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Original Article

# The Protective Effect of The Interleukin 1 Receptor Antagonist on Chronic Thromboembolic Pulmonary Hypertension Model

## Wpływ ochronny antagonisty receptora interleukiny 1 w modelu przewlekłego zakrzepowo-zatorowego nadciśnienia płucnego

**Authors' Contribution:**
A Study Design
B Data Collection
C Statistical Analysis
D Data Interpretation
E Manuscript Preparation
F Literature Search
G Funds Collection

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## Summary

**Aim:** Chronic thromboembolic pulmonary hypertension (CTEPH) is one of the main reasons of severe pulmonary hypertension and has significantly higher morbidity and mortality rates. The pathogenesis of the disease is characterized by the incomplete resolution of acute embolisms. The elevated inflammatory conditions after the acute embolism are one of the critical factors. Therefore, we aimed to investigate whether or not anakinra is an option for treating CTEPH in an animal model.

**Material/Methods:** We studied twenty-one rats in this study They were randomly divided into three groups containing seven animals: the control group: saline-treated control; the embolism group: CTEPH + normal saline; the anakinra group: CTEPH + anakinra.

**Results:** We have observed that the layers of the segmental arteries and the alveolar were normal in the control group. In the cardiac tissue it was observed that muscular tissues and connective tissue were normal in the right ventricle. In embolism group, we detected a widening of the alveolar septum, a surrounding the alveolar infiltrates and a thickening of the segmental arteries in the muscular layer and a hypertrophy in the right ventricle tissues. We have determined that the lung and cardiac tissue specimens in the anakinra group are similar to control group.

**Conclusions:** We have showed that anakinra was useful option for the CTEPH model in rats. Anakinra may be considered as protective effect and the regression of the increased inflammation in CTEPH. The effectiveness of anakinra will continue to be subject to the further experimental and clinical studies.

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## INTRODUCTION

Chronic thromboembolic pulmonary hypertension (CTEPH) is a life-threatening result of acute pulmonary embolism. Although the incidence of CTEPH after pulmonary embolism is very low, there is a high risk of morbidity or mortality [11, 25]. Without treatment or complete resolution of acute pulmonary embolism, the pulmonary arteries become clogged with progressive organized fibrotic materials that cause pulmonary artery hypertension and progressive right heart failure and death [3]. Although the mechanisms related to the incomplete resolution of acute embolisms and progression of CTEPH are not yet fully understood, several responsible mechanisms are considered.

CTEPH is correlated with the high prevalence of inflammatory conditions and inflammatory diseases [6]. After the embolism appears, the physiologic response to a thrombus is organized by leukocyte migration into the embolic area through fibrinolysis and angiogenesis [18]. It has been shown that inflammatory cells (e.g. CD45+) increased in the area of an incomplete embolism, and collagen-secreting cells were found in the vascular walls of patients with CTEPH [33]. In addition, it has been reported that tumor necrosis factor-alfa, monocyte chemoattractant protein (MCP)-1 and C-reactive protein levels platelet EC adhesion molecule 1 (PECAM-1) are increased in that area [18, 25]. A significant positive correlation between MCP-1 and pulmonary vascular resistance is certain [25]. Zabini et al. [34] reported high levels of inflammatory factors in the pulmonary thromboendarterectomy materials and serums. Bonderman et al. [4] showed the relationship between delayed thrombosis resolution and staphylococcus infections.

Pulmonary thromboendarterectomy is a curative surgical choice for the treatment of CTEPH [11]. However, there is limited access to surgical centers, experienced staff and equipment, even if the patient can undergo the operation [24]. Though many medical treatment alternatives have been utilized for inoperable patients, it has been proven that only the soluble guanylate cyclase stimulator, known as riociguat, is beneficial in the treatment [12]. Previously, in patients with pulmonary arterial hypertension, it has been reported that serum levels of interleukin (IL) 1beta were increased and correlated with a worse outcome, and anakinra (IL-1 beta receptor antagonist) has been shown beneficial in the treatment of pulmonary hypertension in a patient with Adult-Onset Still's Disease (AOSD) [8, 27]; however, the efficacy of the IL-1 beta receptor antagonist (anakinra) on CTEPH remains unknown [7]. Our aim is to evaluate the effectiveness of anakinra on an experimental rat model of CTEPH.

## MATERIAL AND METHODS

### Experimental Design

We studied twenty-one rats in this study. They were two-month-old males weighing 200–300 gr. The rats were obtained from the Dokuz Eylul University Clinical Experimental Animals Multi-Disciplinary Laboratory. The animals were given standard rat chow food and water ad libitum. All animals were caged at 18°C to 20°C in 65% to 70% relative humidity and on a 12h light-dark cycle. All experimental procedures complied with the requirements of the Animal Care and Ethics Committee of the Dokuz Eylul University (03/15/2017, protocol no: 05/2017) before the experiments. All participants were certified for the care and use of laboratory animals. Following a week adaptation period, the animals were weighed and allocated into three groups; control, embolism and anakinra groups (total 21, n = 7/group). No deaths were encountered during the study.

The rat model of CTEPH was modified from those previously described [23]. The animals were anesthetized with an intraperitoneal application of ketamine/xylazine (40 mg/kg+5 mg/kg). To generate an embolism, 2 ml of blood was collected from the tail veins of each animals. After keeping the blood samples in a petri dish for 1 day in 37$^0$C, 15 thromboses/petri dishes were generated (dimensions of thromboses: 1x3 mm). Fifteen thromboses were placed into a serum physiologic solution.

In the control group, the rats were injected with an intraperitoneal serum physiologic solution through

the tail veins on day 0 and 14 of the experiment. In the embolism group, 15 thromboses in 2 ml physiologic saline solutions were injected through the tail veins on day 0 and 14 of the experiment. Additionally, physiologic saline solution was applied intraperitoneally all days, expect on day 0 and 14. In anakinra group, 15 thromboses in 2 ml physiologic saline solutions were injected through the tail veins on day 0 and 14 of the experiment, as was the case in the embolism group. Additionally, anakinra (10 mg/kg/day) [16] was applied intraperitoneally on all days, expect on day 0 and 14. In the control group, an intraperitoneal serum physiologic solution was injected to rats through the tail veins at the initiation of the experiment and on day 14 of the experiment. In the embolism group, 15 thromboses in 2 ml physiologic saline solutions were injected through the tail veins at the beginning of the study and on day 14 of the experiment. Additionally, physiologic saline solution was applied intraperitoneally on all days, expect at the initiation of the experiment and on day 14. In the anakinra group, 15 thromboses in 2 ml physiologic saline solutions were injected through the tail veins at initiation of the experiment and on day 14 of the experiment like in the embolism group. Additionally, anakinra (10 mg/kg/day) [16] was applied intraperitoneally all days expect the beginning of the study and on day 14.

The injection of thrombus was well-tolerated in this study. Cyanosis and shortness of breath developed in rats and disappeared after a short time. Additionally, the flow rate was measured (0.5 ml/min). The experiment ended day 28. The rats were sacrificed after four weeks, and the lung and cardiac tissue samples were harvested.

### Histological Analysis

After the tissues were harvested from the animals, the lung and cardiac tissues were fixed in a 10% buffered formalin for 48 h and subsequently embedded into the paraffin. The paraffin blocks were placed into a rotary microtome (RM 2255, Leica, Germany). Sections of 5 µm thickness were obtained. After deparaffinization and rehydration, all sections were stained with hematoxylin-eosin (H-E) and Masson's trichrome stain.

Masson's Trichrome Stain: Lung and cardiac tissues were deparaffinized and rehydrated with 100% alcohol, 95% alcohol 70% alcohol, respectively. The tissues were rinsed under running tap water for 5–10 minutes to remove the yellowish color. The tissues were stained with Wiegert's iron hematoxylin working solution for 10 minutes. After washing, the tissues were stained with Biebrich scarlet-acid fuchsin solution for 10–15 minutes. The sections were washed in distilled water again and differentiated in phosphomolybdic-phosphotungstic acid solution for 10-15 minutes. The tissues were transferred into the aniline blue solution for 5–10 minutes. After rinsing, they were differentiated in 1% acetic acid solution for 2–5minutes. The tissues were dehydrated using the increasing alcohol series. They were mounted with the help of a mounting medium (Nova Ultra Special Stain Kits).

All sections from cardiac and lung tissues were evaluated by the same pathologist, blinded from groups.

### Statistical Analysis

All analyses were performed using SPSS 16.0 for Windows (v16.0, SPSS Inc., Chicago, Illinois). The data was expressed as categorical classification based on the functional classification of World Health Organization (WHO-FC), unless otherwise stated. Fisher's Exact test and McNemar's test were used to classify the groups before and after the experiments separately. $P < 0.05$ was considered as statistically significant.

**Table 1.** The evaluation of symptoms of all animals is made according to WHO-FC

|  | Groups | | | |
|---|---|---|---|---|
|  | Control | Embolism | Anakinra | p |
| Before Experiments | | | | |
| Class 1 | 7(100) | 7(100) | 7(100) | |
| Class 2 | 0(0) | 0(0) | 0(0) | |
| Class 3 | 0(0) | 0(0) | 0(0) | N/A |
| Class 4 | 0(0) | 0(0) | 0(0) | |
| After Experiments | | | | |
| Class 1 | 7(100) | 2(28.57) | 6(85.71) | |
| Class 2 | 0(0) | 0(0) | 0(0) | |
| Class 3 | 0(0) | 4(57.14) | 1(14.29) | 0.017 |
| Class 4 | 0(0) | 1(100) | 0(0) | |

Fisher's Exact test is used. WHO–FC = World Health Organization functional class.

**Table 2.** Comparison of percentile of all animals is made according to WHO-FC

| Class 1 | Before Experiment | | | After Experiment | | | |
|---|---|---|---|---|---|---|---|
| | Class 2 | Class 3 | Class 4 | Class 1 | Class 2 | Class 3 | Class 4 |
| Control | 7(100) | 0(0) | 0(0) | 0(0) | 7(100) | 0(0) | 0(0) | 0(0) |
| Embolism | 7(100) | 0(0) | 0(0) | 0(0) | 2(28.57) | 0(0) | 4(57.14) | 1(14.29) |
| Anakinra | 7(100) | 0(0) | 0(0) | 0(0) | 6(85.71) | 0(0) | 1(14.29) | 0(0) |

McNemar test is used.

## RESULTS

We evaluated all the animals according to WHO-FC for pulmonary hypertension symptoms both at the initiation of the experiments and at day 28. In all groups, none of the animals showed any symptoms related to WHO-FC at the beginning of the experiments.

At the beginning of the study, all the animals were accepted as Class-I. In the control group, none of the animals had pulmonary hypertension symptoms at the end of day 28. Additionally, all the control animals were accepted class-I at the end of the experiment. In the embolism group, 4 rats of 7 were classified as class-3, one of them classified as class-4 and the others were classified as class-1. In the anakinra group, only 1 rat of 7 was classified as class-3 while the others were classified as class-1 at the end of experiments (Table 1).

After harvesting the tissues and preparing the sections, a gross light microscopy evaluation was made by one same pathologist, blinded from groups. Changes in histological findings between groups were listed in Table 2.

In sections of the embolism group, thickening of the segmental arteries muscular layer, atrophy of endothelial structure, hypertrophy in the right ventricular tissue, and an increase in the interstitial tissue samples were reported in all animals (p <0.01). The light microscopic examinations of the control lung sections revealed that the alveolar structures and segmental arterial layers were intact and not disrupted (Fig. 1A). Additionally, the muscular and connective tissue structures were intact and not disrupted in the evaluation of the right ventricular sections of the heart (Fig. 2A, B). The specimens of both lung and cardiac tissue in the anakinra group yielded similar findings as those in the control group (Fig. 1C, 2E, 2F).

However, the light microscopic examinations of the embolism group sections revealed that the alveolar septum was widened in the lung sections due to the increase in the interstitial tissue and widespread
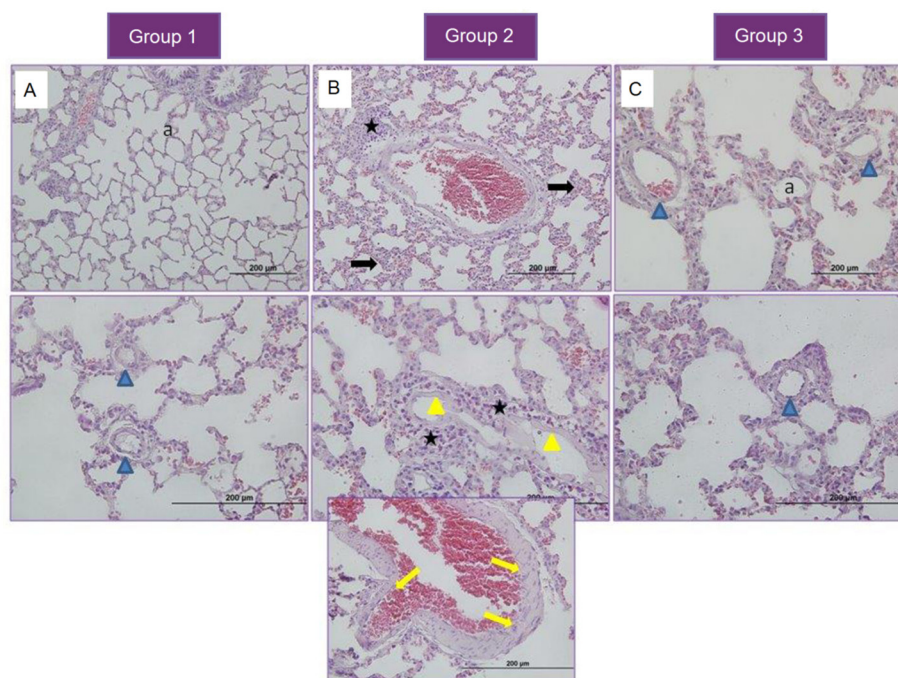


**Fig. 1.** The H&E staining of the lung is shown as following; a: Alveolus; the black arrow: the thickened alveolar septum; the star: mononuclear leukocytes infiltration; the yellow arrow: Hypertrophic segmental arteries; the tip of the blue arrow: segmental arterial branches of a normal structure; the tip of the yellow arrow: irregular segmental arterial branches

polymorph nuclear leukocytes infiltration; the muscular layer of the segmental arteries were thickened, endothelial cell structure were atrophic (Figure 1B); the right ventricular tissue was hypertrophic; inflammatory cells were infiltrated and the connective tissue fibrils were increased (Fig. 2C, 2D).

## DISCUSSION

In this study, we have assessed the effect of anakinra on CTEPH model in rats. The animals were evaluated at the beginning and the end of experiment according to WHO-FC for pulmonary hypertension. After the experiments, the lung and cardiac tissues of the rats were collected and analyzed histologically among groups for the purpose of evaluating the effects of anakinra on the tissue level. The findings in tissue evaluation and symptoms of embolism in embolism group were significantly different than those in the control and anakinra groups. Additionally, we have shown that histological parameters were decreased in the anakinra group compared to those of the embolism group. There are only a few studies performed on animal models (dog and porcine) for CTEPH. In these models, prepared thromboses are directly applied through pulmonary arteries in animals in order to incite an embolism, so that the CTEPH model can progress in animals [9, 15]. Li et al. [21] has used an animal model in which 15 thromboses (each measuring 1x3 mm) were injected through the jugular veins of rats to mimic embolism conditions. They also reported that the rats showed cyanosis and shortness of breath after the application [21]. We developed an animal model with CTEPH by inducing of 15 thromboses. Our results confirmed the presence of the model with showing WHO-FC class 3 to 4 individuals in embolism group.

Zabini et al. [34] and Mercier et al. [22] have demonstrated the critical role of many inflammatory cytokines in the formation of obstructive structures in the pulmonary arteries of CTEPH and reported the high levels of growth factors (fibroblast growth factor 2) and inflammatory cytokines (IL 1 beta, IL 6, monocyte chemoattractant protein 1), and cell adhesion molecules in CTEPH [22, 34]. They also showed the effects of these factors on the development of CTEPH [22, 34].

IL-1 is a pro-inflammatory cytokine that has wide range of effects on metabolism and such mechanisms as pain sensitivity and tissue damage. The stress exposure of cell triggers a deployment of inflammasomes, which are molecular platforms that lead to caspase-1 activation. These signals are essential for the activation of IL 1-beta [26]. It has been shown that chronic inflammation leads to cystic fibrosis and lung fibrosis in the animal models and the indication of treatment with IL-1 receptor antagonists was revealed [7, 16]. There is considerable clinical evidence and studies suggesting that the blockade of IL-1 decreases the severity of diseases and reverses inflammation related loss of organ function in autoinflammatory syndromes [10]. Thus, blockade of IL-1 can be an opportunity for the treatment of those auto-inflammatory syndromes. Anakinra is an approved recombinant form of IL-1 receptor antagonist and a new treatment option for a wide range of diseases associated with inflammatory conditions, such as rheumatoid arthritis [1], familial Mediterranean fever [2], and diabetes [20]; however, the usage of anakinra is not limited with those diseases. The effectiveness of anakinra has been described in the case of pulmonary arterial hypertension [8] as well as in a series of AOSD patients [26]. Elevated IL-1 beta serum levels in PAH correlate with worse outcomes [27]. However, to our knowledge, there
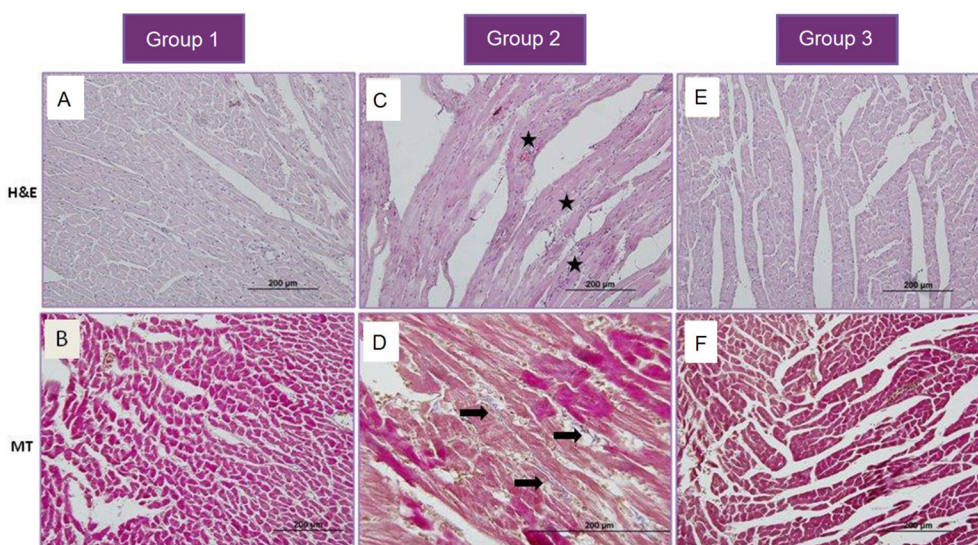


**Fig. 2.** The Masson's trichrome staining of cardiac tissue is shown as following: the star: polymorph nuclear leukocytes infiltration; the black arrow: fiber increase in connective tissue; H&E: Hematoxylin & Eosin stain; MT: Masson's trichrome stain

are not any studies about the use of anakinra in Group 4 pulmonary artery hypertension or CTEPH. In our study, we aimed to investigate whether or not anakinra is an option for the treatment of CTEPH in an animal model.

CTEPH is an intimal disease of organized thrombus and vascular remodeling, with progressive intimal thickening (collagenous, inflammatory, atherosclerotic, hemosiderosis and calcific) resulting in right ventricular pressure overload and remodeling with ultimately decompensation, right ventricle failure and death [19]. Normally, acute pulmonary embolisms resolve within 6 months. However, in 0.1-3.8 % of cases, the transition of thrombosis to a fibrous tissue leads to pulmonary hypertension [3]. Other risk factors for chronic pulmonary thromboembolism are elevated Factor VIII, antiphospholipid antibody, lupus anticoagulants, splenectomy and cancer [5, 6, 32]. In addition, ventricular-atrial (VA) shunts, which are used in the treatment of hydrocephaly, may result in CTEPH. In some cases there are no risk factors for venous thromboembolism, and there may be a relationship between vascular remodeling and staphylococcal infections [4]. In addition to the up-regulation of transforming growth factors and IL-1 beta, the staphylococcal infections can lead to a delay in thrombosis resolution [28].

For a period of 16 weeks, the dual endothelin antagonist, bosentan was evaluated in 157 patients with inoperable CTEPH or persistent/recurrent pulmonary hypertension after a pulmonary endarterectomy [17]. This study indicated that the decrease in pulmonary vascular resistance and increase in the 6-minute walking capacity could not reach the ultimate goal [17]. However, in the MERIT-1 study, patients with inoperable CTEPH were treated with another endothelin receptor antagonist (macitentan), and macitentan significantly improved pulmonary vascular resistance [14]. In addition, for a 16-week period,

an oral cyclic guanosine monophosphate stimulator, riociguat was applied to patients with non-operable CTEPH or persistent/recurrent pulmonary hypertension after a pulmonary endarterectomy. Riociguat was observed to be useful via enhanced 39 meters the 6-minute walking test. A CTEPH team, which included at least one experienced surgeon, has recommended riociguat for the treatment of symptomatic patients with inoperable CTEPH or with persistent or recurrent CTEPH after surgical treatment [13].

The pathogenesis of CTEPH has not yet been fully understood. Despite this fact, its relation to venous thromboembolism raises the following question: Is CTEPH a disorder of misguided vascular remodeling after pulmonary thromboembolism? In some cases, genetic factors can play a significant role leading to a larger thrombus (non-O- blood groups, rare thrombophilia, dysfibrinogenemia) in combination with elevated factor VIII and platelet activation [16]. Recently, these factors should take into consideration about disease phatogenesis. These conditions suppress the two key pathways: thrombosis angiogenesis and innate immune cell function. It is not clear whether similar mechanisms contribute to secondary pulmonary arteriopathy. Alleviating the major vessel obstruction in CTEPH can provide an understanding of the mechanisms of regression of secondary pulmonary arteriopathy and can lead to new treatment objectives.

## CONCLUSION

We have showed that anakinra was a useful option for the CTEPH model in rats. Anakinra may be considered to have a protective effect and can contribute to the regression of the increased inflammation in CTEPH. The effectiveness of anakinra will continue to be subject to the further experimental and clinical studies.

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