Quality ID #337: Psoriasis: Tuberculosis (TB) Prevention for Patients with Psoriasis, Psoriatic Arthritis and Rheumatoid Arthritis on a Biological Immune Response Modifier – National Quality Strategy Domain: Effective Clinical Care

- Meaningful Measure Area: Preventive Care

## 2020 COLLECTION TYPE: MIPS CLINICAL QUALITY MEASURES (CQMS)

# MEASURE TYPE:

Process

# **DESCRIPTION:**

Percentage of patients, regardless of age, with psoriasis, psoriatic arthritis and/or rheumatoid arthritis on a biological immune response modifier whose providers are ensuring active tuberculosis prevention either through negative standard tuberculosis screening tests or are reviewing the patient's history to determine if they have had appropriate management for a recent or prior positive test

## **INSTRUCTIONS:**

This measure is to be submitted a minimum of <u>once per performance period</u> for patients with psoriasis and/or psoriatic arthritis and/or rheumatoid arthritis seen during the performance period. This measure may be submitted by Merit-based Incentive Payment System (MIPS) eligible clinicians who perform the quality actions described in the measure based on the services provided and the measure-specific denominator coding.

## Measure Submission Type:

Measure data may be submitted by individual MIPS eligible clinicians, groups, or third party intermediaries. The listed denominator criteria are used to identify the intended patient population. The numerator options included in this specification are used to submit the quality actions as allowed by the measure. The quality-data codes listed do not need to be submitted by MIPS eligible clinicians, groups, or third party intermediaries that utilize this modality for submissions; however, these codes may be submitted for those third party intermediaries that utilize Medicare Part B claims data. For more information regarding Application Programming Interface (API), please refer to the Quality Payment Program (QPP) website.

## **DENOMINATOR:**

All patients, regardless of age, with a diagnosis of psoriasis and/or psoriatic arthritis and/or rheumatoid arthritis who are on a biologic immune response modifier

## **Definition:**

## Biologic Immune Response Modifier -

- 1. TNF-alpha inhibitors, to include, but not limited to Infliximab (Remicade), Adalimumab (Humira), Etanercept (Enbrel), or Golimumab (Simponi), Certolizumab (Cimzia).
- 2. Inhibitors of IL-12 and/or IL-23 or their receptors to include but not limited to Ustekinumab(Stelara).
- 3. B7 inhibitors, to include but not limited to Abatacept (Orencia).
- 4. Inhibitors of IL-17 family members or their receptors.

**DENOMINATOR NOTE:** A patient would be considered denominator eligible for Measure #337 for submission purposes, if the patient meets the denominator criteria with diagnosis of psoriasis or psoriatic arthritis or rheumatoid arthritis AND is on a biologic immune response modifier PRESCRIBED BY THE PROVIDER BEING EVALUATED FOR THE MEASURE.

\*Signifies that this CPT Category I code is a non-covered service under the Medicare Part B Physician Fee Schedule (PFS). These non-covered services should be counted in the denominator population for MIPS CQMs.

Examples of Applicable Medications				
Brand Name	Chemical Name	Mechanism of Action/ Type of Biologic		
Orencia	abatacept	Blocks T cell activation		
Rituxan	rituximab	CD 20 directed antibody		
Kineret	anakinra	Interleukin (IL)-1 inhibitors		
Actemra	tocilizumab	IL-6 inhibitor		
Kevzara	Sarilumab	IL-6 inhibitor		
Stelara	ustekinumab	IL-12 and IL-23 inhibitors		
Cosentyx	secukinumab	IL-17 inhibitors		
Siliq	brodalumab	IL-17 inhibitors		
Taltz	ixekizumab	IL-17 inhibitors		
Tremfya	guselkumab	IL-23 inhibitors		
Skyrizi	risankizumab-rzaa	IL-23 inhibitors		
Ilumya	Tildrakizumab	Selective (IL)-23p19 inhibitor		
Amjevita	adalimumab-atto	TNF-alpha inhibitors		
Cimzia	certolizumab	TNF-alpha inhibitors		
Cyltezo	adalimumab-adbm	TNF-alpha inhibitors		
Enbrel	etanercept	TNF-alpha inhibitors		
Erelzi	etanercept-szzs	TNF-alpha inhibitors		
Humira	adalimumab	TNF-alpha inhibitors		
Inflectra	infliximab-dyyb	TNF-alpha inhibitors		
Remicade	infliximab	TNF-alpha inhibitors		
Renflexis	infliximab-abda	TNF-alpha inhibitors		
Simponi and Simponi Aria	golimumab	TNF-alpha inhibitors		

# Denominator Criteria (Eligible Cases):

All patients, regardless of age

<u>and</u>

Diagnosis for psoriasis (ICD-10-CM): L40.0, L40.1, L40.2, L40.3, L40.4, L40.8, L40.9 OR

Diagnosis for psoriatic arthritis (ICD-10-CM): L40.50, L40.51, L40.52, L40.53, L40.54, L40.59 OR

Diagnosis for rheumatoid arthritis (ICD-10-CM): M05.10, M05.111, M05.112, M05.119, M05.121, M05.122, M05.129, M05.131, M05.132, M05.139, M05.141, M05.142, M05.149, M05.151, M05.152, M05.159, M05.161, M05.162, M05.169, M05.171, M05.172, M05.179, M05.19, M05.20, M05.211, M05.212, M05.219, M05.221, M05.222, M05.229, M05.231, M05.232, M05.239, M05.241, M05.242, M05.249, M05.251, M05.252, M05.259, M05.261, M05.262, M05.269, M05.271, M05.272, M05.279, M05.29, M05.30, M05.311, M05.312, M05.319, M05.321, M05.322, M05.329, M05.331, M05.332, M05.339, M05.341, M05.342, M05.349, M05.351, M05.352, M05.359, M05.361, M05.362, M05.369, M05.371, M05.372, M05.379, M05.39, M05.40, M05.411, M05.412, M05.419, M05.421, M05.422, M05.429, M05.431, M05.432, M05.439, M05.441, M05.442, M05.449, M05.451, M05.452, M05.459, M05.461, M05.462, M05.469, M05.471, M05.472, M05.479, M05.49, M05.50, M05.511, M05.512, M05.519, M05.521, M05.522, M05.529, M05.531, M05.532, M05.539, M05.541, M05.542, M05.549, M05.551, M05.552, M05.559, M05.561, M05.562, M05.569, M05.571, M05.572, M05.579, M05.59, M05.60, M05.611, M05.612, M05.619, M05.621, M05.622, M05.629, M05.631, M05.632, M05.639, M05.641, M05.642, M05.649, M05.651, M05.652, M05.659, M05.661, M05.662, M05.669, M05.671, M05.672, M05.679, M05.69, M05.70, M05.711, M05.712, M05.719, M05.721, M05.722, M05.729, M05.731, M05.732, M05.739, M05.741, M05.742, M05.749, M05.751, M05.752, M05.759, M05.761, M05.762, M05.769, M05.771, M05.772, M05.779, M05.79, M05.80, M05.811, M05.812, M05.819, M05.821, M05.822, M05.829, M05.831, M05.832, M05.839, M05.841, M05.842, M05.849, M05.851, M05.852, M05.859, M05.861, M05.862, M05.869, M05.871, M05.872, M05.879, M05.89, M05.9, M06.00, M06.011, M06.012, M06.019, M06.021, M06.022, M06.029, M06.031,

M06.032, M06.039, M06.041, M06.042, M06.049, M06.051, M06.052, M06.059, M06.061, M06.062, M06.069, M06.071, M06.072, M06.079, M06.08, M06.09, M06.1, M06.20, M06.211, M06.212, M06.219, M06.221, M06.222, M06.229, M06.231, M06.232, M06.239, M06.241, M06.242, M06.249, M06.251, M06.252, M06.259, M06.261, M06.262, M06.269, M06.271, M06.272, M06.279, M06.28, M06.29, M06.30, M06.311, M06.312, M06.319, M06.321, M06.322, M06.329, M06.331, M06.332, M06.339, M06.341, M06.342, M06.349, M06.351, M06.352, M06.359, M06.361, M06.362, M06.369, M06.371, M06.372, M06.379, M06.38, M06.39, M06.80, M06.811, M06.812, M06.819, M06.821, M06.822, M06.829, M06.831, M06.832, M06.839, M06.841, M06.842, M06.849, M06.851, M06.851, M06.852, M06.861, M06.862, M06.869, M06.871, M06.872, M06.879, M06.87, M06.89, M06.89, M06.89, M06.871, M06.872, M06.879, M06.88, M06.89, M06.89, M06.89, M06.871, M06.872, M06.879, M06.88, M06.89, M06.89, M06.89, M06.871, M06.872, M06.879, M06.88, M06.89, M06.89, M06.89, M06.89, M06.89, M06.89, M06.871, M06.872, M06.879, M06.88, M06.89, M06.89, M06.89, M06.89, M06.871, M06.872, M06.879, M06.88, M06.89, M06.89, M06.89, M06.871, M06.872, M06.879, M06.88, M06.89, M06.89, M06.89, M06.89, M06.871, M06.872, M06.879, M06.88, M06.89, M06.8

#### <u>and</u>

Patient encounter during performance period (CPT or HCPCS): 99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99241\*, 99242\*, 99243\*, 99244\*, 99245\*, G0402 <u>WITHOUT</u> Telehealth Modifier: GQ, GT, 95, POS 02 <u>AND</u> Biologic immune response modifier prescribed: G9506

#### NUMERATOR:

Patients who have a documented negative TB screening or have documentation of the management of a positive TB screening test with no evidence of active tuberculosis, confirmed through use of radiographic imaging (i.e., chest x-ray, CT) prior to treatment with a biologic immune response modifier

#### Definition:

Documentation - To satisfy this measure, documentation will incorporate the following:

- Initials of the clinician that prescribed the biologic
- Initials of clinician that administered the TB test
- Initials of the clinician that performed the skin test evaluating the results note in the medical record that results are negative OR that results are positive and is confirmed via results from radiographic imaging (chest x-ray, CT)

If TB test is administered by a provider other than the biologic prescriber, confirmation of the above mentioned must be obtained via letter or other written means from the administering provider.

**NUMERATOR NOTE:** For Denominator Exception(s): patients are ineligible for this measure if at the time of follow-up there are patient reason(s) for not documenting TB test results (e.g. patient did not return for Mantoux (PPD) skin test evaluation).

	Numerator Options:	
00	Performance Met:	Documentation of negative or managed positive TB screen with further evidence that TB is not active prior to treatment with a biologic immune response modifier (G9359)
<u>OR</u>	Denominator Exception:	Documentation of patient reasons(s) for not having records of negative or managed positive TB screen (e.g., patient does not return for Mantoux (PPD) skin test evaluation) (G9932)
<u>OR</u>	Performance Not Met:	No documentation of negative or managed positive TB screen (G9360)

# RATIONALE:

Pretreatment test for latent TB is essential for appropriate patient care and safety. Prior to the initiation of a biologic immune response modifier, it's important to know the TB status as to prevent tuberculosis reactivation or any type of harm to the patient.

# **CLINICAL RECOMMENDATION STATEMENTS:**

When planning to initiate treatment of a patient with psoriasis with a biologic it is important to obtain an age appropriate history and physical examination along with an updated medication list. In addition, it is also important to obtain a reliable set of baseline laboratory studies that will allow the clinician to detect and be aware of any underlying conditions or risk factors. This is particularly important because after patients have been initiated on a biologic treatment, they are likely to be treated with other biologics or systemic therapies and it may be useful to have reliable baseline laboratory studies. Tuberculosis testing (PPD) should be performed on all patients who will be treated with TNF inhibitors as there are reports of tuberculosis reactivation in patients treated with this class of drug (AAD).

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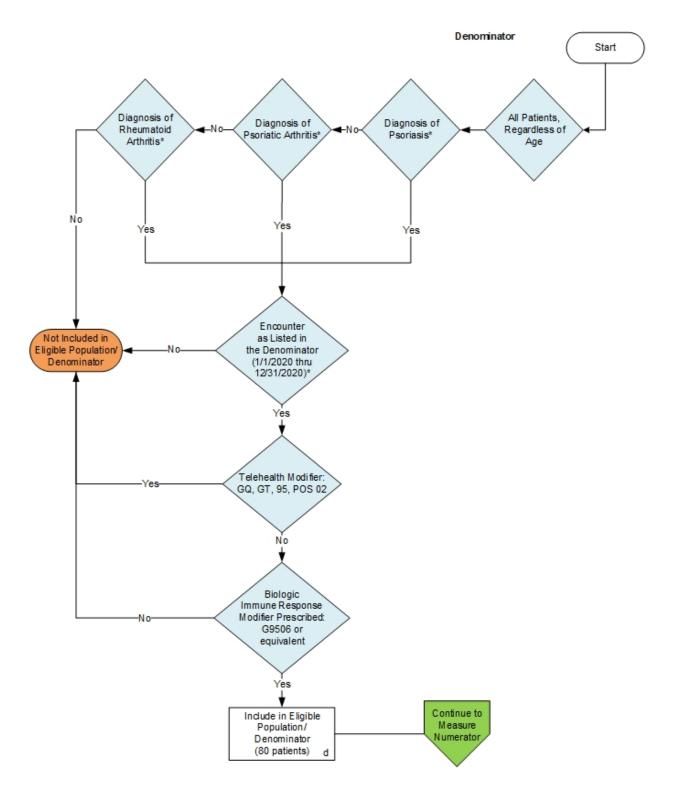
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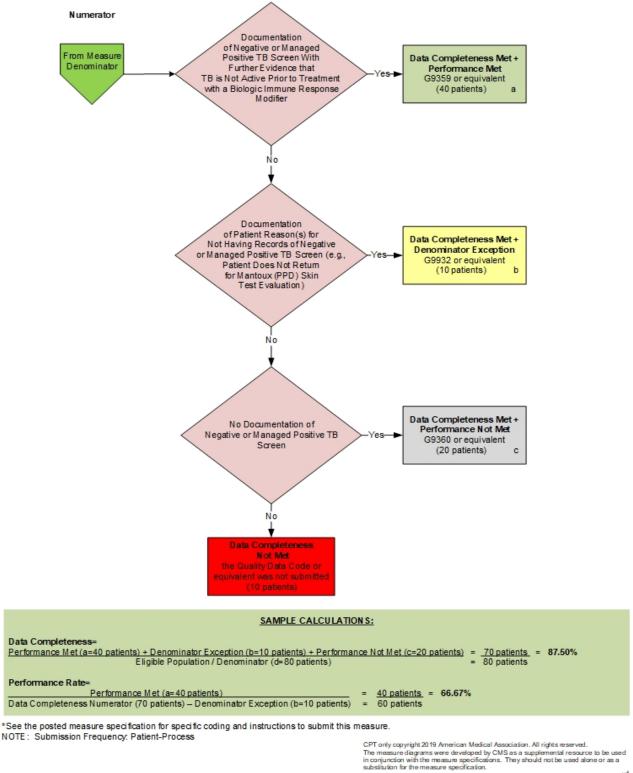
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# 2020 Clinical Quality Measure Flow for Quality ID #337: Psoriasis: Tuberculosis (TB) Prevention for Patients with Psoriasis, Psoriatic Arthritis and Rheumatoid Arthritis on a Biological Immune Response Modifier

Disclaimer: Refer to the measure specification for specific coding and instructions to submit this measure.





# 2020 Clinical Quality Narrative Flow for Quality ID #337: Psoriasis: Tuberculosis (TB) Prevention for Patients with Psoriasis, Psoriatic Arthritis and Rheumatoid Arthritis on a Biological Immune Response Modifier

**Disclaimer:** Refer to the measure specification for specific coding and instructions to submit this measure.

- 1. Start with Denominator
- 2. All Patients, Regardless of Age
- 3. Check Diagnosis of Psoriasis:
  - a. If Diagnosis of Psoriasis equals No, proceed to check Diagnosis of Psoriatic Arthritis.
  - b. If Diagnosis of Psoriasis equals Yes, proceed to check Encounter.
- 4. Check Diagnosis of Psoriatic Arthritis:
  - a. If Diagnosis of Psoriatic Arthritis equals No, proceed to check Diagnosis of Rheumatoid Arthritis.
  - b. If Diagnosis of Psoriatic Arthritis equals Yes, proceed to check Encounter
- 5. Check Diagnosis of Rheumatoid Arthritis:
  - a. If Diagnosis of Rheumatoid Arthritis equals No, do not include in Eligible Population. Stop Processing.
  - b. If Diagnosis of Rheumatoid Arthritis equals Yes, proceed to check Encounter.
- 6. Check Encounter Performed:
  - a. If Encounter as Listed in the Denominator equals No, do not include in Eligible Population. Stop Processing.
  - b. If Encounter as Listed in the Denominator equals Yes, proceed to check Telehealth Modifier.
- 7. Check Telehealth Modifier:
  - a. If Telehealth Modifier equals Yes, do not include in Eligible Population. Stop Processing
  - b. If Telehealth Modifier equals No, proceed to check Biologic Immune Response Modifier Prescribed.
- 8. Check Biologic Immune Response Modifier Prescribed:
  - a. If Biologic Immune Response Modifier Prescribed equals No, do not include in Eligible Population. Stop Processing.
  - b. If Biologic Immune Response Modifier Prescribed equals Yes, include in Eligible Population.
- 9. Denominator Population:
  - a. Eligible population or Denominator is all Eligible Patients in the Denominator. Denominator is represented as Denominator in the Sample Calculation listed at the end of this document. Letter d equals 80 patients in the Sample Calculation.
- 10. Start Numerator

- 11. Check Documentation of Negative or Managed Positive TB Screen With Further Evidence that TB is Not Active prior to treatment with a biologic immune response modifier
  - a. If Documentation of Negative or Managed Positive TB Screen With Further Evidence that TB is Not Active prior to treatment with a biologic immune response modifier equals Yes, include in Data Completeness Met and PerformanceMet.
  - b. Data Completeness Met and Performance Met letter is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter a equals 40 patients in the Sample Calculation.
  - c. If Documentation of Negative or Managed Positive TB Screen With Further Evidence that TB is Not Active prior to treatment with a biologic immune response modifier equals No, proceed to check Documentation of Patient Reasons(s) for Not Having Records of Negative or Managed Positive TB Screen (e.g., Patient Does Not Return for Mantoux (PPD) Skin Test Evaluation).
- 12. Check Documentation of Patient Reasons(s) for Not Having Records of Negative or Managed Positive TB Screen (e.g., Patient Does Not Return for Mantoux (PPD) Skin Test Evaluation):
  - a. If Documentation of Patient Reasons(s) for Not Having Records of Negative or Managed Positive TB Screen (e.g., Patient Does Not Return for Mantoux (PPD) Skin Test Evaluation) equals Yes, include in Data Completeness Met and Denominator Exception.
  - b. Data Completeness Met and Denominator Exception letter is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter b equals 10 patients in the Sample Calculation.
  - c. If Documentation of Patient Reasons(s) for Not Having Records of Negative or Managed Positive TB Screen (e.g., Patient Does Not Return for Mantoux (PPD) Skin Test Evaluation) equals No, proceed to check No Documentation of Negative or Managed Positive TB Screen.
- 13. Check No Documentation of Negative or Managed Positive TB Screen:
  - a. If No Documentation of Negative or Managed Positive TB Screen equals Yes, include in Data Completeness Met and Performance Not Met.
  - b. Data Completeness Met and Performance Not Met letter is represented in the Data Completeness in the Sample Calculation listed at the end of this document. Letter c equals 20 patients in the Sample Calculation.
  - c. If No Documentation of Negative or Managed Positive TB Screen equals No, proceed to check Data Completeness Not Met.
- 14. Check Data Completeness Not Met:
  - a. If Data Completeness Not Met, the Quality Data Code or equivalent was not submitted. 10 patients have been subtracted from the Data Completeness Numerator in the Sample Calculation.

SAMPLE CALCULATIONS:				
Data Completeness=    Performance Met (a=40 patients) + Denominator Exception (b=10 patients) + Performance Not Met (c=20 patients)    Eligible Population / Denominator (d=80 patients)	<u>ents</u> = 87.50% ents			
Performance Rate=				