Policies Repository



Policy Title	Fluticasone Furoate (Veramyst®) Nasal Spray FS.CLIN.30		
Policy Number			
	determined by benefits and contracts. Benefits may vary based on product line, group or he subject to precertification, age, gender or quantity edits. Individual member benefits must		
was developed. Since that time, ne routes, safety or FDA approval ma subject of this document, please p new FDA approved indications, wi whether this Policy should be upda	escribes the status of pharmaceutical information and/or technology at the time the document ew information relating to drug efficacy, interactions, contraindications, dosage, administration y have changed. If the Medical/Pharmacy Reviewer is aware of any new information on the rovide it promptly to the Medical/Pharmacy Policy Department. This information may include thdrawals or other FDA alerts. This type of information is relevant not only when considering ated, but also when applying it to current requests for coverage.		
Policy	Fluticasone furoate (Veramyst®) nasal spray is indicated for the treatment of symptoms of seasonal and perennial allergic rhinitis in adults and children 2 years of age and older.		
	The use of fluticasone furoate (Veramyst®) requires prior authorization (ie, clinical pharmacy and/or Medical Director review).		
Policy Description	Fluticasone furoate (Veramyst®) nasal spray is a synthetic trifluorinated corticosteroid with potent anti-inflammatory activity. The precise mechanism through which fluticasone furoate (Veramyst®) affects rhinitis symptoms is unknown, but corticosteroids have been shown to have a wide range of actions on multiple cell types (eg, mast cells, eosinophils, neutrophils, macrophages, lymphocytes) and mediators (eg, histamine, eicosanoids, leukotrienes, cytokines) involved in inflammation.		
Policy Guideline Inclusion	Fluticasone furoate (Veramyst®) is approved when there is documentation of a diagnosis of seasonal or perennial allergic rhinitis and one of the following:		
	 Documentation that the individual is 2 or 3 years of age, with documentation of trial and failure of or intolerance/contraindication/allergy to mometasone furoate monohydrate (Nasonex®) and triamcinolone acetonide (Nasacort AQ) 		
	 Documentation that the individual is 4 years of age or older, with documentation of trial and failure of or intolerance/contraindication/allergy to fluticasone propionate containing nasal product and one of the following: Mometasone furoate monohydrate (Nasonex®) Triamcinolone acetonide (Nasacort® AQ) 		

- The individual is 4 years of age or older, with no documentation of trial and failure of or intolerance/contraindication/allergy to fluticasone propionate containing nasal product and either mometasone furoate monohydrate (Nasonex[®]) or triamcinolone acetonide (Nasacort[®] AQ)
- The individual is younger than 2 years of age

Policy List of Applicable Drugs	Brand Name	Generic Name	
	Veramyst	fluticasone furoate	
Dosing and Administration	Refer to the specific manufacturer's prescribing information for administration and dosage details for each specific agent.		
Policy References	Facts & Comparisons. Veramyst [®] . [Facts & Comparisons Web site]. Available at: http://online.factsandcomparisons.com [via subscription only]. Accessed September 1, 2008.		
	Micromedex. Veramyst [®] . [Micromedex web site]. Available at: http://www.micromedex.com [via subscription only]. Accessed September 1, 2008.		
	Veramyst® [package insert]. Triangle Park, NC: GlaxoSmithKline, Inc.; 2007. Also available online at: http://us.gsk.com/products/assets/us_veramyst.pdf. Accessed October 16, 2008.		
Policy Link to Related Policies			
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criteria are present:

Policy Guideline Exclusion