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Drug recall notice for valsartan and valsartan-hydrochlorothiazide (HCTZ) tablets

Some manufacturers have voluntarily recalled generic valsartan and valsartan-HCTZ tablets due to an impurity used in the manufacturing of these medications. The drugs that have been recalled are identified by certain national drug codes (NDCs) and lot numbers.

This recall is intended for the following products:

Manufacturer	National drug	Product	Lot numbers	Expiration	Route
	code (NDC)				
Major Pharmaceuticals	00904-6594-61	valsartan 80 mg	T01795; T01807; T01712; T01625; T01596; T01500; T01466; T01270	July 2018 through May 2019	Oral
	00904-6595-61	valsartan 160 mg	T01646; T01788; T01668; T01524; T01269		Oral
	43547-0367-03	valsartan 40 mg			
	43547-0368-09	valsartan 80 mg			
Solco Healthcare LLC	43547-0369-09	valsartan 160 mg			
	43547-0370-09	valsartan 320 mg			
	43547-0311-09	valsartan/HCTZ 80 mg/12.5 mg		July 2018	
	43547-0312-09	valsartan/HCTZ 160 mg/12.5 mg	All lots	through January 2020	Oral
	43547-0313-09	valsartan/HCTZ 160 mg/25 mg			
	43547-0314-09	valsartan/HCTZ 320 mg/12.5 mg			
	43547-0315-09	valsartan/HCTZ 320 mg/25 mg			
Teva Pharmaceuticals USA	00591-2167-19 00591-2167-30	valsartan 40mg	1196934M; 1238462M; 1268429A; 1196936A; 1238463A; 1270617A	July 2018 through October 2019	Oral
	00591-2168-10 00591-2168-19	valsartan 80mg	1177114A; 1219360M; 1250706A; 1175947M; 1175948M; 1177115A; 1219361A;		



		1240434M; 1250704M	
00591-2169-10 00591-2169-19	valsartan 160 mg	1175922M; 1220826M; 1236294M;	
		1240427M; 1270616A;	
		1177880A; 1220831A; 1263941A	
00591-2170-05 00591-2170-19	valsartan 320 mg	1208000M; 1208001M; 1240425A;	
		1208002A; 1247282M; 1263944M	
00591-2315-19	valsartan/HCTZ 80mg/12.5mg	1191191M; 191192M; 1191193M; 191194M; 1191195M;	
		238466M; 1238467M; 253261M; 1256125M; 1277709M	
00591-2316-19	valsartan/HCTZ 160mg/12.5mg	1191160M; 1191161M; 1191162A; 1219363M; 1219364M; 1219365A; 1225613A; 1233944M; 1233945M; 1253253M;	
00591-2317-19	valsartan/HCTZ 160mg/25mg	1253254M 1191164M; 1191165M; 1191166M; 1191167A; 1225612M; 1250717M; 1256111M; 1288798M	
00591-2318-19	valsartan/HCTZ 320mg/12.5mg	1191185M; 1191186M; 1225615M; 1233948M; 1250718M; 1253257M	

00591-2319-19	valsartan/HCTZ	1191188M;	
	320mg/25mg	1191189M;	
	3 3	1191190M;	
		1199220M;	
		1217576M;	
		1217577M;	
		1217578M;	
		1220832M;	
		1220833M;	
		1247283M;	
		1247284M;	
		1247285M;	
		1247286M;	
		1247287A;	
		1280632M;	
		1280633M	

What this means for you

- Check the product name, manufacturer name and NDC on your prescription. If the information is not listed (NDC or lot number), please contact the pharmacy that filled your prescription.
- If your medicine has been recalled, contact your doctor or pharmacist to discuss treatment options. You may be able to get the same medicine that is **not** part of the recall or switch to another medicine.
- It is important that you contact your doctor to discuss continuing to take the recalled medication until you have a replacement product.
- Adverse reactions or quality problems experienced with the use of this product may be reported to the Food and Drug Administration's (FDA's) MedWatch Adverse Event Reporting program either online, by regular mail or by fax.
 - Online: Complete and submit the report: www.fda.gov/medwatch/report.htm
 - Regular mail or fax: Download form www.fda.gov/MedWatch/getforms.htm or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178.

If you have questions about this medicine or the recall, please talk to your doctor or pharmacist. You may also call the number on the back of your Humana member ID card. Our automated phone system may answer your call on Saturdays, Sundays, and some public holidays. Please leave your name and telephone number and we'll call you back by the end of the next business day. For 24-hour service you can sign in to MyHumana, your personal, secure online account on **Humana.com**, to search for other medicines that your plan covers.