## Indications

- Pain or limited mobility due to osteoarthritis (arthrosis), post-traumatic conditions or joint and tendon and tendon lesions.

## Application

- In cases of acute and chronic tendinopathies and/or associated joint disability,
- In the process of tendon repair, even after surgical interventions,
- For the integration of joint fluid, allowing the physiological and rheological properties of joints and tendons affected by osteoarthritis to be restored.

## Warnings

SINOVIAL 50 2.5 ml must only be injected by a physician.
The contents of the ampoule-syringe are sterile.
The syringe is packaged in a sealed blister.
The outer surface of the syringe is not sterile.
Do not use after the expiry date stated on the package.
Do not use if the packaging is open or damaged.
The injection site must be on healthy skin.
Do not inject into vessels.
Do not inject outside the joint cavity, into the synovial tissue or joint capsule. Do not inject in case of abundant intra-articular effusion.
Do not re-sterilise.
The device is intended for single use only.
Do not reuse to avoid risk of contamination.
Once opened, consume immediately and discard after use.
Keep out of reach and sight of children.
After an intramuscular injection or into the synovial membrane of a tendon, the patient should be advised to avoid
any intense physical activity and return to normal activity only after a few days.
The possible presence of air bubbles does not affect the properties of the product.
Do not mix SINOVIAL 50 2.5 ml with disinfectants such as quaternary ammonium salts or chlorhexidine, as a precipitate may form.
Extra-articular infiltration of SINOVIAL 50 2.5 ml may cause local side effects.
When using SINOVIAL 50 2.5 ml, symptoms such as pain, warmth, redness or swelling may occur at the injection site.
These secondary symptoms can be relieved by applying ice to the treated area.
They usually disappear after a short time. It is the doctor's responsibility to ensure that patients inform him/her of any side effects occurring after treatment.
SINOVIAL 50 2.5 ml must not be injected if there is an infection or severe inflammation of the joint or if the patient has a skin disease or infection at the injection site.

The product should not be used if there is a known hypersensitivity to sodium hyaluronate or to any other ingredient in the preparation.
To date, no interactions of SINOVIAL 50 2.5 ml with other drugs are known.
Do not mix with disinfectants such as quaternary ammonium salts or chlorhexidine as a precipitate may form.
Do not inject the drug if there is infection or severe inflammation of the joint, or if the patient has skin disease or infection at the injection site.
The product should not be used if there is a known hypersensitivity to sodium hyaluronate or any other ingredient in the preparation.

## Warnings

SINOVIAL 32 2 ml must only be injected by a doctor.
The contents of the ampoule-syringe are sterile.
The syringe is packaged in a sealed blister.
The outer surface of the syringe is not sterile.
Do not use after the expiry date stated on the packaging.
Use on healthy skin.
Do not inject into vessels.
Do not inject outside the joint cavity, into synovial tissue or joint capsule.
Do not inject in case of profuse intra-articular exudate.
Do not re-sterilise.
The device is intended for single use only.
Do not reuse to avoid risk of contamination.
Once opened, consume immediately and discard after use.
Keep out of reach and sight of children.
After injection intramedullary or into the synovial membrane of the tendon, the patient should be advised to avoid all intense physical activity and only return to normal activity after a few days.
Do not mix SINOVIAL 32 2 ml with disinfectants such as quaternary ammonium salts or chlorhexidine, as a precipitate may form.
Extra-articular infiltration of SINOVIAL 32 2 ml may cause local side effects.
When using SINOVIAL 32 2 ml, symptoms such as pain, warmth, redness or swelling may occur at the injection site.
These secondary symptoms can be relieved by applying ice to the treated area. They usually disappear after a short time.
It is the doctor's responsibility to ensure that patients inform him/her of any side effects occurring after treatment.
SINOVIAL 32 2 ml must not be injected if there is an infection or severe inflammation of the joint or if the patient has a skin condition or infection at the injection site.
The product should not be used if there is a known hypersensitivity to sodium hyaluronate or to any other ingredient in the preparation.
To date, there are no known interactions between SINOVIAL 32 2 ml and other drugs.
Do not mix with disinfectants such as quaternary ammonium salts or chlorhexidine, as a precipitate may form.
Do not inject the drug if there is infection or severe inflammation of the joint or if the patient has skin disease or infection at the injection site.
The product should not be used if there is a known hypersensitivity to sodium hyaluronate or any other ingredient in the product.