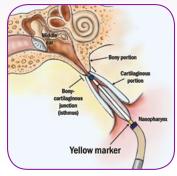
ACCLARENT AERATM EUSTACHIAN TUBE BALLOON

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



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

Designed to relieve symptoms of persistent ETD

ETD is a condition caused by the inadequate opening 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 between 0.9% and 5.0%.^{2,3}

ACCLARENT AERA™ is the only balloon indicated for persistent ETD in the United States





Proven safety and efficacy

ACCLARENT AERA™ is backed by the only prospective, multicenter, randomized clinical trial to demonstrate¹:

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



References:

1. ACCLARENT AERA™ Eustachian Tube Balloon Dilation System Instructions for Use. Irvine, CA: Acclarent, Inc; 2016. **2.** Browning GG, Gatehouse S. The prevalence of middle ear disease in the adult British population. *Clin. Otolaryngol Allied Sci.* 1992;17(4):317-321. **3.** Ockermann T, Reineke U, Upile T, et al. Balloon dilatation eustachian tuboplasty: a clinical study. *Laryngoscope.* 2010;120(7):1411-1416.

Caution: Federal (US) law restricts the sale, distribution or use of the ACCLARENT AERATM by or on the order of a physician who is trained in the use of 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.

33 Technology Drive, Irvine, CA 92618 Customer Support: 877.775.2789 www.acclarent.com

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