

National Cardiogenic Shock Initiative

Case Report Form

(Version 2.0)

Please complete the entirety of the worksheet. Upon completion, please email this worksheet [SECURE] to: NationalCSI@hfhs.org. Please email/call if there are any questions or concerns.

Demographics

Date of Impella Insertion _____

Implanting Physician _____

Hospital: Name _____

Hospital: City, State _____

Age of Patient _____

Gender (please circle) Male Female

Race (please circle) White Black Hispanic Other

Medical History

| | | | |
|---|-----|----|-----|
| Does the patient have a history of Diabetes? | Yes | No | N/A |
| Does the patient have a history of TIA/CVA? | Yes | No | N/A |
| Does the patient have a history of ESRD? | Yes | No | N/A |
| Does the patient have a history of CKD? | Yes | No | N/A |
| Does the patient have a known LVEF <50%? | Yes | No | N/A |
| Has the patient had a prior CABG? | Yes | No | N/A |
| Has the patient had a prior PCI? | Yes | No | N/A |
| Has the patient had a prior Myocardial Infarction? | Yes | No | N/A |

FOR HFH USE ONLY

NCSI #: _____

Admission Characteristics

| | | | |
|---|------|--------------|-----|
| Was the patient transferred from another hospital? | Yes | No | N/A |
| If yes, was the patient on support prior to transport? | Yes | No | N/A |
| What support device was used? | IABP | Other: _____ | |
| | | | |
| Was cardiogenic shock present on admission to your institution? | Yes | No | N/A |
| | | | |
| Did the patient experience any of the following (<i>prior</i> to arrival in the Cath Lab): | | | |
| Anoxic Brain Injury? | Yes | No | N/A |
| Cardiac Arrest (In Hospital)? | Yes | No | N/A |
| Cardiac Arrest (Out of Hospital)? | Yes | No | N/A |
| | | | |
| Did the patient require CPR prior to Impella implant? | Yes | No | N/A |
| | | | |
| Was the patient undergoing active CPR at the time of Impella implantation? | Yes | No | N/A |
| | | | |
| Was the patient treated with medically-induced hypothermia? | Yes | No | N/A |

**** Important Timings **** Please estimate if exact timings are unavailable.

Please do not leave blank.

Arrival to Hospital (date and time) Date _____ Time _____

Onset of AMI (date and time) Date _____ Time _____

Onset of Shock (date and time) Date _____ Time _____

Time of Impella Insertion (date and time) Date _____ Time _____

Using the above timings, please calculate the following times:

Door to Support Time (minutes): _____

Door to Balloon Time (minutes): _____

| Procedural Characteristics (please circle the best choice, if answer is not known please write "N/A") | | | | | | | | |
|--|---|--|---|---|--|---------------------|--|--|
| Impella Placement: | | 1. Prior to PCI | | 2. Post PCI | | 3. Intra-procedural | | |
| RHC Placement: | | 1. Prior to Impella | | 2. Post Impella | | 3. No RHC obtained | | |
| Impella Used: 2.5 CP 5.0 RP Other: _____ | | Impella Access: <input type="checkbox"/> Femoral <input type="checkbox"/> Axillary <input type="checkbox"/> Other: _____ | | AMI Type? STEMI NSTEMI | | | | |
| <u>PCI Attempted?</u> YES NO | TIMI FLOW <u>Pre-PCI</u> <u>Post-PCI</u> 0 0 | | Evidence of Thrombus Pre-PCI? YES NO | # of Diseased Vessels? 0 1 2 3 | # Vessels Treated? 0 1 2 3 | # of Stents? | Culprit Lesion(s) Location(s) 1. LM 5. Ramus 2.LAD 6. SVG 3. LCx 7. LIMA 4. RCA 8. RIMA | Other Lesion(s) Location(s) 1. LM 5. Ramus 2.LAD 6. SVG 3. LCx 7. LIMA 4. RCA 8. RIMA |
| <u>Successful?</u> YES NO | 1 1 2 2 3 3 | | | | | | | |
| <u>Access for PCI?</u> 1. Radial 2. Femoral | Was complete revascularization performed? YES NO | Thrombectomy used? YES NO | Atherectomy used? YES NO | PCI Complications? <input type="checkbox"/> NO <input type="checkbox"/> YES <u>If YES:</u> <input type="checkbox"/> Stent Thrombosis <input type="checkbox"/> Evidence of Residual Thrombus <input type="checkbox"/> OTHER: _____ | | | | |
| Was patient taking antiplatelet medication at home? <input type="checkbox"/> No <input type="checkbox"/> Aspirin <input type="checkbox"/> Clopidogrel <input type="checkbox"/> Ticagrelor <input type="checkbox"/> Prasugrel <input type="checkbox"/> N/A <input type="checkbox"/> OTHER: _____ | | | | | | | | |
| Was patient loaded with antiplatelet prior to PCI: <input type="checkbox"/> No <input type="checkbox"/> Aspirin <input type="checkbox"/> Clopidogrel <input type="checkbox"/> Ticagrelor <input type="checkbox"/> Prasugrel <input type="checkbox"/> Cangrelor <input type="checkbox"/> Abciximab <input type="checkbox"/> Eftifibatide <input type="checkbox"/> Tirofiban <input type="checkbox"/> N/A <input type="checkbox"/> OTHER: _____ | | | | | | | | |
| If oral antiplatelets were given, what route were they given? <input type="checkbox"/> By Mouth <input type="checkbox"/> NG/OG Tube <input type="checkbox"/> Rectal <input type="checkbox"/> N/A <input type="checkbox"/> Other: _____ | | | | | | | | |
| Was the oral antiplatelet crushed? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A | | | | | | | | |
| When was antiplatelet administered? <input type="checkbox"/> EMS <input type="checkbox"/> ER <input type="checkbox"/> Pre-PCI <input type="checkbox"/> Post-PCI <input type="checkbox"/> N/A <input type="checkbox"/> Other: _____ | | | | | | | | |
| Anticoagulation used during PCI: <input type="checkbox"/> Heparin <input type="checkbox"/> Bivalirudin <input type="checkbox"/> Other: _____ | | | | | | | | |

Please give a brief description of the patient admission: _____

HEMODYNAMIC & LABORATORY VALUES (Pre-procedure & Prior to starting Vasoactive Medications)

These values should represent the “worst hemodynamics” that demonstrate level of shock

| | | | |
|----|-----|-----|-----|
| HR | SBP | DBP | MAP |
|----|-----|-----|-----|

HEMODYNAMIC & LABORATORY VALUES: (Pre-Impella)

These values represent the hemodynamics prior to Impella Insertion, at the beginning of the PCI procedure

| | | | | | | | | |
|---------|----------------------|---|----------------|---|-------------|-------------|------------|-----------|
| HR | SBP | DBP | MAP | Troponin | Cr | AST | Hgb | Lactate |
| RA/CVP | RV | PA | PCWP | CO <input type="checkbox"/> Fick <input type="checkbox"/> TD <input type="checkbox"/> CCO | CI | CPO | PAPI | LVEDP |
| PA Sat. | Admission Glucose | <u>VASOACTIVE AGENTS:</u> <u>(DOSE):</u> | Norepinephrine | Dopamine | Epinephrine | Vasopressin | Dobutamine | Milrinone |

HEMODYNAMIC & LABORATORY VALUES: Post-PCI, in the Cath Lab (with Impella running)

These values represent the Cath Lab hemodynamics post PCI, at the end of the procedure

| | | | | | | | | |
|---------|---|----------------|----------|---|-------------|------------|-----------|--------|
| HR | SBP | DBP | MAP | | | | | |
| RA/CVP | RV | PA | PCWP | CO <input type="checkbox"/> Fick <input type="checkbox"/> TD <input type="checkbox"/> CCO | CI | CPO | PAPI | LVEDP |
| PA Sat. | <u>VASOACTIVE AGENTS:</u> <u>(DOSE):</u> | Norepinephrine | Dopamine | Epinephrine | Vasopressin | Dobutamine | Milrinone | Other: |

| HEMODYNAMIC & LABORATORY VALUES: 12 hours Post-Impella Implant (ONLY if Impella is running) | | | | | | | | |
|---|---|----------------|----------|---|-------------|------------|-----------|---------|
| HR | SBP | DBP | MAP | Troponin | Cr | AST | Hgb | Lactate |
| RA/CVP | RV | PA | PCWP | CO <input type="checkbox"/> Fick <input type="checkbox"/> TD <input type="checkbox"/> CCO | CI | CPO | PAPI | LVEDP |
| PA Sat. | <u>VASOACTIVE AGENTS:</u> <u>(DOSE):</u> | Norepinephrine | Dopamine | Epinephrine | Vasopressin | Dobutamine | Milrinone | Other: |

| HEMODYNAMIC & LABORATORY VALUES: 24 hours Post-Impella Implant (ONLY if Impella is running) | | | | | | | | |
|---|---|----------------|----------|---|-------------|------------|-----------|---------|
| HR | SBP | DBP | MAP | Troponin | Cr | AST | Hgb | Lactate |
| RA/CVP | RV | PA | PCWP | CO <input type="checkbox"/> Fick <input type="checkbox"/> TD <input type="checkbox"/> CCO | CI | CPO | PAPI | LVEDP |
| PA Sat. | <u>VASOACTIVE AGENTS:</u> <u>(DOSE):</u> | Norepinephrine | Dopamine | Epinephrine | Vasopressin | Dobutamine | Milrinone | Other: |

POST PROCEDURAL FOLLOW UP

Time & Date of Impella Explant: Time: _____ Date: _____

WEANING OF HEMODYNAMIC SUPPORT:

Support was weaned according to: PA Sat Echo RHC Hemodynamics
 Other: _____

Total hospital duration (Days): _____

Left Ventricle Ejection Fraction (Pre-Impella): _____ (Prior to discharge): _____

DID THE PATIENT SURVIVE THE INDEX PROCEDURE? Yes No N/A

Was the patient transferred to VAD/Transplant Center Yes No N/A
If "Yes", DID THE PATIENT SURVIVE TO TRANSFER? Yes No N/A

Did the patient have any additional support devices implanted post-index procedure? Yes No N/A
If "Yes", which device was used: _____

DID THE PATIENT SURVIVE TO DISCHARGE? Yes No N/A

Was the patient discharged to hospice? Yes No N/A

Did the patient experience any significant Impella-related complications?
 NO Vascular Complications Hemolysis
 OTHER – Please explain: _____

Did the patient require any blood transfusions? No Yes – # of transfusions: _____

Was any external form of vascular bypass performed to provide lower extremity perfusion during Impella (i.e. antegrade access, "up and over" perfusion catheter, etc.)?
 No Yes – Please Explain: _____

Please provide a brief description of the patient's hospital course (including significant complications and discharge circumstances):
Please Explain: _____

If the patient did not survive, please indicate the major cause of death (fatal bleeding, anoxic brain injury, worsening cardiogenic shock, patient/family wishes?)
Please Explain: _____

Cardiac Medications on Discharge (name only): _____

- END OF FORM -