



RenehaVis™

For knee dynamic needs

Sterile Sodium Hyaluronate Solution

- > Dual molecular weight
- > Dual concentration



The sequential injection of 2 HA

Cushioning ②

①
Lubricating



For the relief of the symptoms of osteoarthritis of the knee by providing support and lubrication to the knee joint.





RenehaVis™ is two clear solutions of sterile sodium hyaluronate in a phosphate buffered saline contained in a two chamber pre-filled syringe for single intra-articular injection into the synovial space of the joint.

RenehaVis™ LMW Sodium hyaluronate 15.4mg/0.7ml and HMW Sodium hyaluronate 7.0mg/0.7ml For the relief of the symptoms of osteoarthritis of the knee by providing support and lubrication to the knee joint.

RenehaVis™ 0.7ml LMW and 0.7ml HMW, terminally sterilised by moist heat, is enclosed within a glass, ready

to use, disposable syringe. The syringe is packed within a blister pack and an outer cardboard carton.

RenehaVis™ is a sterile pre-filled two-chamber glass ready to use disposable syringe containing:

Chamber 1 > Sodium hyaluronate **Low Molecular Weight (LMW)**: 0.7ml sterile **2.2%** sodium hyaluronate **1x10⁶ Da** molecular weight.

Chamber 2 > Sodium hyaluronate **High Molecular Weight (HMW)**: 0.7ml sterile **1.0%** sodium hyaluronate **2 x10⁶ Da** molecular weight.

Dosage and Administration

Injection of **RenehaVis™** should only be made by a Healthcare Professional trained in the technique.

The dosage regimen is injection into the affected synovial joint space once a week for up to three injections depending on the severity of the degenerative change to the knee joint.

Clean the skin around the injection site with antiseptic and allow to dry before injection is given. If joint effusion is present it should be aspirated before injection of **RenehaVis™**.

Aspiration of a small amount of synovial fluid as part of the injection procedure to ensure the correct positioning of the needle is possible. Before proceeding, ensure that the plunger rod is tightly screwed into the plunger stopper.

The contents of the syringe are sterile and should be injected using a sterile needle of an appropriate size (25 gauge needle is recommended). The syringe is fitted with a Luer lock (6%).

Discard the syringe and needle after single use.

Uses

For the relief of pain and stiffness of the knee joint in patients with degenerative changes to the synovial joint. The duration of effect in patients with grade 1 to 3 medial compartment osteoarthritis has been demonstrated to be up to twelve months.

The performance of **RenehaVis™** is due to its biocompatibility and physicochemical properties. The LMW and HMW sodium hyaluronate contained in **RenehaVis™** is a biopolymer composed of repeating disaccharide units of N-acetylglucosamine and glucuronic acid and though it is biosynthesised by the bacterium *Streptococcus equi* it has been shown to be the same as the sodium hyaluronate which is found in the human body. **RenehaVis™** supplements the endogenous Sodium Hyaluronate found naturally in the synovium but which has been depleted by degenerative and traumatic changes to the synovial joint.

Contra-indications

Patients with known sensitivity to sodium hyaluronate.

Warnings and Precautions

Do not inject **RenehaVis™** if the area of the injection is infected or where there is evidence of skin disease. **RenehaVis™** pre-filled syringe is single use. The contents of the syringe should be

used for one injection only. Any remaining sodium hyaluronate should be discarded. If a syringe is retained for a subsequent injection there is a risk of contamination resulting in the possible infection of the patient and/or foreign body reaction.

RenehaVis™ should not be re-sterilised as the device performance may be compromised which could cause serious harm to the patient's health and safety. Sodium hyaluronate is manufactured by fermentation of *Streptococcus equi* and rigorously purified. However, the physician should consider the immunological and potential risks that can be associated with the injection of any biological material. Do not use for children. There is no evidence concerning the safety of **RenehaVis™** in human pregnancy and lactation. Administration during pregnancy and lactation is at the discretion of the orthopaedic surgeon. Do not use if sterile packaging has been damaged. Do not use after the expiry date. Follow national or local guidelines for the safe use and disposal of needles. Obtain prompt medical attention if injury occurs.

Adverse Reactions

Transient pain and swelling may occur with intra-articular injections. Transient increases in inflammation in the injected synovial joint following injection of **RenehaVis™** may occur in patients with inflammatory osteoarthritis. Rarely an inflammatory reaction could occur which may or may not be associated with **RenehaVis™**.

Incompatibilities

RenehaVis™ has not been tested for compatibility with other substances for intra-articular injection. Therefore the mixing or simultaneous administration with other intra-articular injectables is not recommended.

Storage

Store between 2°C and 25°C. Do not freeze. Protect from light.

Do not use if sterile packaging has been damaged. Do not use after expiry date.



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