

TennCare Episodes of Care: Detailed description of episode risk adjustment for Wave 2 episodes

Chronic obstructive pulmonary disease acute exacerbation, screening and surveillance colonoscopy, outpatient and non-acute inpatient cholecystectomy, acute percutaneous coronary intervention (PCI), non-acute PCI

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The state of Tennessee has implemented a bundle-based approach to reimburse providers for the care delivered to patients enrolled in the state's Medicaid program. Bundled payments cover all of the services provided to a patient for treatment of a specific condition during a defined episode of care, including services related to diagnosing, managing and treating that condition. The actual provision of services to a specific patient for a specific condition is herein called an "episode," while the grouping for payment of episode-related services normally used to treat the condition is called a "bundle." This distinction is useful because the state may choose, as a matter of policy, to exclude from the bundle some of the services in an episode. For each of these patients and episodes, a provider will be determined to have overall responsibility (the episode "quarterback"). The total cost of care for each quarterback in delivering all bundled services will be measured and compared with targets and thresholds to determine overall performance.

The comparison of bundle costs for a provider is based on the average *risk-adjusted* cost of the provider's episodes with the targets and thresholds established by the state for payment purposes. The health care services required to deliver a bundle of care can vary greatly across patient episodes. Risk adjustment quantifies the part of this variation in cost that can be explained by clinical factors, such as disease progression, comorbidities and other patient attributes, that correlate with clinical need, including age and gender. A higher risk score for an episode means a higher expected cost relative to other episodes of the same type due to the clinical or demographic factors. Risk adjusting bundle costs enables more equitable comparisons across providers and with targets and thresholds.

The first phase of this new payment initiative included 3 bundle types: Asthma – acute exacerbation, perinatal and total joint replacement. An earlier document, that includes several detailed examples of episode risk adjustment, describes the risk adjustment approach used for these 3 bundles. This earlier document may provide useful background to those new to bundled payment.

The present document provides details on the approach used by UnitedHealthcare to compute episode risk and to risk-adjust episode costs for the following bundles: COPD acute exacerbation (COPD); screening and surveillance colonoscopy (COLO); and outpatient and non-acute inpatient cholecystectomy (CHOLE); acute PCI (PCI-A); and non-acute PCI (PCI-N). It describes the general approach used to measure risk across all bundle types, followed by a description of the specific risk markers used for each type of bundle.

I. Overview: Measuring episode risk

Episode risk models are designed to predict the total *expected cost* for an episode of care — those costs that are expected given the clinical characteristics of the patient and the episode. These costs include the payments for all services received by a patient during the course of an episode. Given a measure of the expected cost or relative risk for an episode, actual episode costs can be risk-adjusted. Risk-adjusted costs can then be compared across all quarterbacks and combined with targets to determine performance under the program. Example 1 illustrates this concept.

As shown in Example 1, all episodes for the quarterback are assessed to determine their relative risk and the quarterback's average risk-adjusted cost is computed.

A unique *risk model* was developed for each bundle type based on clinical and demographic variables that would influence the potential cost of those specific episodes.

Episode risk models use 2 key features: episode *risk markers* and episode *risk weights*. *Risk markers* describe those unique clinical characteristics of an episode that were found statistically to affect episode costs. *Risk weights* describe a risk marker's incremental relative contribution to expected episode costs or risk.

As noted above, a separate risk model was developed for each bundle type. As a result, the risk markers and risk weights included in the models differ by bundle type. This is to be expected, given that different clinical factors will have a different impact on bundle costs, depending upon the type of episode.

When assigning a risk score to a bundle, 5 major steps are followed:

1. Identify clinical risk markers using clinical input
2. Assign demographic risk markers
3. Apply risk weights to each risk marker
4. Compute an episode risk score
5. Adjust preliminary risk scores for *risk score neutrality*

Example 1: CHOLE episode risk adjustment

- A surgeon serves as the quarterback for 10 CHOLE episodes during calendar year 2019
- The total cost for each of those episodes is calculated using costs for all services included in the episode (for example medications, imaging and testing, evaluation and management, etc.)
- The characteristics of the 10 patients and their episodes are used to assign a risk score to each individual episode. This risk score represents the relative expected costs of each episode based on clinical and patient factors such as age, gender, diagnoses and disease comorbidities.
- Episode risk is expressed as a relative score. A risk score of 1.000 represents the average risk of episodes for a given set of covered lives. An individual cholecystectomy episode that, based on its clinical and patient factors, is expected to have a 10% higher cost than average would be assigned a risk score of 1.100.
- The actual total cost for each of the surgeon's episodes is risk-adjusted to compute risk-adjusted total cost. Actual cost is divided by episode risk score, so that higher-risk episodes will have costs adjusted down while lower-risk episodes will have costs adjusted up, allowing episodes with different risk to be fairly compared. For example, an episode with a total cost of \$33,000 and a risk score of 1.100 would have a risk-adjusted total cost of \$30,000.
- The quarterback's overall performance is based on average risk-adjusted cost for the 10 episodes. This amount can be compared with that of other providers and with targets to determine performance under the program.

Each of these steps is described below.

II. Assigning clinical risk markers to an episode

The following steps are used to assign clinical risk markers to an episode:

1. Identify qualified services that can contribute diagnoses to risk marker identification
2. Identify the set of initial risk markers using clinical criteria
3. Assign clinically appropriate service timing to risk markers
4. Reduce to a minimum necessary set of risk markers per bundle using statistical criteria

1. Identify qualified services

Only diagnoses from *qualified* service records are considered when identifying risk markers. Qualified services include services such as office visits, consultations, ER visits, surgeries and inpatient stays. Non-qualified services include services such as lab or radiology or services delivered by a durable medical equipment (DME) or ambulance provider. In this way, the methodology does not consider diagnoses from ancillary services or “rule-out” tests. Only services with diagnoses confirmed and assigned by a clinician or facility are used. Qualified services are determined by examining the procedure and revenue codes on an individual service record.

2. Identify initial risk markers

Based on the diagnoses observed on qualified services, 2 sets of clinical risk markers are considered for use in risk-adjusting episodes. First, the diagnoses associated with qualified services are grouped into Episode Treatment Groups® (ETGs®). ETGs are then selected for evaluation as a risk marker based on their clinical relevance to the episode and their prevalence in the episodes.¹

In addition, the state of Tennessee defines risk makers using both Clinical Classifications Software (CCS) groups and their own specific definitions. The second set of risk makers consists of those markers that are specified by the state that meet minimum requirements regarding frequency of occurrence. (The CCS groups are not used since they tend to duplicate information captured by ETGs.)

3. Assign service timing

Service timing is also important when setting initial clinical risk markers. Three windows of service timing, based on clinical appropriateness, were specified for all ETG-based risk markers: (1) risk marker occurred in the 365 days prior to the episode start through 30 days prior to the episode start (*comorbidity risk marker, prior window*); (2) risk marker occurred in the 30 days prior to the episode — start through end of the episode (*episode risk marker window*); (3) risk marker occurred in the 365 days prior to the episode — start through the episode end (*comorbidity risk marker, full window*).

- *Episode risk marker window* – Used to identify risk markers that occurred in the context of the episode itself. The episode risk marker window begins 30 days prior to episode start and extends through the end of the episode.
- *Comorbidity risk marker, full window* – Used to identify risk markers for other conditions not directly related to the episode that increase the complexity and risk associated with its delivery. This window includes a longer period of time — 365 days prior to the episode start through the episode end.
- *Comorbidity risk marker, prior window* – Used to identify risk markers for other conditions not directly related to the episode that increase the complexity and risk associated with its delivery. This window covers the 365 days prior to the episode start through 30 days prior to the episode start. This approach allows for recognition of patient comorbidities that might be considered complications of the episode itself, if first observed during the episode risk marker window.

In general, risk markers defined by the state include their own criteria with regard to service timing. Following this step, all initial clinical risk markers have been assigned to the episode.

4. Reduce to the minimum necessary set of risk markers per bundle

After the initial clinical review, the selected set of clinical risk markers are analyzed statistically to determine their impact on costs for the episode being evaluated. Risk factors for inclusion in the final model are determined based on their clinical relevance to the episode and their impact on costs.

III. Assigning demographic risk markers to a bundle

Demographic characteristics of patients can also affect risk, either because age and gender can affect coverage decisions or because they serve as proxies for unmeasured clinical attributes. For this reason, the statistical evaluation of potential risk markers also evaluates the extent to which the models should distinguish among patients based on age and gender. One of the 5 bundle types (COPD) includes 2 or more demographic risk markers in the final risk model — based on an individual's age and gender at the time of the trigger event. Age and gender did not have a statistically meaningful effect on the cost of the other 4 bundles, which means that all individuals are assigned the same base risk weight that corresponds to an uncomplicated episode.

IV. Apply risk weights to each marker

Each risk marker is assigned a risk weight. This *risk weight* describes a marker's incremental contribution to bundle risk for that bundle type. Model risk weights were estimated using historical data describing a large number of bundles. The risk weights for each risk model by episode type are described below in Tables 1–5. For each episode, all of the demographic and clinical risk markers are captured along with the corresponding *risk weights*. All identified *risk weight* values are then added together to achieve the preliminary risk score for that individual episode.

V. Preliminary risk score

The preliminary risk score for each individual episode is calculated as the sum of individual risk weight values that apply to that episode. Preliminary risk scores for each episode are then adjusted to achieve risk score neutrality across all episodes.

VI. Adjust preliminary risk for risk score neutrality

The preliminary risk score for an episode is multiplied by an episode-specific risk neutrality factor. This factor was based on the adjustment needed to help ensure that the average risk score for each episode was equal to 1.00 for UnitedHealthcare. Risk neutrality factors are calculated at the beginning of each performance period. These values are held constant through the performance period to help ensure that providers are measured against constant risk-adjusted thresholds. The final risk score after this adjustment is then used to risk adjust the cost of the individual episode.

Example 2: Applying risk neutrality factors

- All risk factors associated with an episode are identified and the corresponding risk weight values (clinical and demographic) are added together to achieve the preliminary risk score for an individual episode
- Preliminary risk scores are then multiplied by a risk neutrality factor to help ensure that the average risk score for UnitedHealthcare is 1.00
- The application of the risk neutrality factor will make the final risk score different than the sum of risk weights listed in Tables 1–5 below
- For example, if the risk neutrality factor of an Acute PCI episode was 0.987, then a 58-year-old man without other clinical risk factors would have a final risk score of 0.7341 ($0.987 * 0.7438 = 0.7341$)

Please go to the UnitedHealthcare Provider Portal at UHCprovider.com. Click Sign In in the top-right corner to find the most recent TennCare Episodes of Care risk neutrality factors.

Tables 1–5 below show the risk weights for COPD Acute Exacerbation (COPD); Screening and Surveillance Colonoscopy (COLO); and Outpatient and Non-Acute Inpatient Cholecystectomy (CHOLE); Acute PCI (PCI-A); and Non-Acute PCI (PCI-N). The risk weights shown in these tables were used to risk-adjust the cost of the individual episodes. The preliminary risk score for each episode is the sum of the risk weights for all risk markers observed. The final risk score will be the preliminary risk score for an episode multiplied by an episode-specific risk neutrality factor.

Table 1* COPD Acute exacerbation	
Description of risk marker	Risk weight
All ages, 18–44	0.0372
All ages, 45–54	0.0948
All ages, 55–64	0.1996
Nutritional deficiency (comorbidity risk marker, full window)	0.1578
Bacterial lung infections (comorbidity risk marker, full window)	0.3624
Other metabolic disorders (comorbidity risk marker, full window)	0.1961
Obesity (comorbidity risk marker, full window)	0.0571
Severe presentation of COPD (episode risk marker window)	0.4015
Substance abuse: Alcohol dependence (comorbidity risk marker, full window) or opioid or barbiturate dependence (comorbidity risk marker, full window)	0.0658
Ischemic heart disease (episode risk marker window)	0.1712
Heart failure, diastolic (comorbidity risk marker, full window)	0.1482
Hypertension (comorbidity risk marker, full window)	0.0750
Other inflammatory lung diseases (comorbidity risk marker, full window)	0.3566
Acute respiratory distress syndrome (comorbidity risk marker, full window)	0.5943

*In 2022, the COPD risk model was updated to test new risk markers and incorporate 2022 episode design and configuration file maintenance changes.

Table 2 Colonoscopy	
Description of risk marker	Risk weight
All ages	0.7158
Non-malignant neoplasm of intestines and abdomen (comorbidity risk marker, full window)	0.2132
Acute pancreatitis (episode risk marker window)	0.1677
Malignant neoplasm of large intestine (comorbidity risk marker, full window)	0.1375

Table 2 (cont.) Colonoscopy	
Description of risk marker	Risk weight
Bowel obstruction (comorbidity risk marker, full window)	0.0919
Non-malignant neoplasm of rectum or anus (comorbidity risk marker, full window)	0.0872
Hemorrhoids (comorbidity risk marker, full window)	0.0852
Substance abuse (comorbidity risk marker, full window): cocaine or amphetamine dependence, acute alcohol intoxication, alcohol dependence, opioid or barbiturate dependence or other drug dependence	0.0626
Other inflammation of intestines and abdomen (comorbidity risk marker, full window) or other infectious diseases of intestines and abdomen (comorbidity risk marker, full window)	0.0555
Hiatal hernia (comorbidity risk marker, full window)	0.0519
Diverticulitis and diverticulosis (comorbidity risk marker, full window)	0.0407
Ulcer (comorbidity risk marker, full window)	0.0277
Nutritional deficiency (comorbidity risk marker, full window)	0.0217
Asthma (comorbidity risk marker, full window) or chronic obstructive pulmonary disease (comorbidity risk marker, full window)	0.0101

Table 3 CHOLE	
Description of risk marker	Risk weight
All ages	0.8920
Appendicitis (comorbidity risk marker, full window)	0.1084
Bowel obstruction (comorbidity risk marker, full window)	0.0957
Bacterial lung infections (episode risk marker window)	0.0735
Hernias, except hiatal (comorbidity risk marker, full window)	0.0614
Dehydration (comorbidity risk marker, full window)	0.0492
Heart failure (comorbidity risk marker, full window)	0.0485
Chronic renal failure (comorbidity risk marker, full window)	0.0427
Infection of stomach and esophagus (comorbidity risk marker, full window)	0.0422
Other inflammation of intestines and abdomen (comorbidity risk marker, full window)	0.0368
Obesity (comorbidity risk marker, full window)	0.0358
Hiatal hernia (comorbidity risk marker, full window)	0.0233
Cardiovascular conditions (comorbidity risk marker, full window): Ischemic heart disease, pulmonary heart disease, cardiomyopathy, aortic aneurysm, valvular disorder, other conduction disorders, atrial fibrillation and flutter, hypertension or cardiac congenital disorder	0.0144

Table 4 Acute PCI	
Description of risk marker	Risk weight
All ages	0.7438
Immunodeficiencies (comorbidity risk marker, full window)	0.0663
Obesity (comorbidity risk marker, full window)	0.0683
Cocaine or amphetamine dependence (comorbidity risk marker, full window)	0.0049
Acute renal (comorbidity risk marker, full window)	0.0681
Subendocardial infarction (episode risk marker window)	0.0180
Multiple vessel or staged PCI (episode risk marker window)	0.1846
Heart failure (comorbidity risk marker, full window)	0.0694
Behavioral health disorders (comorbidity risk marker, full window): Mood disorder, depressed; mood disorder, bipolar; psychotic and schizophrenic disorders; dementia; mental disorders, organic and drug-induced; personality disorder, eating disorder; anxiety disorder or phobias; psychosexual disorder; intellectual disability; other neuropsychological or behavioral disorders	0.0456
Gastrointestinal disorders (comorbidity risk marker, full window): Inflammation of esophagus; gastritis and/or duodenitis; ulcer	0.0639

Table 5 Non-Acute PCI	
Description of risk marker	Risk weight
All ages	0.7752
Obesity (comorbidity risk marker, full window)	0.0513
Cocaine or amphetamine dependence (comorbidity risk marker, full window)	0.0170
Aortic aneurysm (comorbidity risk marker, full window)	0.0078
Acute renal (comorbidity risk marker, full window)	0.0495
Multiple vessel or staged PCI (episode risk marker window)	0.1844
Heart failure (comorbidity risk marker, full window)	0.0651
Behavioral health disorders (comorbidity risk marker, full window): Mood disorder, depressed; mood disorder, bipolar; psychotic and schizophrenic disorders; Dementia; mental disorders, organic and drug-induced; personality disorder, eating disorder; anxiety disorder or phobias; psychosexual disorder; intellectual disability; other neuropsychological or behavioral disorders	0.0382
Gastrointestinal disorders (comorbidity risk marker, full window): Inflammation of esophagus; gastritis and/or duodenitis; ulcer	0.0502

¹ The methodology described here uses the clinical constructs of Episode Treatment Groups® (ETGs®) to categorize diagnosis codes into clinically meaningful groups. The clinical constructs within the ETG methodology are defined in terms of both ICD-9-CM and ICD-10-CM/PCS.