



CLIA '88 AND GRADING

The Clinical Laboratory Improvement Amendments of 1988 (CLIA '88) were established by the federal government (CMS) to regulate clinical laboratories and proficiency test providers like API. One of the subjects regulated by CLIA '88 is the way proficiency test results are graded. The method by which an analyte is graded depends on whether the test is qualitative (e.g., a blood sample reactive / nonreactive for Syphilis), semi-quantitative (e.g., color comparison tests on a urine dipstick), or quantitative (e.g., the amount of Cholesterol in a blood sample).

HOW YOUR RESULTS ARE GRADED (QUALITATIVE)

Qualitative results are graded based on 80% agreement (called "consensus") of participants giving a particular answer or group of clinically correct answers. The only exception to this is Immunohematology results which are based on 95% consensus. In cases where participants do not reach consensus, a consensus of referee laboratories will be used. If the necessary consensus is not reached, the sample is not graded.

Microbiology results are graded as described above, but scores for regulated tests are reported at the sub-specialty level (e.g., bacteriology or virology).

HOW YOUR RESULTS ARE GRADED (SEMI-QUANTITATIVE)

Semi-quantitative results are graded based on 80% agreement (called "consensus") of participants giving a group of clinically correct answers. For most analytes, this group of clinically correct answers is generally the most common response and one response lower and higher, to accommodate differences in interpretation that will not affect patient care. Both negative and positive responses will not be accepted on the same sample. For regulated titer analytes (Rheumatoid Factor, Rubella, and Anti-Streptolysin O), CLIA requires the most common response \pm two dilutions be considered acceptable. Regardless of how the group of acceptable answers is determined, the consensus rule is applied to establish whether a sample can be graded. In cases where participants do not reach a consensus, a consensus of referee laboratories may be used. If the proper consensus is not reached, the sample is not graded.

HOW YOUR RESULTS ARE GRADED (QUANTITATIVE)

The criteria used to evaluate your quantitative results are based on a target value (mean) \pm a fixed amount. This fixed amount is determined by CLIA '88 for regulated analytes and is expressed as a percentage, a specific quantity, or a number of standard deviations. The grading criteria are listed below. The target value is the mean (average) value of your comparison group. A comparison group (peer group) may consist of those laboratories that use the same instrument or reagent as you, or those laboratories that use the same method principle. A comparison group of "All Participants" (all facilities performing that test) may also be used as well as a group of referee laboratories. The comparison group must contain at least 10 laboratories. In cases where an appropriate comparison group cannot be found, no grading criteria is applied. In addition, only samples with at least 80% consensus are graded.

GRADING CRITERIA for CMS Regulated Analytes

CHEMISTRY

Allowed variance from the Target Value

Albumin	\pm 10 percent
Alkaline Phosphatase	\pm 30 percent
ALT/SGPT	\pm 20 percent
Amylase	\pm 30 percent
AST/SGOT	\pm 20 percent
Bilirubin, Total	\pm 0.4 mg/dL or \pm 20 percent, whichever is greater
Calcium	\pm 1 mg/dL

CHEMISTRY

Chloride	± 5 percent
Cholesterol	± 10 percent
Creatine Kinase (CK), total	± 30 percent
Creatine Kinase, Isoenzyme (CK-MB)	± 3 SD or ± 3 (ng/mL or U/L), whichever is greater
Creatinine	± 0.3 mg/dL or ± 15 percent, whichever is greater
Glucose	± 6 mg/dL or ± 10 percent, whichever is greater
HDL, Cholesterol	± 30 percent
Iron	± 20 percent
LDH	± 20 percent
Magnesium	± 25 percent
pCO ₂	± 5 mmHg or ± 8 percent, whichever is greater
pH	± 0.04
pO ₂	± 3 SD
Potassium	± 0.5 mmol/L
Sodium	± 4 mmol/L
Total Protein	± 10 percent
Triglycerides	± 25 percent
Urea Nitrogen (BUN)	± 2 mg/dL or ± 9 percent, whichever is greater
Uric Acid	± 17 percent

ENDOCRINOLOGY

Cortisol	± 25 percent
Free Thyroxine (Free T4)	± 3 SD
HCG - Quantitative	± 3 SD or ± 10 mIU/mL, whichever is greater
T-Uptake	± 3 SD
Thyroid Stimulating Hormone	± 3 SD
Thyroxine (T4)	± 1 µg/dL or ± 20 percent, whichever is greater
Triiodothyronine (T3)	± 3 SD

HEMATOLOGY and COAGULATION

White Blood Cell Automated Differential (includes Neutrophils, Lymphocytes, Monocytes, Eosinophils, Basophils)	± 3 SD or ± 1, whichever is greater
Erythrocyte (Red Blood Cell) Count	± 6 percent
Fibrinogen	± 20 percent
Hematocrit	± 6 percent
Hemoglobin	± 7 percent
Leukocyte (White Blood Cell) Count	± 15 percent
Partial Thromboplastin Time (APTT)	± 15 percent
Platelet Count	± 25 percent
Prothrombin Time (PT)	± 15 percent

Allowed variance from the Target Value

Allowed variance from the Target Value

Allowed variance from the Target Value

IMMUNOLOGY

Alpha-1-Antitrypsin	± 3 SD
Alpha-fetoprotein	± 3 SD
Complement C3	± 3 SD
Complement C4	± 3 SD
IgA	± 3 SD
IgE	± 3 SD
IgG	± 25 percent
IgM	± 3 SD

Allowed variance from the Target Value

TOXICOLOGY

Alcohol	± 10 mg/dL or ± 25 percent, whichever is greater
Blood Lead	± 4 μ g/dL or ± 10 percent, whichever is greater
Carbamazepine	± 25 percent
Digoxin	± 0.2 ng/mL or ± 20 percent, whichever is greater
Gentamicin	± 25 percent
Lithium	± 0.3 mmol/L or ± 20 percent, whichever is greater
Phenobarbital	± 20 percent
Phenytoin	± 25 percent
Theophylline	± 25 percent
Tobramycin	± 25 percent
Valproic Acid	± 25 percent

Allowed variance from the Target Value

All other analytes are considered “Not Scored” by CMS and are graded using the Target Value ± 2 SD. **Exceptions** are noted below:

NOT SCORED for CMS

Acetaminophen	± 3 SD or ± 2.5 μ g/mL, whichever is greater
ACT	± 3 SD
Allergen Specific IgE	± 3 SD
Ammonia	± 2 SD or ± 10 μ mol/L, whichever is greater
Anti-CCP	± 3 SD or ± 5 U/mL, whichever is greater
Anti-HBs (Quant)	± 3 SD or ± 3 mIU/mL
Anti-Streptolysin O, quant	± 2 SD or ± 25 IU/mL, whichever is greater
Antithrombin III Activity	± 3 SD
Anti-Xa (Hybrid, LMWH, UFH curves)	± 3 SD
Aspirin Induced Inhibition	± 3 SD or ± 50 ARU, whichever is greater
Bacteria (Urine Microscopic System)	± 3 SD or ± 5 cells/ μ L, whichever is greater
Basophils (CSF/Body Fluid)	± 3
Beta-Hydroxybutyrate	± 3 SD or ± 0.2 mmol/L, whichever is greater
Bilirubin, Direct	± 2 SD or ± 0.4 mg/dL, whichever is greater

Allowed variance from the Target Value

NOT SCORED for CMS

BNP
Body Fluid Albumin
Body Fluid ALP
Body Fluid Amylase
Body Fluid Bilirubin, Total
Body Fluid Calcium
Body Fluid Chloride
Body Fluid Cholesterol
Body Fluid Creatinine
Body Fluid Glucose
Body Fluid LDH
Body Fluid Lipase
Body Fluid pH (color comparison)
Body Fluid Potassium
Body Fluid Sodium
Body Fluid Total Protein
Body Fluid Triglycerides
Body Fluid Urea Nitrogen
Body Fluid Uric Acid
C-Reactive Protein (hs)
C-Reactive Protein (quant)
Calcium, Ionized
Carboxyhemoglobin
Casts (Urine Microscopic System)
CEA
CO₂ (serum)
Conductivity
Creatinine (UAD quant)
Crystals (Iris-quant)
CSF Glucose
CSF Lactic Acid
CSF Total Protein
Cystatin C
D-Dimer
Eosinophils (CSF/Body Fluid)
Epithelial Cells (Urine Microscopic System)
Estriol (unconjugated)
Factor II, IX, V, VII, VIII, X, XI
Ferritin

Allowed variance from the Target Value

± 3 SD or ± 10 pg/mL, whichever is greater
 ± 3 SD or ± 10 percent, whichever is greater
 ± 30 percent
 ± 30 percent
 ± 0.4 mg/dL or ± 20 percent, whichever is greater
 ± 1 mg/dL
 ± 2 SD or ± 5 percent, whichever is greater
 ± 3 SD
 ± 0.3 mg/dL or ± 15 percent, whichever is greater
 ± 6 mg/dL or ± 10 percent, whichever is greater
 ± 20 percent
 ± 2 SD or ± 30 percent, whichever is greater
 ± 1
 ± 0.5 mmol/L
 ± 4 mmol/L
 ± 3 SD or ± 10 percent, whichever is greater
 ± 3 SD or ± 10 mg/dL, whichever is greater
 ± 2 mg/dL or ± 9 percent, whichever is greater
 ± 17 percent
 ± 2 SD or ± 0.2 mg/dL, whichever is greater
 ± 2 SD or ± 0.3 mg/dL, whichever is greater
 ± 3 SD or ± 0.05 mmol/L, whichever is greater
 ± 3 SD or ± 3 , whichever is greater
 ± 3 SD or ± 1 / μ L, whichever is greater
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 2 SD or ± 20 percent, whichever is greater
 ± 10
 ± 6 mg/dL or ± 10 percent, whichever is greater
 ± 3 SD or ± 0.4 mmol/L whichever is greater
 ± 3 SD or ± 20 percent, whichever is greater
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 6
 ± 3 SD or ± 1 cell/ μ L, whichever is greater
 ± 2 SD or ± 0.5 ng/mL, whichever is greater
 ± 3 SD
 ± 3 SD

NOT SCORED for CMS

Folate	± 3 SD
Free PSA	± 3 SD or ± 0.4 ng/mL, whichever is greater
Fructosamine	± 2 SD or ± 0.2 mmol/L, whichever is greater
FSH	± 3 SD
GGT	± 20 percent
Glucose (Whole Blood - waived)	± 10 mg/dL or ± 20 percent, whichever is greater
Glycohemoglobin (HbA1c)	± 3 SD or ± 20 percent, whichever is greater
Growth Hormone	± 3 SD or ± 0.2 ng/mL, whichever is greater
Haptoglobin	± 3 SD
HBV Viral Load	± 3 SD
HCV Viral Load	± 3 SD
Hematocrit (Waived)	± 6 percent
Hemoglobin (Waived)	± 7 percent
Hemoglobin F, quantitative	± 2 SD or ± 0.6 , whichever is greater
HIV-1 RNA Viral Load	± 3 SD
Homocysteine	± 3 SD
IG absolute	± 3 SD
IG percent	± 3 SD
Immature Platelet Fraction	± 3 SD
INR	± 3 SD
Insulin	± 3 SD
IRF	± 3 SD
Lactate (Lactic Acid)	± 3 SD or ± 0.4 mmol/L, whichever is greater
Lamellar Body Count	± 25 percent
Lidocaine	± 3 SD or ± 10 percent, whichever is greater
Lipase	± 2 SD or ± 30 U/L, whichever is greater
Luteinizing Hormone	± 3 SD
Lymphocytes (CSF/Body Fluid)	± 20
Magnesium, Ionized (Blood Gas)	± 3 SD or ± 0.1 mmol/L, whichever is greater
MCH	± 3 SD
MCHC	± 3 SD
MCV	± 3 SD or ± 3 fL, whichever is greater
Methemoglobin	± 2
Microalbumin	± 3 SD or ± 30 percent, whichever is greater
Monocytes (CSF/Body Fluid)	± 20
Mononuclear (CSF/Body Fluid)	± 25
MPV	± 3 SD
Myoglobin	± 2 SD or ± 15 ng/mL, whichever is greater
Neutrophils (CSF/Body Fluid)	± 20

Allowed variance from the Target Value

NOT SCORED for CMS

Nitrite
Non-Squamous Epithelial Cells (Iris, quant)
NT-pro-BNP
NUCL (Body Fluid - Iris)
Nucleated RBCs (Hem - 5C)
Nucleated RBCs (Hem - 5D & Heme - 5S)
Osmolality
Oxidants
Oxyhemoglobin
P2Y12 Inhibition
PAP
Plasminogen
Polymorphonuclear (CSF/Body Fluid)
Prealbumin
Procalcitonin
Protein C Activity
Protein S- Activity, Free Antigen, and Total Antigen
Progesterone
Prolactin
PSA
RBC (Body Fluid automated)
RDW (including CV and SD)
Reticulocyte Count
Reticulocyte Hgb
Rheumatoid Factor, quant
Rubella, quant
Salicylates
Sedimentation Rate
Sex Hormone Binding Globulin
Specific Gravity
Sperm Count (Post- Vasectomy) (manual and automated)
Sperm Morphology
Sperm Viability
TCO₂
Testosterone
Thrombin Time
Thromboelastometry (all analytes)
Thyroglobulin Ab (Anti-TG)
Thyroid Microsomal Ab (Anti-TPO)

Allowed variance from the Target Value

± 2 SD or ± 30 $\mu\text{g/mL}$, whichever is greater
 ± 10
 ± 2 SD or ± 10 pg/mL , whichever is greater
 ± 3 SD
 ± 3 SD or ± 1 , whichever is greater
 ± 3 SD or $\pm 0.01 \times 10^9/\text{L}$, whichever is greater
 ± 3 SD
 ± 2 SD or ± 30 $\mu\text{g/mL}$, whichever is greater
 ± 3 SD or ± 3 , whichever is greater
 ± 3 SD or ± 30 PRU, whichever is greater
 ± 2 SD or ± 30 percent, whichever is greater
 ± 3 SD
 ± 25
 ± 5 mg/dL or ± 25 percent, whichever is greater
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 3 SD or ± 0.4 ng/mL , whichever is greater
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 3 SD or ± 20 IU/mL, whichever is greater
 ± 3 SD or ± 10 IU/mL, whichever is greater
 ± 3 SD or ± 2.8 mg/dL , whichever is greater
 ± 2 SD or ± 3 mm/hr , whichever is greater
 ± 3 SD
 ± 0.01
 ± 3 SD
 ± 20 percent normal
 ± 2 SD or ± 2 percent viable, whichever is greater
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 2 SD or ± 20 IU/mL, whichever is greater
 ± 2 SD or ± 20 IU/mL, whichever is greater

NOT SCORED for CMS

TIBC (all)
Total Nucleated Cell Count
Transferrin
Troponin I
Troponin T
Urine Amylase
Urine Calcium
Urine Chloride
Urine Creatinine
Urine Glucose
Urine Magnesium
Urine Microalbumin
Urine Osmolality
Urine Phosphorus
Urine Potassium
Urine Sodium
Urine Total Protein
Urine Urea (BUN)
Urine Uric Acid
Vancomycin
Vitamin B-12
WBC (Body Fluid automated)

Allowed variance from the Target Value

± 20 percent
 ± 3 SD
 ± 20 percent
 ± 3 SD or ± 0.3 ng/mL, whichever is greater
 ± 2 SD or ± 0.1 ng/mL, whichever is greater
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 20 percent
 ± 3 SD
 ± 25 percent
 ± 3 SD or ± 30 percent, whichever is greater
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 3 SD
 ± 2 μ g/mL or ± 20 percent, whichever is greater
 ± 3 SD
 ± 3 SD