Morphological Assessment of Changes in the Macro- and Microvascular System of Lung Tissue in Diabetes Mellitus

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Abstract: Diabetes mellitus is a chronic metabolic disease that affects various organs, including the lungs. The morphological changes of the macro- and microvascular systems of lung tissue in diabetes mellitus remain largely unknown, which affects our understanding of the pathogenesis of diabetic lung diseases. This article aims to review recent research on morphological assessments of changes in the macro- and microvascular system of lung tissue in diabetes mellitus, with a focus on the available techniques and their limitations, as well as their potential applications in preventing and treating diabetic lung diseases.

Keywords: Diabetes mellitus, lung tissue, macrovascular system, microvascular system, morphological assessment.

Introduction

Diabetes mellitus (DM) is a common metabolic disease characterized by hyperglycemia resulting from insulin insufficiency or resistance, which can cause various complications such as neuropathy, retinopathy, and nephropathy. The respiratory system is also affected by DM, leading to diabetic lung diseases such as pulmonary infections, obstructive airway diseases, and interstitial lung diseases (1, 2). However, despite the growing awareness of lung complications in DM, the precise mechanisms are still poorly understood.

One of the potential ways DM affects the lungs is through vascular changes. DM has been associated with vascular complications in various organs, including the heart, kidneys, and eyes, due to the damage done to the macro- and microvascular systems. Similarly, lung tissue can experience significant structural changes in the macro- and microvasculature in DM, leading to impaired lung function and respiratory diseases (1). These morphological alterations can include thickening and fibrosis of the pulmonary vessel walls, an increase in the number and size of alveolar capillaries, and a reduced number of peripheral pulmonary vessels.

To better understand the morphological changes in the macro- and microvascular systems of lung tissue in DM, various imaging techniques have been utilized, including computed tomography (CT) scans, magnetic resonance imaging (MRI), and histopathologic assessments. CT scans and MRI provide non-invasive methods to assess lung morphology and function but may lack the ability to detect finer features of the vasculature. Microscopic examination of lung tissue is still the gold standard for morphological analysis, but it is invasive and subject to sampling bias due to the heterogeneity of the lung (1). Practical non-invasive imaging techniques are therefore needed to understand and diagnose diabetic lung disease.
Several studies have attempted to investigate the morphological changes of the macro- and microvascular systems of lung tissue in DM using these imaging techniques. In a study using contrast-enhanced CT scans, researchers found that diabetic patients had significant vascular abnormalities, including vascular obstruction, segmental pulmonary artery enlargement, and increased bronchial artery diameters, compared to non-diabetic individuals. Similarly, a study using endobronchial ultrasound (EBUS) found that diabetic patients had significantly increased thickness of the bronchial wall and the submucosa of the central and peripheral bronchi, which may indicate morphological changes in the small airways consistent with diabetic lung disease (2). These findings suggest that morphological assessment of the macro- and microvascular system of lung tissue could provide valuable diagnostic information in DM and contribute to the early detection and treatment of diabetic lung disease.

However, while these imaging techniques can be informative, they still have limitations. For instance, CT scans and MRI may be affected by motion artifacts or respiratory motion, which may lead to image degradation and inaccuracies in the analysis. There may also be variations in the MRI signal intensity depending on the field strength and pulse sequences used (3). Histopathologic analysis, on the other hand, is invasive and may not be feasible for some patients, particularly those with severe respiratory disease. Furthermore, the potential confounding effect of other comorbidities on lung morphology in DM needs to be considered (8).

Despite these limitations, morphological assessment of the macro- and microvascular system of lung tissue in DM is a growing field of study that has the potential to improve our understanding of the pathogenesis of diabetic lung disease. This article aims to review recent research on morphological assessments of changes in the macro- and microvascular system of lung tissue in DM, with a focus on technique limitations and their potential applications in diagnosing and treating diabetic lung diseases.

**Methods**

Diabetes mellitus is a metabolic disorder characterized by hyperglycemia that affects various organs, including the lungs. The lung tissue macro- and microvascular systems undergo structural and functional alterations in diabetes mellitus. Morphological assessment of these changes in the lung tissue micro- and macrovasculature is essential to understand the mechanisms underlying the development of lung complications in diabetes mellitus.

A histomorphometric study could help to evaluate the lung tissue's morphological alterations in patients with diabetes mellitus. This study examines structural changes in the lung tissue microvascular system using various techniques to determine the alveolar surface area density, alveolar septal thickness, vascular density, and the arterial wall thickness. Alveolar septal thickness and alveolar surface area density could be measured using morphometric analysis by the ImageJ program (10, 11). Vascular density and arterial wall thickness could be determined by hematoxylin and eosin (H&E) staining and quantitative analysis by the ImageJ program. This technique has been widely used to evaluate morphometric alterations in the lung tissue of rats with diabetes mellitus (4).

Transmission electron microscopy (TEM) could also help to assess ultrastructural changes in the lung tissue microvascular system. TEM can visualize the morphology of lung tissue cells and extracellular matrix, enabling researchers to describe and quantify morphological changes in the basement membrane and capillary endothelium of the lung tissue. TEM has been used to evaluate ultrastructural changes in the lung tissue of rats with diabetes mellitus (5).

Moreover, vascular corrosion casting is a method used to produce a three-dimensional replica of the lung microvasculature. This technique utilizes a transparent resin injection that fills the vascular network, followed by digestion of the tissue to expose the vascular structures. The resin cast of the microvasculature shows the branching patterns and structure of the lung microvasculature, and the morphology of the alveolar capillaries, which provide gas exchange between alveolar air and blood. This technique allows researchers to analyze the effects of diabetes mellitus on the lung tissue macrovasculature (4, 7).

Animal models have been widely used to investigate the mechanisms, pathologies, and therapeutic strategies in diabetes mellitus. Transgenic mouse models of diabetes mellitus, such as streptozotocin-induced diabetes, could be utilized to conduct preclinical studies on lung tissue morphology. These models show the pathological changes, mortality rates, and complications akin to human diabetes mellitus (6). Animal models
could be used to study changes in the lung tissue’s micro- and macrovascular systems in diabetes mellitus through histomorphometric analysis, TEM, vascular corrosion casting, and confocal microscopy.

Finally, in addition to morphological assessments, functional assessments could be carried out to complement these analyses. Pulmonary function tests could be used to evaluate the lung's functional changes resulting from the morphological alterations. This assessment could also help to determine the effects of diabetes mellitus on lung function and evaluate the efficacy of the therapeutic agents deployed to reduce lung complications in diabetes mellitus.

To conclude, morphological assessment of changes in the macro- and microvascular system of lung tissue in diabetes mellitus could provide critical insights into the mechanisms underlying diabetic lung complications. These morphological assessments could help researchers to identify new therapeutic agents or modify existing therapies. Studies targeting lung complications in diabetes mellitus could benefit from the application of various techniques of lung tissue morphometry, such as histomorphometric analysis, TEM, vascular corrosion casting, confocal microscopy, and pulmonary function tests (9). These techniques could contribute to the generation of knowledge with the potential to reduce significant morbidity and mortality rates in diabetes mellitus patients.

Conclusion

In conclusion, diabetes mellitus is a disease that has significant and diverse effects on various organ systems in the body, including the lung macro- and microvascular systems. These changes manifest in the form of alterations in the lung parenchyma and airway architecture, and impaired blood supply to the lung tissue. These changes ultimately lead to lung damage, respiratory failure and other pulmonary complications in diabetic patients.

Morphological analysis of the lung tissue, involving the use of high-resolution imaging techniques, has provided crucial insights into the pathophysiological mechanisms underlying diabetic lung disease. The studies reviewed in this article have highlighted the complex interplay between metabolic imbalances, systemic inflammation, and oxidative stress in the development of diabetic lung disease. Understanding these mechanisms can help in the development of targeted therapeutics that can prevent, delay or even reverse the onset and progression of diabetic lung disease.

Moreover, the assessment of morphological changes in the macro- and microvascular structure of lung tissue is a valuable diagnostic and prognostic tool for diabetic patients with respiratory symptoms. The use of advanced imaging techniques, such as CT scans and MRI, can help in the early detection and monitoring of lung damage in diabetes mellitus. Further studies are needed to determine the optimal imaging protocols and biomarkers for the precise identification and quantification of lung disease in diabetic patients.

In conclusion, morphological assessment of the lung tissue in diabetes mellitus is a rapidly evolving field with significant implications for early diagnosis, prognosis, and targeted therapy. The application of advanced imaging techniques and the integration of morphological data with clinical and biochemical parameters can offer new insights into the pathophysiology of diabetic lung disease, and provide a rational basis for the development of effective intervention strategies.

References