FDA WARNING ALERT

Regarding EpiPen, EpiPen Jr, and Mylan's Generic Versions

FDA Alert Regarding Epinephrine Devices

[3/24/2020] FDA is alerting patients, caregivers and health care professionals that EpiPen 0.3mg and EpiPen Jr 0.15mg auto-injectors, and the authorized generic versions, may potentially have delayed injection or be prevented from properly injecting

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1. **Device failure from spontaneous activation caused by using sideways force to remove the blue safety release**
2. **Device failure from inadvertent or spontaneous activation due to a raised blue safety release**
3. **Difficulty removing the device from the carrier tube**
4. **User errors**

Please click link for important FDA warning regarding EpiPen, EpiPen Jr. and generic varieties of both