URGENT MEDICINE RECALL

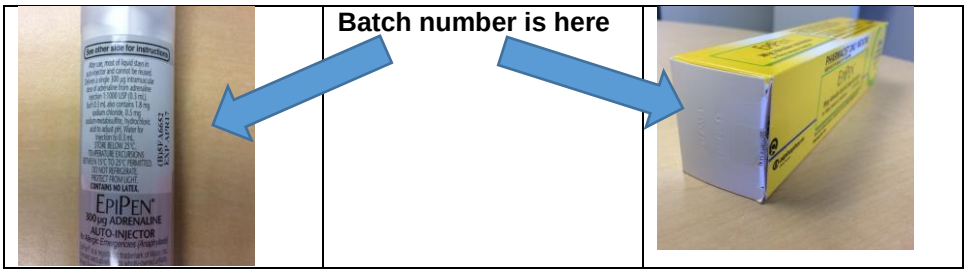
EPIPEN 300 microgram (µg) Adrenaline Injection Syringe Auto-Injector
Batches 5FA665, 5FA6651, 5FA6652, 5FA6653

Alphapharm Pty Ltd, following consultation with the Therapeutic Goods Administration is recalling the following batches of EpiPen® 300 microgram (µg) Adrenaline Injection Syringe Auto-Injectors (used for the treatment of allergic emergencies (anaphylaxis)) due to the potential that these devices may contain a defective part that may result in the device failing to activate or requiring increased force to activate. If the Epipen fails to activate it may result in life threatening adverse events as the underlying anaphylaxis will not be treated.

Batch number	Expiry
5FA665	Apr 17
5FA6651	Apr 17
5FA6652	Apr 17
5FA6653	Apr 17

IF YOU HAVE AN EPIPEN:

1. **Check if you have a 300 µg EpiPen® (yellow carton and label) and if you do, check the batch number and expiry. The batch number and expiry can be found on the label of the pen or on the end of the carton.**



2. **If your EpiPen® 300µg has the following batch numbers 5FA665, 5FA6651, 5FA6652 or 5FA6653 and an expiry of Apr 17, you need to replace it with a new one as soon as possible by returning to your pharmacist.**
3. **Your pharmacist will replace the Epipen® 300µg from the affected batch with an EpiPen® 300µg from a different batch FREE OF CHARGE.**
4. **You must keep your current Epipen® until you get a replacement and use it if required.**

If your EpiPen® 300µg is not from a batch listed above or is a green Epipen® Jr 150µg Adrenaline Injection Syringe Auto Injector, your product is not affected by this recall and no action is required.

Alphapharm Pty Ltd sincerely regrets any inconvenience to its customers