

CPT® Proprietary Laboratory Analyses (PLA) Codes: Long Descriptors

It is important to note that further CPT Editorial Panel (Panel) or Executive Committee actions may affect these codes and/or descriptors. For this reason, code numbers and/or descriptor language in the CPT code set may differ at the time of publication. In addition, further Panel actions may result in gaps in code number sequencing.

| Proprietary Name and Clinical Laboratory and/or Manufacturer | Code | Long Code Descriptor | Released to AMA Website | Effective Date | Publication |
|---|----------------|--|-------------------------------|--------------------|-----------------------|
| Praxis Optical Genome Mapping, Praxis Genomics LLC | ⊕0264U | Rare diseases (constitutional/heritable disorders), identification of copy number variations, inversions, insertions, translocations, and other structural variants by optical genome mapping ▶ (For additional PLA code with identical clinical descriptor, see 0260U. See Appendix O or the most current listing on the AMA CPT website to determine appropriate code assignment) ◄ | July 1, 2021 | October 1, 2021 | CPT [®] 2022 |
| Praxis Whole Genome Sequencing, Praxis Genomics LLC | ●0265U | Rare constitutional and other heritable disorders, whole genome and mitochondrial DNA sequence analysis, blood, frozen and formalin-fixed paraffinembedded (FFPE) tissue, saliva, buccal swabs or cell lines, identification of single nucleotide and copy number variants | July 1, 2021 | October 1, 2021 | CPT [®] 2022 |
| Praxis Transcriptome, Praxis Genomics LLC | ●0266U | Unexplained constitutional or other heritable disorders or syndromes, tissue-specific gene expression by whole-transcriptome and next-generation sequencing, blood, formalin-fixed paraffinembedded (FFPE) tissue or fresh frozen tissue, reported as presence or absence of splicing or expression changes | July 1, 2021 | October 1, 2021 | CPT [®] 2022 |
| Praxis Combined Whole Genome Sequencing and Optical Genome Mapping, Praxis Genomics LLC | ●0267U | Rare constitutional and other heritable disorders, identification of copy number variations, inversions, insertions, translocations, and other structural variants by optical genome mapping and whole genome sequencing | July 1, 2021 | October 1, 2021 | CPT [®] 2022 |