## **Dobutamine**

			<u>Dobutamine</u>													
Form	250mg per 20ml vial															
Reconstitution	Already in solution															
	Dilute further before administration															
Compatibility	Sodium Chloride	0.9%														
& Stability	Glucose 5%															
_	Hartmann's Solution															
Administration	Continuous IV infusion															
	Peripheral administration via a large vein:															
	Recommended concentration: 1mg/ml.															
	If clinically indicated a concentration of 2mg/ml may be used.															
	To prepare a <b>1mg/ml</b> solution: Withdraw 40mls from a 500ml infusion bag of compatible fluid.															
	Add 500mg (40mls) Dobutamine to the remaining 460ml.															
	Infusion rate in ml per hour (ml/hr) is:															
	Dose Weight (Kg)															
	microgram/															
	kg/min															
		45	50	55	60	65	70	75	80	85	90	95	100			
	2.5	6.8	7.5	8.3	9	9.8	10.5	11.3	12.0	12.8	13.5	14.3	15.0			
	5.0	13.5	15.0	16.5	18.0	19.5	21.0	22.5	24.0	25.5	27.0	28.5	30.0			
	7.5 10	20.3	22.5 30.0	24.8 33.0	27.0 36.0	29.3 39.0	31.5 42.0	33.8 45.0	36.0 48.0	38.3 51.0	40.5 54.0	42.8 57.0	45.0 60.0			
	LU															
	To prepare a <b>2mg/ml</b> solution: Withdraw 80mls from a 500ml infusion bag of compatible.															
	Add 1000mg (80mls) Dobutamine to the remaining 420ml.															
	Infusion rate in ml per hour <b>(ml/hr)</b> is:															
	Dose Weight (Kg)															
	microgram/															
	kg/min	45	50	55	60	65	70	75	80	85	90	95	100	1		
	2.5	3.4	3.8	4.1	4.5	4.9	5.3	5.6	6.0	6.4	6.8	7.1	7.5	-		
	5.0	6.8	7.5	8.3	9.0	9.8	10.5	11.3	12.0	12.8	13.5	14.3	15.0	1		
	7.5	10.1		12.4		14.6		16.9				21.4				
	10	13.5	15.0	16.5	18.0	19.5	21.0	22.5	24.0	25.5	27.0	28.5	30.0	_		
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	Central administ	ration	See	HDIL	المانانة	nes fo	r adm	inistra	tion v	ia cen	tral ve	nous	20000	device		
	Central administration: See HDU guidelines for administration via central venous access device using a syringe pump.															
	asing a synnige p	amp.														
Extravasation	Administer via a	cent	ral ve	nous	acces	s devi	ce if	possih	le. Th	ne inf	usion	has a	low	pH and		
	extravasation is I							•						-		
	peripheral ischae	•				-	•							····		
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This information has been summarised to act as a guide for those administering IV medication. The monograph should be used in conjunction with the drug data sheet and BNF for information on dose, adverse effects, cautions and contra-indications. Further information is available from pharmacy on 22142 or 22546.

## **Notes**

- It is recommended that treatment with dobutamine should be discontinued gradually.
- Solution may turn pink due to oxidation of the drug. There is no significant loss of potency during recommended storage and administration periods.
- Continuous ECG and BP monitoring is recommended
- Monitor heart rate and rhythm, arterial blood pressure, blood glucose, urine output, serum potassium and infusion rate.

Information provided relates to Dobutamine manufactured by Mercury Pharma.