



**URGENT RELAYHEALTH NOTIFYRx™ DRUG RECALL NOTICE:**  
**APOTEX – RETAIL RECALL – 05/13/10**

Apotex is initiating a voluntary **RETAIL LEVEL** recall that addresses an issue with the product due to a potential GMP violation, and is asking all retail pharmacies to now inspect for and remove **BELOW LISTED LOT #'s** for the following product (No significant adverse health consequences are expected with these lots):

**It is necessary to reformat the typical notification into tables due to the size of this recall.**

Pharmacies are advised to quarantine and discontinue distribution of these lots. Further instructions will come directly from your distributor.

RelayHealth has responded by implementing a NotifyRx™ safety alert informing pharmacies of this recall at point-of-dispense when the claim is processed through the Intelligent Network.

Beginning today, all pharmacies that participate in the NotifyRx messaging program will see the following messages for the listed drugs:

Recall #	Product Name	Lot #s	Drug Code	Message
1	CARBIDOPA/LEVO ER 25/100MG 1000-CT	HX8239	60505-0131-00	*NRX* Voluntary Recall of CARBIDOPA/LEVO ER 25/100mg (Apotex)-Lots HX8239, HX8241 and JF4544. Safety issue: Potential GMP Violation. Set aside.
		HX8241		
		JF4544		
2	CARBIDOPA/LEVO ER 50/200MG 1000-CT	HW8683	60505-0132-00	*NRX* Voluntary Recall of CARBIDOPA/LEVO ER 50/200mg (Apotex)-Lots HW8683, HW8684, HZ5773, HZ5774, HZ5775, JE1118. Safety issue: Potential GMP Violation. Set aside.
		HW8684		
		HZ5773		
		HZ5774		
		HZ5775		
		JE1118		
3	CARBIDOPA/LEVO 25/100MG 100-CT	HW8674	60505-0131-01	*NRX* Voluntary Recall of CARBIDOPA/LEVO ER 25/100mg (Apotex)-Lots HW8674, HX8240, HX8243, HZ5776, JD0810. Safety issue: Potential GMP Violation. Set aside.
		HX8240		
		HX8243		
		HZ5776		
		JD0810		
4	CARBIDOPA/LEVO ER 50/200MG 100-CT	HW8679	60505-0132-01	*NRX* Voluntary Recall of CARBIDOPA/LEVO ER 50/200mg (Apotex)-See list for affected lots. Safety issue: Potential GMP Violation. Set aside.
		HW8680		
		HW8681		
		HW8682		
		HZ5766		
		HZ5767		



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Recall #	Product Name	Lot #s	Drug Code	Message
		HZ5768		
		HZ5769		
		HZ5770		
		HZ5771		
		JD0814		
		JD0815		
		JE1115		
		JE1116		
		JE1117		
5	CITALOPRAM TAB 40MG 100-ct	JH2204	60505-2520-01	*NRX* Voluntary Recall of Citalopram 40mg (Apotex)-Lot JH2204 only. Safety issue: Potential GMP Violation. Set aside.
6	CLONAZEPAM TAB 0.5MG 100-ct	HX8305 JE7102	60505-0066-01	*NRX* Voluntary Recall of Clonazepam 0.5mg (Apotex)-Lots HX8305 and JE7102. Safety issue: Potential GMP Violation. Set aside.
7	CLONAZEPAM TAB 0.5MG 500-ct	HX8306 JE7105 JE7406	60505-0066-03	*NRX* Voluntary Recall of Clonazepam 0.5mg (Apotex)-Lots HX8306, JE7105 nad JE7406. Safety issue: Potential GMP Violation. Set aside.
8	CLONAZEPAM TAB 2MG 100-ct	HY6874	60505-0068-01	*NRX* Voluntary Recall of Clonazepam 2mg (Apotex)-Lot HY6874 only. Safety issue: Potential GMP Violation. Set aside.
9	CLONAZEPAM TAB 2MG 500-ct	HY6870 HY6871 JE7525	60505-0068-03	*NRX* Voluntary Recall of Clonazepam 2mg (Apotex)-Lots HY6870, HY6871 and JE7525. Safety issue: Potential GMP Violation. Set aside.
10	CYCLOSPORINE CAP 100MG 30-ct	HP9576 HW5445 HY6968 HY6969 JD6291 JE8040	60505-0134-00	*NRX* Voluntary Recall of Cyclosporine 100mg Caps (Apotex)-Lots HP9576, HW5445, HY6968, HY6969, JD6291 and JE8040. Safety issue: Potential GMP Violation. Set aside.
11	CYCLOSPORINE CAP 25MG 30-ct	HW5450 HY6964	60505-0133-00	*NRX* Voluntary Recall of Cyclosporine 25mg (Apotex)-Lots



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		JC0283 JD6290		HW5450, HY6964, JC0283, JD6290. Safety issue: Potential GMP Violation. Set aside.
12	DICLOFENAC POTASSIUM TAB 50MG 100-ct	HX3581 HZ4734 HZ4735 JD1676	60505-0135-00	*NRX* Voluntary Recall of Diclofenac Potassium (Apotex)-Lots HX3581, HZ4734, HZ4735 and JD1676. Safety issue: Potential GMP Violation. Set aside.
13	DILTIAZEM CD CAP 120MG 500-ct	HZ8662 JA4978 JF3578	60505-0007-08	*NRX* Voluntary Recall of Diltiazem CD 120mg (Apotex)-Lots HZ8662, JA4978, JF3578. Safety issue: Potential GMP Violation. Set aside.
14	DILTIAZEM CD CAP 180MG 500-ct	HZ2918 JA0227 JA4985 JD1889 JE9293 JF3580	60505-0008-08	*NRX* Voluntary Recall of Diltiazem CD 180mg (Apotex)-Lots HZ2918, JA0227, JA4985, JD1889, JE9293, JF3580. Safety issue: Potential GMP Violation. Set aside.
15	DILTIAZEM CD CAP 240MG 500-ct	JA0330 JA2104 JA4989 JA4991 JD5696 JF3590	60505-0009-08	*NRX* Voluntary Recall of Diltiazem CD 240mg (Apotex)-Lots JA0330, JA2104, JA4989, JA4991, JD5696, JF3590. Safety issue: Potential GMP Violation. Set aside.
16	DILTIAZEM CD ER CAP 300MG 500-ct	HZ8691 JA4980	60505-0010-08	*NRX* Voluntary Recall of Diltiazem CD ER 300mg (Apotex)-Lots HZ8691 and JA4980. Safety issue: Potential GMP Violation. Set aside.
17	DILTIAZEM CD CAP 120MG 90-ct	HZ8661 JA4977 JE9298 JF3577	60505-0007-04	*NRX* Voluntary Recall of Diltiazem CD 120mg (Apotex)-Lot HZ8661, JA4977, JE9298, JF3577. Safety issue: Potential GMP Violation. Set aside.
18	DILTIAZEM CD CAP 180MG 90-ct	HY9659 HZ2919 JA0229 JA4982 JD1453	60505-0008-04	*NRX* Voluntary Recall of Diltiazem CD 180mg (Apotex)-Lot HY9659, HZ2919, JA0229, JA4982, JD1453, JE3048, JE9296, JE9297, JF3588. Safety



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Recall #	Product Name	Lot #s	Drug Code	Message
		JE3048 JE9296 JE9297 JF3588		issue: Potential GMP Violation. Set aside.
19	DILT-XR CAP 180MG 100-ct	HY3576	60505-0015-06	*NRX* Voluntary Recall of Dilt-XR 180mg (Apotex)-Lot HY3576 only. Safety issue: Potential GMP Violation. Set aside.
20	DILT-CD CAP 240MG 90-ct	JA0332 JA2105 JA4987 JA4988 JD1489 JD5695 JE3034 JE3035 JE9288 JE9289 JF1473 JF3591	60505-0009-04	*NRX* Voluntary Recall of DILT-CD 240mg (Apotex)-See list for affected lot numbers. Safety issue: Potential GMP Violation. Set aside.
21	DILT-CD ER CAP 300MG 90-ct	HZ8690 JC1281 JE9279 JE9280	60505-0010-04	*NRX* Voluntary Recall of DILT-CD ER 300mg (Apotex)-Lots: HZ8690, JC1281, JE9279, JE9280. Safety issue: Potential GMP Violation. Set aside.
22	DILTZAC ER CAP 120MG 30-ct	JD3976 JF3643	60505-0210-03	*NRX* Voluntary Recall of DILTZAC-ER 120mg (Apotex)-Lots: JD3976 and JF3643. Safety issue: Potential GMP Violation. Set aside.
23	DILTZAC ER CAP 120MG 90-ct	JA4854 JD3977 JE3609 JF0150	60505-0210-09	*NRX* Voluntary Recall of DILTZAC-ER 120mg (Apotex)-Lots: JA4854, JD3977, JE3609, JF0150. Safety issue: Potential GMP Violation. Set aside.
24	DILTZAC ER CAP 180MG APX 30	JD3978 JE3604	60505-0211-03	*NRX* Voluntary Recall of DILTZAC-ER 180mg (Apotex)-Lots: JD3978 and JE3604. Safety issue: Potential GMP Violation. Set aside.
25	DILTZAC ER CAP	JA4862	60505-0211-09	*NRX* Voluntary Recall of



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	180MG 90-ct	JD3979 JE3589		DILTZAC-ER 180mg (Apotex)- Lots: JA4862, JD3979, JE3589. Safety issue: Potential GMP Violation. Set aside.
26	DILTZAC ER CAP 240MG 30-ct	JA8448 JD3980 JF0143	60505-0212-03	*NRX* Voluntary Recall of DILTZAC-ER 240mg (Apotex)- Lots: JA8448, JD3980 and JF0143. Safety issue: Potential GMP Violation. Set aside.
27	DILTZAC ER CAP 240MG 90-ct	JA4865 JD3981 JD3982 JE3573	60505-0212-09	*NRX* Voluntary Recall of DILTZAC-ER 240mg (Apotex)- Lots: JA4865, JD3981, JD3982, JE3573. Safety issue: Potential GMP Violation. Set aside.
28	DILTZAC ER CAP 300MG 30-ct	JD3984 JE3579	60505-0213-03	*NRX* Voluntary Recall of DILTZAC-ER 300mg (Apotex)- Lots: JD3984 and JE3579. Safety issue: Potential GMP Violation. Set aside.
29	DILTZAC ER CAP 300MG 90-ct	JC2005 JD3985 JE3583	60505-0213-09	*NRX* Voluntary Recall of DILTZAC-ER 300mg (Apotex)- Lots: JC2005, JD3985, JE3583. Safety issue: Potential GMP Violation. Set aside.
30	DILTZAC ER CAP 360MG 30-ct	JA8450 JD3986 JE3586 JF0157	60505-0214-03	*NRX* Voluntary Recall of DILTZAC-ER 360mg (Apotex)- Lots: JA8450, JD3986, JE3586, JF0157. Safety issue: Potential GMP Violation. Set aside.
31	DILTZAC ER CAP 360MG 90-ct	JA4867 JD3987 JD3988 JE3584 JE3585 JE4351 JF0158 JF0161 JF3678	60505-021409	*NRX* Voluntary Recall of DILTZAC-ER 360mg (Apotex)-see list for affected lots. Safety issue: Potential GMP Violation. Set aside.
32	DILT-XR CAP 240MG 100-ct	HW4437 HY3540	60505-0016-06	*NRX* Voluntary Recall of DILT- XR 240mg (Apotex)-Lots: HW4437 and HY3540. Safety issue: Potential GMP Violation. Set aside.



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33	DILT-XR CP 240MG 500-ct	HW4435	60505-0016-08	*NRX* Voluntary Recall of DILT-XR 240mg (Apotex)-Lots: HW4435 and HY3541. Safety issue: Potential GMP Violation. Set aside
		HY3541		
34	DIVALPROEX SOD DR TAB 125MG 100-ct	HZ4212	60505-3065-01	*NRX* Voluntary Recall of Divalproex DR 125mg (Apotex)-Lot HZ4212 only. Safety issue: Potential GMP Violation. Set aside.
35	DIVALPROEX SOD DR TAB 250MG 100-ct	HW0431	60505-3066-01	*NRX* Voluntary Recall of Divalproex DR 250mg (Apotex)-See list for affected lots. Safety issue: Potential GMP Violation. Set aside.
		HW0433		
		HW0434		
		HW0435		
		HW0436		
		HW0438		
		HW0439		
		HW0441		
		HW7458		
		HW7460		
		HW7466		
		HW7467		
		HZ1063		
		HZ4447		
		HZ4448		
		HZ4449		
		HZ4450		
		HZ4451		
		JC9956		
		JC9957		
JC9958				
JC9959				
JD6584				
JD6585				
JD6586				
JD6587				
JE1856				
JE1857				
36	DIVALPROEX SOD DR	HW0445	60505-3066-05	*NRX* Voluntary Recall of



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	TAB 250MG 500-ct	HW7509 HW7510 HW7513 HY9238 HZ4442 JC9955 JD6588 JD6589 JE1860 JE9516		Divalproex DR 250mg (Apotex)- See list for affected lots. Safety issue: Potential GMP Violation. Set aside.
37	EPLERENONE TAB 25MG 30-ct	HW0774 JC6593 JC9429 JH2193	60505-2651-03	*NRX* Voluntary Recall of Eplerenone 25mg (Apotex)-Lots: HW0774, JC6593, JC9429, JH2193. Safety issue: Potential GMP Violation. Set aside.
38	EPLERENONE TAB 25MG 90-ct	HY7640 JH2190	60505-2651-09	*NRX* Voluntary Recall of Eplerenone 25mg (Apotex)-Lots: HY7640 and JH2190. Safety issue: Potential GMP Violation. Set aside.
39	EPLERENONE TAB 25MG 500-ct	JC9428 JF4992	60505-2651-05	*NRX* Voluntary Recall of Eplerenone 25mg (Apotex)-Lots: JC9428 and JF4992. Safety issue: Potential GMP Violation. Set aside.
40	EPLERENONE TAB 50MG 30-ct	JC6601 JC9477	60505-2652-03	*NRX* Voluntary Recall of Eplerenone 50mg (Apotex)-Lots: JC6601 and JF9477. Safety issue: Potential GMP Violation. Set aside.
41	EPLERENONE TAB 50MG 90-ct	JC6604	60505-2652-09	*NRX* Voluntary Recall of Eplerenone 50mg (Apotex)-Lots: JC6604 only. Safety issue: Potential GMP Violation. Set aside.
42	GEMFIBR TAB 600MG APX 500	JD4444 JD4448	60505-0034-08	*NRX* Voluntary Recall of Gemfibrozil 600mg (Apotex)-Lots: JD4444 and JD4448. Safety issue: Potential GMP Violation. Set aside.
43	LORATADINE TAB 10MG 100-ct	HX8178 JA9708	60505-0147-01	*NRX* Voluntary Recall of Loratadine 10mg (Apotex)-Lots: HX8178 and JA9708. Safety



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				issue: Potential GMP Violation. Set aside.
44	LORATADINE TAB 10MG 1000-ct	HY3556 JA9709	60505-0147-08	*NRX* Voluntary Recall of Loratadine 10mg (Apotex)-Lots: HY3556 and JA9709. Safety issue: Potential GMP Violation. Set aside.
45	MOEXIPRIL TAB 15MG 100-ct	JD7716	60505-0272-01	*NRX* Voluntary Recall of Moexipril 15mg (Apotex)-Lots: JD7716. Safety issue: Potential GMP Violation. Set aside.
46	MOEXIPRIL TAB 7.5MG 100-ct	JD7717	60505-0271-01	*NRX* Voluntary Recall of Moexipril 7.5mg (Apotex)-Lots: JD7717. Safety issue: Potential GMP Violation. Set aside.
47	OMEPRAZOLE DR CAP 10MG 30-ct	HN5167 HW1580 HX9342 HZ1742 JD5422	60505-0145-00	*NRX* Voluntary Recall of Omeprazole 10mg (Apotex)-Lots: HN5167, HW1580, HX9342, HZ1742, JD5422. Safety issue: Potential GMP Violation. Set aside.
48	OMEPRAZOLE DR CAP 10MG 100-ct	HN3625 HR8424	60505-0145-02	*NRX* Voluntary Recall of Omeprazole 10mg (Apotex)-Lots: HN3625 and HR8424. Safety issue: Potential GMP Violation. Set aside.
49	OMEPRAZOLE DR CAP 10MG 1000-ct	HN5142 HR9367 HW1583 HZ1743 JD5421	60505-0145-01	*NRX* Voluntary Recall of Omeprazole 10mg (Apotex)-Lots: HN5142, HR9367, HW1583, HZ1743, JD5421. Safety issue: Potential GMP Violation. Set aside.
50	OMEPRAZOLE DR CAP 20MG 30-ct	HL2492 HL4244 HL7315 HN6311 HP3824 HP3827 HR5045 HT2183 HV0102 HV0103 HW4355	60505-0065-00	*NRX* Voluntary Recall of Omeprazole 20mg (Apotex)-See list for affected lots. Safety issue: Potential GMP Violation. Set aside.



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		HX2483		
		HX8525		
		HY3752		
		HZ2554		
		JA1868		
		JC3915		
		JC3938		
		JD5405		
		JE8165		
		JE8700		
		JE8757		
		JF7544		
51	OMEPRAZOLE DR CAP 20MG 100-ct	HK2931	60505-0065-02	*NRX* Voluntary Recall of Omeprazole 20mg (Apotex)-See list for affected lots. Safety issue: Potential GMP Violation. Set aside.
		HL2477		
		HL3181		
		HL3182		
		HL7314		
		HM3346		
		HM3347		
		HM3348		
		HM9677		
		HM9678		
		HN4405		
		HN4408		
		HN5465		
		HN5826		
		HN6243		
		HN6310		
		HN6312		
		HN6313		
		HN6314		
		HP9353		
		HP9354		
		HR2532		
		HR2533		
		HR2537		
		HV0107		
		HV0108		



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Recall #	Product Name	Lot #s	Drug Code	Message
		HV8054		
		HV8055		
		HV8056		
		HW3108		
		HW4358		
		HW4360		
		HW4361		
		HW4362		
		HW4363		
		HX2479		
		HX2480		
		HX2481		
		HX2482		
		HX8523		
		HX8524		
		HY3750		
		HY3751		
		HZ2540		
		HZ2542		
		HZ5287		
		JA1870		
		JA1871		
		JA1873		
		JA6887		
		JA6888		
		JA6889		
		JC4897		
		JC5709		
		JC9698		
		JD0113		
		JD0117		
		JD3992		
		JD5387		
		JD5388		
		JD5389		
		JD5390		
		JD7326		
		JE4646		



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Recall #	Product Name	Lot #s	Drug Code	Message
		JE8752		
		JE8753		
		JE8756		
		JF1851		
52	OMEPRAZOLE DR CAP 20MG 1000-ct	HJ2268	60505-0065-01	*NRX* Voluntary Recall of Omeprazole 20mg (Apotex)-See list for affected lots. Safety issue: Potential GMP Violation. Set aside.
		HL2499		
		HL2500		
		HL2501		
		HL2502		
		HL2503		
		HL3171		
		HL3174		
		HL3176		
		HL3177		
		HL3178		
		HL3179		
		HL7316		
		HL7317		
		HL7318		
		HM3350		
		HM3351		
		HM3352		
		HM7180		
		HM8609		
		HM9679		
		HM9680		
		HM9681		
		HN4453		
		HN5481		
		HN6330		
		HP2381		
		HP3832		
		HP3833		
		HP3836		
		HP9359		
		HP9360		
		HR4982		
		HR4986		



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Recall #	Product Name	Lot #s	Drug Code	Message
		HR5073		
		HR5075		
		HR5084		
		HR5683		
		HT2230		
		HT2232		
		HT2233		
		HT2234		
		HT2235		
		HT2236		
		HT2237		
		HT2238		
		HT2239		
		HT6557		
		HT6559		
		HT6560		
		HT6561		
		HT8020		
		HT8021		
		HT8152		
		HV0109		
		HV0110		
		HV0111		
		HV0112		
		HV0118		
		HV0121		
		HV8076		
		HV8078		
		HV8079		
		HV8082		
		HW4373		
		HW4374		
		HW4378		
		HW4387		
		HW4397		
		HW4398		
		HW4399		
		HW4400		



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		HW4401		
		HW4402		
		HW4408		
		HW5311		
		HX2484		
		HX2485		
		HX2486		
		HX2487		
		HX7608		
		HX7609		
		HX7610		
		HX7612		
		HX7613		
		HX7614		
		HX7615		
		HX7616		
		HX7617		
		HX7618		
		HX8544		
		HX8545		
		HX8546		
		HX8547		
		HX8548		
		HX8549		
		HX8550		
		HX8551		
		HX8552		
		HX8553		
		HX8554		
		HX8556		
		HX8558		
		HX8559		
		HY3753		
		HY3754		
		HY3755		
		HY3756		
		HY3758		
		HY3760		



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Recall #	Product Name	Lot #s	Drug Code	Message
		HY3763		
		HY3766		
		HY3768		
		HY3771		
		HY3773		
		HY3775		
		HZ0065		
		HZ0066		
		HZ2534		
		HZ2535		
		HZ2536		
		HZ2538		
		HZ2544		
		HZ7137		
		HZ7138		
		HZ7139		
		JA2142		
		JA2143		
		JA2144		
		JA2145		
		JA2147		
		JA2148		
		JA2149		
		JA2151		
		JA6891		
		JA6892		
		JA6893		
		JA6894		
		JA6895		
		JC3481		
		JC3482		
		JC3483		
		JC3484		
		JC3485		
		JC3486		
		JC3487		
		JC3488		
		JC3489		



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Recall #	Product Name	Lot #s	Drug Code	Message
		JC3490		
		JC3491		
		JC8807		
		JC9695		
		JC9696		
		JC9701		
		JC9702		
		JC9703		
		JC9705		
		JD0108		
		JD0109		
		JD0110		
		JD0111		
		JD0112		
		JD0114		
		JD0115		
		JD0118		
		JD5391		
		JD5392		
		JD5393		
		JD5394		
		JD5395		
		JD8787		
		JD9105		
		JD9110		
		JE0294		
		JE0779		
		JE0780		
		JE0781		
		JE0782		
		JE0783		
		JE0784		
		JE2956		
		JE2958		
		JE4472		
		JE4783		
		JE5323		
		JE5324		



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Recall #	Product Name	Lot #s	Drug Code	Message
		JE5341		
		JE8758		
		JE8759		
		JE8760		
		JE8762		
		JE8767		
		JE8768		
		JE8769		
		JE8771		
		JF1119		
		JF1120		
		JF1121		
		JF1122		
		JF1147		
		JF1857		
		JF1859		
		JF1860		
		JF1861		
		JF1981		
		JF2220		
		JF5842		
		JF5843		
		JF5845		
		JF5846		
		JF5847		
		JF7160		
		JF8021		
		JF8022		
		JF8023		
		JF8024		
		JF8025		
		JF9662		
		JF9663		
		JG1912		
		JG1913		
		JG1914		
53	OMEPRAZOLE DR CAP 20MG UD 100-ct	HY9656 JC3942	60505-0065-07	*NRX* Voluntary Recall of Omeprazole 20mg (Apotex)- Lots:



**URGENT RELAYHEALTH NOTIFYRx™ DRUG RECALL NOTICE:**  
**APOTEX – RETAIL RECALL – 05/13/10**

Recall #	Product Name	Lot #s	Drug Code	Message
		JD5404 JE4100		HY9656, JC3942, JD5404, JE4100. Safety issue: Potential GMP Violation. Set aside.
54	OMEPRAZOLE DR CAP 40MG 30-ct	JC2939 JC4538 JC4539 JD8861 JD8873 JF2262 JF2984 JF8082 JF8634 JF8637	60505-0146-00	*NRX* Voluntary Recall of Omeprazole 40mg (Apotex)-See list for affected lots. Safety issue: Potential GMP Violation. Set aside.
55	OMEPRAZOLE DR CAP 40MG 100-ct	JC2826 JC4536 JD1258 JD8872 JE8813 JF2268 JF8087 JG1988	60505-0146-02	*NRX* Voluntary Recall of Omeprazole 40mg (Apotex)-See list for affected lots. Safety issue: Potential GMP Violation. Set aside.
56	OMEPRAZOLE DR CAP 40MG 500-ct	JC2827 JC7602 JD0672 JD2869 JD2870 JD8858 JD8859 JD8860 JE1350 JF2261 JF3153 JF3154 JG1987	60505-0146-01	*NRX* Voluntary Recall of Omeprazole 40mg (Apotex)-See list for affected lots. Safety issue: Potential GMP Violation. Set aside.
57	OMEPRAZOLE SA CAP 20MG 5000-ct	HZ8676 HZ8680 HZ8681 HZ8685	60505-0065-08	*NRX* Voluntary Recall of Omeprazole 20mg (Apotex)-See list for affected lots. Safety issue: Potential GMP Violation. Set



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**APOTEX – RETAIL RECALL – 05/13/10**

Recall #	Product Name	Lot #s	Drug Code	Message
		JA2120		aside.
		JA2121		
		JA2127		
		JA2136		
		JA6930		
		JA6931		
		JA6932		
		JA6933		
		JA6934		
		JA6935		
		JA6936		
		JA6937		
		JC3492		
		JC3493		
		JC3494		
		JC3495		
		JC3496		
		JC3497		
		JC3498		
		JC9699		
		JC9700		
		JC9709		
		JD0123		
		JD0125		
		JD0127		
		JD0128		
		JD0130		
		JD5398		
		JD5399		
		JD5400		
		JD5401		
		JD5402		
		JD7334		
		JE0795		
		JE0798		
		JE5630		
		JE5631		
		JF2231		



**URGENT RELAYHEALTH NOTIFYRx™ DRUG RECALL NOTICE:**  
**APOTEX – RETAIL RECALL – 05/13/10**

Recall #	Product Name	Lot #s	Drug Code	Message
		JF2232		
		JF2234		
		JF5849		
		JG1909		
		JG1910		
Recall #	Product Description	Lot #s	Drug Code	Message:
58	OMEPRAZOLE DR CAP 40MG 90 BTL	JG9727	60505-0146-09	*NRX* Voluntary Recall of Omeprazole 40mg (Apotex)-Lot JG9727 only. Safety issue: Potential GMP Violation. Set aside.
59	ONDANSETRON FCT 4MG 30 BTL	HP9522	60505-1311-03	*NRX* Voluntary Recall of Ondansetron 4mg (Apotex)-Lots: HP9522, HX5296, JA3653. Safety issue: Potential GMP Violation. Set aside.
		HX5296		
		JA3653		
60	ONDANSETRON FCT 8MG 30 BTL	HP9549	60505-1312-03	*NRX* Voluntary Recall of Ondansetron 8mg (Apotex)-Lots: HP9549 and HP9552. Safety issue: Potential GMP Violation. Set aside.
		HP9552		

If you have any questions about NotifyRx or are interested in signing up for the program, please contact your RelayHealth Sales Executive.

[www.relayhealth.com](http://www.relayhealth.com)

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