

Electronic Claims Submission: Clinical Laboratory Improvement Amendments (CLIA)

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The Centers for Medicare & Medicaid Services (CMS) regulates all laboratory testing in the U.S. through the Clinical Laboratory Improvement Amendments (CLIA). The objective of the CLIA program is to ensure quality laboratory testing.

A valid CLIA Certificate Identification number is required and must be included on each electronic claim billed for laboratory services, subject to CLIA legislation. You may not receive reimbursement for your electronic claims if the required certification number is missing.

How to apply for a CLIA Certificate

[cms.gov/Regulations-and-Guidance/Legislation/CLIA/How_to_Apply_for_a_CLIA_Certificate_International_Laboratories](https://www.cms.gov/Regulations-and-Guidance/Legislation/CLIA/How_to_Apply_for_a_CLIA_Certificate_International_Laboratories)

This CMS mandate went in to effect on May 1, 2020. Please work with your software vendor or clearinghouse to ensure that the required information is included in your electronic files to avoid EDI claim rejections.

For detailed information on the tests subject to CLIA, please refer to the CMS link below:
[cms.gov/Regulations-and-Guidance/Legislation/CLIA/](https://www.cms.gov/Regulations-and-Guidance/Legislation/CLIA/)

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URL: <https://providernews.anthem.com/georgia/article/electronic-claims-submission-clinical-laboratory-improvement-amendments-clia>

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