Australian Government

Department of Health and Aged Care

Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry:	170905 Pfizer Australia Pty Ltd - Medicine administration kit, percutaneous, medicated, single-use
ARTG entry for	Medical Device Included Class IIa
Sponsor	Pfizer Australia Pty Ltd
Postal Address	Level 17 151 Clarence Street, Sydney, NSW, 2000 Australia
ARTG Start Date	27/04/2010
Product Category	Medical Device Class IIa
Status	Active
Approval Area	Medical Devices

Conditions

- The inclusion of the kind of device in the ARTG is subject to compliance with all conditions placed or imposed on the ARTG entry. Refer Part 4-5, Division 2 (Conditions) of the Therapeutic Goods Act 1989 and Part 5, Division 5.2 (Conditions) of the Therapeutic Goods (Medical Devices) Regulations 2002 for relevant information.
- Breaching conditions of the inclusion related to the device of the kind may lead to suspension or cancellation of the ARTG entry; may be a criminal offence; and civil penalties may apply.

Manufacturers

Name	Address
Wyeth Farma SA	Autovia del norte A-1 23 Desvio Algete Km 1 San Sebastian , de los Reyes, 28700 Spain

Products

1 . Medicine administration kit, percutaneous, medicated, single-use

Product Type	Procedure Pack	**Effective Date**	27/04/2010
GMDN	45156 Medicine administration kit, percutaneous, medicated, single-use		
Intended Purpose	The kit is a collection of devices used to adminster a medication by injection.		

Specific Conditions

No Specific Conditions included on Record

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