

Supplemental:

<b>TABLE X CLIA Part 493 Laboratory Requirements At A Glance</b>		
<b>Subpart A</b>	<b>General Provisions</b>	Defines test complexities (waived, provider performed microscopy, non-waived (moderate vs high complexity)
<b>Subpart B</b>	<b>Certificate of Waiver</b>	Applies only to labs performing waived tests.
<b>Subpart C</b>	<b>Registration Certificate, Certificate for Provider-performed Microscopy Procedures, and Certificate of Compliance</b>	Applies only to labs performing provider performed microscopy
<b>Subpart D</b>	<b>(D) Certificate of Accreditation</b>	Accreditation requirements.
<b>Subpart E</b>	<b>(E) Accreditation by a Private, Nonprofit Accreditation Organization or Exemption Under an Approved State Laboratory Program</b>	
<b>Subpart F</b>	<b>General Administration</b>	
<b>Subpart H</b> <b>Subpart I</b>	<b>(H) Participation in Proficiency Testing for Laboratories Performing Nonwaived Testing</b>  <b>(I) Proficiency Testing Programs for Nonwaived Testing</b>	General and discipline specific ** requirements for proficiency testing.
<b>Subpart J</b>	<b>Facility Administration for Nonwaived Testing</b>	Facility requirements.
<b>Subpart K</b>	<b>Quality System for Nonwaived Testing</b>	Discipline specific ** requirements for quality nonwaived testing.
<b>Subpart L</b>	<b>General Laboratory Systems</b>	General requirements regarding patient and specimen issues Also listed are specific requirements associated with preanalytic, analytic (general and discipline specific **) and postanalytic phases of laboratory testing.
<b>Subpart M</b>	<b>Personnel for Nonwaived Testing</b>	Personnel requirements for nonwaived testing across test complexity (moderate vs high) and discipline and role (director through testing staff)

<b>Subpart Q</b>	<b>Inspection</b>	Inspection requirements across complexity and discipline
<b>Subpart R</b>	<b>Enforcement Procedures</b>	Details regarding sanctions and penalties.
<b>Subpart T</b>	<b>Consultations</b>	Establishment and function of the Clinical Laboratory Improvement Advisory Committee.
<b>Subparts L, N-P &amp; S</b>	<b>[Reserved]</b>	

**\*\* Bacteriology, mycobacteriology, mycology, parasitology, virology, syphilis serology, general immunology, routine chemistry, urinalysis, endocrinology, toxicology, hematology, immunohematology, histopathology, oral pathology, cytology, clinical cytogenetics, radiobioassay and histocompatibility.**

Supplemental Figure 1: CLIA Requirements at a glance as an example of the extensive detail required for CLIA license from *Title 42—Public Health Chapter IV—Centers for Medicare & Medicaid Services, Department of Health and Human Services*  
<https://www.ecfr.gov/current/title-42/chapter-IV/subchapter-G/part-493/>  
*Subchapter G—Standards and Certification.*