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Follow Up Assessment after Shock Onset – (For serial entry, use the <u>time period from the last assessment</u>) *For the measurement elements below, enter accurate parameters <u>closest to assessment time</u>			
Date/Time of assessment:	/_/ (MM/DD/YYYY HH:MM)	O Unknown	
SCAI Stage:	Shock has resolvedStage BStage C	Stage DStage END/Unable to Determine	
Presence of a Pulmonary Artery Catheter (PAC)	O Yes	O No	
If Yes, Date/Time of the <u>first</u> PAC:	// (MM/DD/YYYY)	O Unknown	
Presence of Mechanical Ventilation	O Yes	O No	
Presence of renal replacement therapy	O Yes	O No	
*BP (Systolic/Diastolic):	/mmHg	O Not Documented	
*Heart Rate:	bpm	O Not Documented	
*CVP/RA:	(mmHg)	 Not Documented 	
*PA Pressure (Systolic/Diastolic):	/ mmHg	O Not Documented	
*PCWP:	(mmHg)	O Not Documented	
*Cardiac Output:	(L/min)	O Not Documented	
MAP (Auto-calculated)	(mmHg)	O Not Documented	
PAPi (<i>Auto-calculated</i>)	(W)	 Not Documented 	
CPO (Auto-calculated)		O Not Documented	
Peak Lactate since the last assessment	(mmol/L)	O Not Documented	
Lowest pH since the last assessment		 Not Documented 	
Peak ALT since the last assessment	(IU/L)	O Not Documented	
Vasoactive Medications at time of assessment (check all that apply)	 □ None □ Dobutamine □ Dopamine □ Epinephrine □ Levosimendan □ Milrinone 	 □ Nitroprusside □ Norepinephrine □ Phenylephrine □ Vasopressin □ Not Documented □ Other (Specify): 	
Presence of MCS Device(s) at assessment (If MCS is present, select all that apply)	□ None □ Impella 2.5 □ Impella CP □ Impella ECP □ Impella 5.0 □ Impella 5.5 □ Impella RP □ ECMO (VA) □ IABP	iVAC □ TandemHeart □ Left □ Right □ Temporary surgical VAD (e.g. CentriMag) □ Left □ Right □ Right □ Other (Specify):	
Was there a device upgrade/escalation since the prior assessment?	O Yes	O No	
If Yes, select reason(s) for device upgrade or escalation since the prior assessment Was there a device de-escalation	 □ None/Not Documented □ Device-related complication or failure □ Inadequate response to vasoactive medications □ Need for escalation to greater hemodynamic support from MCS □ Switch to alternative MCS access site (e.g. fem to axillary) □ Other (Specify): 		
since the prior assessment?	O Yes	O No	

CSRC Cardiogenic Shock Lean CRF: V4 – Follow-up CRF Form

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If Yes, select reason(s) for device downgrade or de-escalation since the prior assessment	 Change in goals of care Durable LVAD or heart transplant MCS no longer needed Transition to central cannulated device (e.g. CentriMag) 		
Vascular complication requiring intervention:	O Yes O No		
If Yes, enter Date:	/ / (MM/DD/YYYY)	O Not Documented	
ii res, enter Date.		O Not Documented	
Other complications of ECMO	 Pulmonary hemorrhage requiring intervention Refractory pulmonary edema Other (Specify): 		