

This is an update about information in the provider manual. For access to the latest provider manual, go online to <https://providers.amerigroup.com>.

Genetic testing services to require prior authorization

Summary: Effective May 1, 2017, genetic testing services for epidermal growth factor receptor (EGFR) testing, prothrombin G20210A (factor II) mutation testing, methylenetetrahydrofolate reductase mutation testing and cell-free fetal DNA-based prenatal testing require prior authorization (PA).

What is the impact of this change?

For dates of service on or after May 1, 2017, PA is required for EGFR testing, prothrombin G20210A (factor II) mutation testing, methylenetetrahydrofolate reductase mutation testing and cell-free fetal DNA-based prenatal testing covered by Amerigroup for STAR and CHIP members. Federal and state law as well as state contract language and CMS guidelines, including definitions and specific contract provisions/exclusions, take precedence over these PA rules and must be considered first when determining coverage. **Noncompliance with new requirements may result in denied claims.**

PA requirements will be added to the following codes:

- 81235
- 81291
- 81420
- 81507
- 0009M

To request PA, contact us by phone (1-800-454-3730), fax (1-800-964-3627) or the provider website (<https://providers.amerigroup.com/TX>).

Not all PA requirements are listed here. Detailed PA requirements are available to contracted providers on the provider self-service website (<https://providers.amerigroup.com/TX> > Provider Resources & Documents > Quick Tools > Precertification Lookup Tool).

What if I need assistance?

If you have questions about this communication or need assistance with any other item, contact your local Provider Relations representative or call Provider Services at 1-800-454-3730.

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