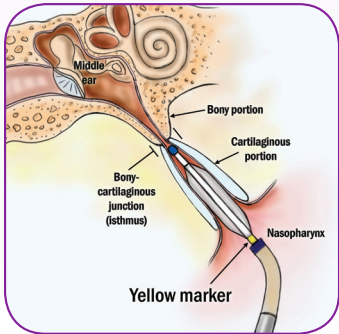


ACCLARENT AERA® EUSTACHIAN TUBE BALLOON DILATION SYSTEM

GO BEYOND SYMPTOMS TO THE SOURCE

The first balloon dilation system intervention indicated for persistent Eustachian Tube Dysfunction (ETD) in the United States



Designed to relieve symptoms of persistent ETD

ETD is a condition caused by the inadequate opening and closing of the Eustachian tubes, and may be causing your patients to experience **chronic ear pressure, pain, and/or clogged or muffled sensations**.¹ Adult prevalence of ETD is estimated to be up to 5%.²

Balloon is inflated to dilate the Eustachian tube to relieve symptoms of persistent ETD.

ACCLARENT AERA® is the only balloon that was specifically designed to treat persistent ETD in the United States

Safe by design, the bulb tip enhances safety by limiting balloon catheter travel to the isthmus.



- 1 Designed specifically for the Eustachian tube anatomy
- 2 Offers the flexibility to adapt to the s-shaped curve of the Eustachian tube
- 3 Preserves natural anatomy with minimally invasive, transnasal access

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Proven safety and efficacy

ACCLARENT AERA® is backed by the only prospective, multicenter, randomized clinical trial to demonstrate:³

99.7%

technical success in dilating the Eustachian tube

51.8% vs 13.9%

higher rate of tympanogram normalization compared to control subjects treated with medical management alone

56.1% vs 8.5%

improvement in the quality of life measure from the Eustachian Tube Dysfunction Questionnaire (ETDQ-7)

We're invested in you

Acclarent offers you the training and support—including guidance on patient selection—to help you successfully treat persistent Eustachian Tube Dysfunction (ETD). Medical management alone cannot always address the source of persistent ETD. With ACCLARENT AERA®, the solution is now in your hands.

Talk to your Acclarent representative to learn more about this new tool in the management of persistent ETD.

ACCLARENT AERA® Eustachian Tube Balloon Dilation System



ACCLARENT AERA® Balloon Catheter



ACCLARENT AERA® Guide Catheter

Description | ACCLARENT AERA® Eustachian Tube Balloon Dilation System

Catalog number | EU061655

References:

1. Llewellyn, A., Norman, G., Harden, M., Coatesworth, A., Kimberling, D., Schilder, A. and McDaid, C. (2014). Interventions for adult Eustachian tube dysfunction: a systematic review. HEALTH TECHNOLOGY ASSESSMENT, 18; 1-180. 2. Ockermann, T., Reineke, U., Upile, T., Ebmeyer, F., Sudhoff, H.H. (2010). Balloon Dilatation Eustachian Tuboplasty: A Clinical Study. Laryngoscope, 120:1411–1416. 3. Poe D, Anand V, Dean M, et al. (2017). Balloon dilation of the Eustachian tube for dilatory dysfunction: A randomized controlled trial. Laryngoscope. Sep 20. doi: 10.1002/lary.26827. Laryngoscope. Sep 20. doi: 10.1002/lary.26827.2010;120(7):1411-1416.

ACCLARENT AERA® Eustachian Tube Balloon Dilation System is intended for use by physicians who are trained on Acclarent Technology. Eustachian tube balloon dilation has associated risks, including tissue and mucosal trauma, infection, or possible carotid artery injury. Prior to use, it is important to read the Instructions for Use and to understand the contraindications, warnings, and precautions associated with these devices. The safety of the device as used under local anesthesia has not been evaluated.

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