Evidence of Compliance:

✓ Records of qualifications including degree or transcript, certification/registration, current license (if required) and work history in related field

REFERENCES
1) Department of Health and Human Services, Centers for Medicare and Medicaid Services. Clinical laboratory improvement amendments of 1988; final rule. *Fed Register.* 2023(Dec 28): [42CFR493.1483].

CYP.07900 Screening Personnel — Phase II

All screening personnel satisfy one or more of the following three criteria.

1. **Pathologist or physician qualified as section director or technical supervisor**
2. **Supervisory level cytotechnologist**
3. **Qualified cytotechnologist**

Evidence of Compliance:

✓ Records of qualifications including degree or transcript, certification/registration, current license (if required) and work history in related field

REFERENCES
1) Department of Health and Human Services, Centers for Medicare and Medicaid Services. Clinical laboratory improvement amendments of 1988; final rule. *Fed Register.* 2023(Dec 28): [42 CFR493.1449(e)], [42CFR493.1469], [42CFR493.1483].

CYP.08100 General Supervisor Qualifications - Cytopathology — Phase II

The cytopathology laboratory has a general supervisor who meets the qualifications defined by CLIA (for laboratories subject to US regulations) and other applicable national, federal, state (or provincial), or local laws and regulations.

NOTE: The supervisor can be a pathologist boarded in anatomic pathology. Alternatively, the supervisor can be qualified as a cytotechnologist, with at least three years of full-time experience as a cytotechnologist within the preceding 10 years. The section director/technical supervisor may also serve as the general supervisor.

For laboratories not subject to US regulations, appropriate national, state or provincial, or local laws and regulations also apply.

Evidence of Compliance:

✓ Records of qualifications including degree or transcript, certification/registration, current license (if required) and work history in related field

REFERENCES
1) Department of Health and Human Services, Centers for Medicare and Medicaid Services. Clinical laboratory improvement amendments of 1988; final rule. *Fed Register.* 1992(Feb 28): [42CFR493.1467].
2) Department of Health and Human Services, Centers for Medicare and Medicaid Services. Clinical laboratory improvement amendments of 1988; final rule. *Fed Register.* 2023(Dec 28): [42CFR493.1469].

CYP.08200 General Supervisor Responsibilities — Phase II

The cytopathology general supervisor fulfills defined responsibilities.

NOTE: The general supervisor, as designated by the laboratory/section director, is responsible for day-to-day supervision or oversight of the laboratory operation and personnel performing testing and reporting test results. This individual must also:

1. *Be accessible to provide consultation to resolve technical problems*
2. *Record the slide interpretation results of each case he or she examined or reviewed*
3. *For each 24-hour period, record the total number of slides he/she examined (screened/ rescreened) or reviewed, as well as ensuring the recording of the total number of slides evaluated by others*
4. *Record the number of hours he/she spent examining slides in each 24-hour period*