H29. 5. 10

韓国 MINISTRY OF FOOD AND DRUG SAFETY からの情報

Reasons for recall	Fraud product	
(Details of defect)	MFDS was reported the criminal investigation report	
	by a Police office for fraud biological product of	
	NABOTA Inj. which produced by illegal	
	manufacturer.	
	These fraud products had been produced total of	
	15,000 Vials, and 2,900 Vials of them were sold.	
	The fraud products have mostly same figure with	
	the real ones. But, there is no API(Botulinum toxin	
	type A) in them.	
	* Information on distribution: Not investigated (May	
	exported to China)	
	The labels on the products and package are	
	written in Korean.	
Company name	DAEWOOG Pharmaceuticals	
(recall obligator)		
	Finished medicinal	NABOTA INJ
Product name	products	
	APIs	Botulinum Toxin Type A
	(INN or Generic name)	
Active ingredients &	Botulinum Toxin Type A / 100 Units	
strength		
Dosage form	Injection (lyophilization)	
Batch number	089139, 091743, 093103	
Expiry date	2019.03.03, 2019.07.21, 2019.10.10	