

TO: MEDICAID PROVIDERS

RE: NEW YORK STATE MEDICAID EXPANDED COVERAGE OF TESTING FOR LYNCH SYNDROME

IMPACTED PLANS: MEDICAID, HIV SNP and HARP

SUMMARY

Effective December 1, 2023, per New York State (NYS) Medicaid guidance update, MetroPlusHealth has included coverage and reimbursement of genetic testing for Lynch Syndrome deoxyribonucleic acid (DNA) mismatch repair (MMR) gene mutations (MLH1, MSH2, MSH6, and PMS2) billed with CPT code **"81288"**.

Billing and Reimbursement Criteria

NYS Medicaid allows reimbursement for initial screening of the MLH1 and MSH2 genes. Genetic testing for Lynch syndrome mutations will be covered when one or more of the following criteria are met:

- Individuals diagnosed with colorectal cancer (CRC) under 70 years of age.
- Individuals 70 years of age or older who meet the Bethesda criteria outlined below as applicable.
- Women who were diagnosed with endometrial cancer at less than 50 years of age.
- Individuals who meet the Amsterdam II criteria.

CPT Code	Description
Effective 12/1/2023	
81288	MLH1 (MUTL HOMOLOG 1, COLON CANCER, NONPOLYPOSIS TYPE 2) (E.G., HEREDITARY NONPOLYPOSIS COLORECTAL CANCER, LYNCH SYNDROME) GENE ANALYSIS; PROMOTER METHYLATION.

Other Screening Codes	Description
81292	MLH1 (MUTL HOMOLOG 1, COLON CANCER, NONPOLYPOSIS TYPE 2) (EG, HEREDITARY NON-POLYPOSIS COLORECTAL CANCER, LYNCH SYNDROME) GENE ANALYSIS; FULL SEQUENCE ANALYSIS
81294	LH1 (MUTL HOMOLOG 1, COLON CANCER, NONPOLYPOSIS TYPE 2) (EG, HEREDITARY NON-POLYPOSIS COLORECTAL CANCER, LYNCH SYNDROME) GENE ANALYSIS; DUPLICATION/DELETION VARIANTS
81295	MSH2 (MUTS HOMOLOG 2, COLON CANCER, NONPOLYPOSIS TYPE 1) (EG, HEREDITARY NON-POLYPOSIS COLORECTAL CANCER, LYNCH SYNDROME) GENE ANALYSIS; FULL SEQUENCE ANALYSIS
81297	MSH2 (MUTS HOMOLOG 2, COLON CANCER, NONPOLYPOSIS TYPE 1) (EG, HEREDITARY NON-POLYPOSIS COLORECTAL CANCER, LYNCH SYNDROME) GENE ANALYSIS; DUPLICATION/DELETION VARIANTS
81298	MSH6 (MUTS HOMOLOG 6 YE, COLI") (EG, HEREDITARY NON-POLYPOSIS COLORECTAL CANCER, LYNCH SYNDROME) GENE ANALYSIS; FULL SEQUENCE ANALYSIS
81300	MSH6 (MUTS HOMOLOG 6 YE, COLI") (EG, HEREDITARY NON-POLYPOSIS COLORECTAL CANCER, LYNCH SYNDROME) GENE ANALYSIS; DUPLICATION/DELETION VARIANTS
81317	PMS2 (POSTMEIOTIC SEGREGATION INCREASED TWO YS. CEREVISIAE") (EG, HEREDITARY NON-POLYPOSIS COLORECTAL CANCER, LYNCH SYNDROME) GENE ANALYSIS; FULL SEQUENCE ANALYSIS
81319	PMS2 (POSTMEIOTIC SEGREGATION INCREASED TWO YS. CEREVISIAE") (EG, HEREDITARY NON-POLYPOSIS COLORECTAL CANCER, LYNCH SYNDROME) GENE ANALYSIS; DUPLICATION/DELETION VARIANTS

Thank you

MetroPlusHealth