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# HHS-OPERATED RISK ADJUSTMENT DATA VALIDATION (HHS-RADV) INTRODUCTION



# OBJECTIVES

- // Introduce the HHS – Operated Risk Adjustment Data Validation (HHS-RADV) Activities for the 2016 Benefit Year
- // Overview of the HHS-RADV Program
- // Identify HHS-RADV Processes
- // Responsibilities of Key Stakeholders
- // Upcoming Timelines



## HHS-RADV DEFINITIONS

- // **HHS** – Governs the HHS-RADV Program
- // **CMS** – Designated by HHS to be responsible for implementing the RA premium stabilization program
- // **Issuer** – Health plan subject to the HHS-RADV Audit – those offering non-grandfathered ACA compliant individual and/or small group health plans both inside and outside the marketplace
- // **IVA** – Entity retained by issuer to perform the Initial Validation Audit (IVA)
- // **SVA** – Entity retained by CMS to perform the Secondary Validation Audit (SVA)



## HHS-RADV AUTHORITY

// Implemented in accordance with the following regulations:

- 45 CFR Section 153.350
- 45 CFR Section 153.620
- 45 CFR Section 153.630
- Premium Stabilization Final Rule
- 2014 Payment Notice Final Rule
- 2015 Payment Notice Final Rule



## HHS-RADV AUTHORITY

- // Section 1343 of the Affordable Care Act (ACA) establishes a permanent Risk Adjustment (RA) program
- // The Premium Stabilization Final Rule requires states to validate a statistically valid sample of data for all issuers that submit for risk adjustment every year and provide an appeals process
- // Finalized in 2018 Payment Notice, HHS implemented a materiality threshold of \$15M in total premiums, beginning for the 2017 benefit year HHS-RADV program





## RISK ADJUSTMENT OVERVIEW

- // **What:** Budget neutral program that transfers funds from plans with lower risk enrollees to plans with higher risk enrollees in a state market risk pool
- // **Who Participates:** ACA-compliant non-grandfathered individual and small group market plans, inside and outside the Marketplace (Issuers)
- // **How:** Data validation in an audit function ensuring integrity and data provided by issuers



## HHS-RADV PROCESSES – SAMPLE SELECTION

- // CMS provides a sample size of enrollees so that the estimated risk score errors will be statistically sound and the enrollee-risk level risk score distributions will reflect enrollee characteristics for each issuer
- // 200 enrollees per issuer for each state in which the issuer offers plans that are HHS-RADV eligible will be sampled for the IVA – a sample less than 200 enrollees may be selected for small enrollee populations





## HHS-RADV PROCESSES – SAMPLE SELECTION

// A sample of 200 enrollees will have up to one (1) or more **Hierarchical Condition Categories (HCCs)**

- CMS requires documentation that supports the presence of the condition and indicate the provider's assessment and/or plan for management of the condition. This must occur at least once each calendar year in order for CMS to recognize the individual continues to have the condition.



## HIERARCHICAL CONDITION CATEGORIES

- Hierarchy logic is imposed on certain disease groups
- The HCC model is cumulative – a patients with more than one HCC are factored into the member’s risk profile
- Disease groups are clinically related diagnoses that have similar Medicare cost implications
- Each disease group relates to a **specific ICD-10-CM** medical condition. Some HCCs are age-related, such as breast malignancies



## HHS-RADV PROCESSES: IVA REGISTRATION

- // Issuer identifies IVA Entity – independent auditor to validate demographic and enrollment data and health status information for the enrollee sample(s)
- // CMS accepts or rejects the IVA entity
- // IVA review of enrollee health status must be conducted by certified coders by a nationally recognized agency



## INITIAL VALIDATION AUDIT – DEMOGRAPHICS AND ENROLLMENT – HEALTH STATUS VALIDATION

- // Source enrollment documentation from the claims processor to the transactions (claims) with the issuer
- // Issuer provides medical record documentation
- // IVA entity validates the risk score of each enrollee in the sample(s)
- // IVA entity provides CMS with final results and supporting documentation



## HHS-RADV PROCESSES - SVA

- // Following the IVA, the SVA is conducted by a CMS auditor to verify accuracy of the IVA findings
- // Issuers may appeal the SVA results and/or accept the error estimation
- // CMS determines an adjustment factor and prepares for payment adjustments for the benefit year based on error estimation



## HHS-RADV – ISSUER RESPONSIBILITIES

// Identify a “Senior Official” to communicate with CMS regarding audit activities

// Confirm completion of results prior to submission to CMS

// Submit appeals on behalf of the Issuer for the SVA

// Provides all claims, medical records, and enrollment documentation to the IVA entity for the sampled enrollees





## HHS-RADV – IVA ENTITY RESPONSIBILITIES

- // Be free of conflicts with the issuer
- // Attend 2016 benefit year HHS-RADV trainings
- // Maintain appropriate personnel to conduct the IVA
  - Claims, demographic, enrollment, finance
  - Ensure certified coders have and maintain current certifications
  - Register in the Audit Tool
  - Perform IVA, **IRR** and submit results to CMS, timely



## HHS-RADV TIMELINE FOR BENEFIT YEAR 2016

Date	Description
February 15, 2017	HHS-RADV Training Begins
February 2017 – April 2017	Issuers Select IVA Entity
April 17 – 24, 2017	Issuers submit IVA Entity to CMS for Acceptance
May 1, 2017	HHS-RADV 2016 Benefit Year Protocols
May 1, 2017	2016 Data Submission Deadline
May 2017	Sample Released to Issuers
June 2017-January 8, 2018	IVA is Conducted
January 8, 2018	IVA Results and Submissions Due
January 18, 2018	IVA Entity Submits SVA Subsample to CMS
January 2018 – April 2018	SVA Conducted
May 2018 – June 2018	CMS Releases Error Rates to Issuers

## CODING AND DOCUMENTATION - HCCS

- // All relevant diagnosis codes should be reported at least once per year for each member (preferably every six months)
- On January 1 each year, the member's diagnosis information is reset in preparation for a new year of diagnosis encounter data
- 2015 initial validation audits completed and will be starting the 2016 audits this spring 2017
- **2015 was a pilot year.** Penalties will begin in 2017 based on results of 2016 audits



## CODING AND DOCUMENTATION – COMMON HCCS

// COPD

// Congestive heart failure

// Acute or chronic renal failure

// Malignancies

// Diabetes with manifestations (neuropathy)

// Newborns with problems

// Complicated deliveries

// Complicated pregnancies



# DOCUMENTATION REQUIREMENTS

## // The Basics

Each page of a note **MUST** include:

- ✓ Patient's full name
- ✓ Date of Birth
- ✓ Date of Service – including year

## **The Provider's signature must be legible –**

- ✓ Must also include provider's credentials
- ✓ Electronic signatures should include the date and time of authentication, the service provider's name and credentials and include a statement such as “electronically signed by....” or “authenticated by....”



## DIAGNOSIS (HCC) ABSTRACTION - VALIDATION

- // Medical record source must be hospital inpatient, **outpatient**, or professional medical treatment (office visits)
- // Face-to-face encounters only
- // Follow the **“MEAT”** documentation criteria – **M**onitored, **E**valuated, **A**ssessed, **T**reated
- // Approved provider types – MD, DO, PA, APRN, Clinical Psychologist, PT, OT, Audiologist, DPM, etc.





## LESSONS LEARNED FROM AN ENTITY'S PERSPECTIVE

- // Lack of documentation to support the MEAT criteria for correctly capturing the HCCs
- // Illegible provider signatures
- // Signatures dated the date the documentation request was made
- // No birth date on progress note
- // Some enrollees had multiple HCCs



## LESSONS LEARNED FROM AN ENTITY'S PERSPECTIVE

- // Failure to capture HCCs once every 12 months
- // Copy and pasted “problem lists” from one encounter to another that could not be used to support the HCC due to not meeting the MEAT criteria
- // Newly identified HCCs - an error but has potential result in a positive impact to the Issuer
- // Audited a year’s worth of documentation, requested HHS to allow audit be based **at claim level** in future



## LESSONS LEARNED FROM AN ENTITY'S PERSPECTIVE

- // Use of quantifying language in the outpatient setting, such as “consistent with, probable, possible....”
- // **Historical status** of a diagnosis unclear, especially with malignancies
- // Chronic or coexisting conditions are not documented or are left out of the clinical documentation of an office visit
- // Coders did not follow Official Coding Guidelines



## MOST AUDITED ENCOUNTER TYPES

// Hospital Anesthesiologist Pre-Evaluations

// Hospital outpatient department records

// Hospital emergency room records

// Hospital inpatient records

// Physician practice office visits – most often

// Oncology and urology coding worst



## PROVIDER IMPLICATIONS

- If coding is accurate and complete, ROI processes are minimally disrupted, allowing greater focus on patient care and other business operations
- If coding is incomplete, higher likelihood of more medical record requests by an Issuer with ROI disruption and cost
- Follow MEAT criteria and be practically audit-proof
- Risk adjustment is an expanding arena, started in 2004 with Medicare Advantage



## RESOURCES: LINKS

Resource	Resource Link
Affordable Care Act (ACA) HHS-Operated Risk Adjustment Data Validation (RADV) Process White Paper, June 22, 2013	<a href="https://www.regtap.info/uploads/library/ACA_HHS_OperatedRADVWhitePaper_062213.pdf">https://www.regtap.info/uploads/library/ACA_HHS_OperatedRADVWhitePaper_062213.pdf</a>
CCIIO ACA RA Data Validation Email Address	CCIIOACARADDataValidation@cms.hhs.gov
The Center for Consumer Information & Insurance Oversight (CCIIO) web page	<a href="https://www.cms.gov/cciiio">https://www.cms.gov/cciiio</a>
Patient Protection and Affordable Care Act (ACA)	<a href="http://www.gpo.gov/fdsys/pkg/PLAW-111publ148/content-detail.html">http://www.gpo.gov/fdsys/pkg/PLAW-111publ148/content-detail.html</a>





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