



Inpatient Admissions Criteria

Effective July 20, 2022, HAP will be transitioning to the latest version of the following medical screening modalities:

- InterQual® 2022
- The CMS Inpatient Only List 2022
- HAP Criteria 2022 (formerly *HAP Criteria for Inpatient Admissions*). Note: This document was renamed for continuity. No other changes were made to the current version.

HAP Criteria for Inpatient Admissions 2022



If the inpatient criteria are not met as follows for each of the medical conditions, observation is the expected level of care.

DIAGNOSIS	CRITERIA FOR INPATIENT ADMISSION
ACUTE KIDNEY INJURY	<ul style="list-style-type: none"> — Creatinine level exceeds twice the patient’s baseline and the upper reference range following an observation period during which intravenous fluids were administered OR — Creatinine level continues to rise following an observation period during which intravenous fluids were administered OR — Acute renal failure with persistent nausea, vomiting or diarrhea following an observation period during which intravenous fluids were administered
ASTHMA	<ul style="list-style-type: none"> — Failed observation period — Concurrent pneumonia confirmed on imaging with systemic toxicity — Ventilatory support (invasive or non-invasive) — Peak expiratory flow (PEF) $\leq 25\%$, forced expiratory volume at one second (FEV1) $\leq 25\%$, arterial PCO2 $\geq 42\text{mmHg}$ and pH ≤ 7.24, or venous PCO2 $\geq 42\text{mmHg}$ and pH ≤ 7.24 — Unresolved documented wheezing following at least 3 doses of a short acting beta-agonist and at least 3 hours of outpatient management along with at least one criteria point and at least one accompanying risk factor: Criteria: Accessory muscle usage, pulsus paradoxus $> 10\text{mmHg}$, PEF 26-69%, FEV1 26-69%, plasma glucose level > 300 Risk factor: History of sudden or severe exacerbation, intubation or critical care admission, or severe and persistent mental health or substance use disorder
ATRIAL FIBRILLATION	<ul style="list-style-type: none"> — Persistent atrial fibrillation confirmed on ECG that remains uncontrolled (rate exceeds 110) despite an observation period OR — The initiation of one of the following anti-arrhythmic agents: amiodarone, disopyramide, dofetilide, sotalol, dronedarone, mexiletine or quinidine
BOWEL OBSTRUCTION	<ul style="list-style-type: none"> — Bowel obstruction confirmed by imaging and unresolved following treatment during an observation period
CELLULITIS	<ul style="list-style-type: none"> — A documented increase in area of involvement or lymph involvement despite an observation period during which intravenous anti-infectives were administered OR — Orbital cellulitis and treatment with intravenous anti-infective OR — Immunocompromised patient OR — Cellulitis in proximity with an indwelling medical device OR — Systemic toxicity

CHOLECYSTITIS	<ul style="list-style-type: none"> — Persistent systemic toxicity despite supportive care and definitive procedure that occurred in an observation period OR — Gangrenous gallbladder OR — Perforated gallbladder
CHRONIC OBSTRUCTIVE PULMONARY DISEASE	<ul style="list-style-type: none"> — Failed observation period OR — Concurrent pneumonia or heart failure confirmed on imaging OR — Ventilatory support (invasive or non-invasive) OR — Persistent documented dyspnea following 3 doses of a short acting beta-agonist and at least one criteria point: O2 sat lower than 90% and the patient's baseline, PaO2 lower than 50mmHg, accessory respiratory muscle usage, paradoxical chest wall movement or acute or progressive central cyanosis
CHRONIC PANCREATITIS	<ul style="list-style-type: none"> — Failed observation period
DEEP VEIN THROMBOSIS	<ul style="list-style-type: none"> — Confirmed by imaging with at least one criteria point: Pregnancy, malignancy, pulmonary embolism, active bleeding, major surgery within the last 6 weeks, history of heparin-induced thrombocytopenia, iliofemoral vein thrombosis, creatinine level exceeds 2.5, platelet count less than 50,000, severe sustained hypertension (SBP exceeds 220 or DBP exceeds 110), bilateral DVTs, GI bleed within the last 6 months or morbid obesity (BMI exceeds 40)
DEHYDRATION OR GASTROENTERITIS	<ul style="list-style-type: none"> — Persistent volume depletion with active vomiting or diarrhea following an observation period OR — Orthostatic changes following an observation period
DIABETIC FOOT ULCER	<ul style="list-style-type: none"> — Ischemia OR — Gangrene OR — Systemic toxicity
DIABETIC KETOACIDOSIS	<ul style="list-style-type: none"> — Plasma glucose level exceeds 250 with at least one criteria point: pH lower than 7.25, serum bicarbonate level lower than 15 or hydroxybutyrate level exceeds 4
DIABETES MELLITUS - HYPERGLYCEMIA	<ul style="list-style-type: none"> — Persistent elevated blood sugar exceeding 450 following an observation period
DIVERTICULITIS	<ul style="list-style-type: none"> — Diverticulitis confirmed on imaging with at least one criteria point: Systemic toxicity, perforation confirmed on imaging or failed observation period
DYSPNEA	<ul style="list-style-type: none"> — Mechanical ventilatory support (invasive or non-invasive) without suspected or actual diagnosis of asthma, COPD, heart failure or pneumonia
FRACTURE	<ul style="list-style-type: none"> — Fracture requiring an inpatient designated surgery

HYPERCALCEMIA	— Diagnosis of hypercalcemia along with a calcium level of 12.1-13.9 with persistent symptoms (abdominal pain, mental status changes, Glasgow Coma Scale (GCS) 9-14, nausea, nephrolithiasis, or vomiting) following an observation period
HYPERKALEMIA	— Potassium level exceeds 6.5 OR — Potassium level remains from 5.5 to 6.5 despite treatment in an observation period OR — Potassium level exceeds 5.4 with associated ECG changes including AV dissociation, loss of P wave, multifocal PVCs, ventricular fibrillation, ventricular tachycardia or widening QRS
HYPERMAGNESEMIA	— Diagnosis of hypermagnesemia along with a magnesium level greater than 3.0 with persistent symptoms (mental status changes, Glasgow Coma Scale (GCS) 9-14, muscle weakness, or vomiting) following an observation period
HYPERNATREMIA	— Diagnosis of hypernatremia along with a sodium level of 151-160 with persistent symptoms (hyperreflexia, irritability, mental status changes, Glasgow Coma Scale (GCS) 9-14, muscle weakness, or restlessness) following an observation period
HYPERTENSION	— Elevated BP (SBP exceeds 180 and/or DBP exceeds 120) along with at least one criteria point: Associated acute neurological symptoms, acute coronary syndrome, acute heart failure, pregnancy, aortic dissection, recent vascular surgery, papilledema or acute kidney injury
HYPOCALCEMIA	— Diagnosis of hypocalcemia along with a calcium level of 5.0-7.4 with persistent symptoms (carpopedal spasm, flaccid paralysis, muscle weakness, paresthesia, perioral numbness, or tetany) following an observation period during which treatment included calcium repletion
HYPOKALEMIA	— Diagnosis of hypokalemia along with a potassium level less than 2.5 and without electrocardiogram changes following an observation period during which treatment included potassium repletion
HYPOMAGNESEMIA	— Diagnosis of hypomagnesemia along with a magnesium level of 1.0-1.4 with persistent symptoms (carpopedal spasm, clonus, hyperreflexia, malaise, nausea, tetany, or weakness) following an observation period during which treatment included magnesium repletion
HYPONATREMIA	— Sodium level less than 130 following a period of observation OR — Sodium level less than 121
HYPOPHOSPHATEMIA	— Diagnosis of hypophosphatemia along with a phosphate level of 1.0-1.4 with persistent muscle weakness following an observation period during which treatment included phosphate repletion
HYPOVOLEMIA OR HYPOTENSION	— SBP lower than 90 <u>without</u> tachycardia, tachypnea, oliguria, mental status changes, acidosis or elevated serum lactate despite adequate fluid resuscitation OR — SBP lower than 90 <u>with</u> tachycardia, tachypnea, oliguria, mental status changes, acidosis or elevated serum lactate
ILEUS	— Ileus confirmed by imaging and unresolved following treatment during an observation period
LOWER GASTROINTESTINAL BLEED	— Lower gastrointestinal bleed with hematochezia or melena along with either a hematocrit level < 30% and at least 20% less than their baseline OR a hemoglobin level < 10 and at least 2 grams less than their baseline and at least one criteria point: INR level exceeds 2.0, mental status changes, non-vitamin K oral anticoagulant, a platelet count less than 60,000 or exceeds 1,000,000, PT level exceeds one and a half the upper reference range, PTT level exceeds one and a half the upper reference range, or orthostatic changes

NEPHROLITHIASIS	— Nephrostomy tube placement
NSTEMI	— Diagnosis of NSTEMI with positive cardiac biomarkers along with a cardiac catheterization performed within 24 hours of presentation OR scheduled to be performed within 24 hours of presentation
NEW ONSET NEUROLOGICAL DISORDER	— Unresolved ataxia, blindness, diplopia, visual field loss, nystagmus, or paresis or paralysis of an extremity following treatment in an observation period
OSTEOMYELITIS	— Confirmed osteomyelitis on imaging with systemic toxicity
RHABDOMYOLYSIS	— CPK level exceeds 5000 despite treatment during an observation period OR — CPK level lower than 5000 and trending up despite treatment during an observation period OR — CPK level exceeds 1500 despite treatment during an observation period and the patient is unable to achieve adequate oral hydration
STROKE	— Diagnosis of stroke with confirmation on imaging of ischemia, thrombus or hemorrhage AND at least one new neurological deficit: aphasia, ataxia, blindness, diplopia, dysarthria, dysphagia, mental status changes (excludes coma, stupor or obtundation), Glasgow Coma Scale (GCS) 9-14, paralysis, paresis, partial or total gaze palsy, sensory deficit, or visual field loss
SYNCOPE	— Documented episode of syncope with at least one criteria point: Acute myocardial ischemia, acute aortic dissection, decompensated heart failure, severe pulmonary hypertension, cardiac rhythm pauses of 3 seconds or more, pre-excitation syndromes such as Wolff-Parkinson White, suspected arrhythmogenic right ventricular cardiomyopathy, aortic or mitral stenosis, 2 nd or 3 rd degree AV block or primary arrhythmia syndrome such as long QT syndrome, Brugada syndrome, idiopathic ventricular tachycardia or short QT syndrome
TIA	— Patient's diagnosis changes to stroke during the observation period for the TIA work-up
UPPER GASTROINTESTINAL BLEED non-variceal	— Upper gastrointestinal bleed non-variceal with coffee ground emesis, hematemesis, hematochezia or melena along with either a hematocrit level < 30% and at least 20% less than their baseline OR a hemoglobin level < 10 and at least 2 grams less than their baseline and at least one criteria point: INR level exceeds 2.0, mental status changes, non-vitamin K oral anticoagulant, a platelet count less than 60,000 or exceeds 1,000,000, PT level exceeds one and a half the upper reference range, PTT level exceeds one and a half the upper reference range, or orthostatic changes
VAGINAL BLEEDING	— Acute severe menorrhagia with a hemoglobin level lower than 6.1 with hemodynamic instability and treatment including blood product transfusion OR — Ruptured ectopic pregnancy OR — Uterine rupture OR — Genitourinary trauma OR — Uterine arteriovenous malformation

Baseline – When no baseline is provided, baseline is considered within the standard reference range. *LOS* - Length of stay.

HAP CLINICAL SURGICAL CRITERIA

WHEN THE SURGICAL PROCEDURE IS NOT ON THE CMS (CENTERS FOR MEDICARE & MEDICAID SERVICES) INPATIENT LIST OR IF THE PROCEDURE IS NOT ON THE INTERQUAL INPATIENT LIST OR IF IT IS CATEGORIZED AS AN ASTERISK PROCEDURE

ADULT

Three or more of the following criteria:

- Cardiovascular disease - cardiomyopathy, unstable coronary syndromes (i.e., unstable or severe angina [Canadian Class III or IV])
- Uncompensated chronic heart failure [CHF] [NYHA class III or IV]
- BMI (Body Mass Index) greater than or equal to 40
- Diabetes mellitus uncontrolled despite optimal medical management with a documented A1C greater than 9.0%
- Hypertension which is poorly controlled despite optimal medical management (described as: systolic blood pressure ≥ 180 mmHg or diastolic blood pressure ≥ 110 mmHg)
- Thrombocytopenia or clotting factor disorders (hemophilia/uncontrolled coagulopathy with anticipated need for transfusions)
- Current treatment of a malignancy
- Prior documented complication with anesthesia or post-operative complications
- ESRD (end stage renal disease) requiring dialysis.
- One of the following criteria:
 - Advanced Liver Disease (MELD score >8)
 - Individual is awaiting a lung or heart transplant
 - MI (myocardial infarction) within the last 3 months
 - CVA (cerebrovascular accident) or TIA (transient ischemic attack) within the last 3 months.

PEDIATRIC

One of the following criteria:

- BMI (Body Mass Index) greater than or equal to 40
- Brittle diabetics or patients who are not well controlled
- Major Cardiac risk factors including transposition of the Great Vessels, Pulmonic stenosis, hypoplastic left heart syndrome and single ventricle
- Respiratory Disease (Cystic fibrosis, Uncontrolled Asthma, requires preoperative oxygen)
- Chronic Kidney Disease
- Neurologic Disease (Cerebral palsy, CNS disease, Poorly controlled epilepsy, Muscular Dystrophy)
- Hematologic Disease
- Post-conceptual age < 60 weeks

Per UpTo Date: The MELD score, American Society of Anesthesiologists (ASA) class, and age predicted mortality in a study of 772 patients with cirrhosis who underwent major digestive, orthopedic, or cardiovascular surgery [41]. The MELD score was the best predictor of 30- and 90-day mortality. Mortality at 30 days ranged from 6 percent (MELD score, <8) to more than 50 percent (MELD score, >20) and correlated linearly with the MELD score.

This information is not intended to represent the level of benefits covered by HAP. Please refer to the Member's Subscriber Contract, Certificate of Coverage and/or applicable Benefit Rider(s). For more information, contact HAP Customer Service at 800-801-1766.

This notification applies to all lines of HAP business and is effective until revised.

Observation: An observation level of care is defined by CMS as "a well-defined set of specific, clinically appropriate services, which include ongoing short-term treatment, assessment, and reassessment before a decision can be made regarding whether patients will require further treatment as hospital inpatients or if they are able to be discharged from the hospital." This determination should be rendered within 48 hours; therefore, observation stays of up to 48 hours will be reimbursed without prior authorization. Exceptional circumstances may require extension of the observation period, of which these may be retrospectively reviewed for appropriate level of care.