

LAB VALUE CHEAT SHEET

VITAL SIGNS

BLOOD PRESSURE	SYSTOLIC	120 mmHg
	DIASTOLIC	80 mmHg
HEART RATE	60 – 100 bpm	
RESPIRATIONS	12 – 20 breaths/min	
TEMPERATURE	97.8 – 99°F (36.5 – 37.2°C)	
OXYGEN	95 – 100%	
OXYGEN IN CORD PT.	as low as 88%	



CORD pts are expected to have low O₂ levels

COMPLETE BLOOD COUNT (CBC)

WBCs	4,500 – 11,000 / μ L
RBCs	4.5 – 5.5 $\times 10^6$ / μ L
PLTs	150,000 – 450,000 / μ L
HEMOGLOBIN (HGB)	FEMALE: 12 – 16 g/dL
	MALE: 13 – 18 g/dL
HEMATOCRIT (HCT)	FEMALE: 36% – 48%
	MALE: 39% – 54%



HBA1C

NON-DIABETIC	4 – 5.6%
PRE-DIABETIC	5.7 – 6.4%
DIABETIC	> 6.5%
Goal for diabetic:	< 7%

LIVER FUNCTION TEST (LFT)

ALT	7 – 56 U/L
AST	5 – 40 U/L
ALP	40 – 120 U/L
BILIRUBIN	0.1 – 1.2 mg/dL

BMI

UNDERWEIGHT	<18.5
HEALTHY WEIGHT	18.5 – 24.9
OVERWEIGHT	25.0 – 29.9
OBESITY	> 30.0

ABGs

PH	7.35 – 7.45
P _a CO ₂	35 – 45 mmHg
P _a O ₂	80 – 100 mmHg
HCO ₃	22 – 26 mEq/L



LIPID PANEL

TOTAL CHOLESTEROL	<200 mg/dL
TRIGLYCERIDE	<150 mg/dL
LDL	<100 mg/dL
HDL	>60 mg/dL



LDL bad cholesterol
- we want **LOW** levels



HDL Happy cholesterol
- we want **HIGH** levels

RENAL

CALCIUM	9 – 11 mg/dL
MAGNESIUM	1.5 – 2.5 mg/dL
PHOSPHORUS	2.5 – 4.5 mg/dL
SPECIFIC GRAVITY	1.010 – 1.030
GFR	90 – 120 mL/min/1.73 m ²
BUN	7 – 20 mg/dL
CREATININE	0.6 – 1.2 mg/dL



PANCREAS

AMYLASE	30 – 110 U/L
LIPASE	0 – 150 U/L



BASIC METABOLIC PANEL (BMP)

SODIUM	135 – 145 mEq/L
POTASSIUM	3.5 – 5.0 mEq/L
CHLORIDE	95 – 105 mEq/L
CALCIUM	9 – 11 mg/dL
BUN	7 – 20 mg/dL
CREATININE	0.6 – 1.2 mg/dL
ALBUMIN	3.4 – 5.4 g/dL
TOTAL PROTEIN	6.2 – 8.2 g/dL

COAGs

PT	10 – 13 sec
PTT	25 – 35 sec
aPTT	NOT ON HEPARIN: 30–40 sec
	ON HEPARIN: 47–70 sec
INR	NOT ON WARFARIN: < 1 sec
	ON WARFARIN: 2 – 3 sec



OTHER

MAP (mean arterial pressure)	70 – 100 mmHg
ICP (intracranial pressure)	5 – 15 mmHg
GLASGOW COMA SCALE	BEST = 15
MILD: 13–15	MODERATE: 9–12
SEVERE: 3–8	

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Cheat Sheets For Reviewing Lab Results

Mike Chapple



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The FDA and Worldwide Current Good Manufacturing Practices and Quality System Requirements Guidebook for Finished Pharmaceuticals José Rodríguez-Pérez, 2014-08-15 This guidance book is meant as a resource to manufacturers of pharmaceuticals providing up to date information concerning required and recommended quality system practices It should be used as a companion to the regulations standards themselves and texts on the specific processes and activities contained within the QMS This book includes chapters on US current Good Manufacturing Practice GMP international GMP global GMP guides and harmonization detailed analysis of the requirements and guidances missing subparts what inspectors are looking for and the price of noncompliance It also includes an appendix with two tabulated comparisons the first compares US European PIC S Canadian and WHO cGMPs while the second compares US cGMPs with effective quality system elements The companion CD contains cGMP regulations for sterile products produced by aseptic processing it also includes updated data of statistical enforcement by the FDA both domestically and abroad a detailed glossary and dozens of FDA guidance documents as well as international regulations EU and Canada and harmonization documents WHO PIC S and ICH A very comprehensive checklist for a cGMP audit that is based on risk management criteria is also included Finally a comprehensive GMP exam is also included

The American Architect and the Architectural Review, 1921

Quality Control in the age of Risk Management, An Issue of Clinics in Laboratory Medicine James O. Westgard, Sten Westgard, 2013-03-28 In October of 2011 CLSI published a new guideline EP23A on Laboratory Quality Control Based on Risk Management In March 2012 CMS announced its intention to incorporate key concepts from EP23A into its Interpretative Guidelines and QC policy for Individualized Quality Control Plans Thus begins a new era of Quality Control in the Age of Risk Management This issue is intended to help laboratories with the transition between traditional QC practices and the new risk management approach Laboratories face a steep learning curve to apply risk analysis for identifying and prioritizing failure modes developing and implementing control mechanisms to detect those failure modes and assessing the acceptability of the residual risks that exist after implementation of a QC Plan One of the main benefits of the new risk analysis based QC Plans should be an integration of all the control mechanisms that are needed to monitor the total testing process including pre analytic analytic and post analytic controls One of the main risks of the new approach is an expectation that Statistical QC is no longer important even though SQC still remains the most useful and flexible approach for monitoring the quality of the analytic process The key to the future is the successful integration of all these control mechanisms to provide a cost effective quality system that monitors all phases of the total testing process This issue should help laboratories understand the evolution of QC practices to include risk management but also to recognize the need to maintain traditional techniques such as Statistical QC especially during the transition to well designed and carefully validated QC Plans Risk analysis may be risky

business unless laboratories proceed carefully and cautiously

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Table of Contents Cheat Sheets For Reviewing Lab Results

1. Understanding the eBook Cheat Sheets For Reviewing Lab Results
 - The Rise of Digital Reading Cheat Sheets For Reviewing Lab Results
 - Advantages of eBooks Over Traditional Books
2. Identifying Cheat Sheets For Reviewing Lab Results
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Cheat Sheets For Reviewing Lab Results
 - User-Friendly Interface
4. Exploring eBook Recommendations from Cheat Sheets For Reviewing Lab Results
 - Personalized Recommendations
 - Cheat Sheets For Reviewing Lab Results User Reviews and Ratings
 - Cheat Sheets For Reviewing Lab Results and Bestseller Lists
5. Accessing Cheat Sheets For Reviewing Lab Results Free and Paid eBooks
 - Cheat Sheets For Reviewing Lab Results Public Domain eBooks
 - Cheat Sheets For Reviewing Lab Results eBook Subscription Services
 - Cheat Sheets For Reviewing Lab Results Budget-Friendly Options

6. Navigating Cheat Sheets For Reviewing Lab Results eBook Formats
 - ePub, PDF, MOBI, and More
 - Cheat Sheets For Reviewing Lab Results Compatibility with Devices
 - Cheat Sheets For Reviewing Lab Results Enhanced eBook Features
7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Cheat Sheets For Reviewing Lab Results
 - Highlighting and Note-Taking Cheat Sheets For Reviewing Lab Results
 - Interactive Elements Cheat Sheets For Reviewing Lab Results
8. Staying Engaged with Cheat Sheets For Reviewing Lab Results
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Cheat Sheets For Reviewing Lab Results
9. Balancing eBooks and Physical Books Cheat Sheets For Reviewing Lab Results
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Cheat Sheets For Reviewing Lab Results
10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
11. Cultivating a Reading Routine Cheat Sheets For Reviewing Lab Results
 - Setting Reading Goals Cheat Sheets For Reviewing Lab Results
 - Carving Out Dedicated Reading Time
12. Sourcing Reliable Information of Cheat Sheets For Reviewing Lab Results
 - Fact-Checking eBook Content of Cheat Sheets For Reviewing Lab Results
 - Distinguishing Credible Sources
13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
14. Embracing eBook Trends
 - Integration of Multimedia Elements

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