Givosiran: Latex Content

The following information is provided in response to your unsolicited inquiry. It is intended to provide you with a review of the available scientific literature and to assist you in forming your own conclusions in order to make healthcare decisions. This document is not for further dissemination or publication without authorization.

The full Prescribing Information for GIVLAARI® (givosiran) is provided here. Alnylam Pharmaceuticals does not recommend the use of its products in any manner that is inconsistent with the approved Prescribing Information. This resource may contain information that is not in the approved Prescribing Information.

If you are seeking additional scientific information related to Alnylam medicines, you may visit the Alnylam US Medical Affairs website at RNAiScience.com.

INDEX

DRUG PRODUCT CONTAINER

The givosiran vial closure is a fluoropolymer-coated rubber stopper. The stopper is not made with natural rubber latex or dry natural rubber.[1]

GIVLAARI PRESCRIBING INFORMATION – RELEVANT CONTENT

The DESCRIPTION section provides the following information[2]:

> *GIVLAARI is supplied as a sterile, preservative free, 1-mL colorless-to-yellow solution for subcutaneous injection containing 189 mg givosiran in a single dose, 2-mL Type 1 glass vial with a fluoropolymer-coated rubber stopper and a flip-off aluminum seal. GIVLAARI is available in cartons containing one single-dose vial each. GIVLAARI is formulated in Water for Injection. Sodium hydroxide and/or phosphoric acid may have been added for pH adjustment during product manufacturing.*

ABBREVIATIONS

mg = milligram; mL= milliliter

Updated 27 June 2024

REFERENCES

1. Alnylam Pharmaceuticals. Data on File. MED-ALL-AS1-2200041.
2. GIVLAARI (givosiran) Prescribing Information. Cambridge, MA: Alnylam Pharmaceuticals, Inc.