Pelosi Medical Center

Preoperative Bloodwork Mandatory Testing 2011

To all concerned:

Please perform all of the following tests on behalf of the patient:

Panel: CBC: complete blood count with platelet count & differential

Complete metabolic panel: must include liver function tests (AST or SGOT, ALT or SGPT)

PT/PTT: prothrombin time/partial thromboplastin time

Screening for HIV, Hepatitis B, Hepatitis C

Thank you for your cooperation.

Marco A. Pelosi, III, MD, FACOG, FACS, FICS, FAACS

Please forward all results and address any questions regarding these tests to:

Marco A. Pelosi, III, MD

Pelosi Medical Center

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Email: mpelosi3@PelosiMedicalCenter.com

Please note: I have attached an example of the required bloodwork testing results for reference.

Test Name	In Range Out of Range	Reference Range	Lab
CHEM-SCREEN PANEL			≠ DD
GLUCOSE	92	65-139 mg/dL	TBR
The glucose ref	erence range is based on a non-fa	eting etato	
SODIUM	137	135-146. mmol/L	
POTASSIUM	4.9	3.5-5.3 mmol/L	
	***	3.3-3.3 MMOI/L	
Results slightl	y increased due to hemolysis.		
CHLURIDE	104	98-110 mmol/L	
CARBON DIOXIDE	24	21-33 mmol/L	
UREA NITROGEN	13		
CREATININE	0.71	7-25 mg/dL	
BUN/CREATININE RATIO	NOTE	0.59-1.07 mg/dL	
	ratio is not reported when the Bu	6-22	
Creatinine value	es are within normal limits.	n and	
URIC ACID	5.0	2 5 7 0 (4)	
PHOSPHORUS	3.7	2.5-7.0 mg/dL	
CALCIUM	9.1	2.5-4.5 mg/dL	
CHOLESTEROL, TOTAL	158	8.6-10.2 mg/dL	
TRIGLYCERIDES	119	125-200 mg/dL	
PROTEIN, TOTAL	8.0	<150 mg/dL	
ALBUMIN	4.4	6.2-8.3 g/dL	
GLOBUL IN, CALCULATED		3.6-5.1 g/dL	
A/G RATIO	3.6	2.2-3.9 g/dL	
BILIRUBIN, TOTAL	1.2	1.0-2.1	
DILINGBIN, IOTHL	0.4	0.2-1.2 mg/dL	
Results slightly	decreased due to hemolysis.		
BILIRUBIN, DIRECT	0.2	4-00 4	
ALKALINE PHOSPHATASE	46	< = 0.2 mg/dL	
GGT	22	33-115 U/L	
AST	44	3-55 U/L	
Test not perform	ed. Specimen unsuitable for testi	10-35 U/L	
hemolysis.	ow. opecimen unsultable for testi	ng due to	
ALT	10	6 40 U/I	
LD	10	6-40 U/L	
Took wat a		100-200 U/L	

Test Name		In Range Out of	Range Reference Range	Lab
EGFR NON EGFR AFR	AFR AMERICAN ICAN AMERICAN	102 118	>=60 mL/min/1.73m2 >=60 mL/min/1.73m2	
CBC (INCLU	DES DIFF/PLT) Test not performed. S	Test Not Performed Specimen received clot	ted.	TBR
PROTHROMB IN	N TIME W/ INR Reference Range Moderate-intensity Wa Higher-intensity Warf	1.1 0.9 - rfarin Therapy 2.0 - arin Therapy 3.0 -	3.0	TBR
PROTHROME		11.8 9.0 - 11.5 seconds	H Seconds	
PTT See foot	mote 1	27	22-34 Seconds	TBR
HBS AG W/R HBS AG	EFLEX CONF	Non Reactive	Non Reactive	TBR
HEPATITIS C HCV RATIO HEPATITIS	C AB A Non Reactive to	0.10 Non Reactive test result does not e of HCV infection sinc oversion is variable.	<1.0 Ratio Non Reactive exclude te the	TBR
HIV 1/2,EII HIV 1/2 E	A SCR W/REFL IA A Nonreactive HIV 1/2 infection since the ti variable. If acute HIV retesting and nucleic testing is recommended (CDC,MMWR, September 2	Nonreactive antibody result does me frame for seroconv infection is suspect acid amplification (H	ersion is ed, antibody IV DNA/RNA)	TBR

FOOTNOTE(S):

The therapeutic range for unfractionated heparin therapy is 1.5-2.5 times the mean of the reference interval.

In patients in whom there is an apparent heparin resistance, a heparin level by an anti-Xa method, test 30292X, is available.

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Government regulations require the assurance of patient confidentiality.

PERFORMING LABORATORY INFORMATION:

TBR Quest Diagnostics One Malcolm Avenue Teterboro NJ 07608
Laboratory Director: William E. Tarr, M.D. CLIA No: 31D0696246
LIST OF RESULTS PRINTED IN THE OUT OF RANGE COLUMN:

PROTHROMBIN TIME 11.8 H Seconds
Reference Range-PT: 9.0 - 11.5 seconds