

AUSTRALIAN PRODUCT INFORMATION – OTODEX (FRAMYCETIN SULFATE, GRAMICIDIN AND DEXAMETHASONE (AS SODIUM METASULFOBENZOATE))

1 NAME OF THE MEDICINE

Otodex (framycetin sulfate, gramicidin and dexamethasone (as sodium metasulfobenzoate))
5 mg/50 µg/500 µg per mL, ear drops.

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each mL contains framycetin sulfate 5mg, gramicidin 50µg and dexamethasone (as sodium metasulfobenzoate) 500µg.

For full list of excipients, see Section 6.1 List of excipients.

3 PHARMACEUTICAL FORM

Otodex is a clear bright colourless aqueous solution.

4 CLINICAL PARTICULARS

4.1 THERAPEUTIC INDICATIONS

Inflammatory and allergic conditions of the ear, e.g. otitis externa. Eczema of the auditory meatus is often present and causes inflammation, exudation and pruritus, which are all rapidly relieved by dexamethasone. Infection, often secondary to scratching, is generally due to staphylococci, *E. coli*, *Pseudomonas* and *Proteus spp.* which respond rapidly to framycetin sulfate.

4.2 DOSE AND METHOD OF ADMINISTRATION

2 or 3 drops should be instilled into the ear three or four times daily; alternatively, a gauze wick kept saturated with the drops may be inserted into the external auditory meatus.

4.3 CONTRAINDICATIONS

Known hypersensitivity to framycetin sulfate or any of the excipients listed in Section 6.1 List of excipients; viral and tubercular lesions; varicella, vaccinia; eardrum perforation.

4.4 SPECIAL WARNINGS AND PRECAUTIONS FOR USE

In patients known to be allergic to Streptomyces-derived antibiotics (neomycin, paromomycin, kanamycin), cross-sensitisation to framycetin sulfate may occur, but not invariably so.

Aminoglycoside antibiotics may cause irreversible, partial or total deafness when applied topically to open wounds or damaged skin. This effect is aggravated by renal or hepatic impairment and by prolonged duration of treatment. The treatment should not be continued after resolution of symptoms.

Visual disturbance may be associated with systemic and topical corticosteroid use. If a patient presents with symptoms such as blurred vision or other visual disturbances, the patient should be considered for referral to an ophthalmologist for evaluation of possible causes which may include cataract, glaucoma or rare diseases such as central serous chorioretinopathy (CSCR) (see Section 4.8 Adverse effects (undesirable effects))

Use in the elderly

No data available.

Paediatric use

No data available.

Effects on laboratory tests

No data available.

4.5 INTERACTIONS WITH OTHER MEDICINES AND OTHER FORMS OF INTERACTIONS

No data available.

4.6 FERTILITY, PREGNANCY AND LACTATION

Effects on fertility

No data available.

Use in pregnancy

Category D

Gentamicin and other aminoglycosides cross the placenta. There is evidence of selective uptake of aminoglycosides by the fetal kidney resulting in damage (probably reversible) to immature nephrons. Eighth cranial nerve damage has also been reported following *in utero* exposure to some of the aminoglycosides.

Because of their chemical similarity, all aminoglycosides must be considered potentially nephrotoxic and ototoxic to the foetus. It should also be noted that therapeutic blood concentrations in the mother do not equate with safety for the foetus.

Use in lactation

No data available.

4.7 EFFECTS ON ABILITY TO DRIVE AND USE MACHINES

The effects of this medicine on a person's ability to drive and use machines were not accessed as part of its registration.

4.8 ADVERSE EFFECTS (UNDESIRABLE EFFECTS)

Local allergic reactions of the hypersensitivity type have rarely been reported.

Eye disorders:

Not known: blurred vision, chorioretinopathy

Reporting suspected adverse reactions after registration of the medicinal product is important. It allows continued monitoring of the benefit-risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions at www.tga.gov.au/reporting-problems (Australia).

4.9 OVERDOSE

For information on the management of overdose, contact the Poisons Information Centre, telephone number 13 11 26 (Australia).

5 PHARMACOLOGICAL PROPERTIES

5.1 PHARMACODYNAMIC PROPERTIES

Mechanism of action

Framycetin sulfate is a bactericidal antibiotic active against a wide variety of Gram-positive and Gram-negative bacteria commonly found in superficial infections; staphylococci (including strains resistant to other antibiotics), *Pseudomonas aeruginosa*, coliforms and pneumococci.

Gramicidin reinforces the action of framycetin sulfate against streptococci.

Dexamethasone is a highly potent topical corticosteroid. Its topical superiority is particularly apparent in cases in which other corticosteroids have failed.

Clinical trials

No data available.

5.2 PHARMACOKINETIC PROPERTIES

No data available.

5.3 PRECLINICAL SAFETY DATA

Genotoxicity

No data available.

Carcinogenicity

No data available.

6 PHARMACEUTICAL PARTICULARS

6.1 LIST OF EXCIPIENTS

It also contains polysorbate 80, industrial methylated spirit, citric acid monohydrate, sodium citrate, lithium chloride, sodium hydroxide, hydrochloric acid, purified water and is preserved with phenethyl alcohol.

6.2 INCOMPATIBILITIES

Incompatibilities were either not assessed or not identified as part of the registration of this product.

6.3 SHELF LIFE

In Australia, information on the shelf life can be found on the public summary of the Australian Register of Therapeutic Goods (ARTG). The expiry date can be found on the packaging

Discard 4 weeks after opening.

6.4 SPECIAL PRECAUTIONS FOR STORAGE

Store below 25°C. Do not refrigerate.

6.5 NATURE AND CONTENTS OF CONTAINER

8mL bottles.

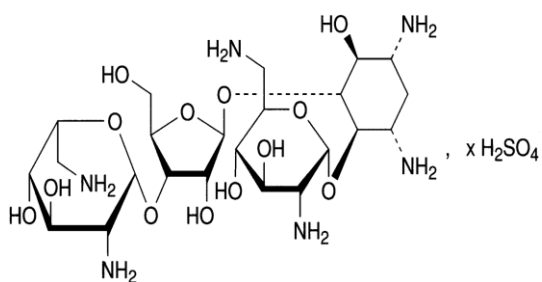
6.6 SPECIAL PRECAUTIONS FOR DISPOSAL

In Australia, any unused medicine or waste material should be disposed of by taking to your local pharmacy.

6.7 PHYSICOCHEMICAL PROPERTIES

Chemical structure

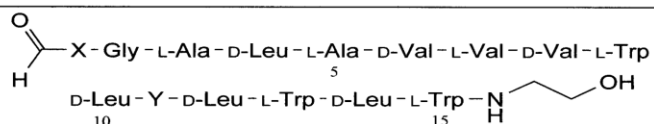
Framycetin sulfate



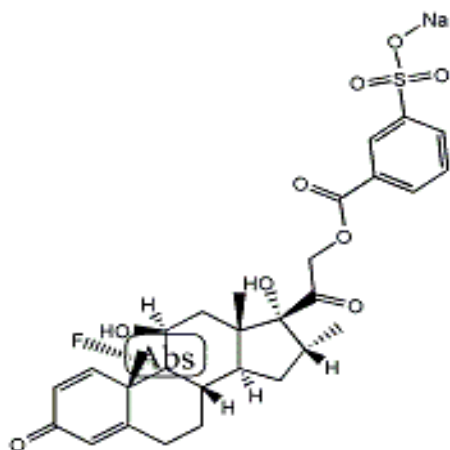
Molecular weight – 712.72

Gramicidin

Gramicidin	X	Y	Mol. formula	M_r
A1	L-Val	L-Trp	$C_{99}H_{140}N_{20}O_{17}$	1882
A2	L-Ile	L-Trp	$C_{100}H_{142}N_{20}O_{17}$	1896
B1	L-Val	L-Phe	$C_{97}H_{139}N_{19}O_{17}$	1843
C1	L-Val	L-Tyr	$C_{97}H_{139}N_{19}O_{18}$	1859
C2	L-Ile	L-Tyr	$C_{98}H_{141}N_{19}O_{18}$	1873



Dexamethasone sodium metasulfobenzoate



Molecular Weight – 598.6

CAS number

Framycetin sulphate – 4146-30-9

Gramicidin – 1405-97-6

Dexamethasone sodium metasulfobenzoate – 3936-02-5

7 MEDICINE SCHEDULE (POISONS STANDARD)

S4

8 SPONSOR

sanofi-aventis australia pty ltd
12-24 Talavera Road
Macquarie Park NSW 2113

9 DATE OF FIRST APPROVAL

01 April 1993

10 DATE OF REVISION

6 April 2018

SUMMARY TABLE OF CHANGES

Section Changed	Summary of new information
All	Reformat to align with new PI format, including additional text to align with the new PI requirements.
4.3	Contraindications expanded and reworded.
4.4	Additional precautions added.
4.8	Additional adverse events added.