



Premium Specialty: Rheumatology
 Credentialed Specialties include: Rheumatology

Use this document with the UnitedHealth Premium® Program Methodology document at UnitedHealthPremium.UHC.com. Please review all of the methodology documents to understand the entire Premium methodology.

We evaluate quality and cost efficiency using national standardized measures and attribution methods. Quality measures are attributed to physicians in applicable specialties. The following chart lists the measures we use to evaluate physicians in the rheumatology Premium specialty by condition or procedure. These measures apply to our UnitedHealthcare commercial, UnitedHealthcare Medicare Advantage and UnitedHealthcare Community Plan patient populations, unless otherwise noted.

Please view the [Quality Evaluation Example](#) and [Attribution Methods](#) documents to learn more.

| Condition/Procedure | Measure | Compliance Criteria | Measure Type | Attribution Method | Source |
|--|---|---|----------------------|--------------------|--|
| Back Pain | Patient(s) with uncomplicated low back pain that did not have imaging studies | Patient with uncomplicated low back pain did not have imaging studies | Acute Condition Care | Rendering | Contact National Committee for Quality Assurance |
| | Patient with a lumbar spine MRI and low back pain diagnosis had antecedent conservative therapy | Patient with a lumbar spine MRI and low back pain diagnosis had antecedent conservative therapy | Diagnostic Care | Ordering | Contact National Quality Forum or Centers for Medicare & Medicaid Services |
| Carotid Imaging for Syncope | Patient(s) with syncope that had a carotid imaging test. | Patient with syncope did not have a carotid imaging test | Diagnostic Care | Ordering | Synopsis |
| Cardiac imaging for Preoperative Risk Assessment for Non-Cardiac; Low-Risk Surgery | Patients of low-risk who did not receive cardiac imaging 30 days prior to a non-cardiac, low-risk surgery | Patient of low-risk did not have cardiac imaging 30 days prior to a non-cardiac, low-risk surgery | Diagnostic Care | Ordering | Contact National Quality Forum or Centers for Medicare & Medicaid Services |

| Condition/Procedure | Measure | Compliance Criteria | Measure Type | Attribution Method | Source |
|---|---|--|----------------------|------------------------|--|
| Concurrent Use of Opioids and Benzodiazepines | Patient(s) who did not have concurrent use of prescription opioids and benzodiazepines | Patient did not have prescription opioid and benzodiazepine medications concurrently dispensed | Patient Safety Care | Prescribing | Contact National Quality Forum or Pharmacy Quality Alliance |
| Head Imaging for Syncope | Patient(s) with syncope that had a head imaging test | Patient with syncope did not have a head imaging test | Diagnostic Care | Ordering | Synopsis |
| Head Imaging for Uncomplicated Headache | Patient(s) with a headache that had a head imaging test | Patient with a headache did not have a head imaging test | Diagnostic Care | Ordering | Synopsis |
| Medication Safety Monitoring | Elderly patient(s) with dementia who did not take an antipsychotic; benzodiazepine; tricyclic antidepressant; H2 receptor antagonist; nonbenzodiazepine hypnotic or anticholinergic agent after the earliest record of dementia | Patient with dementia did not have an antipsychotic, benzodiazepine, tricyclic antidepressant, H2 receptor antagonist, nonbenzodiazepine hypnotic or anticholinergic agent dispensed after the earliest record of dementia | Patient Safety Care | Patient or Prescribing | Contact National Quality Forum or National Committee for Quality Assurance |
| | Elderly patients who had an accidental fall or hip fracture who did not take an anticonvulsant; nonbenzodiazepine hypnotic; SSRI; antiemetic; antipsychotic; benzodiazepine; or tricyclic antidepressant after the incident | Patient with an accidental fall or hip fracture did not have an anticonvulsant, nonbenzodiazepine hypnotic, SSRI, antipsychotic, benzodiazepine, or tricyclic antidepressant medication dispensed after the incident | Patient Safety Care | Patient or Prescribing | Contact National Quality Forum or National Committee for Quality Assurance |
| | Elderly patients with chronic kidney disease who did not take a Cox-2 selective or nonaspirin nonsteroidal anti-inflammatory drug (NSAID) after the earliest record of chronic kidney disease | Patient with chronic kidney disease did not have a Cox-2 selective or nonaspirin nonsteroidal anti-inflammatory drug (NSAID) dispensed after the earliest record of chronic kidney disease | Patient Safety Care | Patient or Prescribing | Contact National Quality Forum or National Committee for Quality Assurance |
| Osteoporosis Management | Patient(s) compliant with prescribed oral bisphosphonate (minimum compliance 80%) | Patient was 80% or more compliant with prescribed oral bisphosphonate medication | Chronic Disease Care | Patient | Synopsis |

| Condition/Procedure | Measure | Compliance Criteria | Measure Type | Attribution Method | Source |
|---|--|---|----------------------|------------------------|--|
| Osteoporosis Management | Women 67 - 85 years of age who were treated or tested for osteoporosis within six months of a fracture | Patient was treated or tested for osteoporosis within six months of a fracture | Chronic Disease Care | Patient | Contact National Quality Forum or National Committee for Quality Assurance |
| Rheumatoid Arthritis | Patient(s) compliant with prescribed methotrexate (minimum compliance 80%) | Patient was 80% or more compliant with prescribed methotrexate medication | Chronic Disease Care | Patient | Synopsis |
| | Patient(s) compliant with prescribed hydroxychloroquine (minimum compliance 80%). | Patient was 80% or more compliant with prescribed hydroxychloroquine medication | Chronic Disease Care | Patient | Synopsis |
| | Patient(s) compliant with prescribed subcutaneous tumor necrosis factor inhibitor (minimum compliance 80%) | Patient was 80% or more compliant with prescribed subcutaneous tumor necrosis factor inhibitor medication | Chronic Disease Care | Patient | Synopsis |
| | Patient(s) taking methotrexate sulfasalazine or leflunomide that had serum alanine aminotransferase (ALT) or aspartate aminotransferase (AST) test in last 3 reported months | Patient taking methotrexate sulfasalazine or leflunomide medication had a serum ALT or AST test | Chronic Disease Care | Patient | Synopsis |
| | Patient(s) taking methotrexate, sulfasalazine, gold, or leflunamide that had a complete blood count (CBC) in last 3 reported months | Patient taking methotrexate, sulfasalazine, gold, or leflunamide medication had a complete blood count (CBC) test | Chronic Disease Care | Patient | Synopsis |
| | Patient(s) who had a prescription dispensed for a disease modifying anti-rheumatic drug (DMARD) during the report period | Patient had a disease modifying anti-rheumatic drug (DMARD) dispensed | Chronic Disease Care | Patient | Contact National Quality Forum or National Committee for Quality Assurance |
| Use of High-Risk Medications in the Elderly | Patient(s) 66 years of age and older who did not receive two or more of the same high-risk medication in the last 12 reported months | Patient did not have two or more of the same high-risk medications dispensed | Patient Safety Care | Patient or Prescribing | Contact National Quality Forum or National Committee for Quality Assurance |
| | Patients 66 years of age and older who did not receive one or more high-risk medications in the elderly in the last 12 reported months | Patient did not have a high-risk medication dispensed | Patient Safety Care | Patient or Prescribing | Contact National Quality Forum or National Committee for Quality Assurance |

| Condition/Procedure | Measure | Compliance Criteria | Measure Type | Attribution Method | Source |
|---------------------------|---|---|---------------------|--------------------|--|
| Use of Opioid Medications | Patient(s) 18 years and older who did not have an average morphine equivalent dose (med) > 120 mg/day during the treatment period | Patient did not have an average morphine equivalent dose greater than 120 mg/day | Patient Safety Care | Prescribing | Contact National Quality Forum or National Committee for Quality Assurance |
| | Patient did not have opioid medications from four or more different prescribers dispensed | Patient did not have opioid medications from four or more different prescribers dispensed | Patient Safety Care | Prescribing | Contact National Committee for Quality Assurance |
| | Patient did not have opioid medication for 15 or more days during the first 30 days following initial opioid treatment | Patient did not have access to opioid medication for 15 or more days during the first 30 days following first opioid treatment initiation | Patient Safety Care | Prescribing | Contact National Committee for Quality Assurance |
| | Patient did not have opioid medication for 31 or more days during the first 62 days following initial opioid treatment | Patient did not have access to opioid medication for 31 or more days during the first 62 days following first opioid treatment initiation | Patient Safety Care | Prescribing | Contact National Committee for Quality Assurance |

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