



# DRUG ALERT

## CLASS 2 MEDICINES RECALL

Action Within 48 Hours  
Pharmacy/Wholesaler Level Recall

Date: 24 August 2020

EL (20)A/40

Our Ref: MDR 091-08/20

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.

NHS regional teams are asked to forward this alert to community pharmacists and dispensing general practitioners for information.

Yours faithfully

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