Policy Statement

One-time genotypic or phenotypic analysis of the thiopurine methyltransferase (TPMT) enzyme may be considered medically necessary in either of the following situations:

- Patients beginning therapy with azathioprine (AZA), mercaptopurine (6-MP), or thioguanine (6-TG)
- Patients on thiopurine therapy with abnormal complete blood count (CBC) results that do not respond to dose reduction

Genotypic and/or phenotypic analysis of the TPMT enzyme is considered investigational in all other situations.

Analysis of the metabolite markers azathioprine (AZA) and mercaptopurine (6-MP), including 6-methyl-mercaptopurine ribonucleotides (6-MMRP) and 6-thioguanine nucleotides (6-TGN), is considered investigational.

Policy Guidelines

Thiopurine methyltransferase (TPMT) testing cannot substitute for complete blood count (CBC) monitoring in patients receiving thiopurines. Early drug discontinuation may be considered in patients with abnormal complete blood count results. Dosage reduction is recommended in patients with reduced TPMT activity. Alternative therapies may need to be considered for patients who have low or absent TPMT activity (homozygous for nonfunctional alleles). Accurate genotyping and phenotyping results are not possible in patients who received recent blood transfusions. TPMT genotyping and phenotyping would only need to be performed once.

Genetic Counseling

Genetic counseling is primarily aimed at patients who are at risk for inherited disorders, and experts recommend formal genetic counseling in most cases when genetic testing for an inherited condition is considered. The interpretation of the results of genetic tests and the understanding of risk factors can be very difficult and complex. Therefore, genetic counseling will assist individuals in understanding the possible benefits and harms of genetic testing, including the possible impact of the information on the individual’s family. Genetic counseling may alter the utilization of genetic testing substantially and may reduce inappropriate testing. Genetic counseling should be performed by an individual with experience and expertise in genetic medicine and genetic testing methods.

Coding

Effective April 1, 2020, there is a new CPT/PLA code specific for the NT (NUDT15 and TPMT) genotyping panel, which is a pharmacogenetic blood/saliva test recommended to guide proper dosing of thiopurines (azathioprine, mercaptopurine, and thioguanine):

- 0169U: NUDT15 (nudix hydrolase 15) and TPMT (thiopurine S-methyltransferase) (e.g., drug metabolism) gene analysis, common variants

The following CPT code is specific for the analysis of common variants of the thiopurine methyltransferase (TPMT) gene:

- 81335: TPMT (thiopurine S-methyltransferase) (e.g., drug metabolism), gene analysis, common variants (e.g., *2, *3)
The analysis of common variants of the thiopurine methyltransferase (TPMT) gene may also be reported with the following CPT code:

- **81401**: Molecular pathology procedure, Level 2 (e.g., 2-10 single nucleotide polymorphisms [SNPs], 1 methylated variant, or 1 somatic variant [typically using nonsequencing target variant analysis], or detection of a dynamic mutation disorder/triplet repeat)

There is a CPT code specific for the analysis of common variants of the nudix hydrolase 15 (NUDT15) gene:

- **81306**: NUDT15 (nudix hydrolase 15) (e.g., drug metabolism) gene analysis, common variant(s) (e.g., *2, *3, *4, *5, *6)

There are no specific CPT codes for metabolite markers of azathioprine, mercaptopurine, or thioguanine.

Genotypic, phenotypic, and metabolite markers are specialized laboratory tests typically performed in reference laboratories.

### Description

The thiopurine class of drugs—which include azathioprine (a pro-drug for mercaptopurine), mercaptopurine, and thioguanine—are used to treat a variety of diseases; however, it is recommended the use of thiopurines be limited due to a high rate of drug toxicity. Mercaptopurine and thioguanine are directly metabolized by the thiopurine S-methyltransferase (TPMT) enzyme. Susceptibility to drug toxicity is linked to the level of TPMT activity. The variation in TPMT activity has been related to three distinct TPMT variants. Pharmacogenomic analysis of TPMT status is proposed to identify patients at risk of thiopurine drug toxicity and adjust medication doses accordingly; measurement of metabolite markers has also been proposed.

### Related Policies

- **N/A**

### Benefit Application

Benefit determinations should be based in all cases on the applicable contract language. To the extent there are any conflicts between these guidelines and the contract language, the contract language will control. Please refer to the member's contract benefits in effect at the time of service to determine coverage or non-coverage of these services as it applies to an individual member.

Some state or federal mandates (e.g., Federal Employee Program [FEP]) prohibits plans from denying Food and Drug Administration (FDA)-approved technologies as investigational. In these instances, plans may have to consider the coverage eligibility of FDA-approved technologies on the basis of medical necessity alone.

### Regulatory Status

Clinical laboratories may develop and validate tests in-house and market them as a laboratory service; laboratory-developed tests must meet the general regulatory standards of the Clinical Laboratory Improvement Amendments. Several thiopurine genotype, phenotype, and metabolite tests are available under the auspices of the Clinical Laboratory Improvement Amendments. Laboratories that offer laboratory-developed tests must be licensed by the
Clinical Laboratory Improvement Amendments for high-complexity testing. To date, the U.S. Food and Drug Administration has chosen not to require any regulatory review of this test.

Prometheus, a commercial laboratory, offers thiopurine genotype, phenotype, and metabolite testing for those on thiopurine therapy. The tests are referred to as Prometheus® TPMT Genetics, Prometheus® TMPT Enzyme, and Prometheus® thiopurine metabolites, respectively. Other laboratories that offer TPMT genotyping include: Quest Diagnostics (TPMT genotype); ARUP Laboratories (TPMT DNA); Specialty Laboratories (TPMT GenoTypR®); Prevention Genetics (TPMT Deficiency via the TPMTGene); Genelex (TPMT); Fulgent Genetics (TPMT); and LabCorp (TPMT enzyme activity and genotyping).

**Rationale**

**Background**

**Thiopurines**

Thiopurines or purine analogues are immunomodulators. They include azathioprine (Imuran), mercaptopurine (6-MP; Purinethol), and thioguanine (6-TG; Tabloid). Thiopurines are used to treat malignancies, rheumatic diseases, dermatologic conditions, and inflammatory bowel disease, and are used in solid organ transplantation. They are considered an effective immunosuppressive treatment of inflammatory bowel disease, particularly in patients with corticosteroid-resistant disease. However, the use of thiopurines is limited by both long onset of action (3-4 months) and drug toxicities, which include hepatotoxicity, bone marrow suppression, pancreatitis, and allergic reactions.

**Pharmacogenomics**

Thiopurines are converted to 6-MP in vivo, where it is subsequently metabolized to 2 active metabolites: either 6-thioguanine nucleotides (6-TGN) by the inosine-5′-monophosphate dehydrogenase enzyme; or to 6-methyl-mercaptopurine ribonucleotides by the thiopurine methyltransferase (TPMT) enzyme. TPMT also converts 6-MP into an inactive metabolite, 6-methylmercaptopurine. The 6-TGN metabolites are considered cytotoxic and thus are associated with bone marrow suppression, while the 6-methyl-mercaptopurine ribonucleotides are associated with hepatotoxicity. In population studies, the activity of the TPMT enzyme has been shown to be trimodal, with 90% of subjects having high activity, 10% intermediate activity, and 0.3% with low or no activity. In patients with intermediate-to-low activity, the metabolism of 6-MP is shunted toward the inosine-5′-monophosphate dehydrogenase pathway with greater accumulation of 6-TGN; these patients are considered at risk for myelotoxicity (i.e., bone marrow suppression).

This variation in TPMT activity has been related to 3 distinct TPMT variants and has permitted the development of TPMT genotyping using a polymerase chain reaction. For example, patients with high TPMT activity are found to have 2 normal (wild-type) TPMT alleles; those with intermediate activity are heterozygous (i.e., have a variant on 1 chromosome), while those with low TPMT activity are homozygous for TPMT variants (i.e., have a variant on both chromosomes). Genetic analysis has been explored as a technique to identify patients at risk for myelotoxicity. Patients with high TPMT activity may be treated with standard doses of thiopurines, patients with intermediate TPMT activity may be initially treated with lower doses of thiopurines, while those with low TPMT activity may not be good candidates for thiopurine therapy.

TPMT activity can also be measured by phenotypic testing. Phenotyping determines the level of thiopurine nucleotides or TPMT activity in erythrocytes. Caution must be taken with phenotyping, because some co-administered drugs can influence the measurement of TPMT activity in blood, and recent blood transfusions will misrepresent a patient’s actual TPMT activity.

Prospective TPMT genotyping or phenotyping may help identify patients at increased risk of developing severe, life-threatening myelotoxicity.
The genotypic analysis of the TPMT gene is based on well-established polymerase chain reaction technology to detect 3 distinct variants. Currently, 3 alleles (TPMT*2, TPMT*3A, TPMT*3C) account for about 95% of subjects with reduced TPMT enzyme activity. Subjects homozygous for these alleles are TPMT-deficient and those heterozygous for these alleles have variable TPMT (low or intermediate) activity. A study by Hindorf and Appell (2012) addressed the concordance between TPMT genotyping and phenotyping. The investigators evaluated data from 7195 unselected and consecutive TPMT genotype and phenotype tests. The genotyping tests examined the 3 most common TPMT variants, previously noted. TPMT genotyping identified 6454 (89.7%) as TPMT wild-type, 704 (9.8%) as TPMT heterozygous, and 37 (0.005%) as TPMT homozygous. The overall agreement between genotyping and phenotyping was 95%. Genotyping alone would have misclassified 3 (8%) of 37 homozygous patients as heterozygous; these 3 subjects were found to have uncommon variants. All three had low TPMT activity. The phenotype test would have misclassified 4 (11%) of 37 of homozygous patients because they had test results above the cutoff level for low TPMT activity (<2.5 U/mL red blood cells).

**Metabolite Markers**

Monitoring of thiopurine therapy has been based on clinical assessment of response in addition to monitoring blood cell counts, liver function, and pancreatic function tests. However, there has been interest in monitoring intracellular levels of thiopurine metabolites (i.e., 6-TGN, 6-methylmercaptopurine ribonucleotides) to predict response and complications, with the ultimate aim of tailoring drug therapy to each patient.

Metabolite markers have been assessed using high-performance liquid chromatography technology. It would be optimal to assess metabolite markers in peripheral leukocytes because they reflect the status of bone marrow precursors. However, it is technically easier to measure metabolites in red blood cells than in leukocytes.

While genotyping and phenotyping of TPMT would only be performed once, metabolite markers might be tested multiple times during the course of the disease to aid in determining the initial dose and in evaluating any ongoing dosing.

**Literature Review**

The primary goal of pharmacogenomic testing and personalized medicine is to achieve better clinical outcomes compared with the standard of care. Drug response varies greatly between individuals, and genetic factors are known to play a role. However, in most cases, the genetic variation only explains a modest portion of the variance in the individual response because clinical outcomes are also affected by a wide variety of factors including alternate pathways of metabolism and patient- and disease-related factors that may affect absorption, distribution, and elimination of the drug. Therefore, a assessment of clinical utility cannot be made by a chain of evidence from clinical validity data alone. In such cases, evidence evaluation requires studies that directly demonstrate that the pharmacogenomic test alters clinical outcomes; it is not sufficient to demonstrate that the test predicts a disorder or a phenotype.

Evidence reviews assess the clinical evidence to determine whether the use of technology improves the net health outcome. Broadly defined, health outcomes are the length of life, quality of life, and a ability to function, including benefits and harms. Every clinical condition has specific outcomes that are important to patients and managing the course of that condition. Validated outcome measures are necessary to ascertain whether a condition improves or worsens; and whether the magnitude of that change is clinically significant. The net health outcome is a balance of benefits and harms.

To assess whether the evidence is sufficient to draw conclusions about the net health outcome of technology, two domains are examined: the relevance, and quality and credibility. To be relevant, studies must represent one or more intended clinical use of the technology in the intended population and compare an effective and appropriate alternative at a comparable intensity. For some conditions, the alternative will be supportive care or surveillance. The quality
and credibility of the evidence depend on study design and conduct, minimizing bias and confounding that can generate incorrect findings. The randomized controlled trial (RCT) is preferred to assess efficacy; however, in some circumstances, nonrandomized studies may be adequate. RCTs are rarely large enough or long enough to capture less common adverse events and long-term effects. Other types of studies can be used for these purposes and to assess generalizability to broader clinical populations and settings of clinical practice.

Genotype Analysis

Clinical Context and Test Purpose

The purpose of testing for thiopurine methyltransferase (TPMT) genotype markers in patients treated with thiopurines is:

- to identify individuals likely or unlikely to be at high-risk of adverse drug reactions (ADRs) from thiopurines; or
- to optimize dose selection or frequency by identifying individuals who are likely to require higher or lower doses of a drug.

The question addressed in this evidence review is: Does a genotypic analysis of TPMT gene variants improve the net health outcomes in patients who are treated with thiopurines?

The following PICOs were used to select literature to inform this review.

Patients

The relevant population of interest are patients treated with thiopurines. Thiopurines are administered for a wide variety of clinical conditions such as malignancies, rheumatic diseases, dermatologic conditions, inflammatory bowel disease (IBD), and those undergoing solid organ transplants.

Interventions

Commercial testing for TPMT genotype is available from multiple labs and companies. The genotypic analysis of the TPMT gene is based on polymerase chain reaction technology to detect three distinct variants. Currently, 3 alleles (TPMT*2, TPMT*3A, TPMT*3C) account for about 95% of subjects with reduced TPMT enzyme activity. Subjects homozygous for these alleles are TPMT-deficient and those heterozygous for these alleles have variable TPMT (low or intermediate) activity.

Thiopurines are used in a wide variety of clinical conditions and therefore may be prescribed by a wide variety of specialists such as rheumatologists, gastroenterologists, oncologists, dermatologists, and transplant surgeons (or team). Most patients are likely to be tested in an outpatient setting.

Comparators

The following practice is currently being used to treat malignancies, rheumatic diseases, dermatologic conditions, IBD, and those undergoing solid organ transplants: standard management without TPMT genotype analysis, which may be administered by a wide variety of specialists such as rheumatologists, gastroenterologists, oncologists, dermatologists, and transplant surgeons (or team). Most patients are likely to be tested in an outpatient setting.

Outcomes

Potential beneficial outcomes of interest for individuals undergoing TPMT testing are positive changes in symptoms and disease status and reduction or elimination of morbid events such as side effects from steroid use or incidence of toxicity associated with thiopurines (e.g., bone marrow toxicity, hepatotoxicity, pancreatitis, gastric intolerance, skin reaction).

The potential harmful outcomes are those resulting from a false test result. False-positive or false-negative test results can lead to under- or overtreatment with thiopurines including the potential
loss of therapeutic benefit from undertreatment or adverse events from overtreatment or possibly from an alternative treatment other than thiopurines.

Testing is typically done prior to initiation of therapy with thiopurines but may also be done during treatment with thiopurines.

**Technically Reliable**
Assessment of technical reliability focuses on specific tests and operators and requires a review of unpublished and often proprietary information. Review of specific tests, operators, and unpublished data are outside the scope of this evidence review and alternative sources exist. This evidence review focuses on the clinical validity and clinical utility.

**Clinically Valid**
A test must detect the presence or absence of a condition, the risk of developing a condition in the future, or treatment response (beneficial or adverse).

**Study Selection Criteria**
For the evaluation of the clinical validity of this test, studies that meet the following eligibility criteria were considered:
- Reported on the accuracy of the marketed version of the technology (including any algorithms used to calculate scores)
- Included a suitable reference standard (describe the reference standard)
- Patient/sample clinical characteristics were described
- Patient/sample selection criteria were described.

Several systematic reviews of studies on the diagnostic performance of TPMT genotyping have been published. Most reviews have provided ranges of diagnostic performance measures, while two also conducted meta-analyses. The most recent meta-analysis (Zur et al [2016]) included 27 studies and reported pooled genotyping sensitivity and specificity rates of 90% (95% credible interval, 79% to 99%) and 100%, respectively. Limitations to the evidence included small numbers of homozygous patients and the inability to conduct subgroup analyses by ethnicity. The incidence of TPMT variants differs by ethnicity, which may affect sensitivity and specificity estimates.

**Clinically Useful**
A test is clinically useful if the use of the results informs management decisions that improve the net health outcome of care. The net health outcome can be improved if patients receive correct therapy, or more effective therapy, or avoid unnecessary therapy, or avoid unnecessary testing.

**Direct Evidence**
Direct evidence of clinical utility is provided by studies that have compared health outcomes for patients managed with and without the test. Because these are intervention studies, the preferred evidence would be from RCTs.

The use of pharmacogenomic testing creates the possibility of tailoring a drug regimen for each patient, with the ultimate goal of attaining disease remission and eliminating steroid therapy. The preferred study design would compare patient management (e.g., drug choice) and health outcomes in patients managed with and without testing.

**Randomized Controlled Trials**
Two RCTs have compared TPMT genotype testing with no testing and empirical weight-based thiopurine dosing. In both RCTs patients with a normal enzyme and genotype started full-dose thiopurine, while those with intermediate enzymatic activity or heterozygous genotype had a 50% dose reduction. Those with low or absent enzyme activity or homozygous genotype were
not given thiopurine or were given a reduced dose at 0% to 10% of the initial dose. These three RCTs are discussed below.

Coenen et al (2015) published results of the Thiopurine response Optimization by Pharmacogenetic testing in Inflammatory bowel disease Clinics (TOPIC) trial, which randomized 761 patients with IBD across 30 centers to empirical weight-based thiopurine dosing (n=378) or genotype-guided dosing (n=405). The trial did not meet its primary endpoint of showing a statistically significant reduction in hematologic ADR among the group that received genotype-guided thiopurines dosing compared with empirical weight-based dosing. After 20 weeks, the percentage of patients with hematologic ADRs was 7.4% for genotype-based dosing and 7.9% for empirical weight-based thiopurine dosing, with a relative risk of 0.93 (95% confidence interval [CI], 0.57 to 1.52). However, among TPMT carriers, only 1 (2.6%) of 39 patients developed a hematologic ADR compared with 8 (22.9%) of 35 patients in the control group (relative risk, 0.11; 95% CI, 0.01 to 0.85). While the results of this secondary analysis were statistically significant, the event rate was low with a wide CI indicating imprecise estimates. Further, there was no statistically significant difference in clinical outcome between the groups in an intention-to-treat analysis at 20 weeks after treatment initiation (p=0.18 for Crohn’s Disease Activity Scale score; p=0.14 for ulcerative colitis). In summary, 200 patients would have to be genotyped to avoid 1 episode of a hematologic ADR (7.4% vs 7.9%; i.e., 0.5% risk difference). The number needed to treat to avoid 1 episode of a hematologic ADR would be 5 for at-risk individuals (risk difference in patients with a genetic variant, 20.3; 2.6% vs 22.9%).

Newman et al (2011), reported results of the RCT of thiopurine methyltransferase genotyping prior to azathioprine treatment (TARGET), which randomized 333 IBD patients to genotype-guided dosing or to empirical weight-based thiopurine dosing. Data were available for 322 (97%) of 333 patients at 4 months. The trial did not meet its primary endpoint of showing a statistically significant reduction in the proportion of patients stopping azathioprine(AZA) treatment due to any ADR in genotype-guided dosing arm compared with empirical weight-based dosing. The respective proportion of patients in both arms who stopped taking AZA because of an ADR was 29% (47/163) and 28% (44/159; p=0.74), respectively. The trial included few patients with non-wild-type gene variants (seven heterozygous patients in the genotyping group; two heterozygous patients, one homozygous patient in the non genotyping group) and therefore was underpowered to detect a difference of the impact of TPMT genotyping.

Observational Studies
Several prospective studies have examined variations in the efficacy of medication by patient TPMT status. In a study that involved 131 patients with IBD, Gisbert et al (2006) reported that the choice of AZA or mercaptopurine dose, based on red blood cells TPMT activity, did not prevent myelotoxicity; no patients in this study exhibited low activity. In a study from New Zealand, Gardiner et al (2008) noted that initial target doses to attain therapeutic levels in patients with IBD ranged from 1 to 3 mg/kg/d in intermediate (heterozygous) and normal (wild-type) metabolizers. This conclusion was based on an analysis of 52 patients with IBD who were started on AZA or mercaptopurine and followed for 9 months while 6-thioguanine nucleotide (6-TGN) levels and clinical status were evaluated. This study suggested that knowledge of TPMT activity could assist with initial dosing.

In a study from Europe that included 394 patients with IBD, Gisbert et al (2006) found the probability of myelotoxicity was 14.3% in the TPMT intermediate group compared with 3.5% in groups with high (wild-type) activity. The authors concluded that determining TPMT activity before initiating treatment with AZA could minimize the risk of myelotoxicity.

Section Summary: Genotype Analysis
Several systematic reviews have examined the diagnostic performance of TPMT genotyping. The most recent meta-analysis reported genotyping sensitivity and specificity rates of 90% and 100%, respectively. Two RCTs (total n=1094 patients) compared TPMT genotype testing with no testing and empirical weight-based thiopurine dosing. In these trials, only 0.17% (n=2) were
Pharmacogenomic and Metabolite Markers for Patients Treated With Thiopurines

2.04.19

Page 8 of 20

homozygous. Of the 2 RCTs, only the TOPIC trial (n=761) was adequately powered. Hematologic adverse events and treatment discontinuation were used as surrogate outcomes for the benefits of TPMT testing. There were no significant differences in either outcome based on TPMT testing and treatment discontinuation. Additionally, there was also no significant difference in clinical remission rates in these groups based on TPMT testing in the largest RCT. However, secondary analysis of individuals who were intermediate enzymatic activity (a heterozygous genotype) or low enzymatic activity (a homozygous genotype) showed that TPMT testing to guide dosing was associated with an 89% risk reduction of hematologic adverse events. In conclusion, although the risk of harm from not testing a TPMT level before initiating therapy is minimal (indicated by a large number needed to treat), in most cases there is considerable risk of harm (indicated by a small number needed to harm) in the 0.3% patients who are homozygous genotype or have low or absent TPMT enzymatic activity.

Phenotype Analysis

Clinical Context and Test Purpose

The purpose of testing for TPMT phenotype in patients treated with thiopurines is:

• to identify individuals likely or unlikely to be at high-risk of ADRs from thiopurines; or
• to optimize dose selection or frequency by identifying individuals who are likely to require higher or lower doses of a drug.

The question addressed in this evidence review is: Does a phenotypic analysis of TPMT variants improve the net health outcomes in patients who are treated with thiopurines?

The following PICOs were used to select literature to inform this review.

Patients

The relevant population of interest are patients treated with thiopurines who receive TPMT phenotype analysis. Thiopurines are administered for a wide variety of clinical conditions such as malignancies, rheumatic diseases, dermatologic conditions, IBD, and those undergoing solid organ transplants.

Interventions

Commercial testing for TPMT phenotype is available from multiple labs and companies. Thiopurines are used in a wide variety of clinical conditions and therefore may be prescribed by a wide variety of specialists such as rheumatologists, gastroenterologists, oncologists, dermatologists, and transplant surgeons (or team). Most patients are likely to be tested in an outpatient setting.

Comparators

The following practice is currently being used to treat malignancies, rheumatic diseases, dermatologic conditions, IBD, and those undergoing solid organ transplants: standard management without TPMT phenotype analysis, which may be administered by a wide variety of specialists such as rheumatologists, gastroenterologists, oncologists, dermatologists, and transplant surgeons (or team). Most patients are likely to be tested in an outpatient setting.

Outcomes

Potential beneficial outcomes of interest for individuals undergoing TPMT testing are positive changes in symptoms and disease status and reduction or elimination of morbid events such as side effects from steroid use or incidence of toxicity associated with thiopurines (e.g., bone marrow toxicity, hepatotoxicity, pancreatitis, gastric intolerance, skin reaction). The potential harmful outcomes are those resulting from a false test result. False-positive or false-negative test results can lead to under- or overtreatment with thiopurines including the potential loss of therapeutic benefit from undertreatment or adverse events from overtreatment or possibly from an alternative treatment other than thiopurines.

Reproduction without authorization from Blue Shield of California is prohibited
Testing is typically done prior to initiation of therapy with thiopurines but may also be done during treatment with thiopurines.

**Technically Reliable**
Assessment of technical reliability focuses on specific tests and operators and requires a review of unpublished and often proprietary information. Review of specific tests, operators, and unpublished data are outside the scope of this evidence review and alternative sources exist. This evidence review focuses on the clinical validity and clinical utility.

**Clinically Valid**
A test must detect the presence or absence of a condition, the risk of developing a condition in the future, or treatment response (beneficial or adverse).

**Study Selection Criteria**
The eligibility criteria for the evaluation of the clinical validity of this test are described in the first indication.

In addition to reporting on genotype testing (see indication 1), the systematic review by Zur et al (2016) reported that in 27 studies, pooled sensitivity of phenotyping to detect TPMT deficiency vs others was 75.9% (95% credible interval [CrI], 58.3% to 87.0%), and specificity was 98.9% (95% CrI, 96.3% to 100%). The pooled sensitivity of phenotype testing to deficient or intermediate TPMT activity vs others was 91.3% (95% CrI, 86.4% to 95.5%), and specificity was 92.6% (95% CrI, 86.5% to 96.6%).

**Clinically Useful**
A test is clinically useful if the use of the results informs management decisions that improve the net health outcome of care. The net health outcome can be improved if patients receive correct therapy, or more effective therapy, or avoid unnecessary therapy, or avoid unnecessary testing.

**Direct Evidence**
Direct evidence of clinical utility is provided by studies that have compared health outcomes for patients managed with and without the test. Because these are intervention studies, the preferred evidence would be from RCTs.

The use of pharmacogenomic testing creates the possibility of tailoring a drug regimen for each patient, with the ultimate goal of attaining disease remission and eliminating steroid therapy. The preferred study design would compare patient management (e.g., drug choice) and health outcomes in patients managed with and without testing.

**Randomized Controlled Trials**
One RCT, by Sayani et al (2005), assessed testing for TPMT phenotype enzymatic activity before treatment of IBD with AZA, looking specifically at cost and adverse events. The results showed that testing TPMT enzymatic activity before AZA therapy did not predict AZA-induced toxicity and reduce adverse effects.

**Section Summary: Phenotype Analysis**
One systematic review has been identified that evaluated the diagnostic performance of TPMT phenotyping, reporting high sensitivity and specificity for detecting TPMT deficiency. An RCT that assessed how testing for TPMT phenotype enzymatic activity before treatment of IBD with AZA could reduce adverse events found that such testing did not predict AZA-induced toxicity.

**Metabolite Marker Testing**
**Clinical Context and Test Purpose**
The purpose of testing for TPMT metabolite markers in patients treated with thiopurines are:
- to identify individuals likely or unlikely to be at high-risk of ADRs from thiopurines; or
to optimize dose selection or frequency by identifying individuals who are likely to require higher or lower doses of a drug.

The question addressed in this evidence review is: Does metabolite marker analysis of TPMT improve the net health outcomes in patients who are treated with thiopurines?

The following PICOs were used to select literature to inform this review.

**Patients**
The relevant population of interest are patients treated with thiopurines who receive AZA and/or 6-mercaptopurine metabolite analysis. Thiopurines are administered for a wide variety of clinical conditions such as malignancies, rheumatic diseases, dermatologic conditions, IBD, and those undergoing solid organ transplants.

**Interventions**
Commercial testing for TPMT metabolite marker (TPMT function) is available from multiple labs and companies.

Metabolite markers are measured from red blood cell samples using high-performance liquid chromatography. It would be optimal to assess metabolite markers in peripheral leukocytes because they reflect the status of bone marrow precursors; however, it is technically easier to measure metabolites in red blood cells than in leukocytes.

Thiopurines are used in a wide variety of clinical conditions and therefore may be prescribed by a wide variety of specialists such as rheumatologists, gastroenterologists, oncologists, dermatologists, and transplant surgeons (or team). Most patients are likely to be tested in an outpatient setting.

**Comparators**
The following practice is currently being used to treat malignancies, rheumatic diseases, dermatologic conditions, IBD, and those undergoing solid organ transplants: standard management without TPMT metabolite analysis, which may be administered by a wide variety of specialists such as rheumatologists, gastroenterologists, oncologists, dermatologists, and transplant surgeons (or team). Most patients are likely to be tested in an outpatient setting.

**Outcomes**
Potential beneficial outcomes of interest for individuals undergoing TPMT testing are positive changes in symptoms and disease status and reduction or elimination of morbid events such as side effects from steroid use or incidence of toxicity associated with thiopurines (e.g., bone marrow toxicity, hepatotoxicity, pancreatitis, gastric intolerance, skin reaction).

The potential harmful outcomes are those resulting from a false test result. False-positive or false-negative test results can lead to under- or overtreatment with thiopurines including the potential loss of therapeutic benefit from undertreatment or adverse events from overtreatment or possibly from an alternative treatment other than thiopurines.

Testing is typically done prior to initiation of therapy with thiopurines but may also be done during treatment with thiopurines.

**Clinically Valid**
A test must detect the presence or absence of a condition, the risk of developing a condition in the future, or treatment response (beneficial or adverse).

**Study Selection Criteria**
The eligibility criteria for the evaluation of the clinical validity of this test are described in the first indication.
One systematic review evaluating the diagnostic accuracy of metabolite testing has been identified. The review focused on the association between metabolite levels and disease remission or adverse events. In a literature search through January 2013, Konidari et al (2014) identified 15 studies (total n=1026 children with IBD), none of the studies were RCTs. Reviewers did not pool findings. Metabolite testing among the studies was inconsistent in terms of predicting clinical outcomes and assessing toxicity.

Several studies have considered the optimal therapeutic cutoff level of metabolites, and the use of metabolite levels vs ratios of metabolite levels as predictors of clinical outcomes. Two studies suggested that 235 pmol/8×10^8 is the optimal therapeutic 6-TGN cutoff, and another study suggested a cutoff of 220 pmol/8×10^8 between patients who did and did not stay in remission. Kopylov et al (2014) found that 6-methyl-mercaptopurine (6-MMP)/6-TGN ratios performed better than 6-TGN levels for predicting relapse in pediatric patients with Crohn disease.

**Clinically Useful**

A test is clinically useful if the use of the results informs management decisions that improve the net health outcome of care. The net health outcome can be improved if patients receive correct therapy, or more effective therapy, or avoid unnecessary therapy, or avoid unnecessary testing.

**Direct Evidence**

Direct evidence of clinical utility is provided by studies that have compared health outcomes for patients managed with and without the test. Because these are intervention studies, the preferred evidence would be from RCTs.

The use of thiopurine metabolite testing creates the possibility of tailoring a drug regimen for each patient, with the ultimate goal of attaining disease remission and eliminating steroid therapy. The preferred study design would compare patient management (e.g., drug choice) and health outcomes in patients managed with and without testing.

**Randomized Controlled Trials**

Sayani et al (2005) reported results of a small RCT (n=29) in which IBD patients were randomized to the TPMT assay testing (n=15) or no assay testing (n=14) prior to AZA dosing. All 14 patients who received TPMT assay were found to have normal TPMT levels and therefore commenced AZA at 2.5 mg/kg/d while the individuals in the control arm underwent an upward dose-titration protocol to a target dose of 2.5 mg/kg/d. While the trial was small and did not report power calculations, results showed that 53% (8/15) in the no assay group and 57% (8/14) in the TPMT assay group, withdrew as a result of AZA-induced adverse events.

Friedman et al (2018) conducted a multicenter RCT in which 73 patients with clinically active or steroid-dependent IBD were randomized to 2 different doses of adjunctive allopurinol with thiopurine (AZA or mercaptopurine) therapy. The purpose of the trial was to compare the efficacy of the 2 different doses of allopurinol (50 mg or 100 mg), as the thiopurine dose was modified based on metabolite testing at 4, 12, and 18 weeks. The modifications in dosing were aimed at achieving a therapeutic level of more than 260 pmol/8×10^8 red blood cells. The primary outcome was the proportion of patients in steroid-free clinical remission at 24 weeks. Tables 1 and 2 summarize the trial characteristics and results. Adverse events did not differ between the two groups.

**Table 1. Summary of Randomized Controlled Trial Characteristics**

<table>
<thead>
<tr>
<th>Study</th>
<th>Country</th>
<th>Sites</th>
<th>Dates</th>
<th>Participants</th>
<th>Interventions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sasyani (2005)</td>
<td>Canada</td>
<td>1</td>
<td>2002-2003</td>
<td>Patients with Crohn disease or ulcerative colitis, starting azathioprine</td>
<td>TPMT assay (n=14) No TPMT assay (n=15)</td>
</tr>
</tbody>
</table>
Table 2. Summary of Randomized Controlled Trial Results

<table>
<thead>
<tr>
<th>Study</th>
<th>Number (%) Withdrawing</th>
<th>Reason for Withdrawal (n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sasyani (2005)&lt;sup&gt;18&lt;/sup&gt;</td>
<td>8 (57)</td>
<td>Nausea, vomiting, fatigue, cramps, headache (5); pancreatitis (1); no therapeutic effect (2)</td>
</tr>
<tr>
<td>No TPMT Assay</td>
<td>8 (53)</td>
<td>Nausea, vomiting, fatigue, cramps, headache (5); leukopenia (1); elevated liver enzymes (2)</td>
</tr>
<tr>
<td>Achieving Steroid-Free Remission, %</td>
<td>6-TGN Concentration pmol/8´10^8</td>
<td>p</td>
</tr>
<tr>
<td>Achieving Steroid-Free Remission, %</td>
<td>6-MMP Concentration pmol/8´10^8</td>
<td>p</td>
</tr>
</tbody>
</table>

Friedman et al (2018)<sup>18</sup>

<table>
<thead>
<tr>
<th>Study</th>
<th>Number (% Withdrawing)</th>
<th>Reason for Withdrawal (n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole cohort (n=61)</td>
<td>53</td>
<td>NR</td>
</tr>
<tr>
<td>Allopurinol 50 mg group (n=34)</td>
<td>54</td>
<td>382</td>
</tr>
<tr>
<td>Allopurinol 100 mg group (n=27)</td>
<td>53</td>
<td>420</td>
</tr>
<tr>
<td></td>
<td></td>
<td>NS</td>
</tr>
<tr>
<td></td>
<td></td>
<td>435</td>
</tr>
<tr>
<td></td>
<td></td>
<td>0.03</td>
</tr>
</tbody>
</table>

NR: not reported; 6-MMP: 6-methyl-mercaptopurine; 6-TGN: 6-thioguanine nucleotide; TPMT: thiopurine methyltransferase.

The purpose of the limitations tables (see Tables 3 and 4) is to display notable limitations identified in each study. This information is synthesized as a summary of the body of evidence following each table and provides the conclusions on the sufficiency of evidence supporting the position statement. The limitations stated in these tables are specific to the current review and do not reflect a comprehensive limitations assessment.

Table 3. Relevance Limitations

<table>
<thead>
<tr>
<th>Study</th>
<th>Population&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Intervention&lt;sup&gt;b&lt;/sup&gt;</th>
<th>Comparator&lt;sup&gt;c&lt;/sup&gt;</th>
<th>Outcomes&lt;sup&gt;d&lt;/sup&gt;</th>
<th>Duration of Follow-Up&lt;sup&gt;e&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sasyani (2005)&lt;sup&gt;18&lt;/sup&gt;</td>
<td>3</td>
<td>3. Different doses of allopurinol is not the intervention of interest.</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Friedman et al (2018)&lt;sup&gt;18&lt;/sup&gt;</td>
<td>3</td>
<td>3. Different doses of allopurinol is not the intervention of interest.</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

The study limitations stated in this table are those notable in the current review; this is not a comprehensive limitations assessment.

<sup>a</sup> Population key: 1. Intended use population unclear; 2. Clinical context is unclear; 3. Study population is unclear; 4. Study population not representative of intended use.

<sup>b</sup> Intervention key: 1. Classification thresholds not defined; 2. Version used unclear; 3. Not intervention of interest.

<sup>c</sup> Comparator key: 1. Classification thresholds not defined; 2. Not compared to credible reference standard; 3. Not compared to other tests in use for same purpose.

<sup>d</sup> Outcomes key: 1. Study does not directly assess a key health outcome; 2. Evidence chain or decision model not explicated; 3. Key clinical validity outcomes not reported (sensitivity, specificity, and predictive values); 4. Reclassification of diagnostic or risk categories not reported; 5. Adverse events of the test not described (excluding minor discomforts and inconvenience of venipuncture or noninvasive tests).

<sup>e</sup> Follow-Up key: 1. Follow-up duration not sufficient with respect to natural history of disease (true-positives, true-negatives, false-positives, false-negatives cannot be determined).
### Table 4. Study Design and Conduct Limitations

<table>
<thead>
<tr>
<th>Study</th>
<th>Selection</th>
<th>Blindingb</th>
<th>Delivery of Testc</th>
<th>Selective Reportingd</th>
<th>Data Completenesse</th>
<th>Statisticalf</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sasyani (2005)20</td>
<td>3.57% and 53% withdrawal from each arm of study</td>
<td>3.25% dropout rate in 1 arm of study</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Friedman et al (2018)21</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

The study limitations stated in this table are those notable in the current review; this is not a comprehensive limitations assessment.

a Selection key: 1. Selection not described; 2. Selection not random or consecutive (i.e., convenience).
b Blinding key: 1. Not blinded to results of reference or other comparator tests.
c Test Delivery key: 1. Timing of delivery of index or reference test not described; 2. Timing of index and comparator tests not same; 3. Procedure for interpreting tests not described; 4. Expertise of evaluators not described.
e Data Completeness key: 1. Inadequate description of indeterminate and missing samples; 2. High number of samples excluded; 3. High loss to follow-up or missing data.
f Statistical key: 1. Confidence intervals and/or p values not reported; 2. Comparison with other tests not reported.

### Observational Studies

Garritsen et al (2018) measured thiopurine metabolite levels in patients with atopic dermatitis and/or chronic dermatitis during maintenance (n=32) and dose escalation (n=8). The patient population included both high and intermediate activity genotypes and 6-TGN metabolite levels varied widely, from 42 to 696 pmol/8´10^8 red blood cells. Interpretation of results is limited due to the small sample size and the heterogeneity in patient genotypes and drug doses.

Meijer et al (2017) retrospectively reviewed the charts of 24 patients with 6-MMP-induced leukocytopenia. The authors reported that patients' symptoms resolved on altering the treatment regimens. However, due to the retrospective nature of the study, the altering of treatment regimens cannot be attributed directly to metabolite testing.

Wong et al (2017) reported on the result of a post hoc analysis of the TOPIC trial to address the predictive value of 6-MMP ribonucleotide concentrations 1 week after treatment initiation for development of hepatotoxicity during the first 20 weeks of treatment. They reported that, in more than 80% of patients, hepatotoxicity could be explained by elevated 6-MMP ribonucleotide concentrations and the independent risk factors of age, sex, and body mass index, allowing personalized thiopurine treatment in IBD to prevent early failure. Placing 174 patients on a stable thiopurine dose showed that those exceeding the 6-MMP ribonucleotide threshold of 3615 pmol/8´10^8 erythrocytes were more likely to have hepatotoxicity (odds ratio, 3.8; 95% CI, 1.8 to 8.0).

Goldberg et al (2016) retrospectively reviewed medical records of patients (n=169) with IBD who were treated with thiopurines for at least 4 weeks. Metabolite levels of 6-TGN showed 52% were subtherapeutic, 34% were therapeutic, and 14% were supratherapeutic. Among patients who experienced active disease despite therapy, 86% were managed differently following metabolite testing. Clinical outcomes following the management changes were not reported.

Kennedy et al (2013) retrospectively reviewed medical records of patients who had undergone metabolite testing in South Australia. The analysis reported on 151 patients with IBD who had been taking a thiopurine for at least 4 weeks and underwent at least 1 metabolite test. The 151 patients had a total of 157 tests. Eighty (51%) of 157 tests were done because of flare or lack of medication efficacy, 18 (12%) were for adverse events, and 54 (34%) tests were routine. Forty-four (55%) of the 80 patients who had a metabolite test due to flare or lack of efficacy had
better outcomes after the test was performed. Outcomes also improved after testing for 5 (28%) of 18 patients with an adverse event to a thiopurine. For patients who had routine metabolite tests, 7 (13%) of 54 had better outcomes following testing. The rate of benefit was significantly higher in patients tested because of flare or lack of efficacy compared with those who underwent routine metabolite testing (p<0.001). Changes in patient management included medication dose adjustments, change in medication, and surgical treatment. The study lacked a control group, and thus, outcomes cannot be compared with patients managed without metabolite testing.

Smith et al (2013) retrospectively reviewed medical records of 189 patients with IBD who had 6-TGN metabolite monitoring during thiopurine treatment. When 6-TGN concentrations were below the therapeutic range (n=47), 18 of the patients were given dose increases and 2 patients were given a combination of allopurinol with AZA. When 6-TGN concentrations were above the upper limit of the therapeutic range (n=55), 14 of the patients were given dose reductions. When nonresponders (n=53) were identified, 74% underwent treatment changes including dose increases, switching to a treatment combination of allopurinol and AZA or methotrexate, or surgery. Clinical outcomes related to the management changes were not reported. Armstrong et al (2011) conducted a retrospective chart review of pediatric patients who had a poor clinical response to thiopurine medication for at least 3 months for the treatment of IBD (n=70). Testing of 6-TGN found that 32% of values were within therapeutic levels. Management was changed based on metabolite measurements in 25 (36%) of the patients (lowering dose, increasing dose, or switching to methotrexate). Clinical outcomes following the management changes were not reported.

Chain of Evidence
Indirect evidence on clinical utility rests on clinical validity. If the evidence is insufficient to demonstrate test performance, no inferences can be made about clinical utility.

The current evidence base is insufficient to construct a chain of evidence supporting the use of metabolite testing for this patient population.

Section Summary: Metabolite Marker Testing
One systematic review assessed the diagnostic accuracy of metabolite testing. The review did not pool results due to heterogeneity across studies. Results among the studies were inconsistent in terms of predicting clinical outcomes and assessing toxicity. The evidence for the use of metabolite marker testing to manage patients who are treated with thiopurines is limited to two RCTs and a number of retrospective studies. One small RCT had over 50% withdrawal rate due to the adverse effects of the treatment, limiting the interpretation of results. Another RCT used metabolite testing to adjust thiopurine doses, the purpose of the trial was to compare two different allopurinol doses. Most of the retrospective studies have described changes in management following metabolite testing, but clinical outcomes following the management changes were not reported. Without a control group in these studies, outcomes cannot be compared for patients managed without metabolite testing. It is possible that, in the absence of metabolite testing, patients who were not seeing a benefit or who were experiencing adverse events would have had their treatments adjusted without having metabolite testing.

Summary of Evidence
For individuals who are treated with thiopurines who receive TPMT genotype analysis or TPMT phenotype analysis, the evidence includes studies of diagnostic performance, systematic reviews, and RCTs. The relevant outcomes are symptoms, morbid events, and change in disease status. A large number of studies have assessed the diagnostic performance of TPMT genotyping and phenotyping tests. The most recent meta-analysis reported genotyping sensitivity and specificity of 90% and 100% respectively, and a phenotyping sensitivity and specificity of 76% and 99% respectively, for identifying patients with subnormal enzymatic activity. Three RCTs (total n=1145 patients) have compared TPMT genotype/phenotype testing with no testing and empirical weight-based thiopurine dosing. There were no significant differences in the incidence
of hematologic adverse events, treatment discontinuation rates, or clinical remission rates. However, secondary analysis of a small number of individuals who had intermediate enzymatic activity (a heterozygous genotype) or low enzymatic activity (a homozygous genotype) showed that TPMT testing to guide dosing was associated with statistically significant risk reduction in hematologic adverse events with a wide margin of error. In summary, 200 patients would have to be genotyped to avoid 1 episode of a hematologic adverse drug reaction (7.4% vs 7.9% i.e., 0.5% risk difference). The number needed to treat to avoid 1 episode of a hematologic adverse drug reaction would be 5 for at-risk individuals (risk difference in patients with a genetic variant, 20.3% 2.6% vs 22.9%). In addition, a small, inadequately powered RCT, which assessed phenotype TPMT testing, found no difference in treatment discontinuation rates due to adverse drug reactions between the two arms. The evidence is sufficient to determine that the technology results in a meaningful improvement in the net health outcome.

For individuals who are treated with thiopurines who receive AZA and/or 6-mercaptopurine metabolite analysis, the evidence includes a systematic review as well as prospective and retrospective studies. The relevant outcomes are symptoms, morbid events, and change in disease status. The systematic review, which assessed the diagnostic accuracy of metabolite testing, reported that the ability of the metabolite tests to predict clinical outcomes and toxicity was inconsistent across studies. There is insufficient evidence from prospective studies to determine whether knowledge of metabolite marker status will lead to improved outcomes (primarily improved disease control and/or less adverse drug events). Findings from studies evaluating the association between metabolite markers and clinical remission are mixed, and no prospective comparative trials have compared health outcomes in patients managed using metabolite markers with current approaches to care. The evidence is insufficient to determine the effects of technology on net health outcomes.

Supplemental Information
Practice Guidelines and Position Statements

National Comprehensive Cancer Network
National Comprehensive Cancer Network (v. 2.2019) guidelines on acute lymphoblastic leukemia state:

- “For patients receiving 6-MP [mercaptopurine] consider testing for TPMT [thiopurine methyltransferase] gene polymorphisms, particularly in patients who develop severe neutropenia after starting 6-MP.”
- “Determination of patient TPMT genotype using genomic DNA is recommended to optimize 6-MP dosing, especially in patients who experience myelosuppression at standard doses.”
- “Quantification of 6-MP metabolites can be very useful in determining whether the lack of myelosuppression is due to non-compliance or hypermetabolism.”

North American Society for Pediatric Gastroenterology, Hepatology and Nutrition
The North American Society for Pediatric Gastroenterology, Hepatology and Nutrition (2013) on inflammatory bowel disease (IBD) published consensus recommendations on the role of the TPMT enzyme and thiopurine metabolite testing in pediatric IBD. Recommendations (high and moderate) included:

1. “TPMT testing is recommended before initiation of TPs [thiopurines] to identify individuals who are homozygous recessive or have extremely low TPMT activity....
2. Individuals who are homozygous recessive or have extremely low TPMT activity should avoid use of TPs because of concerns for significant leukopenia.
3. All individuals on TPs should have routine monitoring of CBC [complete blood cell] and WBC [white blood cell] counts to evaluate for leukopenia regardless of TPMT testing results.
4. Metabolite testing can be used to determine adherence to TP therapy.
5. Metabolite testing can be used to guide dosing increases or modifications in patients with active disease....
6. Routine and repeat metabolite testing has little or no role in patients who are doing well and taking an acceptable dose of a TP.”

American Gastroenterological Association Institute
Recommendations from the American Gastroenterological Association Institute (2017) guidelines on therapeutic drug monitoring in IBD are summarized in Table 5.28,29.

Table 5. Evidence-Based Clinical Guidelines on Therapeutic Drug Monitoring in IBD

<table>
<thead>
<tr>
<th>Recommendation</th>
<th>SOR</th>
<th>QOE</th>
</tr>
</thead>
<tbody>
<tr>
<td>In adults with IBD being started on thiopurines, AGA suggests routine TPMT testing (enzymatic activity or genotype) to guide thiopurine dosing</td>
<td>Conditional</td>
<td>Low</td>
</tr>
<tr>
<td>In adults treated with thiopurines with active IBD or adverse effects thought to be due to thiopurine toxicity, AGA suggests reactive thiopurine metabolite monitoring to guide treatment changes</td>
<td>Conditional</td>
<td>Very low</td>
</tr>
<tr>
<td>In adults with quiescent IBD treated with thiopurines, AGA suggests against routine thiopurine metabolite monitoring</td>
<td>Conditional</td>
<td>Very low</td>
</tr>
</tbody>
</table>


U.S. Preventive Services Task Force Recommendations
Not applicable.

Medicare National Coverage
There is no national coverage determination. In the absence of a national coverage determination, coverage decisions are left to the discretion of local Medicare carriers.

Ongoing and Unpublished Clinical Trials
Some currently ongoing and unpublished trials that might influence this review are listed in Table 6.

Table 6. Summary of Key Trials

<table>
<thead>
<tr>
<th>NCTNo.</th>
<th>Trial Name</th>
<th>Planned Enrollment</th>
<th>Completion Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ongoing</td>
<td>Effectiveness of Thiopurine Dose Optimization by NUDT15 R139C on Reducing Thiopurine-Induced Leucopenia in Inflammatory Bowel Disease</td>
<td>400</td>
<td>Aug 2018 (ongoing; last updated May 2018)</td>
</tr>
<tr>
<td></td>
<td>PREemptive Pharmacogenomic Testing for Preventing Adverse Drug REactions (PREPARE)</td>
<td>6892</td>
<td>Dec 2019</td>
</tr>
<tr>
<td></td>
<td>A Prospective Trial to Assess Cost and Clinical Outcomes of a Clinical Pharmacogenomic Program at Eskenazi Hospital (INGenious)</td>
<td>4465</td>
<td>May 2018 (active, not recruiting; updated Aug 2019)</td>
</tr>
</tbody>
</table>

NCT: national clinical trial.

References


Documentation for Clinical Review

Please provide the following documentation:

- History and physical and/or consultation notes including:
  - Tests required
  - Purpose of testing
  - Treatment plan
- Laboratory report(s)

Post Service (in addition to the above, please include the following):

- Results/reports of tests performed

Coding

This Policy relates only to the services or supplies described herein. Benefits may vary according to product design; therefore, contract language should be reviewed before applying the terms.
of the Policy. Inclusion or exclusion of codes does not constitute or imply member coverage or provider reimbursement.

**MN/IE**

The following services may be considered medically necessary in certain instances and investigational in others. Services may be considered medically necessary when policy criteria are met. Services may be considered investigational when the policy criteria are not met or when the code describes application of a product in the position statement that is investigational.

<table>
<thead>
<tr>
<th>Type</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0169U</td>
<td>NUDT15 (nudix hydroxylase 15) and TPMT (thiopurine S-methyltransferase) (e.g., drug metabolism) gene analysis, common variants (Code effective 4/1/2020)</td>
</tr>
<tr>
<td></td>
<td>81306</td>
<td>NUDT15 (nudix hydroxylase 15) (e.g., drug metabolism) gene analysis, common variant(s) (e.g., *2, *3, *4, *5, *6)</td>
</tr>
<tr>
<td></td>
<td>81335</td>
<td>TPMT (thiopurine S-methyltransferase) (e.g., drug metabolism), gene analysis, common variants (e.g., *2, *3)</td>
</tr>
<tr>
<td></td>
<td>81401</td>
<td>Molecular Pathology Procedure Level 2</td>
</tr>
</tbody>
</table>

**Policy History**

This section provides a chronological history of the activities, updates and changes that have occurred with this Medical Policy.

<table>
<thead>
<tr>
<th>Effective Date</th>
<th>Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>12/07/2006</td>
<td>New Policy Adoption</td>
</tr>
</tbody>
</table>
| 01/07/2011     | Policy title change from Pharmacogenomic and Metabolite Markers for Patients Treated with Azathioprine (6-MP)  
Policy revision with position change |
| 02/22/2013     | Coding update                                                                                   |
| 06/30/2015     | Coding update                                                                                   |
| 02/01/2016     | Policy title change from Pharmacogenomic and Metabolite Markers in Inflammatory Bowel Disease  
Policy revision without position change |
| 02/01/2017     | Policy revision without position change                                                          |
| 02/01/2018     | Policy revision without position change                                                                 |
| 05/01/2018     | Coding update                                                                                   |
| 01/01/2019     | Policy revision without position change                                                                 |
| 02/01/2020     | Annual review. No change to policy statement. Literature review updated.                          |
| 07/01/2020     | Coding update                                                                                   |

**Definitions of Decision Determinations**

**Medically Necessary:** Services that are Medically Necessary include only those which have been established as safe and effective, are furnished under generally accepted professional...
standards to treat illness, injury or medical condition, and which, as determined by Blue Shield, are: (a) consistent with Blue Shield medical policy; (b) consistent with the symptoms or diagnosis; (c) not furnished primarily for the convenience of the patient, the attending Physician or other provider; (d) furnished at the most appropriate level which can be provided safely and effectively to the patient; and (e) not more costly than an alternative service or sequence of services at least as likely to produce equivalent therapeutic or diagnostic results as to the diagnosis or treatment of the Member’s illness, injury, or disease.

**Investigational/Experimental:** A treatment, procedure, or drug is investigational when it has not been recognized as safe and effective for use in treating the particular condition in accordance with generally accepted professional medical standards. This includes services where approval by the federal or state governmental is required prior to use, but has not yet been granted.

**Split Evaluation:** Blue Shield of California/Blue Shield of California Life & Health Insurance Company (Blue Shield) policy review can result in a split evaluation, where a treatment, procedure, or drug will be considered to be investigational for certain indications or conditions, but will be deemed safe and effective for other indications or conditions, and therefore potentially medically necessary in those instances.

## Prior Authorization Requirements (as applicable to your plan)

Within five days before the actual date of service, the provider must confirm with Blue Shield that the member's health plan coverage is still in effect. Blue Shield reserves the right to revoke an authorization prior to services being rendered based on cancellation of the member's eligibility. Final determination of benefits will be made after review of the claim for limitations or exclusions.

Questions regarding the applicability of this policy should be directed to the Prior Authorization Department at (800) 541-6652, or the Transplant Case Management Department at (800) 637-2066 ext. 3507708 or visit the provider portal at www.blueshieldca.com/provider.

Disclaimer: This medical policy is a guide in evaluating the medical necessity of a particular service or treatment. Blue Shield of California may consider published peer-reviewed scientific literature, national guidelines, and local standards of practice in developing its medical policy. Federal and state law, as well as contract language, including definitions and specific contract provisions/exclusions, take precedence over medical policy and must be considered first in determining covered services. Member contracts may differ in their benefits. Blue Shield reserves the right to review and update policies as appropriate.