HA	HALF-PINT Form 5: Daily ICU				
Sec	Section: Daily Assessment				
1.	Ventilatory support this study day (check all that apply)	<ul> <li>Intubated and mechanically ventilated</li> <li>Tracheostomy and mechanically ventilated</li> <li>Non-invasive ventilation [including BiPAP, Humidified High Flow Nasal Cannula (HHFNC ≥ 5 L/min of Oxygen flow), or CPAP ≥ 5 cm H<sub>2</sub>O]</li> <li>Non-invasive ventilation &lt; 5 cm H<sub>2</sub>O</li> <li>None</li> </ul>			
2.	Devices this study day	Arterial line:     Yes    No Central venous line (CVL):     Yes    No Peripheral IV (PIV):     Yes    No Peripherally inserted central catheter (PICC):     Yes    No Enteral feeding tube (new or pre-existing):     Yes    No Bladder catheter:     Yes    No Chest tube:     Yes    No Surgical wound drain:     Yes    No			
3.	CPR this study day	○ Yes ○ No			
4.	ECMO/VAD this study day	○ Yes ○ No			
Sec	Section: Medications				
5.	Inotropes/vasopressors this study day	<pre>Types If Yes, which inotropes/vasopressors?  Dopamine &gt; 5 mcg/kg/min:     Yes    No Dobutamine &gt; 5 mcg/kg/min:     Yes    No</pre> Ores    No			

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Note: Record the most abnormal (lowest and/or highest) value documented this study day (00:00 to 23:59).		
	tion: Daily PELOD Score	
12.	Total caloric intake this study day	kcal
		Lipid: kcal
		Protein: kcal
		If Yes, enter caloric intake: CHO: kcal
11.	Was parenteral nutrition given this study day?	○ Yes
		Lipid: kcal
		Protein: kcal
		If Yes, enter caloric intake: CHO: kcal
10.	Was enteral nutrition given this study day?	O Yes
Sec	tion: Nutrition	
9.	Diuretics this study day	○ Yes ○ No
8.	Empiric/treatment antibiotics this study day	○ Yes ○ No
7.	Steroids this study day	○ Yes ○ No
6.	Paralytic drip this study day	○ Yes ○ No
		Epinephrine (any):  Yes No Norepinephrine (any): Yes No Milrinone (any): Yes No Vasopressin for hypotension: Yes No No inotropes/vasopressors

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13.	LOWEST Glasgow Coma Score (GCS)  Note: All 3 elements must be collected at the same time.  Only assess subjects with known or suspected acute central nervous system disease/process.	Subject had known or suspected acute central nervous system disease/process this study day:  Yes - If Yes, enter GCS below  No  Eye opening:  (4) Spontaneous  (3) To speech  (2) To pain  (1) None  Not done  Best verbal response (use best estimate for intubated subjects):  (5) Oriented/smiles, fixes and follows  (4) Confused conversation/irritable cries
		(3) Inappropriate words/cries to pain (2) Incomprehensible sounds/moans to pain (1) None Not done
		Best motor response:  (6) Obeys commands, normal spontaneous movement  (5) Localizes/withdraws to touch  (4) Withdraws to pain  (3) Decorticate-abnormal flexion  (2) Decerebrate-abnormal extension  (1) None  Not done
14.	Worst pupillary response to bright light  Note: Do not assess after iatrogenic pupillary dilatation.	<ul><li>Both fixed and &gt; 3 mm</li><li>One fixed and &gt; 3 mm</li><li>Both responsive</li><li>Unknown</li></ul>
15.	Heart rate  Note: Do not assess during crying or iatrogenic agitation.	LOWEST HIGHEST One Measurement Only Not Done bpm bpm
16.	Systolic blood pressure  Note: Do not assess during crying or iatrogenic agitation.	LOWEST HIGHEST One Measurement Only Not Done mmHg mmHg
17.	Creatinine	LOWEST HIGHEST One Draw Only Not Done mg/dL mg/dL
18.	PaO <sub>2</sub> /FiO <sub>2</sub>	LOWEST PF ratio HIGHEST PF ratio

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19.	Note: For PaO <sub>2</sub> , use arterial measurement only. PaO <sub>2</sub> /FiO <sub>2</sub> cannot be assessed in subjects with intracardiac shunts and is considered as normal in children with cyanotic heart disease. PaCO <sub>2</sub> Note: PaCO <sub>2</sub> may be measured from arterial,	PaO <sub>2</sub> PaO <sub>2</sub> One Draw Only Not Done  mmHg FiO <sub>2</sub> FiO <sub>2</sub> LOWEST HIGHEST One Draw Only Not Done  mmHg mmHg
20.	Capillary, or venous samples.  White blood cell count	LOWEST HIGHEST One Draw Only Not Done  K/µL K/µL
21.	Platelet count	LOWEST HIGHEST One Draw Only Not Done K/μL K/μL
22.	AST (SGOT)	LOWEST HIGHEST One Draw Only Not Done  IU/L IU/L
23.	Prothrombin time (PT)	LOWEST HIGHEST One Draw Only Not Done sec
24.	INR	LOWEST HIGHEST One Draw Only Not Done
Sec	tion: Adverse Events	
25.		Hypoglycemia <60 mg/dL Hyperglycemia >250 mg/dL (>6 hours after initiation of insulin infusion) Hypokalemia <2.5 mmol/L New seizure (in subject without a known seizure disorder) Catheter-associated bloodstream infection (CA-BSI) Catheter-associated urinary tract infection (CA-UTI) Surgical site infection (SSI) Ventilator-associated pneumonia (VAP) Other hospital-acquired infection Continuous Glucose Monitor (CGM) related adverse event, bleeding or other (that did not involve hypoglycemia, hyperglycemia, or infection) Bedside glucose meter related adverse event (that did not involve hypoglycemia, hyperglycemia, or infection) VAMP related adverse event Computerized insulin dosing protocol related adverse event (that did not result in a hypoglycemia or hyperglycemia event) Insulin dosing error (that did not result in a hypoglycemia or hyperglycemia event)

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		Other, specify:  None
Section: Study Discharge		
26.	If the subject was on non-invasive ventilation or ventilation via tracheostomy overnight or during sleep prior to this illness, has the subject returned to that baseline level of support?	<ul><li>Yes</li><li>No</li><li>Not on non-invasive ventilation or ventilation via tracheostomy prior to this illness</li></ul>
27.	Was the subject discharged from the HALF-PINT study this study day for reasons related to the primary outcome?  If study discharged, end daily data collection here and continue to Form 6.	Yes If Yes, confirm that all of the following are true for at least 24 consecutive hours: Extubated Off non-invasive ventilation that provides ≥ 5 cm H <sub>2</sub> O or reached ventilatory settings used prior to the illness that led to this ICU admission Not on intravenous vasopressors or inotropes (i.e., dopamine or dobutamine > 5 mcg/kg/min, or any dose of epinephrine, norepinephrine, milrinone, or vasopressin if used to treat hypotension) No
28.	Was the subject discharged from the HALF-PINT study this study day for any other reason?  If study discharged, end daily data collection here and continue to Form 6.	If Yes, other reason for study discharge:     Family/team have decided to limit/redirect from aggressive ICU technological support     Withdrawal of consent     Hospital discharge     Study Day 28     Subject had new cardiac surgery     Other, specify:

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