PRODUCT WITHDRAWAL



Instanyl Multi-dose 50 microgams/dose nasal spray, solution Instanyl Multi-dose 100 microgams/dose nasal spray, solution Instanyl Multi-dose 200 microgams/dose nasal spray, solution

EU/1/09/531/001-009

1st November 2018

Dear Healthcare Professional

As the Marketing Authorisation Holder (MAH) for Instanyl Multi-dose nasal spray, Takeda has made the decision to withdraw the product for all multi-dose strengths in the United Kingdom from the 14th December 2018. This decision has been made for commercial reasons and means that the multi-dose product will no longer be available for use.

Instanyl Multi-dose is indicated for the management of breakthrough pain (BTP) in adults who are already receiving maintenance opioid therapy for chronic cancer pain. The active ingredient in Instanyl Multi-dose is fentanyl citrate.

Alternative fentanyl citrate products indicated for the management of breakthrough pain in adult patients using opioid therapy for chronic cancer pain are available in the UK. Alternative products include, single dose intranasal preparations, sublingual tablets and buccal tablets. Please note that the Instanyl single dose presentation will remain available.

Please note that fentanyl preparations for the treatment of BTP are not interchangeable. Any patient being treated with Instanyl Multi-dose should be referred back to their treating physician for consideration of an alternate preparation. If patients are switched from Instanyl Multi-dose to another fentanyl preparation, a new dose titration is required.

No new patients should be prescribed Instanyl Multi-dose from the date of this notification.

Should you have any questions regarding the use of Instanyl Multi-dose nasal spray or questions about the content of this letter, please contact Takeda:

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We thank you for your attention and understanding.

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Yours sincerely,

Dr Simon Meadowcroft

Medical Director

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