

This information is valid for the Slovak Republic and may differ for other countries.
For detailed information, please contact the relevant foreign representative:

<http://www.imuna.sk/en/o-nas/zahranicne-zastupenia-a-business-development/>

Abridged Drug Information

Name of drug: STAFAL. **Composition:** 1 ml of the drug contains Phagi particulae contra staphylococcus polyvalens with 1×10^7 phage particles at minimum. **Therapeutic indications:** STAFAL is intended exclusively for local application in infections caused by staphylococcal strains. It is intended for destruction of staphylococcal cells at the site of ongoing infection. **Dosage and route of administration:** The medication should be applied in affected sites, surgical wounds and potential reservoirs by lavage, compressing, spraying, or using a bandage soaked in STAFAL. Intranasal administration: 1-2 drops. Application: twice daily according to clinical symptoms. **Contraindications:** Hypersensitivity to any of the components of the drug. **Precautions:** Cloudy preparation is unusable. Store at +2 to +8 °C. Protect from light! No drug or other interactions were observed. **Use in pregnancy/ lactation:** Consideration should be given by the treating physician to using the drug in pregnancy/lactation. Undesirable effects are not anticipated when applied locally. The drug has no influence on the ability to drive vehicles or operate machinery. **Adverse events:** In exceptional cases a burning sensation can develop at the site of application. **Packaging:** One packaging contains 1 glass vial of 10 ml. **Marketing authorisation holder:** IMUNA Ltd., Průmyslová 1472/11, 102 19 Praha 10, Czech Republic. Date of the last revision of the text: September 2012. The drug is available on prescription only. This text does not replace full drug prescribing information. Full drug prescribing information is available upon request.