

# PROVIDER BULLETIN

## PROVIDER INFORMATION

December 3, 2018

### Lab Management CPT® Code Updates for Fully Insured Commercial and Medicare Advantage Members – eviCore Healthcare Specialty Utilization Management (UM) Program

The following CPT® Codes have been deleted by the American Medical Association (AMA) effective **December 31, 2018**:

Code	Description
81211	BRCA1, BRCA2 (breast cancer 1 and 2) (eg, hereditary breast and ovarian cancer) gene analysis; full sequence analysis and common duplication/deletion variants in BRCA1 (ie, exon 13 del 3.835kb, exon 13 dup 6kb, exon 14-20 del 26kb, exon 22 del 510bp, exon 8-9 del 7.1kb)
81213	BRCA1, BRCA2 (breast cancer 1 and 2) (eg, hereditary breast and ovarian cancer) gene analysis; uncommon duplication/deletion variants
81214	BRCA1 (breast cancer 1) (eg, hereditary breast and ovarian cancer) gene analysis; full sequence analysis and common duplication/deletion variants (ie, exon 13 del 3.835kb, exon 13 dup 6kb, exon 14-20 del 26kb, exon 22 del 510bp, exon 8-9 del 7.1kb)

The following new Proprietary Laboratory Analyses CPT® Codes have been added by the AMA and will **require prior authorization (PA) beginning February 1, 2019**:

Code	Description
81163	BRCA1 (BRCA1, DNA repair associated), BRCA2 (BRCA2, DNA repair associated) (eg, hereditary breast and ovarian cancer) gene analysis; full sequence analysis
81164	BRCA1 (BRCA1, DNA repair associated), BRCA2 (BRCA2, DNA repair associated) (eg, hereditary breast and ovarian cancer) gene analysis; full duplication/deletion analysis (ie, detection of large gene rearrangements)
81165	BRCA1 (BRCA1, DNA repair associated) (eg, hereditary breast and ovarian cancer) gene analysis; full sequence analysis
81166	BRCA1 (BRCA1, DNA repair associated) (eg, hereditary breast and ovarian cancer) gene analysis; full duplication/deletion analysis (ie, detection of large gene rearrangements)
81167	BRCA2 (BRCA2, DNA repair associated) (eg, hereditary breast and ovarian cancer) gene analysis; full duplication/deletion analysis (ie, detection of large gene rearrangements)
81173	AR (androgen receptor) (eg, spinal and bulbar muscular atrophy, Kennedy disease, X chromosome inactivation) gene analysis; full gene sequence
81174	AR (androgen receptor) (eg, spinal and bulbar muscular atrophy, Kennedy disease, X chromosome inactivation) gene analysis; known familial variant
81185	CACNA1A (calcium voltage-gated channel subunit alpha1 A) (eg, spinocerebellar ataxia) gene analysis; full gene sequence

Code	Description
81186	CACNA1A (calcium voltage-gated channel subunit alpha1 A) (eg, spinocerebellar ataxia) gene analysis; known familial variant
81189	CSTB (cystatin B) (eg, Unverricht-Lundborg disease) gene analysis; full gene sequence
81190	CSTB (cystatin B) (eg, Unverricht-Lundborg disease) gene analysis; known familial variant (s)
81286	FXN (frataxin) (eg, Friedreich ataxia) gene analysis; full gene sequence
81289	FXN (frataxin) (eg, Friedreich ataxia) gene analysis; known familial variant (s)
81306	NUDT15 (nudix hydrolase 15) (eg, drug metabolism) gene analysis, common variant(s) (eg, *2, *3, *4, *5, *6)
81336	SMN1 (survival of motor neuron 1, telomeric) (eg, spinal muscular atrophy) gene analysis; full gene sequence
81337	SMN1 (survival of motor neuron 1, telomeric) (eg, spinal muscular atrophy) gene analysis; known familial sequence variant(s)
81443	Genetic testing for severe inherited conditions (eg, cystic fibrosis, Ashkenazi Jewish-associated disorders [eg, Bloom syndrome, Canavan disease, Fanconi anemia type C, mucopolipidosis type VI, Gaucher disease, Tay-Sachs disease], beta hemoglobinopathies, phenylketonuria, galactosemia), genomic sequence analysis panel, must include sequencing of at least 15 genes (eg, ACADM, ARSA, ASPA, ATP7B, BCKDHA, BCKDHB, BLM, CFTR, DHCR7, FANCC, G6PC, GAA, GALT, GBA, GBE1, HBB, HEXA, IKBKAP, MCOLN1, PAH)
81518	Oncology (breast), mRNA, gene expression profiling by real-time RT-PCR of 11 genes (7 content and 4 housekeeping), utilizing formalin-fixed paraffin- embedded tissue, algorithms reported as percentage risk for metastatic recurrence and likelihood of benefit from extended endocrine therapy [Breast Cancer Index, Biotheranostics, Inc]
81596	Infectious disease, chronic hepatitis C virus (HCV) infection, six biochemical assays (ALT, A2-macroglobulin, apolipoprotein A-1, total bilirubin, GGT, and haptoglobin) utilizing serum, prognostic algorithm reported as scores for fibrosis and necroinflammatory activity in liver [HCV FibroSURE, FibroTest, BioPredictive S.A.S.]

The following existing codes will now **require PA** through the eviCore Lab Management Program **effective February 1, 2019**:

Code	Description
0002M	Liver disease, ten biochemical assays (ALT, A2-macroglobulin, apolipoprotein A-1, total bilirubin, GGT, haptoglobin, AST, glucose, total cholesterol and triglycerides) utilizing serum, prognostic algorithm reported as quantitative scores for fibrosis, steatosis and alcoholic steatohepatitis (ASH)
0003M	Liver disease, ten biochemical assays (ALT, A2-macroglobulin, apolipoprotein A-1, total bilirubin, GGT, haptoglobin, AST, glucose, total cholesterol and triglycerides) utilizing serum, prognostic algorithm reported as quantitative scores for fibrosis, steatosis and nonalcoholic steatohepatitis (NASH)

eviCore's Lab Management clinical guidelines are available on the Blue Cross website at [providers.bluecrossmn.com](http://providers.bluecrossmn.com)

- To access the link, select “**Medical Policy**” under **Tools and Resources**, read and accept the Blue Cross Medical Policy Statement
- Click on the “+” (plus) sign next to “**Medical and Behavioral Health Policies**” and locate the “**Medical Policy Supporting Documents**” section
- Click on “**eviCore healthcare Specialty Utilization Management Clinical Guidelines**” link
- Scroll down to the **Lab Management** section.
- Click on the “**BCBSMN Lab Resources Page**”

### **Products Impacted**

- This change only applies to the **fully insured commercial** and **Medicare Advantage** members.
- The changes do not impact commercial self-insured health plans, Government Programs lines of business (Families and Children (F&C), MinnesotaCare (MNCare), SecureBlue (MSHO), and Minnesota Senior Care Plus (MSC+) health plans), Federal Employee Program (FEP), Medicare Supplement (Senior Gold, Basic Medicare Blue and Extended Basic Blue), Platinum Blue as those lines of business have separate PA requirements.

### **To submit a PA Request to eviCore**

Providers submit eviCore PA requests via our free [Availity](#) provider portal. Instructions on how to utilize this portal are found on the Availity website.

Providers need to reference the eviCore clinical guideline criteria, submit prior authorization request via Availity, and submit all applicable clinical documentation with the PA request. Failure to submit required information may result in review delays or denial of the request due to insufficient information.

Note: An approved PA does not guarantee coverage under a member's benefit plan. Member benefit plans vary in coverage and some plans may not provide coverage for certain services discussed in the medical policies.

### **Questions?**

If you need to submit a PA by phone or need to speak to an eviCore representative call **844-224-0494**, 7:00 a.m. to 7:00 p.m. CST, Monday - Friday.