

# FILTERLEX - Full-Body Embolic Protection Device for TAVI

## AT A GLANCE

**IP:** PCT submitted in Sep. 2016, received exceptionally positive search report and written opinion

**Status:** Pre-clinical

## UNIQUE FEATURES

- ▮ Provides full-body embolic protection
- ▮ No additional arterial access required
- ▮ Performs deflection, capture and removal of embolic particles
- ▮ Easy and intuitive deployment and retrieval
- ▮ Protects aorta surface
- ▮ Stable and safe device anchoring
- ▮ No interference to procedure devices
- ▮ Applicable for a variety of procedures

## THE MARKET

The worldwide TAVI market is estimated to surpass \$5.5B by 2020. Additional major markets include TMVR (estimated at five times of the TAVI market) and AF Ablation (1.2M procedures annually). The total addressable market for Filterlex is valued at over \$5B.

Filterlex Medical Ltd. was founded in June 2015 to develop a novel, full-body embolic protection device to reduce the risk of stroke and related embolic complications during left-heart interventional procedures.

## THE NEED

During catheter-based, left-heart procedures such as TAVI, embolic particles are often released to the blood flow. Particles migration to the brain may cause a spectrum of neurological deficiencies, from cognitive impairment to debilitating stroke. Emboli released to distal organs may result in acute kidney injury and ischemia.

## THE PRODUCT

The Filterlex device is a next-generation full-body embolic protection device, easily and intuitively deployed and retrieved. The device is securely positioned in the aorta, protects its surface while facilitating a seamless TAVI procedure. Its distinctive, triple action design provides a full-body embolic protection by deflecting, capturing and removing embolic particles. Uniquely, it requires no additional arterial access and does not interfere to the procedure workflow.

## THE COMPETITION

**Claret Medical** – Sentinel, overall funding estimate - \$41M.

FDA clearance 510k De-novo in June 2017, CE since Jan. 2014.

**Keystone Heart** – TriGuard, overall funding estimate - \$25M.

FDA, 510K, undergoing IDE trial, CE since Sept. 2013.

**ICS** – Emblok, early clinical-stage.

**Protembis** – ProtEmbo, early clinical-stage.

## THE TEAM

**Ms. Sigal Eli - CEO:** Entrepreneur, inventor and interventional cardiology imaging expert with extensive experience in TAVI procedures, a former clinical and training specialist at a medical imaging company.

**Mr. Eyal Teichman - CTO:** Entrepreneur, mechanical engineer and CTO with over 25 years of R&D experience with extensive experience in implantable devices in the cardiovascular field.

**Giora Weisz, M.D. - Chairman of Advisory Board:** Director of Interventional Cardiology at Montefiore Medical Center, and faculty of the Cardiovascular Research Foundation in New York, NY.

**Mr. Ron Davidson – Active Board Member:** Seasoned executive with 20 years' experience in managing and accelerating companies in the healthcare field, led two companies to successful exits.

**Dr. Shimon Eckhouse – Chairman:** Founder and investor of numerous medical technology firms. One of Israel's leading medical entrepreneur and the owner and Active Chairman of Alon MedTech Ventures incubator.