

## DRUG ALERT

### CLASS 2 MEDICINES RECALL

Action within 48 Hours

Pharmacy and Wholesaler Level Recall

Dear Healthcare Professional

#### Sanofi

**Fasturtec® 7.5 mg. 1.5 mg/ml powder and solvent for concentrate for solution for infusion**

**EU/1/00/170/002**

Batch Number	Expiry Date	Pack Size	First Distributed
A9306	02 / 2022	7.5mg vial	14/11/2019

Generic Name: rasburicase

#### Brief description of problem

Sanofi has informed us of an Out Of Specification (OOS) result which was detected for Rasburicase enzyme activity according to a specific method and specifications for US market, at 12 months stability time point. Sanofi is recalling Fasturtec® 7.5 mg (Rasburicase) Solution for IV infusion - 7.5 mg/5 ml (Injectable powder in vial packaged with 5 ml solvent in ampoule), batch number A9306 as a precautionary measure.

#### Advice for healthcare professionals

Stop supplying the above batch immediately. Quarantine all remaining batch stock and return it to your supplier using your supplier's approved process.

#### Company contacts for further information

For medical information enquiries please contact [uk-medicalinformation@sanofi.com](mailto:uk-medicalinformation@sanofi.com)  
Med Info Phone : 0800 035 25 25. For stock control enquiries please contact [GB-CustomerServices@sanofi.com](mailto:GB-CustomerServices@sanofi.com) Phone number: 0800 854 430

Recipients of this Drug Alert should bring it to the attention of relevant contacts by copy of this notice.



RQIA should bring this information to the attention of private hospitals/clinics registered with them and any other relevant care facilities

The Business Services Organisation is asked to bring this information to the attention of Community Pharmacists and General Medical Practitioners directly.

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**TO ALL CHEMISTS, DOCTORS ON THE LISTS**

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**24<sup>th</sup> August 2020**

Pharmaceutical website: <http://www.hscbusiness.hscni.net>