

Shock Review & Vasopressors

Type of Shock	Common Examples	Right heart filling	Left heart filling	Cardiac output	Afterload	Treatment
		CVP	PCWP	CI	SVR	
Hypovolemic	<ul style="list-style-type: none"> Hemorrhage Intravascular volume loss 	↓	↓	↓	↑ ↑	Fluids +/- vasopressors to temporize
Cardiogenic	<ul style="list-style-type: none"> Cardiomyopathy MI/ischemia Valvular lesions Arrhythmia 	↑	↑	↓	↑ ↑	inotropes +/- vasopressors
Distributive	<ul style="list-style-type: none"> Sepsis Anaphylaxis Adrenal insuff. Neurogenic shock 	↓	↓	↓ vs. ↑*	↓	Fluids + vasopressors (rarely inotropes)
Obstructive	<ul style="list-style-type: none"> PE** Cardiac tamponade Tension PTX 	↑	PCWP ↑ ↓	↓	↑ ↑	Fluids + vasopressors (occasionally inotropes for R heart failure)

*distributive shock often presents with low cardiac output at the onset, but may become a high cardiac output state once adequately fluid resuscitated

**PE's is the most common example of an obstructive shock and is used to fill out the hemodynamic table.

Shock definition: insufficient blood flow to the tissues

- First: try and identify the type of shock
- Second: tailor your treatment to the PRIMARY PROBLEM (black box) of that particular shock state
 - hypovolemic shock** = treat with **fluids**
 - Crystalloids** (Ringer's Lactate is preferred)
 - SMART-MED and SMART-SURG 2018 → Ringer's Lactate reduced rate of death, need for dialysis, lower rates of AKI when compared to Normal Saline
 - Colloids**
 - Albumin 5%** 500 mL → results in an increase in intravascular volume of ~500 mL x 12-24h (1:1) → more immediate effect
 - Albumin 25%** 100 mL → results in an increase in intravascular volume of ~450 mL x 12-24h (1:4.5) → slower effect, use if fluid overloaded but intravascularly dry
 - Blood → typically reserved for anemia; transfusion trigger of 70 g/L in a stable, non-bleeding patient
 - AVOID starches → increased risk of AKI + need for dialysis
 - cardiogenic shock** = **increase cardiac output** with inotropes (may need vasopressors to **support BP**)
 - inotropes**

- dobutamine 0-20 ug/kg/min
 - milrinone 0.25-0.75 ug/kg/min → slow onset, 6hr duration, more likely to cause hypotension, use with caution in renal failure
 - epinephrine 0-10 **ug/min**
 - digoxin load
- **distributive shock = hypovolemia** due to vasodilation + vasopressors to counter **vasodilation**
 - if using CVP, aim for 8-12 mm Hg if not intubated, 12-15 mm Hg if intubated
 - remember that vasopressors also raise CVP, so if the patient is in their “CVP target” but on vasopressors, they may still benefit from more fluid
 - obstructive shock (**PE**) = fluids (CVP is high, but LV is under filled), vasopressors to support BP, inotropes if RV is failing, consider **thrombolysis** if:
 - hemodynamically UNSTABLE
 - stable patients IF:
 - severe or worsening RV dysfunction
 - cardiac arrest due to PE
 - extensive clot burden
 - free-floating RA or RV thrombus

Vasopressors

Drug	Dose	Notes
Norepinephrine (Levophed)	0-30 ug/kg/min	<ul style="list-style-type: none"> • often 1st line vasopressor • can go up to 1 ug/kg/min in refractory shock
Vasopressin	0-2.4 u/hr	<ul style="list-style-type: none"> • often added as a 2nd vasopressor if on high doses of Norepinephrine • still effective when acidotic and hypothermic whereas other vasopressors may not be
Phenylephrine	0-360 ug/min	<ul style="list-style-type: none"> • useful when you want to avoid b-agonism, e.g. rapid a-fib/flutter and hypotensive • not as potent → can be a significant volume load when running at high doses • watch for hyponatremia when mixed in D5W and running at high doses
Dopamine	0-20 ug/kg/min	<ul style="list-style-type: none"> • 0-3 ug/kg/min = “renal dose”, doesn’t prevent AKI • 3-10 ug/kg/min = primarily b-agonism (inotropy) • > 10 ug/kg/min = progressive alpha-effect (vasoconstriction)
Epinephrine	0-10 ug/min	<ul style="list-style-type: none"> • Drug of choice for anaphylaxis • Watch for tachycardia/arrhythmias