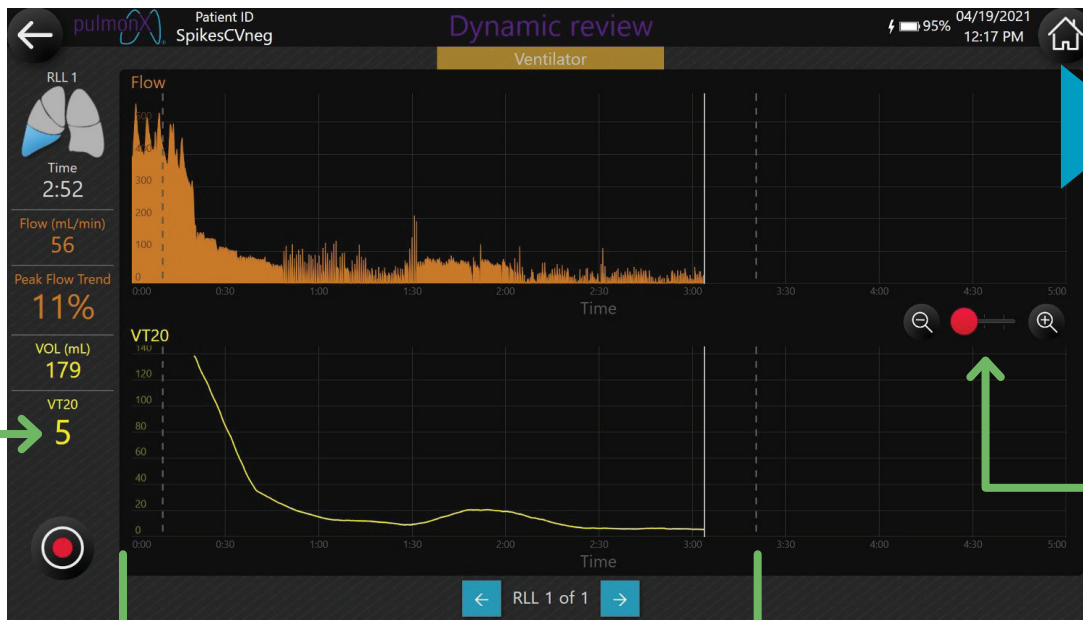




Introducing Chartis[®] V.6.1

Designed For A Simpler & Faster Assessment Experience:



A survey of experienced Chartis physicians reported Chartis V.6.1 is preferred over the previous Chartis software¹

Three level zoom windows enable granular or global view of assessment

1 Simpler

2 Faster

- Chartis V.6.1 software and new VT20 (Volume Trend Over 20 Seconds) feature determines collateral ventilation more easily²
- A study showed that 100% of CV- patients recorded VT20 <6 mL²

- Chartis V.6.1 determines collateral ventilation status faster²
- A study showed assessment times are 60 seconds faster compared to previous Chartis software²

Additional Features:

- Record, export, and share video assessments easily
- Real-time playback and fast-forwarding features

References

1. Physician survey of 10 procedures completed with Chartis V.6.1. Data on File at Pulmonx
2. Koster, TD, Klooster, K, McNamara, H, Shargill, NS, Radhakrishnan, S, Olivera, R, & Slebos, DJ. An adjusted and time-saving method to measure collateral ventilation with Chartis. ERJ Open Res, 2021. doi: <https://doi.org/10.1183/23120541.00191-2021>.

Brief Statement: The Zephyr® Endobronchial Valve is an implantable bronchial valve intended to control airflow in order to improve lung functions in patients with hyperinflation associated with severe emphysema with little to no collateral ventilation, and/or to reduce air leaks. The Zephyr Valve is contraindicated for: Patients for whom bronchoscopic procedures are contraindicated; Evidence of active pulmonary infection; Patients with known allergies to Nitinol (nickel-titanium) or its constituent metals (nickel or titanium); Patients with known allergies to silicone; Patients who have not quit smoking. Use is restricted to a trained physician. Prior to use, please reference the Zephyr Endobronchial System Instructions for more information on indications, contraindications, warnings, all precautions, and adverse events.

zephyr®
valve

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