo Clin Proc* 2012, **87**:853–861.
133. Rana JS, Mittleman MA, Sheikh J, Hu FB, Manson JE, Colditz GA, Speizer FE, Barr RG, Camargo CA Jr: **Chronic obstructive pulmonary disease, asthma, and risk of type 2 diabetes in women.** *Diabetes Care* 2004, **27**:2478–2484.
134. Song Y, Klevak A, Manson JE, Buring JE, Liu S: **Asthma, chronic obstructive pulmonary disease, and type 2 diabetes in the Women's Health Study.** *Diabetes Res Clin Pract* 2010, **90**:365–371.
135. Bolton CE, Evans M, Ionescu AA, Edwards SM, Morris RH, Dunseath G, Luzio SD, Owens DR, Shale DJ: **Insulin resistance and inflammation - A further systemic complication of COPD.** *COPD* 2007, **4**:121–126.
136. Bayliss EA, Blatchford PJ, Newcomer SR, Steiner JF, Fairclough DL: **The effect of incident cancer, depression and pulmonary disease exacerbations on type 2 diabetes control.** *J Gen Intern Med* 2011, **26**:575–581.
137. Spies CM, Strehl C, van der Goes MC, Bijlsma JW, Buttgerit F: **Glucocorticoids.** *Best Pract Res Clin Rheumatol* 2011, **25**:891–900.
138. Niewoehner DE, Erbland ML, Deupree RH, Collins D, Gross NJ, Light RW, Anderson P, Morgan NA: **Effect of systemic glucocorticoids on exacerbations of chronic obstructive pulmonary disease. Department of Veterans Affairs Cooperative Study Group.** *N Engl J Med* 1999, **340**:1941–1947.
139. Burt MG, Roberts GW, Aguilar-Loza NR, Frith P, Stranks SN: **Continuous monitoring of circadian glycemic patterns in patients receiving prednisolone for COPD.** *J Clin Endocrinol Metab* 2011, **96**:1789–1796.
140. Decramer M, Lacquet LM, Fagard R, Rogiers P: **Corticosteroids contribute to muscle weakness in chronic airflow obstruction.** *Am J Respir Crit Care Med* 1994, **150**:11–16.
141. Gallagher CG: **Respiratory steroid myopathy.** *Am J Respir Crit Care Med* 1994, **150**:4–6.
142. Perez T, Becquart LA, Stach B, Wallaert B, Tonnel AB: **Inspiratory muscle strength and endurance in steroid-dependent asthma.** *Am J Respir Crit Care Med* 1996, **153**:610–615.
143. Weiner P, Azgad Y, Weiner M: **The effect of corticosteroids on inspiratory muscle performance in humans.** *Chest* 1993, **104**:1788–1791.
144. Ernst P, Suissa S: **Systemic effects of inhaled corticosteroids.** *Curr Opin Pulm Med* 2012, **18**:85–89.
145. O'Byrne PM, Rennard S, Gerstein H, Radner F, Peterson S, Lindberg B, Carlsson LG, Sin DD: **Risk of new onset diabetes mellitus in patients with asthma or COPD taking inhaled corticosteroids.** *Respir Med* 2012, **106**:1487–1493.
146. Slatore CG, Bryson CL, Au DH: **The association of inhaled corticosteroid use with serum glucose concentration in a large cohort.** *Am J Med* 2009, **122**:472–478.
147. van den Oever IA, Raterman HG, Nurmohamed MT, Simsek S: **Endothelial dysfunction, inflammation, and apoptosis in diabetes mellitus.** *Mediators Inflamm* 2010, **2010**:792393.
148. Jermendy G: **Vascular memory: can we broaden the concept of the metabolic memory?** *Cardiovasc Diabetol* 2012, **11**:44.
149. Guazzi M, Brambilla R, De Vita S, Guazzi MD: **Diabetes worsens pulmonary diffusion in heart failure, and insulin counteracts this effect.** *Am J Respir Crit Care Med* 2002, **166**:978–982.
150. Guazzi M, Oreglia I, Guazzi MD: **Insulin improves alveolar-capillary membrane gas conductance in type 2 diabetes.** *Diabetes Care* 2002, **25**:1802–1806.

151. Hsia CC, Raskin P: The diabetic lung: relevance of alveolar microangiopathy for the use of inhaled insulin. *Am J Med* 2005, **118**:205–211.
152. Ceglia L, Lau J, Pittas AG: Meta-analysis: efficacy and safety of inhaled insulin therapy in adults with diabetes mellitus. *Ann Intern Med* 2006, **145**:665–675.
153. Davis TM, Davis WA: An assessment of eligibility for inhaled insulin (Exubera): the Fremantle Diabetes Study. *Diabetes Care* 2007, **30**:360–361.
154. Becker RH, Sha S, Frick AD, Fountaine RJ: The effect of smoking cessation and subsequent resumption on absorption of inhaled insulin. *Diabetes Care* 2006, **29**:277–282.
155. Ang E, Lawrence MK, Heilmann CR, Ferguson JA, Tobian JA, Webb DM, Berclaz PY: Safety and efficacy of AIR inhaled insulin compared with subcutaneous insulin in patients having diabetes and asthma: A 12-month, randomized, noninferiority trial. *Diabetes Technol Ther* 2009, **11**(Suppl 2):S35–S44.
156. Raskin P, Heller S, Honka M, Chang PC, Boss AH, Richardson PC, Amin N: Pulmonary function over 2 years in diabetic patients treated with prandial inhaled Technosphere Insulin or usual antidiabetes treatment: a randomized trial. *Diabetes Obes Metab* 2012, **14**:163–173.
157. Guvener N, Tutuncu NB, Akcay S, Eyuboglu F, Gokcel A: Alveolar gas exchange in patients with type 2 diabetes mellitus. *Endocr J* 2003, **50**:663–667.
158. Chance WW, Rhee C, Yilmaz C, Dane DM, Pruneda ML, Raskin P, Hsia CC: Diminished alveolar microvascular reserves in type 2 diabetes reflect systemic microangiopathy. *Diabetes Care* 2008, **31**:1596–1601.
159. Minette P, Buyschaert M, Rahier J, Veriter C, Frans A: Pulmonary gas exchange in life-long nonsmoking patients with diabetes mellitus. *Respiration* 1999, **66**:20–24.
160. Scherthaner G, Haber P, Kummer F, Ludwig H: Lung elasticity in juvenile-onset diabetes mellitus. *Am Rev Respir Dis* 1977, **116**:544–546.
161. Sokolov EI, Demidov Iul [Gas exchange function of the lungs in patients with type 1 diabetes mellitus]. *Ter Arkh* 2008, **80**:63–66.
162. Asanuma Y, Fujiya S, Ide H, Agishi Y: Characteristics of pulmonary function in patients with diabetes mellitus. *Diabetes Res Clin Pract* 1985, **1**:95–101.
163. Sandler M, Bunn AE, Stewart RI: Cross-section study of pulmonary function in patients with insulin-dependent diabetes mellitus. *Am Rev Respir Dis* 1987, **135**:223–229.
164. Cooper BG, Taylor R, Alberti KG, Gibson GJ: Lung function in patients with diabetes mellitus. *Respir Med* 1990, **84**:235–239.
165. Ramirez LC, Dal Nogare A, Hsia C, Arauz C, Butt I, Strowig SM, Schnurr-Breen L, Raskin P: Relationship between diabetes control and pulmonary function in insulin-dependent diabetes mellitus. *Am J Med* 1991, **91**:371–376.
166. Niranjan V, McBrayer DG, Ramirez LC, Raskin P, Hsia CC: Glycemic control and cardiopulmonary function in patients with insulin-dependent diabetes mellitus. *Am J Med* 1997, **103**:504–513.
167. Strojek K, Ziora D, Sroczynski JW, Oklek K: Pulmonary complications of type 1 (insulin-dependent) diabetic patients. *Diabetologia* 1992, **35**:1173–1176.
168. Innocenti F, Fabbri A, Anichini R, Tuci S, Pettinà G, Vannucci F, De Giorgio LA, Seghieri G: Indications of reduced pulmonary function in type 1 (insulin-dependent) diabetes mellitus. *Diabetes Res Clin Pract* 1994, **25**:161–168.
169. Schnack C, Festa A, Schwarzmaier-D'Assiè A, Haber P, Scherthaner G: Pulmonary dysfunction in type 1 diabetes in relation to metabolic long-term control and to incipient diabetic nephropathy. *Nephron* 1996, **74**:395–400.
170. Saler T, Cakmak G, Saglam ZA, Ataoglu E, Yesim Erdem T, Yenigun M: The assessment of pulmonary diffusing capacity in diabetes mellitus with regard to microalbuminuria. *Intern Med* 2009, **48**:1939–1943.
171. Scaramuzza AE, Morelli M, Rizzi M, Borgonovo S, De Palma A, Mameli C, Giani E, Beretta S, Zuccotti GV: Impaired diffusing capacity for carbon monoxide in children with type 1 diabetes: is this the first sign of long-term complications? *Acta Diabetol* 2012, **49**:159–164.
172. Sinha S, Guleria R, Misra A, Pandey RM, Yadav R, Tiwari S: Pulmonary functions in patients with type 2 diabetes mellitus & correlation with anthropometry & microvascular complications. *Indian J Med Res* 2004, **119**:66–71.
173. Boulbou MS, Gourgoulisian KI, Petinaki EA, Klisiaris VK, Maniatis AN, Molyvdas PA: Pulmonary function and circulating adhesion molecules in patients with diabetes mellitus. *Can Respir J* 2003, **10**:259–264.
174. Benbassat CA, Stern E, Kramer M, Lebzelter J, Blum I, Fink G: Pulmonary function in patients with diabetes mellitus. *Am J Med Sci* 2001, **322**:127–132.
175. Andersen H: Motor dysfunction in diabetes. *Diabetes Metab Res Rev* 2012, **28**(Suppl 1):89–92.
176. Sinclair AJ, Conroy SP, Bayer AJ: Impact of diabetes on physical function in older people. *Diabetes Care* 2008, **31**:233–235.
177. Ottenheijm CA, Heunks LM, Dekhuijzen RP: Diaphragm adaptations in patients with COPD. *Respir Res* 2008, **9**:12.
178. Fierro B, Bennici S, Raimondo DM: Phrenic nerve conduction in diabetic patients. A preliminary study. *Acta Neurol (Napoli)* 1982, **4**:357–361.
179. Wolf E, Shochina M, Fidel Y, Gonen B: Phrenic neuropathy in patients with diabetes mellitus. *Electromyogr Clin Neurophysiol* 1983, **23**:523–530.
180. Wanke T, Paternostro-Sluga T, Gritsold W, Formanek D, Auinger M, Zwick H, Irsigler K: Phrenic nerve function in type 1 diabetic patients with diaphragm weakness and peripheral neuropathy. *Respiration* 1992, **59**:233–237.
181. White JE, Bullock RE, Hudgson P, Home PD, Gibson GJ: Phrenic neuropathy in association with diabetes. *Diabet Med* 1992, **9**:954–956.
182. Polotsky VY, Wilson JA, Haines AS, Scharf MT, Soutiere SE, Tankersley CG, Smith PL, Schwartz AR, O'Donnell CP: The impact of insulin-dependent diabetes on ventilatory control in the mouse. *Am J Respir Crit Care Med* 2001, **163**:624–632.
183. Ramamurthy B, Höök P, Jones AD, Larsson L: Changes in myosin structure and function in response to glycation. *FASEB J* 2001, **15**:2415–2422.
184. Wanke T, Formanek D, Auinger M, Popp W, Zwick H, Irsigler K: Insulin muscle performance and pulmonary function changes in insulin-dependent diabetes mellitus. *Am Rev Respir Dis* 1991, **143**:97–100.
185. Tang EW, Jardine DL, Rodins K, Evans J: Respiratory failure secondary to diabetic neuropathy affecting the phrenic nerve. *Diabet Med* 2003, **20**:599–601.
186. Rison RA, Beydoun SR: Bilateral phrenic neuropathies in a diabetic patient responsive to intravenous immunoglobulin: a case report. *Clin Neurol Neurosurg* 2011, **113**:788–791.
187. Brannagan TH, Promisloff RA, McCluskey LF, Mitz KA: Proximal diabetic neuropathy presenting with respiratory weakness. *J Neurol Neurosurg Psychiatry* 1999, **67**:539–541.
188. Dharwadkar AR, Dharwadkar AA, Banu G, Bagali S: Reduction in lung functions in type-2 diabetes in Indian population: correlation with glycemic status. *Indian J Physiol Pharmacol* 2011, **55**:170–175.
189. Scano G, Seghieri G, Mancini M, Filippelli M, Duranti R, Fabbri A, Innocenti F, Iandelli I, Misuri G: Dyspnoea, peripheral airway involvement and respiratory muscle effort in patients with type I diabetes mellitus under good metabolic control. *Clin Sci (Lond)* 1999, **96**:499–506.
190. Mancini M, Filippelli M, Seghieri G, Iandelli I, Innocenti F, Duranti R, Scano G: Respiratory muscle function and hypoxic ventilatory control in patients with type I diabetes. *Chest* 1999, **115**:1553–1562.
191. Scano G, Filippelli M, Romagnoli I, Mancini M, Misuri G, Duranti R, Rosi E: Hypoxic and hypercapnic breathlessness in patients with type I diabetes mellitus. *Chest* 2000, **117**:960–967.
192. Fusco L, Pitocco D, Longobardi A, Zaccardi F, Contu C, Pozzuto C, Basso S, Varone F, Ghirlanda G, Antonelli Incalzi R: Reduced respiratory muscle strength and endurance in type 2 diabetes mellitus. *Diabetes Metab Res Rev* 2012, **28**:370–375.
193. Lange P, Parmer J, Schnohr P, Jensen G: Copenhagen City Heart Study: longitudinal analysis of ventilatory capacity in diabetic and nondiabetic adults. *Eur Respir J* 2002, **20**:1406–1412.
194. Walter RE, Beiser A, Givelber RJ, O'Connor GT, Gottlieb DJ: Association between glycemic state and lung function: the Framingham Heart Study. *Am J Respir Crit Care Med* 2003, **167**:911–916.
195. Lawlor DA, Ebrahim S, Smith GD: Associations of measures of lung function with insulin resistance and Type 2 diabetes: findings from the British Women's Heart and Health Study. *Diabetologia* 2004, **47**:195–203.
196. Davis WA, Knuiaman M, Kendall P, Grange V, Davis TM: Glycemic exposure is associated with reduced pulmonary function in type 2 diabetes: the Fremantle Diabetes Study. *Diabetes Care* 2004, **27**:752–757.
197. McKeever TM, Weston PJ, Hubbard R, Fogarty A: Lung function and glucose metabolism: an analysis of data from the Third National Health and Nutrition Examination Survey. *Am J Epidemiol* 2005, **161**:546–556.
198. Litonjua AA, Lazarus R, Sparrow D, Demolles D, Weiss ST: Lung function in type 2 diabetes: the Normative Aging Study. *Respir Med* 2005, **99**:1583–1590.

199. Duvoix A, Dickens J, Haq I, Mannino D, Miller B, Tal-Singer R, Lomas DA: **Blood fibrinogen as a biomarker of chronic obstructive pulmonary disease.** *Thorax* 2012, 2012 Jun 28. [Epub ahead of print].
200. Groenewegen KH, Postma DS, Hop WC, Wielders PL, Schlösser NJ, Wouters EF: **Increased systemic inflammation is a risk factor for COPD exacerbations.** *Chest* 2008, **133**:350–357.
201. Anzueto A: **Impact of exacerbations on COPD.** *Eur Respir Rev* 2010, **19**:113–118.
202. Muller LM, Gorter KJ, Hak E, Goudzwaard WL, Schellevis FG, Hoepelma AI, Rutten GE: **Increased risk of common infections in patients with type 1 and type 2 diabetes mellitus.** *Clin Infect Dis* 2005, **41**:281–288.
203. Philips BJ, Meguer JX, Redman J, Baker EH: **Factors determining the appearance of glucose in upper and lower respiratory tract secretions.** *Intensive Care Med* 2003, **29**:2204–2210.
204. Philips BJ, Redman J, Brennan A, Wood D, Holliman R, Baines D, Baker EH: **Glucose in bronchial aspirates increases the risk of respiratory MRSA in intubated patients.** *Thorax* 2005, **60**:761–764.
205. Lee AS, Huttner B, Harbarth S: **Control of methicillin-resistant *Staphylococcus aureus*.** *Infect Dis Clin North Am* 2011, **25**:155–179.
206. McAlister FA, Majumdar SR, Blitz S, Rowe BH, Romney J, Marrie TJ: **The relation between hyperglycemia and outcomes in 2,471 patients admitted to the hospital with community-acquired pneumonia.** *Diabetes Care* 2005, **28**:810–815.
207. Baker EH, Janaway CH, Philips BJ, Brennan AL, Baines DL, Wood DM, Jones PW: **Hyperglycaemia is associated with poor outcomes in patients admitted to hospital with acute exacerbations of chronic obstructive pulmonary disease.** *Thorax* 2006, **61**:284–289.
208. Chakrabarti B, Angus RM, Agarwal S, Lane S, Calverley PM: **Hyperglycaemia as a predictor of outcome during non-invasive ventilation in decompensated COPD.** *Thorax* 2009, **64**:857–862.
209. Küpeli E, Ulubay G, Ulasli SS, Sahin T, Erayman Z, Gürsoy A: **Metabolic syndrome is associated with increased risk of acute exacerbation of COPD: a preliminary study.** *Endocrine* 2010, **38**:76–82.
210. Kim HJ, Lee JY, Jung HS, Kim DK, Lee SM, Yim JJ, Yang SC, Chung HS, Kim YW, Han SK, Shim YS, Lee CH: **The impact of insulin sensitizers on lung function in patients with chronic obstructive pulmonary disease and diabetes.** *Int J Tuberc Lung Dis* 2010, **14**:362–367.
211. Man SF, Sin DD: **Could oral hypoglycemics be the new therapeutic answer to chronic obstructive pulmonary disease?** *Int J Tuberc Lung Dis* 2010, **14**:262.
212. Kelly AS, Bergenstal RM, Gonzalez-Campoy JM, Katz H, Bank AJ: **Effects of Exenatide vs. Metformin on Endothelial Function in Obese Patients with Pre-Diabetes: A Randomized Trial.** *Cardiovasc Diabetol* 2012, **11**:64. [Epub ahead of print].
213. Lai SW, Liao KF, Chen PC, Tsai PY, Hsieh DP, Chen CC: **Antidiabetes drugs correlate with decreased risk of lung cancer: a population-based observation in Taiwan.** *Clin Lung Cancer* 2012, **13**:143–148.
214. Celli BR: **Chronic obstructive pulmonary disease and lung cancer: common pathogenesis, shared clinical challenges.** *Proc Am Thorac Soc* 2012, **9**:74–79.
215. Memmott RM, Mercadob JR, Maier CR, Kawabata S, Fox SD, Dennis PA: **Metformin prevents tobacco carcinogen-induced lung tumorigenesis.** *Cancer Prev Res (Phila)* 2010, **3**:1066–1076.
216. Wu N, Gu C, Gu H, Hu H, Han Y, Li Q: **Metformin induces apoptosis of lung cancer cells through activating JNK/p38 MAPK pathway and GADD153.** *Neoplasma* 2011, **58**:482–490.
217. Ashinuma H, Takiguchi Y, Kitazono S, Kitazono-Saitoh M, Kitamura A, Chiba T, Tada Y, Kurosu K, Sakaida E, Sekine I, Tanabe N, Iwama A, Yokosuka O, Tatsumi K: **Antiproliferative action of metformin in human lung cancer cell lines.** *Oncol Rep* 2012, **28**:8–14.
218. Tan BX, Yao WX, Ge J, Peng XC, Du XB, Zhang R, Yao B, Xie K, Li LH, Dong H, Gao F, Zhao F, Hou JM, Su JM, Liu JY: **Prognostic influence of metformin as first-line chemotherapy for advanced nonsmall cell lung cancer in patients with type 2 diabetes.** *Cancer* 2011, **117**:5103–5111.
219. Noto H, Goto A, Tsujimoto T, Noda M: **Cancer risk in diabetic patients treated with metformin: a systematic review and meta-analysis.** *PLoS One* 2012, **7**:e33411.
220. Bodmer M, Becker C, Jick SS, Meier CR: **Metformin does not alter the risk of lung cancer: A case-control analysis.** *Lung Cancer* 2012, 2012 Sep 6. [Epub ahead of print].
221. Tsaknis G, Siempos II, Kopterides P, Maniatis NA, Magkou C, Kardara M, Panoutsou S, Kotanidou A, Roussos C, Armaganidis A: **Metformin attenuates ventilator-induced lung injury.** *Crit Care* 2012, **16**:R134.
222. Young T, Palta M, Dempsey J, Skatrud J, Weber S, Badr S: **The occurrence of sleep-disordered breathing among middle-aged adults.** *N Engl J Med* 1993, **328**:1230–1235.
223. Lavie L: **Oxidative stress inflammation and endothelial dysfunction in obstructive sleep apnea.** *Front Biosci (Elite Ed)* 2012, **4**:1391–1403.
224. Hakim F, Gozal D, Kheirandish-Gozal L: **Sympathetic and catecholaminergic alterations in sleep apnea with particular emphasis on children.** *Front Neurol* 2012, **3**:7.
225. Phillips CL, Yee BJ, Marshall NS, Liu PY, Sullivan DR, Grunstein RR: **Continuous positive airway pressure reduces postprandial lipidemia in obstructive sleep apnea: a randomized, placebo-controlled crossover trial.** *Am J Respir Crit Care Med* 2011, **184**:355–361.
226. Ansar S, Koska J, Reaven PD: **Postprandial hyperlipidemia, endothelial dysfunction and cardiovascular risk: focus on incretins.** *Cardiovasc Diabetol* 2011, **10**:61.
227. Toth PP, Simko RJ, Palli SR, Koselleck D, Quimbo RA, Cziraky MJ: **The impact of serum lipids on risk for microangiopathy in patients with type 2 diabetes mellitus.** *Cardiovasc Diabetol* 2012, **11**:109. [Epub ahead of print].
228. Kasai T, Floras JS, Bradley TD: **Sleep apnea and cardiovascular disease: a bidirectional relationship.** *Circulation* 2012, **126**:1495–1510.
229. Jaffe LM, Kjekshus J, Gottlieb SS: **Importance and management of chronic sleep apnoea in cardiology.** *Eur Heart J* 2012, 2012 Mar 16. [Epub ahead of print].
230. Baguet JP, Barone-Rochette G, Tamisier R, Levy P, Pépin JL: **Mechanisms of cardiac dysfunction in obstructive sleep apnea.** *Nat Rev Cardiol* 2012, 2012 Sep 25. doi: 10.1038/nrcardio.2012.141. [Epub ahead of print].
231. Mirrahkhimov AE: **Obstructive sleep apnea and kidney disease: is there any direct link?** *Sleep Breath* 2011, 2011 Dec 15. [Epub ahead of print].
232. Mirrahkhimov AE: **Supine fluid redistribution: should we consider this as an important risk factor for obstructive sleep apnea?** *Sleep Breath* 2012, 2012 Aug 8. [Epub ahead of print].
233. Mannarino MR, Di Filippo F, Pirro M: **Obstructive sleep apnea syndrome.** *Eur J Intern Med* 2012, **23**:586–593.
234. McNicholas WT: **Chronic obstructive pulmonary disease and obstructive sleep apnea: overlaps in pathophysiology, systemic inflammation, and cardiovascular disease.** *Am J Respir Crit Care Med* 2009, **180**:692–700.
235. Nadel JA, Widdicombe JG: **Reflex effects of upper airway irritation on total lung resistance and blood pressure.** *J Appl Physiol* 1962, **17**:861–865.
236. Machado MC, Vollmer WM, Togeiro SM, Bilderback AL, Oliveira MV, Leitão FS, Queiroga F Jr, Lorenzi-Filho G, Krishnan JA: **CPAP and survival in moderate-to-severe obstructive sleep apnoea syndrome and hypoxaemic COPD.** *Eur Respir J* 2010, **35**:132–137.
237. Marin JM, Soriano JB, Carrizo SJ, Boldova A, Celli BR: **Outcomes in patients with chronic obstructive pulmonary disease and obstructive sleep apnea: the overlap syndrome.** *Am J Respir Crit Care Med* 2010, **182**:325–331.
238. Lesser DJ, Bhatia R, Tran WH, Oliveira F, Ortega R, Keens TG, Mittelman SD, Khoo MC, Davidson Ward SL: **Sleep fragmentation and intermittent hypoxemia are associated with decreased insulin sensitivity in obese adolescent Latino males.** *Pediatr Res* 2012, **72**:293–298.
239. Jain SK, Kahlon G, Morehead L, Lieblong B, Stapleton T, Hoeldtke R, Farrington Bass P 3rd, Levine SN: **The Effect of Sleep Apnea and Insomnia on Blood Levels of Leptin, Insulin Resistance, IP-10, and Hydrogen Sulfide in Type 2 Diabetic Patients.** *Metab Syndr Relat Disord* 2012, **10**:331–336.
240. Qian X, Yin T, Li T, Kang C, Guo R, Sun B, Liu C: **High levels of inflammation and insulin resistance in obstructive sleep apnea patients with hypertension.** *Inflammation* 2012, **35**:1507–1511.
241. Pamidi S, Tasali E: **Obstructive sleep apnea and type 2 diabetes: is there a link?** *Front Neurol* 2012, **3**:126.
242. Fredheim JM, Rollheim J, Omland T, Hofso D, Røislien J, Vegsgaard K, Hjelmæsæth J: **Type 2 diabetes and pre-diabetes are associated with obstructive sleep apnea in extremely obese subjects: a cross-sectional study.** *Cardiovasc Diabetol* 2011, **10**:84.
243. Pallayova M, Steele KE, Magnuson TH, Schweitzer MA, Hill NR, Bevans-Fonti S, Schwartz AR: **Sleep apnea predicts distinct alterations in glucose homeostasis and biomarkers in obese adults with normal and impaired glucose metabolism.** *Cardiovasc Diabetol* 2010, **9**:83.
244. Togeiro SM, Carneiro G, Ribeiro Filho FF, Zanella MT, Santos-Silva R, Taddei JA, Bittencourt LR, Tufik S: **Consequences of Obstructive Sleep Apnea on Metabolic Profile: A Population-Based Survey.** *Obesity (Silver Spring)* 2012, 2012 Jun 14. doi: 10.1038/oby.2012.146. [Epub ahead of print].

245. Tamura A, Kawano Y, Watanabe T, Kadota J: **Obstructive sleep apnea increases hemoglobin A1c levels regardless of glucose tolerance status.** *Sleep Med* 2012, **13**:1050–1055.
246. Priou P, Le Vaillant M, Meslier N, Chollet S, Masson P, Humeau MP, Pigeanne T, Bizieux-Thaminy A, Goupil F, Gagnadoux F: **Independent Association Between Obstructive Sleep Apnea Severity and Glycated Hemoglobin in Adults Without Diabetes.** *Diabetes Care* 2012, **35**:1902–1906.
247. Pamidi S, Wroblewski K, Broussard J, Day A, Hanlon EC, Abraham V, Tasali E: **Obstructive Sleep Apnea in Young Lean Men: Impact on insulin sensitivity and secretion.** *Diabetes Care* 2012, 2012 Aug 21. [Epub ahead of print].
248. Botros N, Concato J, Mohsenin V, Selim B, Doctor K, Yaggi HK: **Obstructive sleep apnea as a risk factor for type 2 diabetes.** *Am J Med* 2009, **122**:1122–1127.
249. Lindberg E, Theorell-Haglöw J, Svensson M, Gislason T, Berne C, Janson C: **Sleep apnea and glucose metabolism - a long-term follow-up in a community-based sample.** *Chest* 2012, **142**:935–942.
250. Wang X, Bi Y, Zhang Q, Pan F: **Obstructive sleep apnoea and the risk of type 2 diabetes: a meta-analysis of prospective cohort studies.** *Respirology* 2012, 2012 Sep 18. doi: 10.1111/j.1440-1843.2012.02267.x. [Epub ahead of print].
251. Sharma SK, Agrawal S, Damodaran D, Sreenivas V, Kadiravan T, Lakshmy R, Jagia P, Kumar A: **CPAP for the metabolic syndrome in patients with obstructive sleep apnea.** *N Engl J Med* 2011, **365**:2277–2286.
252. Weinstock TG, Wang X, Rueschman M, Ismail-Beigi F, Aylor J, Babineau DC, Mehra R, Redline S: **A controlled trial of CPAP therapy on metabolic control in individuals with impaired glucose tolerance and sleep apnea.** *Sleep* 2012, **35**:617–625B.
253. Duarte FH, Jallad RS, Amaro AC, Drager LF, Lorenzi-Filho G, Bronstein MD: **The impact of sleep apnea treatment on carbohydrate metabolism in patients with acromegaly.** *Pituitary* 2012, 2012 Sep 16. [Epub ahead of print].
254. Roemmler J, Gutt B, Fischer R, Vay S, Wiesmeth A, Bidlingmaier M, Schopohl J, Angstwurm M: **Elevated incidence of sleep apnoea in acromegaly-correlation to disease activity.** *Sleep Breath* 2012, 2012 Jan 13. [Epub ahead of print].
255. Yang D, Liu Z, Yang H, Luo Q: **Effects of continuous positive airway pressure on glycemic control and insulin resistance in patients with obstructive sleep apnea: a meta-analysis.** *Sleep Breath* 2012, 2012 Mar 13. [Epub ahead of print].
256. Mirrakhimov AE, Polotsky VY: **Obstructive Sleep Apnea and Non-alcoholic Fatty Liver Disease: Is the Liver Another Target?** *Front. Neur* 2012, **3**:149.
257. Paredes AH, Torres DM, Harrison SA: **Nonalcoholic fatty liver disease.** *Clin Liver Dis* 2012, **16**:397–419.
258. Bhatia LS, Curzen NP, Calder PC, Byrne CD: **Non-alcoholic fatty liver disease: a new and important cardiovascular risk factor?** *Eur Heart J* 2012, **33**:1190–1200.
259. Targher G, Chonchol M, Zoppini G, Abaterusso C, Bonora E: **Risk of chronic kidney disease in patients with non-alcoholic fatty liver disease: is there a link?** *J Hepatol* 2011, **54**:1020–1029.

doi:10.1186/1475-2840-11-132

Cite this article as: Mirrakhimov: Chronic obstructive pulmonary disease and glucose metabolism: a bitter sweet symphony. *Cardiovascular Diabetology* 2012 **11**:132.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

