Movair and LUISA

The only vendor partner with a full-featured* home ventilator on the market

Over 20,000 LUISA devices sold in the U.S. with inventory available and immediate shipping from Austin, Texas

WHAT MOVAIR OFFERS

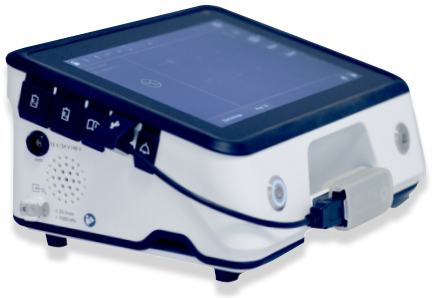
The LUISA® Home Ventilator
PDAC approved for invasive and non-invasive ventilation (E0465 and E0466)

***AUTO RATE:** LUISA is the **only home ventilator with an auto rate feature[1].** LUISA's Auto Rate is the only one that factors in both the patient's respiratory rate and their minute ventilation which is designed to improve patient-ventilator synchrony. With TTV-VAPS-AE, LUISA is **also the only home ventilator offering a combination of Auto Rate, Auto-EPAP and Target Tidal Volume.**

LUISA provides ventilator support and utilizes all standard volume, pressure, and mouthpiece ventilation modes with the added benefit of **High Flow Therapy.** Ventilator modes include LUISA's TTV-VAPS-AE with unique Comfort Settings designed to help patients be more comfortable and compliant.

TTV-VAPS-AE's unique Comfort Settings include:

- Sensitive Inspiratory and Expiratory Triggers with **Inspiratory Lock Out Time**
- Three Target Volume Speeds
- Pressure Increase (Rise) and **Pressure Reduction (Drop)**
- Flow-Based Auto-EPAP
- **Auto-Rate algorithm**



Remote Monitoring NOW Available!

The LUISA home ventilator is an award-winning easy-to-learn and easy-to-use home ventilator – now with remote monitoring!

On-Going Customer Support

- **24/7/365 Clinical Hotline staffed by Movair RTs**
- Clinical device training (in person or virtual) by Movair RTs
- In-person support for sales calls with your clinicians/physicians by Movair RTs
- U.S.-based customer service and repair facility
- Biomed certification training programs

Movair is the exclusive sales, clinical and service partner for LUISA in the U.S.

Find out why LUISA is proven and preferred.

Movair provides competitive pricing, low cost of ownership and financing options.

The LUISA has been authorized by the FDA under an EUA but has not been FDA cleared or approved. The LUISA is authorized only for the duration of the declaration that circumstances exist justifying the authorization of the emergency use under section 564(b)(1) of the Act, 21 U.S.C. § 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner. U.S. federal law restricts this device to sale by or on the order of a physician.

1. Based on search of FDA 510k database for product code NOU as of 1/26/2024

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CE Marked as Class IIb medical device by TÜV Rheinland LGA Products GmbH
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MOVAIR Breath | Life | Empowered