



Nonsafety voluntary recall and withdrawal of a limited portion of Daytrana[®] (methylphenidate transdermal system) patches

Shire has voluntarily recalled one lot and conducted a voluntary market withdrawal of ten lots of the Attention-Deficit/Hyperactivity Disorder (ADHD) patch Daytrana[®]. Shire is taking this action because some Daytrana patches do not meet or in the future may not meet liner removal specifications, and as a result, patients and caregivers could have difficulty removing the liners.

This action is not due to safety issues. All Daytrana patches can continue to be used unless the release liner cannot be removed or the patches are damaged while being opened. The United States Food and Drug Administration (FDA) has been notified of this voluntary action.

If you have questions about Daytrana, call Shire's Daytrana customer service line at **1-800-828-2088, option 1**. Pharmacists should call **1-888-879-8144**.

Daytrana lots affected by this action are

VOLUNTARY RECALL				
Strength	Count	NDC#	Lot Number(s)	Expiry Date
15 mg	30	54092-553-30	40794	11/30/2011

VOLUNTARY MARKET WITHDRAWAL				
Strength	Count	NDC#	Lot Number(s)	Expiry Date
10 mg	30	54092-552-30	40856	11/30/2011
20 mg	30	54092-554-30	37908	7/31/2011
			37910	7/31/2011
			37911	7/31/2011
			38125	7/31/2011
			41465	10/31/2011
			41521	10/31/2011
30 mg	30	54092-555-30	37698	7/31/2011
			37750	7/31/2011
			38128	7/31/2011

Please see [Full Prescribing Information](#), and [Medication Guide](#), including Warning regarding abuse and dependence.