(ST) step therapy (PA) prior auth

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Humana	FMOLHS Plan	UHC	BCBS	Drug Class	Medication (AWP cost / 30 days)	Clinical Points *Adherence to < 2gm Na ⁺ and diabetic diets *Sodium restriction enhances the effect of some anti- hypertensive meds
2-vial	3-vial			Thiazide &	Chlorothiazide	1. Thiazides are less effective as GFR declines but can be
1	1	1	\$0	Thiazide - Like	Chlorthalidone	added to loop diuretics for enhanced diuresis
1	1	1	\$0	Diuretics	Hydrochlorothiazide	2. Chlorthalidone and Indapamide are long-acting
1	1	1	\$0		Indapamide	3. Increases risk of hypokalemia
2	1	1	\$0		Metolazone	
2	1	1	\$0	Loop Diuretics	Bumetanide	1. Equivalency: Bumetanide 1mg = Torsemide 20mg = Fu- rosemide 40mg
1 <i>-tabs</i> 2-soln	1	1	\$0		Furosemide	 2. Torsemide, longer acting, may be preferred 3. Thiazides can be added to loop diuretics for enhanced
2	1	1	\$0		Torsemide	diuresis
	1	2	1	Aldosterone Antagonists	Eplerenone (>\$130)	1. Preferred for primary aldosteronism and resistant hypertension
3 (PA)	3 (PA)	4(PA)			Finerenone (Kerendia >\$790)	2. Risk of hyperkalemia
1	1	1	\$0		Spironolactone	3. Finerenone is FDA approved reduced the risk of CV and
2	1	1	\$0		Spironolactone / HCTZ	 renal outcomes in pts with T2DM & CKD (FIDELIO-DKD) 4. Finerenone use recommended when eGFR ≥25ml/mir albuminuria ≥30mg/g (≥3mg/mmol), and a normal serur potassium concentration
1	1	1	\$0	ACE inhibitors	Benazepril	1. Use for CKD with urine albumin >300mg/24hr, or
3	1	1	1	ACE/ARB can re- duce the risk of	Captopril	CKD/DM pts with urine albumin >30-300mg/24hr
1	1	1	\$0		Enalapril	2. Monitor for hypotension, decreased GFR, and hyper-
1	1	1	\$0	developing micro-	Fosinopril	kalemia
1	1	1	\$0	albuminuria or progression to macroalbuminu- ria	Lisinopril	3. Not recommended for use in combination with ARB or
1	1		1		Moexipril	Renin Inhibitors (increases CV and renal risks)
2	1	2	1		Perindopril	4. Consider use even if GFR <30ml/min due to reno-protec-
1	1	1	\$0		Quinapril	tive properties
1	1	1	\$0		Ramipril	5. Increases risks of hyperkalemia
1	1	1	1		Trandolapril	6. Utilize K ⁺ binders to remain on ACE/ARB therapy
3	1	3	1	ARBs ACE/ARB can re-	Candesartan	 1. Use for CKD with urine albumin >300mg/24hr, or CKD/DM pts with urine albumin >30-300mg/24hr
				duce the risk of	Azilsartan (Edarbi >\$310)	2. Monitor for hypotension, decreased GFR, and hyper-
	1			developing micro- albuminuria or	Eprosartan	kalemia 3. Not recommended for use in combination with Ace In-
1	1	1	\$0	progression to	Irbesartan	hibitors or Renin Inhibitors (increases CV and renal risks)

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	erapy (PA) p			Drug Class	Medication	Clinical Points
ŋ	S			Diug Class	(AWP cost / 30 days)	
าลท	ОГН		S		(AWF LOST / SO days)	*Adherence to < 2gm Na ^{$+$} and diabetic diets
Humana	FMOLHS Plan	инс	BCBS			*Sodium restriction enhances the effect of some anti-
						hypertensive meds
1	1	1	\$0	macroalbuminu- ria	Losartan	4. Consider use even if GFR <30ml/min due to reno-protec-
1	1	2	\$0	110	Olmesartan	 tive properties 5. Increases risks of hyperkalemia
2	1	2	1		Telmisartan	6. Utilize K ⁺ binders to remain on ACE/ARB therapy
1	1	2	\$0		Valsartan	 7. Patients with history of ACE angioedema may try ARB after six-week washout
OTC	OTC	οτο		Vitamin D prope	*Chalacalaifaral (D2)	
ОТС	OTC	OTC		Vitamin D preps *nonactivated	*Cholecalciferol (D3)	1. *Use when evidence of a documented deficiency, use general population guidelines for dosing
2	3	1	1	or activated	*Ergocalciferol (D2)	2. If CKD G4-5 and persistently elevated PTH, use calcitriol
2 - cap 4- soln	2	1	1		Calcitriol (>\$45)	or vitamin D analogs for more direct effect on PTH
4	3		1		Doxercalciferol (Hectorol generic	
	(ST,PA)				>\$350)	
3-vials	1(ST,PA)	1	1		Paricalcitol (Zemplar generic	
4-caps					>\$250)	
	2			Phosphate	Auryxia (>\$1,600)	1. Auryxia may increase serum iron and the risk of alumi-
	1	1-cap	1	Binders	Calcium acetate (>\$130)	num toxicity
ОТС	OTC	OTC	ОТС		Calcium carbonate (>\$10-\$15)	2. Elemental calcium should not exceed 1500mg/24hr
	3		3 (PA)		Lanthanum carbonate (>\$1,200)	3. Renagel may contribute to metabolic acidosis
	1	2	3 (PA)		Sevelamer carbonate (>\$300)	4. Velphoro has minimal increase in serum iron
	2				Sevelamer HCL (Renagel generic >\$650)	
	3	4 (ST)	3 (PA)		Velphoro (>\$1,900)	1
3	1	3 (PA)	2	Potassium	Lokelma (>\$900)	1. Avoid in severe constipation, bowel obstruction, or im-
				Binders		paction
3	1	3	1		Sodium Polystyrene Sulfonate (>\$250)	 Sodium from SPS & Lokelma may exacerbate edema Veltassa may bind magnesium, consider supplementa-
	1	3 (PA)	3		Veltassa (>\$1,200)	tion
			J. J			3. Separation of dosing may be warranted with other med- ications, typically 3hrs before or 3hrs after treatment
					<u> </u>	4. Should not be used as emergency treatments due to de-
						layed onset of action
				SGLT2 inhibi-	Brenzavvy <i>(\$50)</i>	1. SGLT2i class may have more marked effects on de-
3		(ST)		tors	*Invokana (>\$700)	creased hospitalizations for CHF and progression of CKD
		(31)				

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าล	Ŷ			Drug cluss	(AWP cost / 30 days)	*Adherence to < 2gm Na ⁺ and diabetic diets
nar	, C	J	Š			*Sodium restriction enhances the effect of some anti-
Humana	FMOLHS Plan	UHC	BCBS			
						hypertensive meds
3		(ST)		*May delay the progression of di-	Invokamet (>\$700)	2. May increase the risk of mycotic genital infections, in-
3		(ST)		abetic nephropa-	Invokamet XR (>\$700)	- cluding Fournier's gangrene
		(0.7		thy; May reduce		3. To reduce risk of AKI, consider diuretic dose reduction before starting
				CV mortality in	Dapagliflozin (>\$650)	4. Risk of DKA, including euglycemic DKA
			2	pts with estab-	*Farxiga (>\$700)	5. EMP-REG showed Jardiance may prevent new or wors-
				lished CV disease	Dapagliflozin / Metformin (\$>650)	ening nephropathy in 1 out of 16 pts over 3 years. It is FDA
			2	-	Xigduo XR (>\$675)	approved for reducing CV mortality in DM pts with estab-
					Qtern (>\$675)	lished CV disease
3	\$0(PA)	2	2		*Jardiance (>\$730)	6. CREDENCE showed Invokana may prevent the doubling of SCr in 1 out of 31 DM pts over 2.62 years.
3	\$0(PA)	2	2		Synjardy (>\$730)	7. Farxiga when added to ACE or ARB therapy in CKD pts
3	\$0(PA)	2	2		Synjardy XR (>\$730)	may reduce the decline in eGFR of at least 50% and delay the progression to ESRD (DAPA-CKD)
3	\$0(PA)	2	2		Trijardy XR (>\$730)	8. Standards of Diabetes Care 2023 states to use SGLT2i in
3	\$0(PA)	2(ST)	2		Glyxambi (>\$730)	people with an eGFR ≥20ml/min per 1.73m ² to reduce (progression
	\$0(PA)	(ST)			Steglatro (\$>425)	
	\$0(PA)				Segluromet (>\$425)	
					Steglujan (>\$660)	
		4 (ST)		GLP-1 Receptor agonists *May delay the progression of di- abetic nephropa- thy	Adlyxin (>\$800)	 Risk of gallbladder disease or pancreatitis (acute and chronic) C/I in patient or family history of medullary thyroid c
	\$0(PA)	2(PA)			Byetta (<\$1,000)	cer or MEN2 3. Ozempic – monitor for worsening diabetic retinopathy
4	\$0(PA)	2(PA)	2(PA)		Bydureon BCise(>\$1,000)	4. Ozempic: SUSTAIN-6 showed it had a lower incidence of nephropathy (driven by preventing new microalbuminuria) in 1 out of 44 pts over two years. FLOW trial demonstrated
3	\$0(PA)	2(PA)	2(PA)		Mounjaro (>\$1,300)	24% decreased risk of major kidney disease events over 3.4 years in pts with DM & CKD <mark>5. Victoza:</mark> LEADER showed it had a lower incidence of
3	\$0(PA)	2(PA)	2 (PA)		*Ozempic (>\$1,500)	nephropathy (driven by preventing new onset macroall minuria) in 1 out of 67 pts over 4 years; Victoza is FDA

Humana	FMOLHS Plan	UHC	BCBS	Drug Class	Medication (AWP cost / 30 days)	Clinical Points *Adherence to < 2gm Na ⁺ and diabetic diets *Sodium restriction enhances the effect of some anti- hypertensive meds
3	\$0(PA)	2(PA)	2(PA)		Rybelsus (>\$1,150)	approved for reducing the combined endpoints of CV death, MI, or stroke in DM pts with CV disease (LEADER)
3	\$0(PA)	2(PA)	2 (PA)		Trulicity (>\$1,150)	6. Mounjaro - SURPASS-4 post-hoc analysis showed a lower occurrence of the kidney composite (eGFR decline, ESRD,
3		2(PA) (2pk) 3 (PA) (3pk)			*Liraglutide (>\$625 2-pk, >\$950 3- pk)	 death due to kidney failure, & new onset macroalbuminuria) verses insulin glargine 7. Trulicity is FDA approved for reducing the combined endpoints of CV death, MI, or stroke in DM patients with CV disease or at high CV risk (REWIND)

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