ACONITUM BELLADONNA- aconitum belladonna liquid
Uriel Pharmacy Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Aconitum Belladonna

Directions: FOR ORAL USE.

Take the contents of one ampule
under the tongue and hold for
30 seconds, then swallow.

Active Ingredients: Aconitum
e tub. 30X, Atropa belladonna
e rad. 30X, Rhus tox. e fol. 30X

Inactive Ingredients: Water, Salt

Use: Temporary relief of headache.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions
worsen or persist. If pregnant or nursing, consult a doctor before use. Natural ingredients may cause color, scent
and/or taste variation.

Questions? Call 866.642.2858
Uriel, East Troy, WI 53120
www.urielpharmacy.com

ACONITUM BELLADONNA
aconitum belladonna liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-9166
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII:WQZ3G9PF0H)	ATROPA BELLADONNA	4 [hp_X] in 1 mL
APIS MELLIFERA (UNII: 7S82P3R43Z) (APIS MELLIFERA - UNII:7S82P3R43Z)	APIS MELLIFERA	5 [hp_X] in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:48951-9166-1	10 in 1 BOX		
1		1 mL in 1 AMPULE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-9166)

Revised: 3/2014 Uriel Pharmacy Inc.