

AESTHETICS non-CQC - Sculptra Consent

Patient Name: _____ Patient DOB: _____ dd / mm / yyyy

Date: _____ dd / mm / yyyy

LMC

No

☐ No

WHAT IS SCULPTRA?

Sculptra is reconstituted by your physician and injected into the skin to increase the volume of depressed areas, particularly to correct skin depressions, such as in skin creases, wrinkles, folds, scars and for skin aging.

MY TREATING PRACTITIONER HAS EXPLAINED:

- ☐ How and when Sculptra is used.
- ☐ I have been given the opportunity to ask questions and have received satisfactory answers to my questions. In particular, I have received information regarding when treatment with Sculptra should not take place and have also been informed of precautions, warnings for use with these products and common injection related reactions. These reactions include bruising, itching, pain, transient bleeding, redness or swelling in the treatment area. These reactions are usually resolved within a few days to one week. In some cases more serious adverse events have been reported, but these are rare. These reported reactions include papules/nodules, swelling/oedema, pain, granuloma, symptoms of visual disturbance, infection/abscess, mass/induration, paraesthesia and facial nerve paralysis, erythema, inflammation, bruising/ hematoma, discoloration, deformity, scarring/atrophy, hypersensitivity, pruritus, rash, muscle disorders and ischemia/necrosis.
- ☐ In some cases injection site nodules have occurred early within weeks, or with late onset several months to over one year post-injection. Such nodules are occasionally associated with inflammation or discoloration.
- ☐ In some cases, the nodules were reported to resolve spontaneously or following treatment, but may have a prolonged time or may need surgical excision. Isolated rare cases of vision abnormalities including blindness have been reported following injection of Sculptra into the temple area, periorbital areas or cheek.
- ☐ My practitioner has also informed me that topical anaesthetic cream might be used to provide pain relief. I have received information regarding when and what topical anaesthetics will be used, and information regarding contraindications, warnings/ precautions of use of these products and potential side effects.
- ☐ My treating practitioner has also informed me that, depending on the treated area and injection technique, effects from the Sculptra treatment can last up to 25 months, but this period may vary, either longer or shorter. Follow-up treatment can help maintain the desired correction.
- ☐ I have responded to questions pertaining to any hypersensitivity to anaesthetic agents, to Sculptra, to any of the constituents, and to my medical history honestly. I have received a post-treatment advice leaflet as well as information about its contents. I understand the importance of following the advice indicated in the checklist.
- ☐ I have informed my practitioner of all my previous aesthetic treatments, including injections, surgeries, chemical peels etc.
- ☐ I have read and understood the above and hereby consent to treatment with Sculptra
- ☐ I confirm that I have discussed what the procedure is likely to involve, the benefits and risks of any available alternative treatments (including no treatment) and any particular concerns of this patient.

Patient Signature *

Patient Name: _____ Date: _____ dd / mm / yyyy

Practitioner Signature *

Practitioner Name: _____ Date: _____ dd / mm / yyyy