

Local Coverage Determination (LCD): MoIDX: Prometheus IBD sgi Diagnostic Policy (L37313)

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Contractor Information

Contractor Name	Contract Type	Contract Number	Jurisdiction	State(s)
Noridian Healthcare Solutions, LLC	A and B MAC	02101 - MAC A	J - F	Alaska
Noridian Healthcare Solutions, LLC	A and B MAC	02102 - MAC B	J - F	Alaska
Noridian Healthcare Solutions, LLC	A and B MAC	02201 - MAC A	J - F	Idaho
Noridian Healthcare Solutions, LLC	A and B MAC	02202 - MAC B	J - F	Idaho
Noridian Healthcare Solutions, LLC	A and B MAC	02301 - MAC A	J - F	Oregon
Noridian Healthcare Solutions, LLC	A and B MAC	02302 - MAC B	J - F	Oregon
Noridian Healthcare Solutions, LLC	A and B MAC	02401 - MAC A	J - F	Washington
Noridian Healthcare Solutions, LLC	A and B MAC	02402 - MAC B	J - F	Washington
Noridian Healthcare Solutions, LLC	A and B MAC	03101 - MAC A	J - F	Arizona
Noridian Healthcare Solutions, LLC	A and B MAC	03102 - MAC B	J - F	Arizona
Noridian Healthcare Solutions, LLC	A and B MAC	03201 - MAC A	J - F	Montana
Noridian Healthcare Solutions, LLC	A and B MAC	03202 - MAC B	J - F	Montana
Noridian Healthcare Solutions, LLC	A and B MAC	03301 - MAC A	J - F	North Dakota
Noridian Healthcare Solutions, LLC	A and B MAC	03302 - MAC B	J - F	North Dakota
Noridian Healthcare Solutions, LLC	A and B MAC	03401 - MAC A	J - F	South Dakota
Noridian Healthcare Solutions, LLC	A and B MAC	03402 - MAC B	J - F	South Dakota
Noridian Healthcare Solutions, LLC	A and B MAC	03501 - MAC A	J - F	Utah
Noridian Healthcare Solutions, LLC	A and B MAC	03502 - MAC B	J - F	Utah
Noridian Healthcare Solutions, LLC	A and B MAC	03601 - MAC A	J - F	Wyoming
Noridian Healthcare Solutions, LLC	A and B MAC	03602 - MAC B	J - F	Wyoming

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LCD Information

Document Information

LCD ID L37313	Original Effective Date For services performed on or after 01/30/2018
LCD Title MoIDX: Prometheus IBD sgi Diagnostic Policy	Revision Effective Date For services performed on or after 01/30/2018
Proposed LCD in Comment Period N/A	Revision Ending Date N/A
Source Proposed LCD DL37313	Retirement Date N/A
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	Notice Period End Date 01/29/2018

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CMS National Coverage Policy

Title XVIII of the Social Security Act, §1862(a)(1)(A). Allows coverage and payment for only those services that are considered to be reasonable and necessary.

Title XVIII of the Social Security Act, §1833(e). Prohibits Medicare payment for any claim which lacks the necessary information to process the claim.

42 Code of Federal Regulations (CFR) 410.32(a). Diagnostic x-ray tests, diagnostic laboratory tests, and other diagnostic tests: Conditions.

CMS On-Line Manual, Publication 100-02, *Medicare Benefit Policy Manual*, Chapter 15, §§80.0, 80.1.1, 80.2. Clinical Laboratory services.

CMS Internet-Only Manuals, Publication 100-04, *Medicare Claims Processing Manual*, Chapter 16, §50.5 Jurisdiction of Laboratory Claims, 60.12 Independent Laboratory Specimen Drawing, 60.2. Travel Allowance.

CMS Internet Online Manual Pub. 100-04 (*Medicare Claims Processing Manual*), Chapter 23 (Section 10) "Reporting ICD Diagnosis and Procedure Codes".

Coverage Guidance

Coverage Indications, Limitations, and/or Medical Necessity

This is a non-coverage policy for the Prometheus IBD sgi Diagnostic test. The intended use of this test is to aid healthcare providers in the differentiating inflammatory bowel disease (IBD) vs non-IBD, and Crohn's disease (CD) vs UC in a comprehensive blood test. The test includes nine serological markers: ASCA IgA, ASCA IgG, anti-OmpC IgA, anti-CBir1 IgG, anti-A4 Fla2 IgG, anti-FlaX IgG, IBD-specific pANCA auto-antibody, IBD-specific pANCA IFA (perinuclear pattern), IBD-specific pANCA IFA DNase Sensitivity; four genetic immune response markers (SNPs): ATG16L1, STAT3, NKX2-3, and ECM1; and five inflammatory biomarkers: ICAM-1, VCAM-1, VEGF, CRP and SSA. A proprietary Smart Diagnostic Algorithm interprets patterns among the multiple assay values to produce an IBD score. The test results are reported as "consistent with IBD" (consistent with UC; consistent with CD, or inconclusive for UC vs CD) or "not consistent with IBD". In addition to the algorithmic test interpretation, the results of the 17 biomarkers are also individually reported.

Summary of Evidence

CD and UC represent the two main forms of idiopathic chronic IBD. While the etiology remains idiopathic, evidence suggests that the ongoing inflammation in IBD results from persistent overly aggressive inflammatory responses to a subset of commensal microorganisms in a genetically susceptible host with exposure to environmental triggers. CD is characterized by discontinuous, transmural regions of intestinal inflammation most frequently involving the terminal ileum and colon, but can affect any part of the gastrointestinal tract, with symptoms of abdominal pain, weight loss and variable degrees of diarrhea, and complications of intestinal fibrosis, strictures and fistula formation. In contrast, UC is limited to the mucosa and submucosa of the colon, with particular involvement of the rectum. Classic symptoms of active UC include diarrhea, hematochezia, tenesmus and defecatory urgency. Extra intestinal manifestations of IBD occur in up to 25% of patients. Joints, skin and eyes may be affected. In both CD and UC, disease activity is typically relapsing and remitting, although the disease course of CD is typically progressive. Although UC and CD can usually be differentiated on the basis of clinical, radiographic, endoscopic, and histologic findings, these conditions can be difficult to distinguish in about 10% to 15% of IBD patients.

Evolution of IBD Testing

In the mid-2000s, two serologic markers – anti-*Saccharomyces cerevisiae* antibodies (ASCA) and perinuclear anti-neutrophil cytoplasmic antibodies (pANCA) – were used to diagnose IBD, and distinguish between UC and CD. Studies had shown that patients with CD had significantly higher ASCA antibodies than did controls or patients with UC. The reason CD patients have both IgA and IgG-ASCA is unclear. Overall, the sensitivity for either IgA or IgG-ASCA is in the range of 55% with specificity of about 90%. On the other hand, pANCA, a true autoantibody, was observed to be associated with colonic forms of IBD, particularly UC, with a sensitivity of approximately 60-70%. However, these pANCA-positive CD patients typically have a clinical phenotype resembling left-sided UC, so pANCA detection alone is of little value in distinguishing between UC and Crohn's colitis.

A second generation IBD panel (IBD First Step®) was marketed by Prometheus Laboratories (2000) consisting of more sensitive ASCA and pANCA assays and the addition of a second microbial antigen, OmpC. Anti-OmpC was added to increase the sensitivity for CD. Subsequently, a third generation serology panel (IBD Serology 7) was offered by Prometheus Laboratories in 2006. The panel is composed of the following markers: ASCA-IgA, ASCA-IgG, anti-OmpC-IgA, anti-CBir1-IgA, and three ANCA tests: pANCA, ANCA-IgG and DNase-sensitive pANCA. The Smart Diagnostic Algorithm analyzes and correlated test results with patterns known to the database to be associated with IBD. It supposed can predict an IBD diagnosis even when all 7 of the parameters of the IBD Serology 7 panel would be considered normal on the basis of the reference ranges provided.¹ It was reported to identify another 20% or more of otherwise seronegative CD patients.² The IBD Serology 7 panel has a sensitivity of 93%, specificity of 95% and positive predictive value of 96% in population prevalence of 59% according to Prometheus. A positive anti-CBir1 can additionally help distinguish between UC and CD in pANCA positive patients. However, in a comparison study evaluating the predictive IBD Serology 7 with routine blood test (IgA-ASCA, IgG-ASCA) in a pediatric population referred for initial evaluation of suspected IBD, the sensitivity, specificity, positive predictive value, negative predictive value, and k value for the serologic panel was 67%, 76%, 63%, 79% and 47%. The anti-flagellin antibody assay had sensitivity of 50% and specificity of 53%. Despite the inclusion of anti-flagellin in the IBD7 panel, the IBD7 panel had lower predictive values compared with routine laboratory tests in pediatric screening for IBD.³

Concern has been raised about serologic testing for IBD because the data evaluating the role of serologic testing were obtained in individuals with a known diagnosis of either CD or UC. In many of these studies, the controls were normal healthy individuals. The use of the Smart Diagnostic Algorithm based on pattern recognition has not been published in a peer-reviewed journal. Similarly, the characteristics of the validation cohort (age, gender, race, etc.) are not known or whether any of these patient characteristics affect serologic markers. However, the greatest uncertainty pertains to the precise role for serologic testing in the diagnosis of IBD patients. Austin, et al¹ state that "while there are no prospectively validated data on the accuracy of IBD serologic testing in patients with suspected IBD, the presence of positive serologic markers likely does increase the probability that the person has IBD compared with the general population". However, they note that when a physician has a reasonable index of suspicion for IBD, more definitive imaging and endoscopic studies are required to confirm or refute the diagnosis and plan treatment, regardless of the serologic results. When the physician has a low index of suspicion for IBD, a positive serologic test is likely to result in unnecessary evaluation, and a negative serologic test only adds additional expense without benefit. These authors specify that further research is required to develop the evidence that is necessary for rational use of serologic testing.

The American College of Gastroenterology, in its "Ulcerative Colitis Practice Guidelines in Adults"⁴ specifies that serologic testing (ANCA/ASCA) may be useful in the occasional patient in whom no other clinical or pathologic features allow a differential diagnosis between UC and CD. The AGA states that serologic testing may have important consequences in terms of counseling, prognosis, and the choice of medical and surgical therapies.⁵ Additionally, serological studies evaluating antibodies against *Saccharomyces cerevisiae*, antineutrophil

cytoplasmic antibodies, antibodies directed against CBir1, OmpC are evolving to provide adjunctive support for the diagnosis of CD but are not sufficiently sensitive or specific to be recommended for use as a screening tools.

The fourth iteration of Prometheus' IBD testing, IBD sgi Diagnostic test, combines serologic (n=8), genetic (n=4) and inflammatory biomarkers (n=5). In addition to the 7 serologic tests in the IBD Serology 7, two additional serologic markers: anti-Fla-X and anti-A4-FL2; four genetic markers: ATG16L1, ECM1, NKX2.3 and STAT3; and four inflammatory markers: VEGF, ICAM and VCAM, CRP and SAA are marked to increase the discriminatory ability of the assay to be an adjunct in the diagnosis of UC vs CD. The IBD sgi Diagnostic™ product monograph⁶ includes an extensive bibliography that documents associations of the 17 component markers, individually and in combination, with UC and/or CD. Development and performance characteristics of the 17-marker panel are described without citation, and it is unclear what standard criterion was used for diagnosis. Overall sensitivity for IBD, UC, and CD is reported as 74%, 98%, and 89%, respectively; specificity is reported as 90%, 84%, and 81%, respectively; receiver operating characteristic (ROC) analysis showed greater discrimination with the 17-marker panel (area under the curve [AUC], 0.871) compared with any individual marker (greatest AUC=0.690 for IgA anti-Saccharomyces cerevisiae antibodies [ASCA]). Test performance characteristics for distinguishing UC from CD were not provided.

In a 2012 review of the monograph, Shirts et al⁷ observed that serologic tests for ASCA-IgA, ASCA-IgG, and atypical perinuclear anti-neutrophil cytoplasmic antibody are standard of care in the diagnostic workup of IBD although not all investigators include these tests in recommended diagnostic strategies. These 3 markers are included in the 17-marker panel. Based on a meta-analysis of 60 studies (total N=11,608), pooled sensitivity and specificity of the 3-test panel were 63% and 93%, respectively, for diagnosing IBD. Because the product monograph does not include a comparison of the 17-marker panel with the 3- marker panel, incremental improvement in diagnosis with the 17-marker panel is unknown. Shirts et al calculated an AUC for the 3-marker panel of 0.899.

Analysis of Evidence (Rationale for Determination)

Level of Evidence

Quality: Poor

Strength: Moderate

Weight: Moderate

Although manufacturer data supports clinical validity of the test for diagnosing IBD, this evidence is insufficient to support an indirect chain of evidence for clinical utility due to lack of details about study methodology and lack of replication of the findings. For distinguishing UC from CD, clinical validity has not been established. No studies examining the clinical utility of IBD sgi Diagnostic™ have been identified. Furthermore, there are no US Preventive Services Task Force (USPSTF) recommendations for genetic or molecular testing for inflammatory bowel diseases, and no recommendations for multi-marker panels that include genetic tests to facilitate diagnosis or prognosis of CD or UC.^{4, 5} Consequently, this assay does not meet Medicare's reasonable and necessary criteria for coverage. Additionally, each of the individual components that comprise this assay, except ASCA-IgA, ASCA-IgG, and atypical perinuclear anti-neutrophil cytoplasmic antibody, are additionally non-covered for the diagnosis of IBD.

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Coding Information

Bill Type Codes:

Contractors may specify Bill Types to help providers identify those Bill Types typically used to report this service. Absence of a Bill Type does not guarantee that the policy does not apply to that Bill Type. Complete absence of all Bill Types indicates that coverage is not influenced by Bill Type and the policy should be assumed to apply equally to all claims.

N/A

Revenue Codes:

Contractors may specify Revenue Codes to help providers identify those Revenue Codes typically used to report this service. In most instances Revenue Codes are purely advisory. Unless specified in the policy, services reported under other Revenue Codes are equally subject to this coverage determination. Complete absence of all Revenue Codes indicates that coverage is not influenced by Revenue Code and the policy should be assumed to apply equally to all Revenue Codes.

N/A

CPT/HCPCS Codes

Group 1 Paragraph: N/A

Group 1 Codes:

81479 UNLISTED MOLECULAR PATHOLOGY PROCEDURE

82397 CHEMILUMINESCENT ASSAY

83520 IMMUNOASSAY FOR ANALYTE OTHER THAN INFECTIOUS AGENT ANTIBODY OR INFECTIOUS AGENT ANTIGEN; QUANTITATIVE, NOT OTHERWISE SPECIFIED

86140 C-REACTIVE PROTEIN;

ICD-10 Codes that Support Medical Necessity

Group 1 Paragraph: N/A

Group 1 Codes:

ICD-10 Codes Description

XX000 Not Applicable

ICD-10 Codes that DO NOT Support Medical Necessity N/A

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General Information

Associated Information

N/A

Sources of Information

N/A

Bibliography

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6. <https://www.prometheuslabs.com/Resources/IBDDiagnostic/IBD-sgi-Monograph.pdf>
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Revision History Information

Revision History Date	Revision History Number	Revision History Explanation	Reason(s) for Change
01/30/2018	R1	The 5th biomarker, CRP, is added to the listing of biomarkers in the following sentence under Coverage Indications, Limitations and/or Medical Necessity: "...and five inflammatory biomarkers: ICAM-1, VCAM-1, VEGF, CRP and SSA."	<ul style="list-style-type: none">Creation of Uniform LCDs With Other MAC Jurisdiction

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Associated Documents

Attachments N/A

Related Local Coverage Documents Article(s) [A55779 - Response to Comments: MoIDX: Prometheus IBD sgi Diagnostic Policy](#) LCD(s) [DL37313 - MoIDX: Prometheus IBD sgi Diagnostic Policy](#)

Related National Coverage Documents N/A

Public Version(s) Updated on 04/03/2018 with effective dates 01/30/2018 - N/A [Updated on 12/01/2017 with effective dates 01/30/2018 - N/A](#) [Back to Top](#)

Keywords

- 81479
- 82397
- 83520
- 86140
- Prometheus IBD sgi
- MoIDX
- inflammatory
- bowel
- Crohn's
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