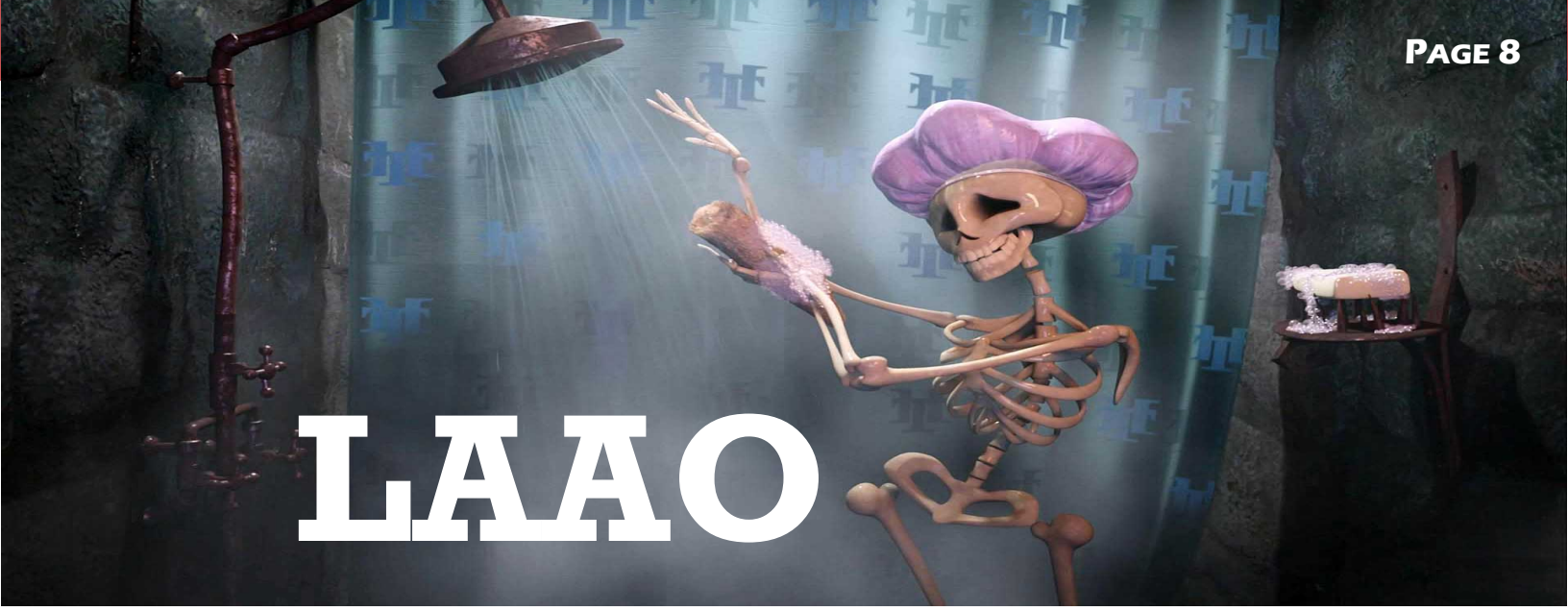

LAAO

Dashboard NEWS…..

New LAAO Device Added

The Abbott's Amplatzer™ Amulet™ Left Atrial Appendage Occluder (LAAO) Device has received U.S. Food and Drug Administration (FDA) approval and is now available for selection in the LAAO Registry device list under Sequence # 7275. The device may be selected for any procedures in which an Amulet Device was implanted on or after 8/14/2021. Procedures that are part of the Catalyst Trial will not be included in the LAAO Registry.

LAAO Registry Added Metrics

The LAAO Registry is pleased to share new metrics

* In-Hospital Informational Data Metrics
    * Patients' average HAS-BLED score [Metric 24]
    * Patients' average CHA2DS2-VASc Score [Metric 25]
* 45-Day Follow-up:
    * Metric 216: Proportion of WATCHMAN procedures where a 45 day follow- up diagnostic imaging was performed.

New Device details

* The Amulet received FDA approval 8/14/2021
* Device is available for selection on the Device list

The Amulet

* If a patient is coded to have received an Amulet device after August 14, 2021, the patient may be entered into the LAAO Registry if not enrolled in the CATALYST study.
    * IMPORTANT!!
        * Patients receiving a Amulet participating in the CATALYST trial will **NOT** be included in the registry (but stay tuned!)