

## Changes to MyTruAdvantage's Formulary Updated 09/27/2024

MyTruAdvantage may immediately remove a brand name drug on our Drug List if we are replacing it with a new generic drug that will appear on the same or lower cost sharing tier and with the same or fewer restrictions. Or, when adding the new generic drug, we may decide to keep the brand name drug on our Drug List, but immediately move it to a different cost-sharing tier or add new restrictions. We may not tell you in advance before we make that change, but we will later provide you with information about the specific change(s) we have made. Also, if the Food and Drug Administration deems a drug on our formulary to be unsafe or the drug's manufacturer removes the drug from the market, we may immediately remove the drug from our formulary and provide notice to members who take the drug.

Before we make other changes during the year to our Drug List that affect members currently taking a drug and that require us to provide advance notice, we will notify affected members of the change at least 30 days before the change becomes effective, or at the time the member requests a refill of the drug, at which time the member will receive a one-month supply of the drug.

If you are affected by a change in drug coverage or restriction, depending on the type of change, there may be different options to consider. For example:

You may be able to use another drug on our Drug List to treat your medical condition. Alternative drug(s) are provided below to help your prescriber to find a covered drug that might work for you. Ask your prescriber if one of the possible alternative drug(s) is right for you.

You, your prescriber, or your authorized representative may also ask for an exception. The notice we provide you will also include information on the steps to request an exception. To learn more about coverage decisions and how to ask for an exception, see your *Evidence of Coverage*, or call us at 1-877-403-6035 (TTY: 711), 24 hours a day, 7 days a week.

The table below outlines changes to our formulary that may impact you.

CMS Formulary ID	Effective Date	Drug Name	Change Description	Reason Description	Alternate Drugs and Tier
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24140	2/1/2024	VOTRIENT 200 MG ORAL TABLET	BRAND DELETION, ADD FRF GENERIC	REMOVAL OF BRAND NAME DRUG FROM FORMULARY DUE TO ADDITION OF NEW GENERIC EQUIVALENT	PAZOPANIB HCL 200 MG ORAL TABLET-5
24140	2/1/2024	ALPHAGAN P 0.1 % OPHTHALMIC DROPS	BRAND DELETION, ADD FRF GENERIC	REMOVAL OF BRAND NAME DRUG FROM FORMULARY DUE TO ADDITION OF NEW GENERIC EQUIVALENT	BRIMONIDINE TARTRATE 0.1 % OPHTHALMIC DROPS-2
24140	4/1/2024	PROLENSA 0.07 % OPHTHALMIC DROPS	BRAND DELETION, ADD FRF GENERIC	REMOVAL OF BRAND NAME DRUG FROM FORMULARY DUE TO ADDITION OF NEW GENERIC EQUIVALENT	BROMFENAC SODIUM 0.07 % OPHTHALMIC DROPS-3
24140	4/1/2024	RISPERDAL CONSTA 37.5MG/2ML INTRAMUSC. VIAL	BRAND DELETION, ADD FRF GENERIC	REMOVAL OF BRAND NAME DRUG FROM FORMULARY DUE TO ADDITION OF NEW GENERIC EQUIVALENT	RISPERIDONE ER 37.5MG/2ML INTRAMUSC. VIAL-5
24140	4/1/2024	RISPERDAL CONSTA 50 MG/2 ML INTRAMUSC. VIAL	BRAND DELETION, ADD FRF GENERIC	REMOVAL OF BRAND NAME DRUG FROM FORMULARY DUE TO ADDITION OF NEW GENERIC EQUIVALENT	RISPERIDONE ER 50 MG/2 ML INTRAMUSC. VIAL-5
24140	4/1/2024	RISPERDAL CONSTA 25 MG/2 ML INTRAMUSC. VIAL	BRAND DELETION, ADD FRF GENERIC	REMOVAL OF BRAND NAME DRUG FROM FORMULARY DUE TO ADDITION OF NEW GENERIC EQUIVALENT	RISPERIDONE ER 25 MG/2 ML INTRAMUSC. VIAL-2
24140	4/1/2024	RISPERDAL CONSTA 12.5MG/2ML INTRAMUSC. VIAL	BRAND DELETION, ADD FRF GENERIC	REMOVAL OF BRAND NAME DRUG FROM FORMULARY DUE TO ADDITION OF NEW GENERIC EQUIVALENT	RISPERIDONE ER 12.5MG/2ML INTRAMUSC. VIAL-2
24140	4/1/2024	FORTEO 20MCG/DOSE SUBCUTANE. PEN INJCTR	BRAND DELETION, ADD FRF GENERIC	REMOVAL OF BRAND NAME DRUG FROM FORMULARY DUE TO ADDITION OF NEW GENERIC EQUIVALENT	TERIPARATIDE 20MCG/DOSE SUBCUTANE. PEN INJCTR- 2
24140	4/1/2024	TRACLEER 125 MG ORAL TABLET	BRAND DELETION, ADD FRF GENERIC	REMOVAL OF BRAND NAME DRUG FROM FORMULARY DUE TO ADDITION OF NEW GENERIC EQUIVALENT	BOSENTAN 125 MG ORAL TABLET-5



				REMOVAL OF BRAND NAME DRUG	
		TRACLEER 62.5 MG ORAL	BRAND DELETION, ADD	FROM FORMULARY DUE TO ADDITION	BOSENTAN 62.5 MG
24140	4/1/2024	TABLET	FRF GENERIC	OF NEW GENERIC EQUIVALENT	ORAL TABLET-5
		LEVONORG-ETH ESTRAD-FE			
		BISGLYC 0.1-0.02MG ORAL	DELETION OF DRUG		
24140	5/1/2024	TABLET	FROM FORMULARY	NOT A PART D COVERED DRUG	
				REMOVAL OF BRAND NAME DRUG	BROMFENAC SODIUM
		BROMSITE 0.075 %	BRAND DELETION, ADD	FROM FORMULARY DUE TO ADDITION	0.075 % OPHTHALMIC
24140	5/1/2024	OPHTHALMIC DROPS	FRF GENERIC	OF NEW GENERIC EQUIVALENT	DROPS-3
	, ,			REMOVAL OF BRAND NAME DRUG	
		KORLYM 300 MG ORAL	DDAND DELETION ADD	FROM FORMULARY DUE TO ADDITION	MIFEPRISTONE 300 MG
24140	5/1/2024	TABLET	BRAND DELETION, ADD FRF GENERIC	OF NEW GENERIC EQUIVALENT	ORAL TABLET-5
24140	3/1/2024	TABLET	FRE GEINERIC	OF NEW GENERIC EQUIVALENT	LOTEPREDNOL
				REMOVAL OF BRAND NAME DRUG	ETABONATE 0.2 %
		ALREX 0.2 % OPHTHALMIC	BRAND DELETION, ADD	FROM FORMULARY DUE TO ADDITION	OPHTHALMIC DROPS
24140	5/1/2024		FRF GENERIC	OF NEW GENERIC EQUIVALENT	SUSP-3
24140	3/1/2024	AZOPT 1 % OPHTHALMIC	THI GLIVLING	OF NEW GENERIC EQUIVALENT	3031-3
24140	7/1/2024	DROPS SUSP	FORMULARY DELETION	FORMULARY DELETION	
24140	7/1/2024	DROI 3 3031	TORRIOLARY DELETION		
				REMOVAL OF BRAND NAME DRUG	NITROGLYCERIN 0.4%
2.1.10	6/1/0001	RECTIV 0.4% (W/W) RECTAL	BRAND DELETION, ADD	FROM FORMULARY DUE TO ADDITION	(W/W) RECTAL OINT. (G)-
24140	6/1/2024	, ,	FRF GENERIC	OF NEW GENERIC EQUIVALENT	2
24442	7/4/2024	MITIGARE 0.6 MG ORAL	500041114014051571001	5000 4111 4 007 0 51 5710 41	
24140	7/1/2024		FORMULARY DELETION	FORMULARY DELETION	
24442	0.14.1000.4	TRUSELTIQ 75 MG/DAY ORAL	50 A MUTU DO AMA		
24140	8/1/2024		FDA WITHDRAWAL		
	0 /4 /0 05 :	TRUSELTIQ 125 MG/DAY			
24140	8/1/2024		FDA WITHDRAWAL		
		FARYDAK 10 MG ORAL	DELETION OF DRUG		
24140	8/1/2024	CAPSULE	FROM FORMULARY	PRODUCT WITHDRAWN FROM MARKET	



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		TRUSELTIQ 50 MG/DAY ORAL			
24140	8/1/2024	CAPSULE	FDA WITHDRAWAL		
		FARYDAK 15 MG ORAL	DELETION OF DRUG		
24140	8/1/2024	CAPSULE	FROM FORMULARY	PRODUCT WITHDRAWN FROM MARKET	
		TRUSELTIQ 100 MG/DAY			
24140	8/1/2024	ORAL CAPSULE	FDA WITHDRAWAL		
		FARYDAK 20 MG ORAL	DELETION OF DRUG		
24140	8/1/2024	CAPSULE	FROM FORMULARY	PRODUCT WITHDRAWN FROM MARKET	
			PAYMENT		
			DETERMINATION		
		MOUNJARO 10MG/0.5ML	ADDED TO PRIOR		
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION		
			PAYMENT		
			DETERMINATION		
		MOUNJARO 2.5 MG/0.5	ADDED TO PRIOR		
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION		
			PAYMENT		
			DETERMINATION		
		OZEMPIC 2MG/0.75ML	ADDED TO PRIOR		
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION		
			PAYMENT		
			DETERMINATION		
		MOUNJARO 12.5MG/0.5	ADDED TO PRIOR		
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION		
			PAYMENT		
			DETERMINATION		
			ADDED TO PRIOR		
24140	10/1/2024	RYBELSUS 7 MG ORAL TABLET	AUTHORIZATION		
			PAYMENT		
24140	10/1/2024	RYBELSUS 3 MG ORAL TABLET	DETERMINATION		



			ADDED TO PRIOR
			AUTHORIZATION
			PAYMENT
		_	DETERMINATION
		MOUNJARO 5 MG/0.5ML	ADDED TO PRIOR
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION
			PAYMENT
			DETERMINATION
		OZEMPIC 1/0.75 (3)	ADDED TO PRIOR
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION
			PAYMENT
			DETERMINATION
		OZEMPIC .25 OR 0.5	ADDED TO PRIOR
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION
			PAYMENT
			DETERMINATION
		TRULICITY 4.5 MG/0.5	ADDED TO PRIOR
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION
			PAYMENT
			DETERMINATION
		TRULICITY 0.75MG/0.5	ADDED TO PRIOR
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION
			PAYMENT
			DETERMINATION
		TRULICITY 3 MG/0.5ML	ADDED TO PRIOR
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION
			PAYMENT
			DETERMINATION
		MOUNJARO 7.5 MG/0.5	ADDED TO PRIOR
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION



			PAYMENT		
			DETERMINATION		
		RYBELSUS 14 MG ORAL	ADDED TO PRIOR		
24140	10/1/2024	TABLET	AUTHORIZATION		
			PAYMENT		
			DETERMINATION		
		OZEMPIC 0.25 OR .5	ADDED TO PRIOR		
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION		
			PAYMENT		
			DETERMINATION		
		TRULICITY 1.5 MG/0.5	ADDED TO PRIOR		
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION		
			PAYMENT		
			DETERMINATION		
		MOUNJARO 15MG/0.5ML	ADDED TO PRIOR		
24140	10/1/2024	SUBCUTANE. PEN INJCTR	AUTHORIZATION		
				REMOVAL OF BRAND NAME DRUG	
		CORLANOR 5 MG ORAL	BRAND DELETION, ADD	FROM FORMULARY DUE TO ADDITION	IVABRADINE HCL 5 MG
24140	10/1/2024	TABLET	FRF GENERIC	OF NEW GENERIC EQUIVALENT	ORAL TABLET-3
				REMOVAL OF BRAND NAME DRUG	
		ENDARI 5 G ORAL POWD	BRAND DELETION, ADD	FROM FORMULARY DUE TO ADDITION	L-GLUTAMINE 5 G
24140	10/1/2024	PACK	FRF GENERIC	OF NEW GENERIC EQUIVALENT	ORAL POWD PACK-5
				REMOVAL OF BRAND NAME DRUG	
		CORLANOR 7.5 MG ORAL	BRAND DELETION, ADD	FROM FORMULARY DUE TO ADDITION	IVABRADINE HCL 7.5
24140	10/1/2024	TABLET	FRF GENERIC	OF NEW GENERIC EQUIVALENT	MG ORAL TABLET-3

<sup>\*</sup>Alternative drug(s) are drugs that you could consider with your prescriber. Only your prescriber can determine alternative drugs that are appropriate for you given the individualized nature of drug therapy. Please consult your prescriber to confirm if this is an appropriate drug for you.