

VEXASYN- dressing, wound and burn, hydrogel w/drug and/or biologic
Sterling-Knight Pharmaceuticals, LLC

Vexasyn Gel
Prescription Only | Rx Only

Vexasyn is a soothing wound gel that promotes a moist wound environment that is ideal for the healing process.

Directions:

Cleanse wound then apply Vexasyn to the wound and skin surrounding the wound 3 - 4 times daily.

Indications:

For the local management of painful skin wounds, including:

- Pressure ulcers
- Venous stasis ulcers
- Superficial wounds and scrapes
- 1st and 2nd degree burns

Warnings:

External use only. Do not use this product if you are allergic to any of the ingredients. Avoid contact with eyes. Sterile unless opened or damaged.

Caution:

Federal law restricts this device to sale by or on the order of a licensed healthcare practitioner.

Storage:

Store at 25°C (77°F); excursions permitted to 15°-30°C (59°-86°F). Protect from freezing [See USP Controlled Room Temperature].

Ingredients:

Ingredients: Water, Lidocaine HCl, Collagen, Aloe Vera gel, Sodium alginate, Glycerin, Hydroxyethylcellulose, Triethanolamine, Benzethonium chloride.

CALL YOUR DOCTOR ABOUT SIDE EFFECTS.

Call your doctor about side effects. You may report side effects to Sterling Knight Pharmaceuticals, LLC at 1-888-460-1531 or the FDA at 1-800-FDA-1088.

How Supplied

Vexasyn wound gel is supplied as a gel in:
1 oz. (28.33g) tube, NDC 69336-831-01

Distributed By:

Sterling-Knight Pharmaceuticals, LLC
Ripley, MS 38663

Rev 121517-1

Principal Display Panel

NDC 69336-831-01 Rx Only

Vexasyn GEL

 **Sterling | Knight**
PHARMACEUTICALS NET WT.
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Manufactured for:
Sterling-Knight Pharmaceuticals, LLC | Ripley, MS 38663
Dispense in Original Container

Item 831
Rev 122017-1


N 3 69336 81301 3

VEXASYN

dressing, wound and burn, hydrogel w/drug and/or biologic

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NHRC:69336-831
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Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYETHYL CELLULOSE (100 MPA.S AT 2%) (UNII: R33S7TK2EP)	
TROLAMINE (UNII: 9O3K93S3TK)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BOVINE TYPE I COLLAGEN (UNII: FHJ3ATL51C)	

Product Characteristics

(SPLSTERILEUSE)	false
(SPLMRISAFE)	true

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NHRIC:69336-831-01	28.33 g in 1 BOTTLE; Type 0: Not a Combination Product	01/02/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
premarket notification	K020540	01/01/2017	

Labeler - Sterling-Knight Pharmaceuticals, LLC (079556942)

Establishment

Name	Address	ID/FEI	Business Operations
Sterling-Knight Pharmaceuticals, LLC		079556942	label(69336-831) , analysis(69336-831)

Revised: 7/2018

Sterling-Knight Pharmaceuticals, LLC