

Best Practice Protocol

# Nephropathy Post-TAVR

Nephropathy after TAVR is associated with increased mortality in MISHC data and in peerreviewed literature.

Predictors of nephropathy post-TAVR in MISHC data include higher STS risk, diabetes, lower body weight, coronary artery disease, higher baseline serum creatinine, and anemia. Procedural characteristics strongly associated with acute kidney injury (AKI) include use of general anesthesia and use of contrast volume greater than 3 times the eGFR.

In addition to avoiding higher doses of iodinated contrast media during the procedure, adequate hydration has been demonstrated to decease the incidence of nephropathy post procedure.

One protocol that has been utilized to manage hydration is the POSIDEON study protocol that is based on end-diastolic pressure measured during the catheterization procedure.

### Definitions of AKI

## VARC2

Creatinine increase to at least any stage 1 of the AKIN classification system in accordance with VARC-2 definitions. Increase in serum creatinine to 150–199% (1.5–1.99 × increase compared with baseline).

#### BMC2

Serum Creatinine increase > 0.5mg/dL over baseline.

## Hydration protocol for TAVR

## **Pre-TAVR**

Normal saline infusion: 3mL/kg x 1 hour

LVEDP-based post-TAVR hydration LVEDP at conclusion of TAVR	Rate of Normal saline infusion X 4 hours post TAVR
< 13	5mL/kg/hr
13-18	3 mL/kg/hr
18-25	1.5 mL/kg/hr
> 25	No additional hydration/implant team to decide

#### **Contrast limits for TAVR procedure**

- eGFR calculated for all TAVR patients before the procedure
- Goal for every TAVR is to limit the dose of contrast
- 2X and 3X the eGFR "contrast limits" are communicated to the implant team in the pre-TAVR time-out
  - When relevant, methods to limit the volume of contrast utilized are discussed prior to the TAVR procedure
  - Efforts are made to limit the volume of contrast delivered to patients during the TAVR procedure, with > 3X eGFR the maximum

## ACKNOWLEDGEMENTS

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#### DISCLAIMER

Michigan TAVR Best Practice Protocols are based on consortium-wide consensus at the time of publication. Protocols will be updated regularly, and should not be considered formal guidance, and do not replace the professional opinion of the treating physician.

#### REFERENCES

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