STUDY DOCUMENT

Official Study Title:
The Effect of Prone Position on Oxygen Saturation, Blood Gas Parameters, and Respiratory Rate in Intensive Care Patients with COVID-19-induced ARDS

NCT Number: NCTXXXXXXXX

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INFORMED CONSENT FORM

1. Study Title

The Effect of Prone Position on Oxygen Saturation, Blood Gas Parameters, and Respiratory Rate in Intensive Care Patients with COVID-19-induced ARDS

2. Introduction

You are being invited to participate in a research study. Please read this form carefully and ask any questions you may have before agreeing to take part.

3. Purpose of the Study

This study aims to investigate whether a short-term prone position can improve oxygen levels and respiratory function in patients with COVID-19-induced ARDS.

4. Procedures

If you agree to participate, you will be randomly assigned to a group. If in the intervention group, you will be placed in a prone position for 30 minutes on two consecutive days. Blood gas levels, oxygen saturation, and respiratory rate will be measured.

5. Voluntary Participation

Your participation in this study is entirely voluntary. You may withdraw at any time without affecting your medical care.

6. Risks and Discomforts

The prone position may cause mild discomfort, including pressure-related discomfort in some areas of the body. All necessary precautions will be taken to minimize risk.

7. Benefits

You may experience improved oxygenation. The information gained from this study may help other patients in the future.

8. Confidentiality

Your information will be kept confidential. Your name will not appear in any published results. Data will be stored securely and used only for research purposes.

9. Contacts

If you have any questions about this study, please contact:
Dr. Veysel Tekin at veyseltekin571453@gmail.com

10. Consent Statement

I have read and understood the information provided above. I have had the opportunity to ask questions and all of my questions have been answered. I voluntarily agree to participate in this study.

Participant Name: _____

Participant Signature: _____ Date: _____

Researcher Name: _____

Researcher Signature: _____ Date: _____