



2020/21 MediGold Coding Guide

We understand the challenges of working with multiple payers and meeting measurements, guidelines and documentation for Medicare beneficiaries. This Coding Guide is intended to make things easier for you and your staff when working with MediGold. The guide includes assistance in understanding:

- Star Ratings and the HEDIS reporting process.
- · Your role in reporting and documenting care.
- Medical record requests (MRR).
- · Star measure guidance and codes.

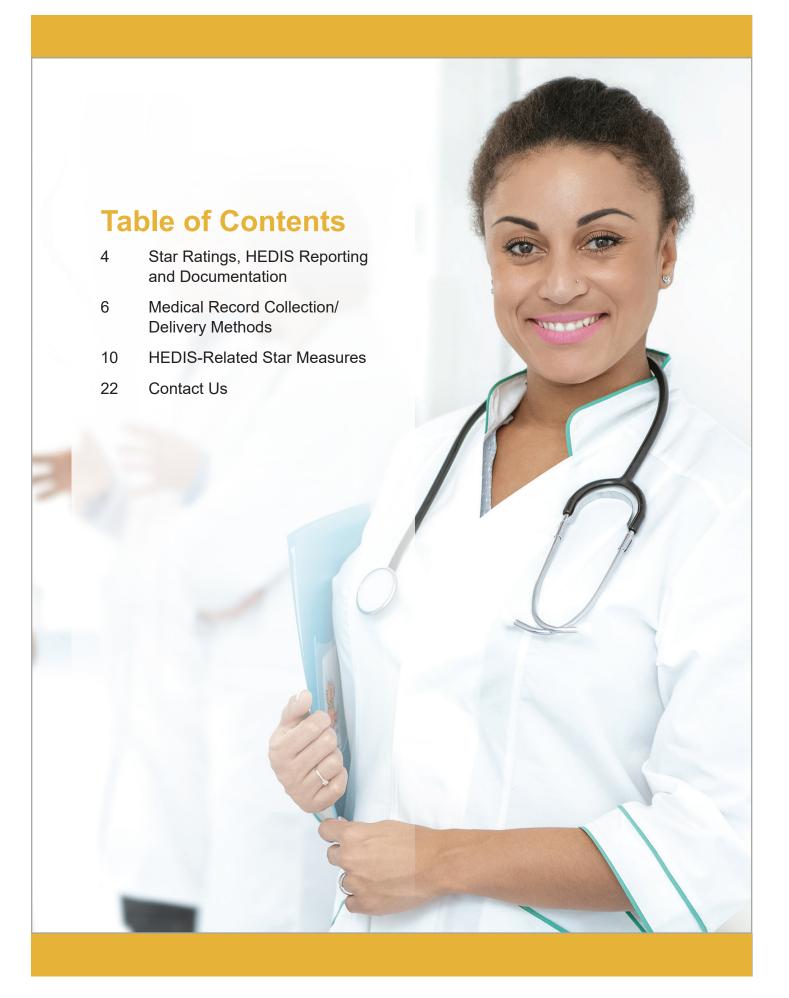
We always welcome your feedback on how we can make this guide better.

Thank you for partnering with MediGold to improve the health and well-being of MediGold members. We sincerely consider you our partner



and recognize that we cannot succeed without the compassionate and high-quality care delivered by the providers in our network. Working together, we can have a positive impact on patient outcomes. **

Greg Wise, MD, FAAFP, Chief Medical Officer, MediGold



Star Ratings, HEDIS Reporting and Documentation

What are Star Ratings?

All Medicare Advantage plans are awarded Star ratings annually by the Centers for Medicare & Medicaid Services (CMS). On a scale of one to five, a 5-Star rating is considered excellent. MediGold's overall Star rating combines rankings of quality and performance, including how well we help our members to stay healthy and manage chronic conditions. This information is gathered from HEDIS® scores, HOS and CAHPS Survey data and CMS administrative data. This guide covers the HEDIS-related Star Measures, and the needed coding and documentation for those measures, used in our HEDIS scores.

HEDIS Reporting and the Role You Play

HEDIS, the acronym for Healthcare Effectiveness Data and Information Set, is a performance measurement tool for health plans, administered by the National Committee for Quality Assurance (NCQA). HEDIS measures are a significant component of Medicare Star Ratings and the NCQA accreditation process. The coding and documentation necessary to meet measures is collected from our claims database and review of medical records. In the eyes of measurement reporting, if it isn't documented, then it didn't happen. To meet requirements, it's important to make every visit count. Useful tips include:

- Promote all patient's health and encourage an annual wellness visit before June 30 each year, when possible.
- Give patients reminder calls 48 hours before their appointments.
- Schedule follow-up visits before patients leave.
- · Accurately code all claims.
- Thoroughly document all care in the patient's chart at the time service is provided, including date and provider's signature.
- Utilize MediGold's Gaps In Care report to close measures and strengthen patient relationships.

Feel free to request a gaps in care report for your office by emailing starsandhedis@mchs.com

What are CPT Category II codes?

Current Procedural Terminology (CPT) Category II codes were developed by the American Medical Association (AMA) as a supplemental performance tracking set of procedural codes in addition to the Category I and III code settings.

- Category I codes are used for tracking and billing common procedures.
- Category III codes are temporary codes for emerging technology.
- Category II codes are optional and intended to be used for measuring performance on quality metrics such as Healthcare Effectiveness Data and Information Set (HEDIS®)

Category II codes are alphanumeric and consist of four digits followed by the letter 'F'.

Category II codes are **NOT** billing codes; they are used to track services on claims for performance measurement.

Category II codes are not to be used as a substitute for Category I codes.

What is the purpose of CPT Category II codes?

Category II codes are intended to facilitate the reporting of services or test results that support quality of care performance measures. MediGold highly encourages (and even incentivizes*) clinical office staff to utilize CPT II codes.

By accurately coding you can decrease the need for manual record abstraction and chart review, minimizing the burden on physicians and office staff to report this information through other methods.

CPT Category II codes are arranged according to the following categories:

Category	Code Range	Category	Code Range
Composite measures	0001F-0015F	Therapeutic, preventive or other interventions	4000F - 4306F
Patient management	0500F - 0575F	Follow-up or other outcomes	5005F - 5100F
Patient history	1000F - 1220F	Patient safety	6005F - 6045F
Physical examination	2000F - 2050F	Structural measures	7010F - 7025F
Diagnostic/screening processes or results	3006F - 3573F		

CPT II codes allow providers to measure and display the quality of care they provide.

CPT® is a registered trademark of the American Medical Association. Copyright 2016 American Medical Association (AMA). All rights reserved. HEDIS® is a registered trademark of the National Committee for Quality Assurance (NCQA).

HEDIS Code	Measure	Category II CPT Code	Incentive
		2022F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	
	Comprehensive Diabetes Care-Retinal Eye Exam (Diabetic members only. One time per year.)	2023F, 2025F (NEW) Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	
CDC		2024F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	\$20
		2026F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	
		3072F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	
	Comprehensive Diabetes Care- HbA1c level less than 7.0 (Diabetic members only.)	3044F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	\$10
	Comprehensive Diabetes Care- HbA1c level greater than 9.0 (Diabetic members only.)	3046F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	\$10
CDC	Comprehensive Diabetes Care- HbA1c level greater than or equal to 7.0 and less than 8.0 (Diabetic members only.)	3051F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	\$10
	Comprehensive Diabetes Care- HbA1c level greater than or equal to 8.0 and less than 9.0 (Diabetic members only.)	3052F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	\$10
		3060F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	
	Comprehensive Diabetes Care-Attention to Nephropathy (Diabetic members only. One time per year.)	3061F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	
CDC		3062F Filed with ICD-10 Diag Codes: ICD-10-E10.9, E10.10-E13.9, O24.011-O24.33, O24.811-O24.83	\$10
		3066F, 4010F Filed with ICD-10 Diag Codes: ICD-10-E08.21- E08.29, E09.21-E09.29,E10.21-E10.29, E11.21- E11.29, E13.21- E13.29,I12.0-I13.2, I15.0-I15.1, N00.0-N08, N14.0-N14.4, N17.0-N19, N25.0- N26.9, Q60.0-Q61.9, R80.0-R80.9,N18.4, N18.5, N18.6, Z91.15, Z94.0, Z99.2	
MRP	Medication Reconciliation Post- Discharge	111F	\$25

^{*}Please note that the codes listed here, when applied correctly, will result in closure of an identified care opportunity. This is not a guarantee of benefits or payment. Benefits are subject to the terms and limitations of the plan.

Documentation Requirements

Correctly documenting patient encounters is critical for quality reporting and accurate reimbursement. This is key as health care reform continues to move toward quality-driven reimbursement.

- Documentation is legible.
- Ensure correct CPT, CPT II and ICD-10 codes are used.
- Blood pressure diagnosis is documented prior to June 30.
- All patient encounters, including telephone, fax and electronic message exchanges are documented.

Common HEDIS Barriers and Obstacles

- Let us know if member attribution is incorrect (patient assigned to wrong PCP.)
- Claim submitted without correct codes will not count toward the measure. This means we
 will be required to ask for the medical record.
- Claim submitted with inaccurate diagnosis code will incorrectly add to a measure.
- Not coding A1c, blood pressure or BMI values/results.
- Services not documented in the patient's medical chart.
- All required components of the measure not provided, e.g., diabetes diagnosis or hypertension without blood pressure reading.
- Records not transferred when patient changed PCP.
- Appointment availability when patient tries to schedule preventive services.
- Practice not seeing new patient in a timely manner.
- PCPs should include documentation received from specialists and other sources in outpatient chart i.e. eye exams, inpatient and discharge summaries, radiology, gastro, gaps summaries from health plan

Medical Record Collection/Delivery Methods

Medical Record Confidentiality

MediGold strictly maintains the confidentiality of any records, which are accessed only by authorized people adhering to the following guidelines. Records are:

- Kept in a safe and secure location.
- Appropriately destroyed when they are no longer needed for the purpose requested.
- Not further disclosed or otherwise distributed.

We are not asking for nor do we want any medical record information related to psychotherapy, HIV, substance abuse or developmental disabilities.

Further, your MediGold Provider Agreement stipulates that copies of members' medical records shall be provided to MediGold, or its respective designees, for quality improvement activities, e.g., HEDIS.

If you have questions concerning this request, please contact: StarsAndHEDIS@mchs.com.

Medical Record Collection/Delivery Methods

Data collection methods include the following, as long as they meet HIPAA guidelines:

- Remote electronic medical record (EMR) system. EMR submissions, which are highly recommended, result in fewer visits and emails from MediGold.
- Fax.
- Hard copy, flash or CD delivered via postal service certified mail, or other signature-required service.
- Email encrypted to HIPAA standards.
- Schedule time with one of our coordinators to come into your office to collect a copy of the records on-site.
- Ask that one of our coordinators come by to pick up the records.

Online Submission of Medical Records for Stars and HEDIS Gaps In Care

- **1.** Access the provider portal at: MediGold.com/For-Providers/Provider-Portal. (For first-time portal users, follow the easy steps at the link to set up an account and log in.)
- 2. On the portal home page, select Close Gaps In Care.

Welcome to the MediGold Provider Portal!

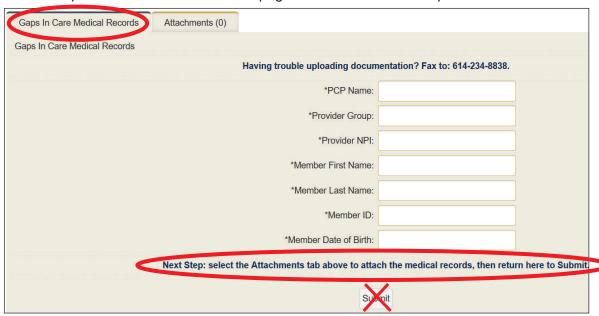
This site will allow you to:

- Verify eligibility and coverage
- View claims history and payment status
- Ask a Claim, Eligibility or Benefit Question
- Special Investigation Unit: upload requested medical records or documents



Close Gaps In Care

3. On the 'Gaps In Care Medical Records' page enter content in all required fields.



Note: do not hit the submit button at this point. Instead, select the Attachments tab above.



Online Submission of Medical Records for Stars and HEDIS Gaps In Care (continued)

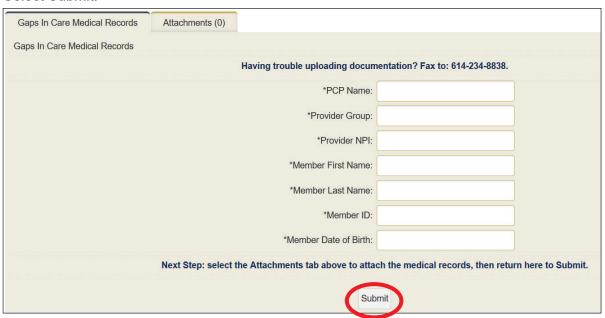
4. Select browse to select the file, then select the Add button.



5. After the file(s) finish uploading it will indicate the number of attachments in the Attachments tab. Now, click the Gaps In Care Medical Records tab.



6. Select Submit.



Frequently Asked Questions

Who reviews the medical records?

MediGold uses our own professionals and/or partners with expert organizations working on our behalf. All professionals reviewing the medical records will treat your patient's protected health information (PHI) with total protection and confidentiality.

Is a review of medical records permitted by HIPAA without a signed member release?

HIPAA allows providers to disclose PHI to another covered entity without a signed release in reference to health care operations. These operations include activities such as quality assessment and improvement and health plan performance evaluations. HEDIS scores are a significant part of these activities.

When will I be asked to provide the records for HEDIS?

Records may be requested throughout the year. However, the majority of records are requested and reviewed between early February to middle May each year.

Is my participation in data collection mandatory and what am I required to do?

Yes. Network participants are contractually required to provide medical record information so we may fulfill our state and federal regulatory obligations. You and your staff are responsible for responding to MediGold's request for medical record documentation in a timely manner. You may provide the records yourself, or schedule time with one of our professionals to come into your office to collect a copy of the records on-site. If a patient included on the list is not part of your practice, you should notify us immediately.

Should I allow a record review for a patient who is no longer with MediGold or a patient who is deceased?

Yes. Medical record reviews may require data collection on the services obtained over multiple years when the patient was receiving benefits from MediGold.

Am I required to provide medical records for a patient who was seen by a provider who has retired, died or moved?

Yes. Data collection includes reviewing medical records as far back as 10 years (including before your patient was a MediGold member). Archived medical records and data may be required to complete data collection.

If you have further questions, please contact: <u>StarsAndHEDIS@mchs.com</u>.

HEDIS-Related Star Measures

Breast Cancer Screening (BCS)

Breast Cancer Screening (BCS)	Percentage of women 50-74 years of age who had a mammogram to screen for breast cancer any time on or between October 1 two years prior to the measurement year, and December 31 of the measurement year. This measure evaluates primary screening, not diagnostic screenings.
Star Weight:	1
Provider Actions:	Mammogram to screen for cancer in the time period listed in measure.
Coding:	
CPT 4	77055 - 77057 77061 - 77063 77065 - 77067
HCPCS	G0202 G0204 G0206
ICD9PCS	87.36 87.37
Revenue	0401 0403
Exclusions:	Members with advanced illness and frailty. Members with a history of bilateral or two unilateral mastectomies. Members in hospice. Members 66 years of age and older as of December 31 of the measurement year who are enrolled in an I-SNP any time during the measurement year or living long term in an institution (LTI).

Colorectal Cancer Screening (COL)

Colorectal Cancer Screening (COL)	Percentage of members 50-75 years of age who had appropriate screening for colorectal cancer.
Star Weight:	1
Provider Actions:	Annual FOBT or FIT during the measurement year.
	Flexible sigmoidoscopy during the measurement year or the four years prior to the measurement year.
	Colonoscopy during the measurement year or the nine years prior to the measurement year.
	CT Colonography during the measurement year or the four years prior.
Coding:	
LOINC	Noninvasive colorectal cancer DNA and occult blood screening [Interpretation] in Stool Narrative – 77353-1
	Noninvasive colorectal cancer DNA and occult blood screening [Presence] in Stool – 77354-9
CPT 4	FOBT – 82270, 82274
	Flexible Sigmoidoscopy – 45330-45335, 45337-45342, 45345-45347, 45349, 45350
	FIT – 81528
	Colonoscopy - 44388-44394, 44397, 44401-44408, 45355, 45378-45393, 45398
	CT Colonoscopy – 74261-74263
HCPCS	FOBT - G0328
	Flexible Sigmoidoscopy – G0104
	FIT – G0464
	Colonoscopy – G0105, G0121
	Colorectal cancer screening; stool-based DNA and fecal occult hemoglobin (e.g., kras, ndrq4 and
	bmp3) -G0464
SNOMED CT US Edition	Stool DNA-based colorectal cancer screening positive (finding) –708699002
ICD-9-CM Procedure	Flexible Sigmoidoscopy – 45.24
Exclusions:	Members with advanced illness and frailty.
	Members with a diagnosis of colorectal cancer or total colectomy are excluded.
	Members in hospice.
	Members 66 years of age and older as of December 31 of the measurement year who are enrolled
	in an I-SNP any time during the measurement year or living long term in an institution (LTI).

Controlling Blood Pressure (CBP)

Controlling Blood Pressure (CBP)	Percentage of members 18-85 years of age who had a diagnosis of hypertension (HTN) and whose BP was adequately controlled (<140/90 mm Hg) during the measurement year.			
Star Weight:	1			
Provider Actions:	The most recent BP reading during th hypertension	The most recent BP reading during the measurement year on or after the second diagnosis of hypertension		
Coding				
CPT 2	Systolic BP <130 mmHg.	3074F		
	Systolic BP 130-139 mmHg.	3075F		
	Systolic BP ≥140 mmHg.	3077F		
	Diastolic BP <80 mmHg.	3078F		
	Diastolic BP 80-89 mmHg.	3079F		
	Diastolic BP ≥90 mmHg.	3080F		
Exclusions:	Members with advanced illness and frailty. Members in hospice. Members with evidence of End-stage Renal Disease (ESRD) or kidney transplant on or prior to December 31 of the measurement year. Members 66 years of age and older as of December 31 of the measurement year who are enrolled in an I-SNP any time during the measurement year or living long term in an institution (LTI).			

Comprehensive Diabetes Care (CDC)

Comprehensive

The percentage of members 18-75 years of age with diabetes (Type 1 and Type 2) who had each of the following:

- Hemoglobin A1c (HbA1c) ≤ 9.0%.
- Retinal Eye Exam performed by an eye care professional.
- Medical attention for nephropathy either evidence of nephrology or a nephropathy screen.
- BP Control (<140/90 mm Hg).

HbA1C Testing Star Weight:

Provider Actions:

Annual documentation of most recent date and result of HbA1c.

Members who were dispensed insulin or hypoglycemics/ antihyperglycemics on an ambulatory basis during the measurement year or the year prior to the measurement year

Diabetes Medications

Description		Prescription	
Alpha-glucosidase inhibitors	Acarbose	Miglitol	
Amylin analogs	Pramlinitide		
Antidiabetic combinations	Alogliptin-metformin Alogliptin-pioglitazone Canagliflozin-metformin Dapagliflozin-metformin Empaglifozin-linagliptin	Empagliflozin-metformin Glimepiride-pioglitazone Glipizide-metformin Glyburide-metformin Linagliptin-metformin	Metformin-pioglitazone Metformin-repaglinide Metformin-rosiglitazone Metformin-saxagliptin Metformin-sitagliptin
Insulin	Insulin aspart Insulin aspart-insulin aspart protamine Insulin degludec Insulin detemir Insulin glargine Insulin glulisine	Insulin isophane human Insulin isophane-insulin r Insulin lispro Insulin lispro-insulin lispro Insulin regular human Insulin human inhaled	
Meglitinides	Nateglinide	Repaglinide	
Glucagon-like peptide-1 (GLP1) agonists	Dulaglutide Exenatide	Albiglutide Liraglutide	
Sodium glucose cotransporter 2 (SGLT2) inhibitor	Canagliflozin	Dapagliflozin	Empagliflozin
Sulfonylureas	Chlorpropamide Glimepiride	Glipizide Glyburide	Tolazamide Tolbutamide
Thiazolidinediones	Pioglitazone	Rosiglitazone	
Dipeptidyl peptidase-4 (DDP-4) inhibitors	Alogliptin Linagliptin	Saxagliptin Sitagliptin	

Note: Glucophage/metformin as a solo agent is not included because it is used to treat conditions other than diabetes; members with diabetes on these medications are identified through diagnosis codes only.

	L	_
Coding:		
CPT 2	Level <7.0%	3044F
	Level >9.0%	3046F
	Level >7.0<8.0%	3051F
	Level <u>></u> 8.0%<9.0%	3052F
CPT4	83036-83037	
Exclusions:	Members with advance	ced illness and frailty for all CDC measures.

Member in hospice.

Members 66 years of age and older as of December 31 of the measurement year who are enrolled in an I-SNP any time during the measurement year or living long term in an institution (LTI).

Retin	ıal	Ey	/e	Ex	am
	St	ar	W	eia	ht:

Dravidar Actions	Annual daguman	tation of most recent ratingles dilete	d ava avam ar	
Provider Actions:		tation of most recent retinal or dilate	,	o.r
	documentation of a negative retinal or dilated eye exam in prior year or chart/photograph of retinal abnormalities indicating date when the fundus photography was			
	performed and evidence it was reviewed by an eye care professional (optometrist or			
	ophthalmologist) in current year.			
Coding:	opritriali fiologist)	in current year.		
CPT 2	Diahetic Retinal S	Screening with Eye Care Profession	al· 2022F	2024F, 2026F
0112	Negative Indicators for Diabetic Retinopathy 2023F, 2025F			
		Screening Negative:	3072F	20201
Exclusions:	Member in hospid		00721	
ZXOIGOIOIIOI	Members 66 years of age and older as of December 31 of the measurement year who are enrolled			
	in an I-SNP any time during the measurement year or living long term in an institution (LTI).			
	,	j	<u> </u>	,
Evidence of Treatment of				
Nephropathy				
Star Weight:	1			
Provider Actions:	Annual documen	tation of one of the following service	s in the measure	ement year:
	 Nephrop 	pathy screening or monitoring test.		
		e of nephropathy.		
		st for albumin or protein		
	 At least 	1 dispensing medication of either ar	n ACE inhibitor of	or ARB.
	ACE Inhibitor and A	ARB Medications		
	Description	Pr	escription	
	Angiotensin	Benazepril	nopril • Peri	ndopril • Ramipril
	converting	Captopril		
	enzyme inhibitors		101 5 0000	
	Angiotensin II	Azilsartan	artan • Teln	nisartan
	inhibitors	Candesartan	nesartan • Vals	artan
	Antihypertensive	Amlodipine-benazepril Azilsartan-ci	hlorthalidone	Hydrochlorothiazide-moexipril
	combinations			Hydrochlorothiazide-
		hydrochlorothiazide- • Candesartar	• 1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-	olmesartan
		valsartan hydrochlorot		 Hydrochlorothiazide-quinapril
		Amlodipine- hydrochlorothiazide- Captopril-hy	drochlorothiazide	Hydrochlorothiazide-
		olmesartan • Enalapril-hy	drochlorothiazide	telmisartan
		Amlodipine-olmesartan Fosinopril-hy	ydrochlorothiazide	Hydrochlorothiazide-valsartan
			thiazide-irbesartan	Sacubitril-valsartan
		Amlodipine-telmisartan Hydrochloro	thiazide-lisinopril	Trandolapril-verapamil
		Amlodipine-valsartan Hydrochloro	thiazide-losartan	
Coding:	-			-
CPT 2	Positive microalb	uminuria test documented/reviewed	3060F	
-		buminuria test documented/reviewe		
		ouminuria test documented/reviewed		
		tment documentation	3066F,	4010F
CPT 4:	81000 - 81005			
	82042 - 81044			
	84156			
Exclusions:	Evidence of stage	e 4 chronic kidney disease, ESRD, o	or kidney transpl	ant. Member in hospice.
		rs of age and older as of December		
		<u>ime during the measurement year o</u>		

Osteoporosis Management in Women Who Had a Fracture (OMW)

Osteoporosis	The percentage of women 67-85 years of age who suffered a fracture and			
Management in	who had either a bo	one mineral density (BMI	D) test or prescription for a drug	
Women Who Had a	to treat or prevent of	steoporosis in the six n	nonths after the fracture. Note:	
Fracture (OMW)			ncluded in this measure.	
	J			
Star Weight:	1			
Provider Action:			edication therapy to treat osteoporosis	
Codings	within 6 months of a fracti	ure. Allowable every 24 months		
CPT 4	Rone Mineral Density Tes	et: 76077 77078 77080 _ 7708	2 77085 - 77086	
HCPCS	Bone Mineral Density Test: 76977, 77078, 77080 – 77082, 77085 - 77086 Bone Mineral Test: G0130			
1.01.00	Injection, calcitonin salmo	on, up to 400 units	J0630	
	Injection, Denosumab, 1 r		J0897	
	Injection, Ibandronate soc		J1740	
	Injection, Teriparatide, 10	mg	J3110	
	Injection, Zoledronic acid		J3487	
	Injection, Zoledronic acid		J3488	
	Injection, Zoledronic acid		J3489	
LODODOO		not otherwise classified, 1 mg	Q2051	
ICD9PCS	Bone mineral density stud		88.98	
ICD10PCS	Ultrasonography of Right Ultrasonography of Left S		BP48ZZ1 BP49ZZ1	
	Ultrasonography of Right		BP4GZZ1	
	Ultrasonography of Left E		BP4HZZ1	
	Ultrasonography of Right		BP4LZZ1	
	Ultrasonography of Left W		BP4MZZ1	
	Ultrasonography of Right		BP4NZZ1	
	Ultrasonography of Left H		BP4PZZ1	
	Plain Radiography of Right Hip, Densitometry Plain Radiography of Left Hip, Densitometry		BQ00ZZ1	
			BQ01ZZ1	
	Plain Radiography of Righ		BQ03ZZ1	
	Plain Radiography of Left Femur, Densitometry Plain Radiography of Cervical Spine, Densitometry Plain Radiography of Thoracic Spine, Densitometry Plain Radiography of Lumbar Spine, Densitometry		BQ04ZZ1	
			BR00ZZ1	
			BR07ZZ1	
	Plain Radiography of Luft Plain Radiography of Who		BR09ZZ1 BR0GZZ1	
Provider Action:				
Trovider Action.	1	otation of the following prescribed medications listed below:		
	Description	P	rescription	
	Biphosphonates	Alendronate	Risedronate	
	International Participation	Alendronate-cholecalcife	erol • Zoledronic acid	
		Aleridionale-cholecalche	• Zoledionic acid	
		 Ibandronate 		
	Other agents	Albandronate	Raloxifene	
	Other agents	Form and court of	0000 00 00000	
		Calcitonin	Teriparatide	
		Denosumab		
Evolucionos		91 17 9		
Exclusions:	Members with advanced		700 days (04 mands -)	
	Episode Start Date (IESD		ne 730 days (24 months) prior to the Index	
			erapy during the 364 days (12 months)	
	prior to the IESD.		or any and our days (12 months)	
	1 •	dispensed prescription or had a	an active prescription to treat osteoporosis	
	during the 365 days (12 n	during the 365 days (12 months) prior to the IESD.		
	Member in hospice.			
			en e e	
	Members 66 years of age		of the measurement year who are enrolled ng long term in an institution (LTI).	

Plan All-Cause Readmission (PCR)

Plan All-Cause Readmissions (PCR)	Those with an acute inpatient stay during the measurement year that were followed-up by an unplanned acute readmission for any diagnosis within 30-days and the predicted probability of an acute readmission.
Star Weight:	No weight, currently display for CY 2019 and 2020
Provider Action:	Outreach to your patient and see them within 7 days of discharge. Reconcile current and discharge medications, when applicable. If medications are prescribed, provide education to the patient, including side effects, importance of adherence, etc.
Exclusions:	None

Medication Reconciliation Post-Discharge (MRP)

Medication Reconciliation Post- Discharge (MRP)	Percentage of discharges from January 1 to December 1 of the measurement year for members 18 years of age and older for whom medications were reconciled on or within 30 days of discharge (31 total days).
Star Weight:	1
Provider Action:	 Documentation in the outpatient medical record must include evidence of medication reconciliation and the date on which it was performed. Any of following meets criteria: Documentation of the current medications with a notation that the provider reconciled the current and discharge medications. Documentation of the current medications with a notation that references the discharge medications (e.g., no changes in medications since discharge, same medications at discharge, discontinue all discharge medications). Documentation of the member's current medications with a notation that the discharge medications were reviewed. Documentation of a current medication list, a discharge medication list and notation that both lists were reviewed on the same date of service. Documentation of the current medications with evidence that the member was seen for post-discharge hospital follow-up with evidence of medication reconciliation or review. Documentation in the discharge summary that the discharge medications were reconciled with the most recent medication list in the outpatient medical record. There must be evidence that the discharge summary was filed in the outpatient chart on the date of discharge through 30 days after discharge (31 total days). Notation that no medications were prescribed or ordered upon discharge. Only documentation in the outpatient record meets the intent of the measure, but an outpatient visit is not required.
Coding:	Luure .
CPT 2	1111F
CPT4	99495 99496
Exclusions:	Member in hospice. Members 66 years of age and older as of December 31 of the measurement year who are enrolled in an I-SNP any time during the measurement year or living long term in an institution (LTI).

Transitions of Care (TRC)

Transitions of Care (TRC)	Percentage of discharges for members 18 and older who had each of the following.			
	Four rates are reported:			
Star Weight:	Display			
Provider Action:	 Notification of Inpatient Admission. Documentation of receipt of notification of inpatient admission on the day of admission or the following day. Receipt of Discharge Information. Documentation of receipt of discharge information on the day of discharge or the following day. At a minimum, the discharge information must include all of the following: The practitioner responsible for the member's care during the inpatient stay. Procedures or treatment provided. Diagnoses at discharge. Current medication list. Testing results, or documentation of pending tests or no test pending. Instructions to the PCP or ongoing care provider for patient care. Discharge instructions provided to the member to follow-up with their PCP does not meet criteria. Patient Engagement After Inpatient Discharge. Documentation of patient engagement provided within 30 days after discharge. Medication Reconciliation Post-Discharge. Documentation of medication reconciliation on the date of discharge through 30 days after discharge (31 total days). 			
Coding:				
CPT 2	1111F			
CPT4	99495 99496			
Exclusions:	Members with advanced illness & frailty. Member in hospice. Members 66 years of age and older as of December 31 of the measurement year who are enrolled in an I-SNP any time during the measurement year or living long term in an institution (LTI).			

Disease-Modifying Anti-Rheumatic Drug Therapy for Rheumatoid Arthritis (ART)

Disease-Modifying Anti-Rheumatic Drug Therapy for Rheumatoid Arthritis (ART)	Percentage of members 18 years of age and older who were diagnosed with rheumatoid arthritis and who were dispensed at least one ambulator prescription for a disease modifying anti-rheumatic drug (DMARD).			ory	
Star Weight:	1				
Provider Action:	Notation of the following prescribed DMARD medications listed below:				
	Description		Prescription		
	5-Aminosalicylates	Sulfasalazine			
	Alkylating agents	Cyclophosphamide			
	Aminoquinolines	Hydroxychloroquine	•		\Box
	Anti-rheumatics	Auranofin Leflunomide	Methotrexate Penicillamine		
	Immunomodulators	Abatacept Adalimumab Anakinra Certolizumab	Certolizumab pegolEtanerceptGolimumabInfliximab	Rituximab Tocilizumab	
	Immunosuppressive agents	Azathioprine	Cyclosporine	Mycophenolate	
	Janus kinase (JAK) inhibitor	Tofacitinib			
	Tetracyclines	Minocycline			
Coding:					
HCPCS	under the direct supervi	sion of a physician, not 20 mg	for Medicare when drug adn for use when drug is self-adn be used for Medicare when c	ministered) J013	35
	Injection, certolizumab pegol, 1 mg (code may be used for Medicare when drug administered under the direct supervision of a physician, not for use when drug is self-administered) Injection, etanercept, 25 mg (code may be used for Medicare when drug administered			Irug is self- ministered J143	38
			for use when drug is self-adr		00
	Injection, gold sodium the Injection, golimumab, 1		2	J160 J160	
	Injection, infliximab, exc			J174	
	Injection, tocilizumab, 1			J326	
	Cyclosporine, oral, 100	mg		J750	
	Cyclosporine, oral, 25 m			J751	
	Cyclosporine, parentera			J751	
	Mycophenolate mofetil,			J751	
	Mycophenolic acid, oral Methotrexate sodium, 5			J751 J925	
	Methotrexate sodium, 5			J925 J926	
	Injection, rituximab, 100			J931	
Exclusions:	Members with advanced Members in hospice.	d illness and frailty.			
	Members 66 years of ag	ge and older enrolled in	an I-SNP or living long-term	in institutional setting	ngs

Statin Therapy for Patients with Cardiovascular Disease

Statin Therapy for Patients with Cardiovascular Disease (SPC)	The percentage of males 21-75 years of age and females 40-75 years of ag with clinical atherosclerotic cardiovascular disease (ASCVD) who receive high or moderate-intensity statin medication during the measurement year			
Star Weight	1	1		
Provider Action:		o adhere at least 80% or more to the gh-intensity or moderate-intensity sta		
	Description	Pres	cription	
	High-intensity statin therapy	 Atorvastatin 40-80 mg Amlodipine-atorvastatin 40-80 mg Ezetimibe-atorvastatin 40-80 mg 	Rosuvastatin 20-40 mgSimvastatin 80 mgEzetimibe-simvastatin 80 mg	
	Moderate-intensity statin therapy	Atorvastatin 10-20 mg Amlodipine-atorvastatin 10-20 mg Ezetimibe-atorvastatin 10-20 mg Rosuvastatin 5-10 mg Simvastatin 20-40 mg Ezetimibe-simvastatin 20-40 mg Niacin-simvastatin 20-40 mg	Sitagliptin-simvastatin 20-40 mg Pravastatin 40-80 mg Lovastatin 40 mg Niacin-lovastatin 40 mg Fluvastatin XL 80 mg Fluvastatin 40 mg bid Pitavastatin 2–4 mg	
Exclusions (When appropriate diagnosis code on claim):	Rhabdomyolysis and End Members dispensed with measurement year or the Members diagnosed with measurement year. Member in hospice. Members 66 years of age	Muscular Pain and Disease to includ l-stage Renal Disease (ESRD). at least one prescription for clomiph year prior to the measurement year Cirrhosis during the measurement y	nene (Estrogen Agonist) during the cyear or the year prior to the e measurement year who are enrolled	

Part D Measures

Medication Adherence - Cholesterol	had at lea	The percentage of Medicare Part D beneficiaries, 18 years or older, who had at least two fills of medication(s) on unique dates of services (DOS) and were 80% or more adherent to their statin medication			
Star Weight	3	3			
Provider Action:			roughout the year for the for Table PDC-STA-A: Statin Medication	Statins	ibed statin
			Statin Medicati	ons	
		fluvastatin	pitavastatin	rosuvastatin	
		atorvastatin (+/- amlodipine, ezetimibe)		pravastatin	3
		simvastatin (+/-ezetimibe, niacin, sitagliptin) lovastatin (+/- niacin)			7
		Note: The active ingredients are limited to oral formulations only.			
Exclusions:		Beneficiaries enrolled in hospice any time during the measurement period Beneficiaries that have ESRD			

Medication Adherence – Diabetes

The percentage of Medicare Part D beneficiaries, 18 years or older, who had at least two fills of medication(s) on unique dates of services (DOS) and were 80% or more to their diabetes medications.

Star Weight

1 3

Provider Action:

Always prescribe 90 days when possible. Encourage patients to adhere to their prescribed drug therapy 80% or more throughout the year for the following medications: Biguanides, Sulfonylureas, Thiazolidinediones, DPP-IV inhibitors, Incretin Mimetics, Meglitinides, and SGLT2 inhibitors:

Table PDC-DR-A: Biguanide Medications

Biguanides

metformin (+/- alogliptin, canagliflozin, dapagliloflozin, empagliflozin, ertugliflozin, glipizide,

glyburide, linagliptin, pioglitazone, repaglinide, rosiglitazone, saxagliptin, sitagliptin)

Note: Active ingredients are limited to oral formulations only.

Excludes nutritional supplement/dietary management combination products

Table PDC-DR-B: Sulfonylureas Medications

Sulfonylurea Medications and Combinations			
chlorpropamide	glipizide (+/- metformin)	tolazamide	
glimepiride (+/- pioglitazone)	glyburide (+/- metformin)	tolbutamide	

Note: Active ingredients are limited to oral formulations only (includes all salts and dosage forms).

Table PDC-DR-C: Thiazolidinediones

Thiazolidinedione Medications and Combinations	
pioglitazone (+/- alogliptin, glimepiride, metformin)	rosiglitazone (+/- metformin)

Note: Active ingredients are limited to oral formulations only.

Table PDC-DR-D: DPP-4 Inhibitors

DPP-4 Medications and Combinations			
alogliptin (+/- metformin, pioglitazone)	saxagliptin (+/- metformin, dapagliflozin)	sitagliptin (+/- metformin, simvastatin)	

Note: Active ingredients are limited to oral formulations only.

Table PDC-DR-E: Incretin Mimetics

DPP-4 Medications and Combinations		
albiglutide	exenatide	lixisenatide
dulaglutide	liraglutide	semaglutide

Note: Active ingredients are limited to oral formulations only.

Table PDC-DR-F: Meglitinides

	Meglinitides and Combinations		
nateglinide	repaglinide (+/-metformin)		

Note: Active ingredients are limited to oral formulations only.

Exclusions:

Beneficiaries who have one or more of the following prescriptions for insulin in the measurement period listed below.

Table PDC-H: Insulin Exclusion

Insulins	
insulin aspart (+/-insulin aspart protamine)	insulin glargine (+/- lixisenatide)
insulin regular (including inhalation powder)	insulin glulisine
insulin lispro (+/- insulin lispro protamine)	insulin degludec (+/- liraglutide)
insulin isophane (+/- regular insulin)	insulin deternir

Note: The active ingredients are limited to inhaled and injectable formulations only.

Beneficiaries enrolled in hospice any time during the measurement period. Beneficiaries that have ESRD

Medication Adherence - Hypertension-RAS Antagonists	The percentage of Medicare Part D beneficiaries, 18 years or older, who had at least two fills of medication(s) on unique dates of services (DOS) and were 80% or more to a RAS antagonist			
Star Weight	Always prescribe 90 days when possible. Encourage patients to adhere to their prescribed AC			
Provider Action:	inhibitors, ARBs, or Direct Renin Inhibitors 80%	urage patients to adhere to their prescribed ACE or more throughout the year.		
	Table PDC-RASA-A: Renir	Angiotensin System (RAS)		
	Antag	gonists		
	Direct Renin Inhibitor Me	edications and Combinations		
	aliskiren (+/- amlodipine, hydrochlorothiazide)			
	ARB Medications	s and Combinations		
	azilsartan (+/- chlorthalidone)	irbesartan (+/- hydrochlorothiazide)		
	candesartan (+/- hydrochlorothiazide)	losartan (+/- hydrochlorothiazide)		
	eprosartan (+/- hydrochlorothiazide)	olmesartan (+/- amlodipine, hydrochlorothiazide)		
	telmisartan (+/- amlopdipine, hydrochlorothiazide)	valsartan (+/- amlodipine, hydrochlorothiazide nebivolol)		
	ACE Inhibitor Medications and Combination Products			
	benazepril (+/- amlodipine, hydrochlorothiazide)	lisinopril (+/- hydrochlorothiazide)		
	captopril (+/- hydrochlorothiazide)	moexipril (+/- hydrochlorothiazide)		
	enalapril (+/- hydrochlorothiazide)	perindopril (+/- amlodipine)		
	fosinopril (+/- hydrochlorothiazide)	quinapril (+/- hydrochlorothiazide)		
	ramipril	trandolapril (+/- verapamil)		
	Note: Active ingredients are limited to oral formulations only. Excludes nutritional supplement/dietary management combination products.			
Exclusions:	Beneficiaries that received one of more prescription claims for Sacubitril/Valsartan.			
	Table PDC-RASA-B Exclusion: Sacubitril/Valsartan			
	ARB/Neprilysin Inhibitor Combination Medication sacubitril/valsartan			
	Beneficiaries enrolled in hospice any time during the measurement period Beneficiaries that have ESRD			

Statin Therapy for Patients with Diabetes (SUPD)	The percentage of Medicare Part D beneficiaries, ages 40-75 years, dispensed at least two diabetes medication fills who received a statin medication fill.				
Star Weight	3				
Provider Action:	<u></u>	Table SUPD-A: Di	abetes Medica	tions	
		Biguanides and Biguan	ide Combination Pr	oducts	
	metformin (+/- alogliptin, car repaglinide, rosiglitazone, si		ozin, ertugliflozin, glipiz	zide, glyburide, linagliptin, pioglitzone	
	8	Sulfonylureas and Sulfony	lurea Combination	Products	
	chlorpropamide	glyburide (+/- metformin)	tolazamide	glipizide (+/- metformin)	
	tollbutamide	glimepiride (+/- pioglitazone)		1	
		Megl	itinides		
	nateglinide	repaglinide (+/- metformin)	Ī	Ï	
		Alpha- Glucos	sidase Inhibitors	1	
	acarbose	miglital		1	
	Th	iazolidinediones and Thiazol	idinedione Combin	ation Products	
	pioglitazone (+/- alogliptin, g		rosiglitazone (+/- me		
			metic Agents		
	albiglutide	exenatide	semaglutide	dulaglutide	
		lec) lixisenatide (+/- insulin glargi	1		
	Amylin Analogs				
	pramlintide	7	- Tallongs	T	
		DPP-4	Inhibitors		
	alogliptin (+/- metformin, pio		linagliptin (+/- empag	diflozin metformin)	
	sitagliptin (+/- metformin, ertugliflozin, simvastatin) saxagliptin (+/-dapagliflozin, metformin)				
			ulins	3	
	insulin aspart (+/-insulin asp		insulin detemir	insulin glulisine	
	insulin lispro (+/- insulin lisp		GATEST RESPONSE IN	liraglutid insulin glargine (+/- lixisena	
	insulin isophane (+/- regular	The second secon	insulin regular (including inhalation powder)		
	mount toophone (17- regular	Sodium Glucose Co-trans			
	canagliflozin (+/- metformin)		Market Street Street		
	empagliflozin (+/- linagliptin,		ertugliflozin (+/- sitag	liptin, metformin)	
	Note: The active ingredients are limited to oral, inhalation and injectable formulations only (includes all dosage forms; excludes nutritional supplement/dietary management combination products).				
Exclusions:	Beneficiaries enrolled in hospice any time during the measurement period. Beneficiaries that have ESRD.				
	Table SUPD-B: Statin Medications				
		Statin Med			
	fluvastatin pitavastatin rosuvastatin			X	
		- amlodipine, ezetimibe)	pravas		
	simvastatin (+/-ezetimibe, niacin, sitagliptin) lovastatin (+/- niacin) Note: The active ingredients are limited to oral formulations only.			tin (+/- niacin)	

Advanced Illness and Frailty

Patients with an advanced illness diagnosis or limited life expectancy may not benefit from recommended services required to meet certain quality measures. Unnecessary tests and treatments may be burdensome or even harmful to these patients. To account for this the National Committee for Quality Assurance (NCQA) updated their specifications to allow exclusions for advanced illness and frailty.

To qualify, patients must have at least one of the following in the measurement year or year prior:

- Two outpatient claims on different dates of service with an advanced illness code
- One inpatient claim with an advanced illness code
- One filled prescription for a dementia medication

AND

At least one claim with a frailty diagnosis or treatment claim in the measurement year.

Exclusions can be applied to the following Star Measures:

Breast Cancer Screening (BCS)

Colorectal Cancer Screening (COL)

Comprehensive Diabetes Care (CDC)

Controlling Blood Pressure (CBP)*

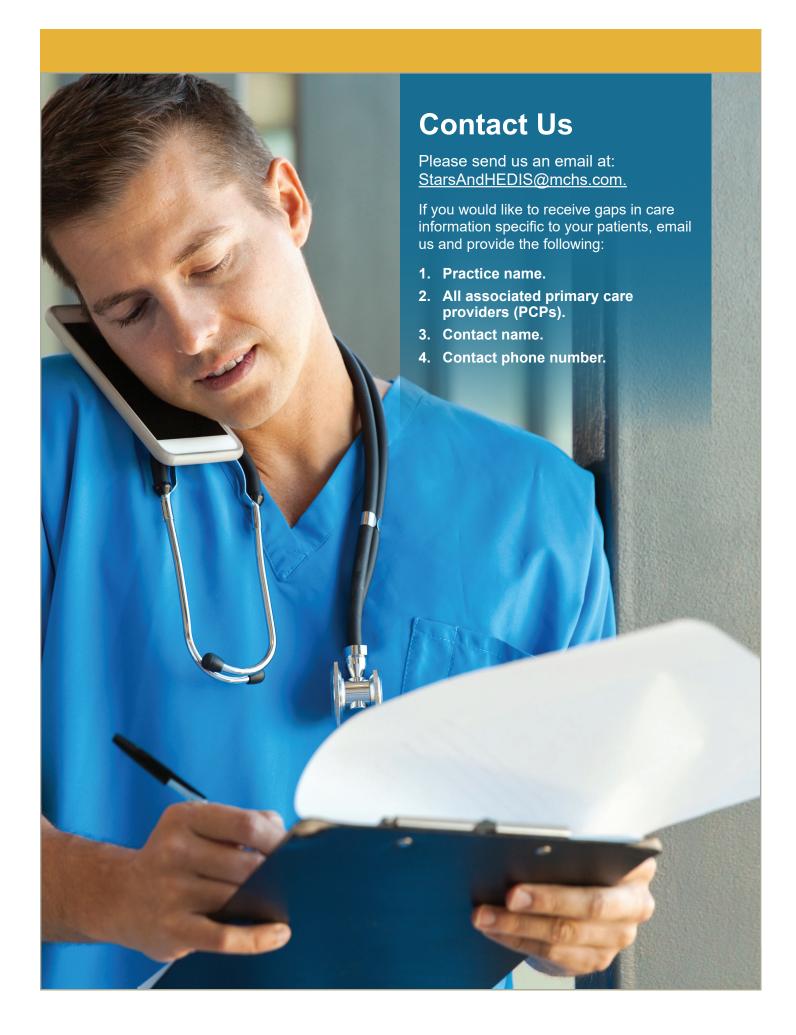
Disease-Modifying Anti Rheumatic Drug Therapy for Rheumatoid Arthritis (ART) Osteoporosis Management in Women with a

Fracture (OMW)*

Statin Therapy for Patients with Cardiovascular Disease (SPC)*

*Patients age 81 and older can be excluded with a frailty diagnosis or treatment alone.

For a complete listing of advanced illness and frailty codes please visit MediGold.com.



Notes:				
·			·	

