CASE STUDY InterVapor®



A Broncus Company

Sana Klinikum Hof



Treating Physician: Dr. med. Wolfgang Hohenforst-Schmidt Hospital / Country: Sana Klinikum Hof/ Germany

Patient Background

Age / Gender:	62/male
Disease:	COPD IV
Clinical History:	Progressive COPD
Referral Pathway:	MD Dr. Rumpf, Hof

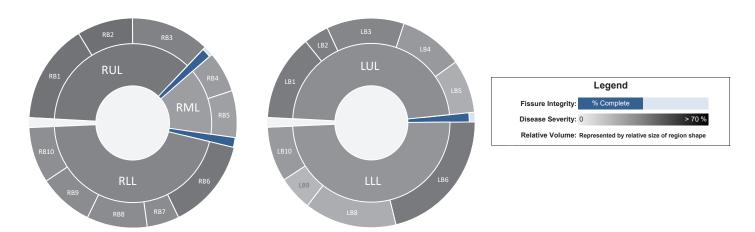
Procedure

Basis of decision to treat with BTVA* (InterVapor®): Incomplete fissure, patient did not want implants. The BTVA treatment has been performed under sedation.

The total treatment time was 12 minutes.

Segments RB1 and RB2 have been treated on a subsegmental and segmental level, respectively.

IMAGES



Pulmonary Function Changes: RV down from 264% to 206%, FEV1 up from 31% to 44%

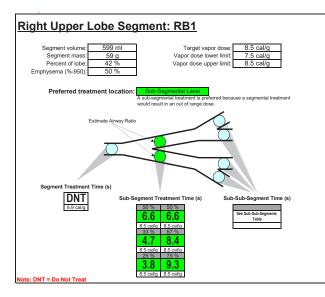
Quality of life changes: Increased activity level, patient feels much better, no oxygen, improved quality of life

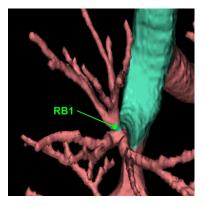
CONCLUSION

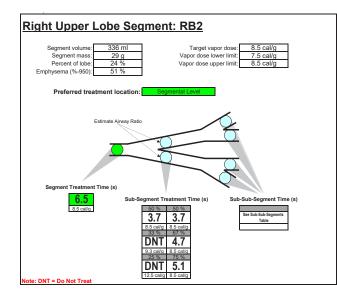
PHYSICIAN PERSPECTIVE ON OUTCOMES:

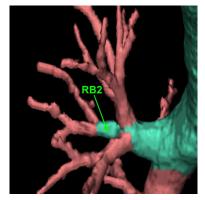
MD Dr. med. Rumpf, the referring physician, indicates that the patient has significant improvement of his pulmonary functions, especially RV and FEV1. BTVA treatment can be used successfully in patients with heterogeneous emphysema with upper lobe predominance. Patients may achieve clinically important ameliorations in physiology, quality of life and exercise tolerance following a single session of unilateral BTVA. The clinical condition of the patient plays a crucial role on the day of admission since he may have no signs of exacerbation.

TREATMENT



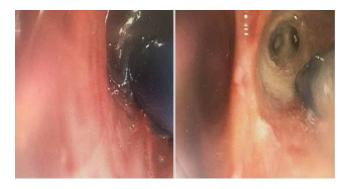






Left picture shows the catheter inside the target lobe with the balloon dilated during the procedure.

The right picture emphasizes the Vapor catheter distally from the target lobe and the ablated area.



Bronchoscopic Thermal Vapor Ablation (BTVA®) treatment included in the

2019 GOLD Guidelines

CONTACT

Uptake Medical[®], a Broncus Company **Phone:** +31 (0) 733032563 **Email:** info@uptakemedical.com or contact your local representative



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