SCHEDULE OF ASSESSMENTS

Trial Activity	Screening	Baseline	6 hours (+/- 1 hour)	12 hours (+/- 4 hours)	24 hours (+/-6 hours)	48 hours (+/-12 hours)	72 hours (+/-12 hours)	+/- 7 days	Hospital Discharge	30 days	90 days
Eligibility – Inclusion/Exclusion*	X	X						X			
Pregnancy test *		X									
Written Informed Consent	X										
Demographics/Medical History/estimated weight/ Frailty score		X									
Vital signs		X	X	X	X	X	X				
Blood results (routine) incl lactate		X	X**	X**	X**	X**	X**				
IMP administration		X	X	X	X	X					
IMP adherence		X			X	X					
Total intravenous fluid volume delivered		X	X	X	X	X	X				
Total dose of norepinephrine delivered		X	X	X	X	X	X				
Total dose of other vasopressors delivered		X	X	X	X	X	X				
Safety outcomes (pulmonary oedema/extravasation)							X		X		
Mortality/interventions/length of stay/readmissions									X	X	X

Adverse Events ***		X	X	X	X	X	X**	X			
EQ-5D-5L ****		X								X	X

**All women of childbearing potential must have a negative urine or serum pregnancy test completed as part of study eligibility checks.*

*** Daily (+/- 12 hours) for any routine bloods collected up to 72 hours. If bloods (or individual parameters) are not requested by the clinical team, this will not be recorded as a deviation.*

**** Adverse Event reporting is a continuous process*

***** EQ-5D-5L should only be completed if the patient has capacity.*