Form no.: HIM-PMR-A15



[PC]: Personal Computer

PATIENT'S INFORMATION (Please Stick Label)

ULTRASONIC ENHANCING AGENT ECHOCARDIOGRAPHY PROCEDURE

Allergic to Sulfur Hexafluoride: YES NO							
1.	SonoVue is a contrast agent used in echocardiography procedure to improve ultrasound image quality of the heart. It contains millions of microbubbles that act as a reflector of the ultrasound have and provide image appropriate.						
2. 3.	beam and provide image enhancement. It is used when the result of the echocardiography study without contrast agent is inconclusive. CVT will consult the echo referring doctor to obtain permission to use SonoVue contrast agent in he/she encounters a patient with poor echo window (ie if ≥ 2 adjacent segments are not visualised).						
1	and for suspected cardiac thrombus.						
4.	Once the approval has been obtained, CVT will proceed with SonoVue administration protocol as per WI/SOP.						
5.	The requesting doctor shall transcribe the SonoVue contrast media ordering in Trackare (in eNCL Investigation Request section) immediately once he/she received a request from CVT or within 24 hours if he/she has no immediate access to Trackcare.						
6.							
7.	The recommended dose of SonoVue for thrombus assessment is 2 ml administered as an intravenous bolus injection. During a single examination, a second injection of 2 ml SonoVue may be administered to prolong contrast enhancement. For patient with poor echo window, the recommended dose of SonoVue is 1ml administered as an intravenous bolus injection. During a single examination, repeat injection of 1 ml SonoVue may be administered to prolong contrast enhancement. Follow each SonoVue injection with an intravenous flush using 5 mL of 0.9% Sodium Chloride injection.						
SonoVue CONTRAST INJECTION:							
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	otal Dose in mg (5mg/ml) ven (√)	5 mg	10 mg	15 mg	20 mg	25 mg	
_							
Prepared by:			Adminis	Administrated by:			
Signature:				Signature:			
Name :				Name :			
Date :			Date Time				
[C\	/T] : Cardiovascular Technologist						

Effective Date: 13 July 2020
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