JUGLANS REGIA- juglans regia pellet Rxhomeo Private Limited d.b.a. Rxhomeo, 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.

ACTIVE INGREDIENT

JUGLANS REGIA HPUS 1X and higher

USES

Facial Acne

INDICATIONS

Condition listed above or as directed by the physician

DOSAGE

Adults- Take 4 or 6 Pellets by mouth, three times daily or as suggested by physician. Children 2 years and older- take 1/2 the adult dose.

WARNINGS

This product is to be used for self-limiting conditions

If symptoms do not improve in 4 days, or worsen, discontinue use and seek assistance of health professional

As with any drug, if you are preganant, or nursing a baby, seek professional advice before taking this product

Keep this and all medication out of reach of children

Do not use if capseal is broken or missing.

Close the cap tightly after use.

INACTIVE INGREDIENTS

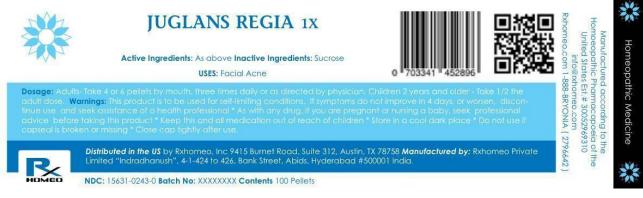
Sucrose

STORAGE

Store in a cool dark place

QUESTIONS OR COMMENTS

www.Rxhomeo.com | 1.888.2796642 | info@rxhomeo.com Rxhomeo, Inc 9415 Burnet Road, Suite 312, Austin, TX 78758



J	UGLANS RE	GIA							
ju	glans regia pellet								
F	Product Informa	tion							
Product T ype			HUMAN OTC DRUG Item Code (Source)		NDC:15631-02		243		
F	Route of Administra	e of Administration ORAL							
A	Active Ingredien	nt/Active Moi	ety						
		Basis of Strength		Strength					
E	NGLISH WALNUT (ENGLISH WALNUT		1 [hp_X]					
I	nactive Ingredie	ents							
		Strength							
s	SUCROSE (UNII: C151H8M554)								
P	ackaging								
#	Item Code		Package Description		Marketing S	Start Date	Marketin	g End Date	
1	NDC:15631-0243-0	100 in 1 PACKA	GE; Type 0: Not a Combination P	roduct	0 1/0 1/20 18				
2	NDC:15631-0243-1	200 in 1 PACKA	GE; Type 0: Not a Combination P	Product	0 1/0 1/20 18				
3	NDC:15631-0243-2	400 in 1 PACKA	GE; Type 0: Not a Combination P	roduct	0 1/0 1/20 18				
4	NDC:15631-0243-3	750 in 1 PACKA	GE; Type 0: Not a Combination P	roduct	0 1/0 1/20 18				
5	NDC:15631-0243-4	2500 in 1 PACK	AGE; Type 0: Not a Combination	Product	0 1/0 1/20 18				
6	NDC:15631-0243-5	12500 in 1 PACK	AGE; Type 0: Not a Combination	n Product	0 1/0 1/20 18				

Marketing Information									
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date						
unapproved homeopathic		09/03/2015							
unapproved homeopathic		09/03/2015							

Labeler - Rxhomeo Private Limited d.b.a. Rxhomeo, Inc (650833994)

Establishment									
Name	Address	ID/FEI	Business Operations						
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc		650833994	manufacture(15631-0243) , label(15631-0243)						

Revised: 3/2020

Rxhomeo Private Limited d.b.a. Rxhomeo, Inc