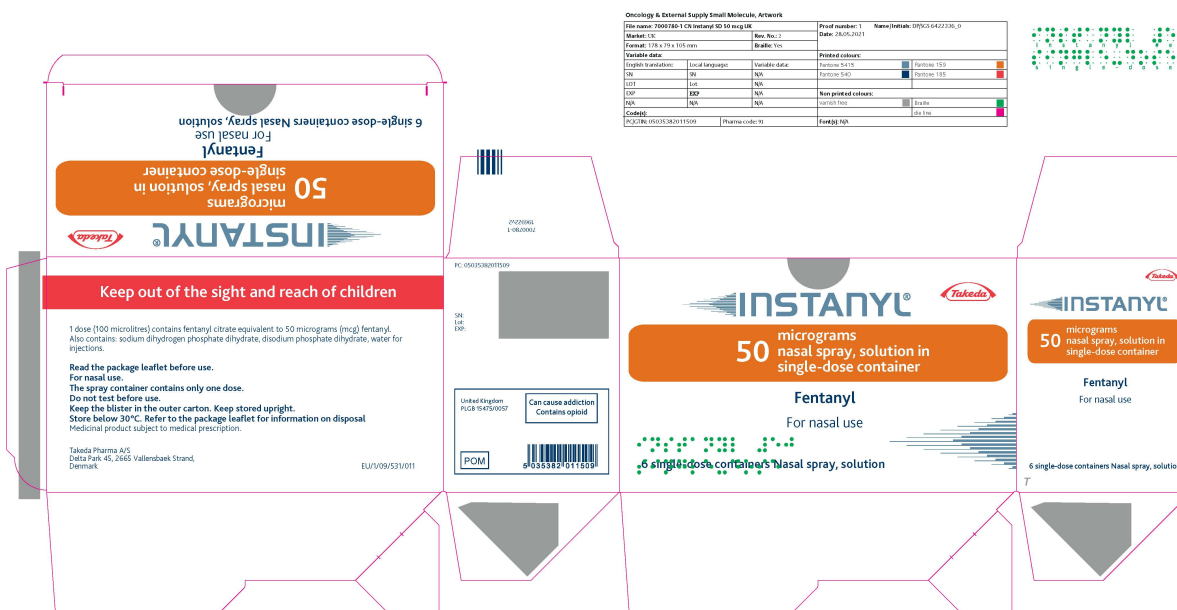


INSTANYL- fentanyl nasal spray spray Renaissance Lakewood LLC

Instanyl Fentanyl Nasal Spray

Instanyl single dose nasal spray carton



INSTANYL

fentanyl nasal spray spray

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:49396-0512
Route of Administration	NASAL	DEA Schedule	CII

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FENTANYL CITRATE (UNII: MUN5LYG46H) (FENTANYL - UNII:UF599785JZ)	FENTANYL	2 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JH2SW)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49396-0512-1	50 in 1 CARTON	06/29/2011	
1		0.125 mL in 1 VIAL; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Export only		06/29/2011	

Labeler - Renaissance Lakewood LLC (077744035)

Revised: 3/2023

Renaissance Lakewood LLC