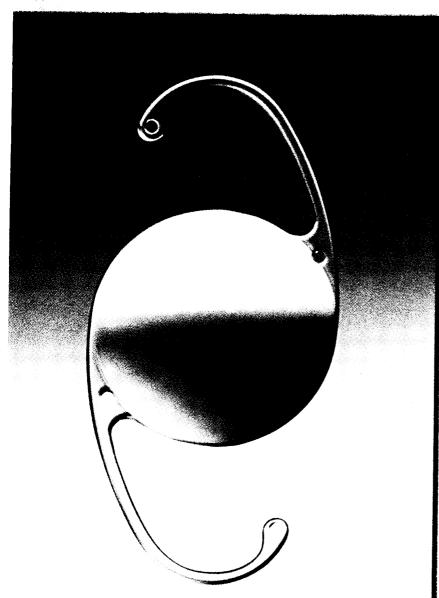
The J-Flex posterior chamberlens from CILCO.



The J-Flex[™] lens is a single-piece posterior chamber lens with loops as well as optic lathe cut from Perspex® CQ polymethylmethacrylate.

This lens is available in uniplaner design (Style SK-1) or with loops angled 10° anteriorly (Style SK-2). Loops measure 0.14 mm imes 0.17 mm. Overall length is 13.0 mm. Lenticular diameter is 6.0 mm.

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TLX: 848 507

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Telephone: (93) 38 06 99
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AN IMPORTA IN THE THE OF HERPES SIMI

ZOVIRAX is a highly selective antiherpes agent with a fundamentally different mode of action, and extremely low toxicity.

A Wellcome discovery, ZOVIRAX is the first antiherpes agent that is activated to any significant extent only when the herpes simplex virus is present. ZOVIRAX is converted to a monophosphate form by a herpes-specific thymidine kinase enzyme. This starts a chain of events resulting in the active

Unique mode of action

compound, the triphosphate form, which inhibits viral replication. In chemically signalling its presence the virus thus seals its own fate.

Because of its unique mode of action, ZOVIRAX can be regarded as an ultra-selective agent. Once "bioactivated" it has a 10 to 30-fold greater affinity for viral DNA polymerase than for cellular polymerase. In tissue culture experiments it was 3,000 times more active against the herpes simplex virus than it was against the host cell?

Because of its ultra-selectivity, ZOVIRAX has extremely low toxicity. A report on this selectivity describes ZOVIRAX as "... a new class of antiviral agent that has extremely low toxicity for normal cells while having an inhibitory activity against HSV which is greater than that of any hitherto known compound." 3

• Highly effective with rapid action

In clinical studies ZOVIRAX has been shown to be superior to idoxuridine. In the trial, healing time was quicker with ZOVIRAX. Antiviral activity has been well demonstrated in *in vitro* studies. ZOVIRAX ... was found to be between 5 and 10 times more active than cytarabine, idoxuridine and trifluorothymidine, and more than 100 times more active than vidarabine."²

An agent of promise

Wellcome take particular pride in introducing ZOVIRAX, a preparation which we believe heralds a new era in antiviral chemotherapy.

1. Collum, L M T et al Brit. J. Ophthalmol., (1980), 64, 766 2. J. Antimicrob. Chemother., (1979), 5, 431 3. Proc Natl Acad Sci USA., (1977), 74/12, 5716

PRESCRIBING INFORMATION Presentation Acyclovir 3 per cent w/w in a white soft paraffin base. Uses Treatment of herpes simplex keratitis. Dosage and administration A 1 cm ribbon of ointment should be placed inside the lower conjunctival sac five times a day at approximately four-hourly intervals. Treatment should continue for at least 3 days after healing is complete. Contra-indications Patients with a known hypersensitivity to acyclovir. Warnings and adverse effects For ophthalmic use only. Transient mild stinging immediately following administration occurs in a small proportion of patients. Superficial punctate keratopathy has been reported but has not resulted in patients being withdrawn from therapy, and healing has occurred without apparent sequelae. (PL 3/0150)

ITADVANCE ATMENT EX INFECTIONS







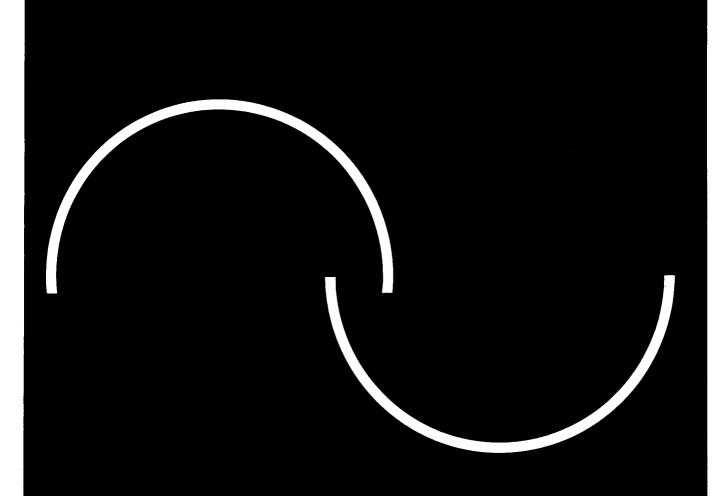
OPHTHALMIC OINTMENT

The effective, selective antiviral









FML, the potent steroid, that's gentle on pressure.

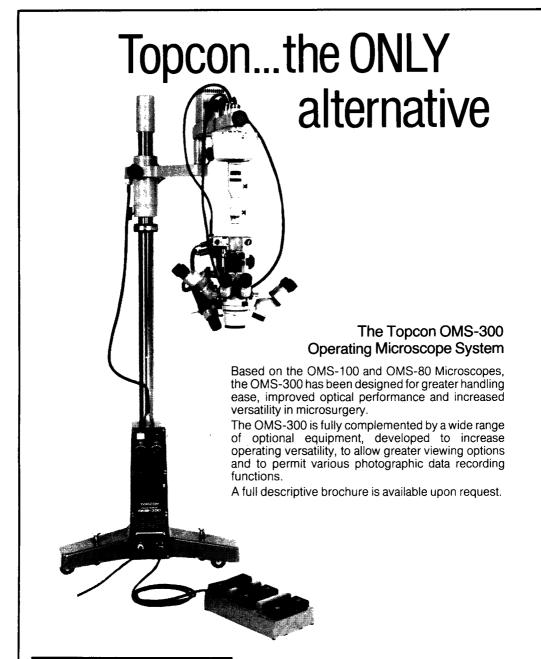
FML (FLUOROMETHOLONE) LIQUIFILM STERILE OPHTHALMIC SUS-PENSION Abbreviated Prescribing Information Uses Topical ophthalmic suspension for steroid responsive inflammation of the papebral and bulbar conjunctiva, cornea, and anterior segment of the globe Dosage and administration 110 2 drops instilled into the conjunctival size two to four times daily Contra-indications, warnings, etc. Contra-indications. Acute superficiency of the segment of the cornea and conjunctival Eucracian Contra-indications, warnings, etc. Contra-indications. Acute superficiency of the eye Warnings. Steroid medication in the treatment of herpes culosis of the eye Warnings. Steroid medication in the treatment of herpes amp microscopy is mandatory. Protonged use may result in glaucorna damage to the optic nerve defects in usual additivation and steroid in the secondary ocular infections from fung or viruses betweated from ocular issue in those diseases causing timining of the cornea or sclera, perforation has been known to occur with use of topical steroids. This preparation contains benzalkonium chiorde and should be used with caution in association with hydrophilic contact lenses. Precautions As fungal infections of the cornea are particularly because the supported in any experience of the cornea are particularly stress on must be supported in any existing the authority of the defects, posterior subcapsular cataract formation, secondary ocular infections from pathogens liberated from ocular issues, perforation of the globe Local side effects of steroid therapy i.e. skin activities and telapietectasia, are especially likely to affect facal skin Legal Category. POM Basic NHS cost. £1 82 for 5mt £2.57 for 10mt Product Lenence, No 3426/0028. References 1. Kuppforman and Lenowitz. Acro. Office and 1986. Statement of the perforation of combine of the policy of the defects, observed on the combine of the policy. Acro. Office and the production of the perforation of the globe Local skin Legal Category. POM Basic NHS cost. £1 82 for 5mt £2.57 for



FML has comparable efficacy to dexamethasone and prednisolone with less tendency to raise IOP than prednisolone, dexamethasone or betamethasone.

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DOSAGE AND ADMINISTRATION Adults: One drop to be instilled into the eye once or twice daily or at the discretion of the physician. Children: At the discretion of the physician.

CONTRA-INDICATIONS, WARNINGS ETC. Ganda 1 - 0.2 should not be used in the case of a narrow angle between the iris and cornea as pupillary dilation may precipitate angle closure. Occasionally, orbital discomfort or red eye (hyperaemia) may occur. Other side effects, such as local irritation and headache are rare. When used in conjunction with miotics, Ganda 1+0.2 should follow the miotic after an interval of 5-10 minutes. Ganda 1+0.2 should not be used if the solution has become dark amber. The contents of the bottle should be discarded one month after the pouch has been opened. Ganda 1+0.2 is fully potent for two years providing the pouch remains unopened

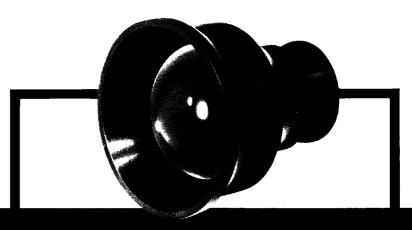
Full prescribing information is available on request

References 1 Romano J., Nagasubramanian S., and Poinoosawmy D. Double-masked cross-over comparison of Ganda 1.02 (Guanethidine 1% and Adrenaline 0.2% mixture) with Guttae Adrenaline 1% (Simplene 1%) and with Pilocarpine 1% (Sno-Pilo 1%). British Journal of Ophthalmology — in press.

2 Mills K. B. Personal communication. 3 Urner-Bloch U., Aeschlimann J. E., and Gloor B. P. (1980). Treatment of Chronic Simple Glaucoma with an

Adrenaline/Guanethidine Combination at Three Different Dosages (Comparative Double-Blind Study) Albrecht v. Graefes Arch. klin. exp. Ophthal. 213, 175-185

A new instrument for wide-field fundus observation



THE RODENSTOCK

PANFUNDOSCOPE

The Panfundoscope is a contact optical system utilizing the practical advantages of indirect ophthalmoscopy. As an accessory to the slit lamp, it provides full synoptic examination of the fundus with its observable area reaching the ocular

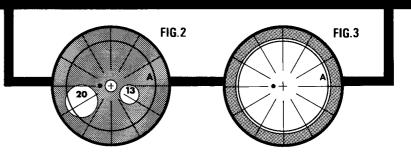
equator. This area can be further extended by movement of the patient's eye, so that even the extreme quadrant peripheries can be readily viewed.

Figures 2 & 3 compare the sizes of the fundus areas which can be simultaneously observed, using different ophthalmoscopic procedures with stationary eye and ophthalmoscope.

In Fig. 2 the white central circle represents the area observed using a conventional ophthalmoscope with upright images. The circle marked 13 corresponds to the synoptic field with inverted image using a lens of 13D; the circle marked 20 indicates the synoptic field with a lens of 20D.

The white circular field in Fig. 3 corresponds to the synoptic field of the Panfundoscope and demonstrates the enormous advantage in field size.

Rodenstock have two types of Panfundoscope available. Type D is a diagnostic contact optical device with a plexiglass sphere weighing only 15 g. Type Th, of greater interest to ophthalmologists, is a therapeutic contact optical device for use in laser coagulation of the fundus. It utilises a silicate glass sphere, weighing 27 g., the glass surface is treated with a non-reflective coating to prevent injury to



observers by reflected laser light.

The advantages of the Panfundoscope for laser coagulation are easy handling, and the ideal synopsis of the working area. Thus the surgeon is not dependent on small keyhole-like observation fields of other optical contact systems. This applies particularly to the treatment of peripheral portions of the medial and lateral fundus areas. Coagulations of the pericentral areas and of the middle periphery as, for example, in diabetic retinopathy, can be performed with an excellent overview of the corresponding area and occur in a considerably shorter time.

As several laser types are mounted to the slit lamp in such a way that the delivery head of the laser beam restricts the working space, the doctor's hand guiding the contact lens is considerably hampered. The Panfunoscope solves this problem, since the fundus image lies in front of the patient's eye. Thus the slit lamp has to be drawn towards the therapeutist providing ample space for the manipulation of the Panfundoscope.

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INDICATIONS Glaucoma. DOSAGE Adults: One capsule at night and in the morning. Contra-indications: Idiopathic renal hyperchloraemic acidosis. Addison's disease or all types of suprarenal gland failure. Long-term administration in patients with chronic congestive angle-closure glaucoma. Precautions: The patient should be cautioned to report any unusual skin rash. Periodic blood cell counts are recommended. The transitory loss of hearing calls for immediate cessation of medication Side-effects: Drowsiness, paraesthesia of extremities and face may occur. Diamox is a sulphonamide derivative and therefore some side-effects similar to those caused by sulphonamides have occasionally been reported. **Drug interactions:** Possible potentiation of the effects of folic acid antagonists, hypoglycaemics and oral anticoagulants may occur. **Package quantities:** Pransparent orange capsules each containing 500mg, of acetazolamide and printed 'L ederle 4 309' in bettles of 30, 100 and 500. **Basic N.H.S.** cost: £21.70 per 100. PL 0095-5074.
*Trademark. **References:** L Garner, L.L. et al. Amer. J. Ophthalmol. 1903. 55, 2-323-327. 2. Lichter, P.R. et al. Amer. J. Ophthalmol. 1978.85, 4-495-502.

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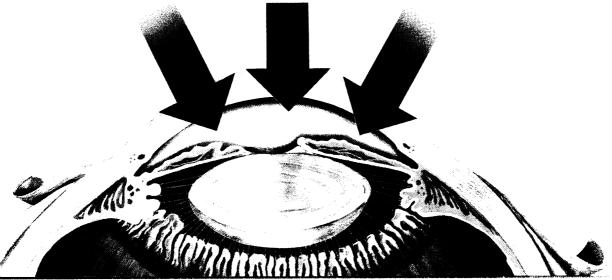
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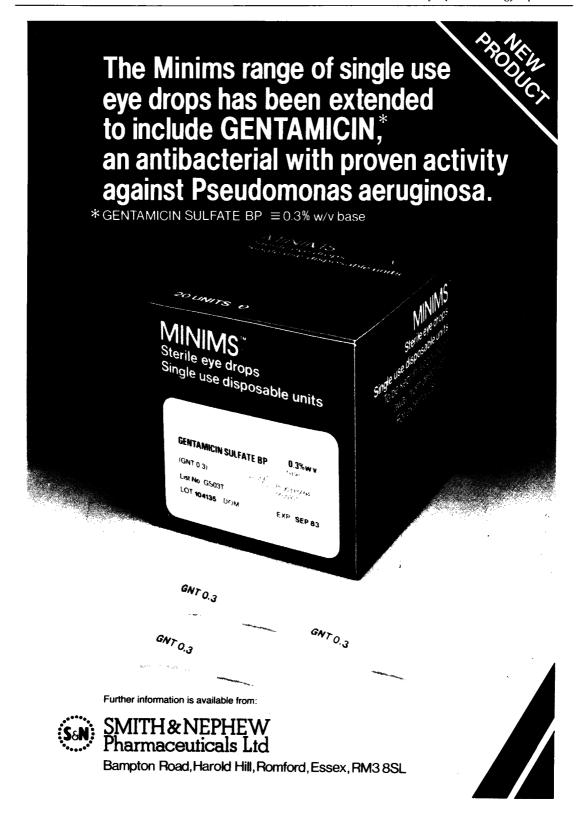
Ophthalmic suspension Dexamethasone 0.1%

fast relief from inflammation



Prescribing information

Presentation Documethasone 3 to in a venicle containing 3.5 in not overoup productions in hydroxycoup methyboillulose integrance oscillations of the anterior segment. A line and or thronic alterior venicles is system between control of the anterior segment. A line and or thronic alterior venicles is system between or the present and productions of the anterior segment. A line and or thronic alterior venicles is system between or standards in the present as a control conjunctivitis, when a conjunctivitis alterior is exercitivated in the conjunctivitis. For unrent manage a negroid of the or alterior of the conjunctivitis and administration of a secretarization of the conjunctivities of the conjunctivities. Advisor unital and so the conjunctivities of the conjunctivities. A days of unital and standards of the conjunctivities of the conjuncti



THE ORIGINAL SINSKEY POSTERIOR CHAMBER LENS

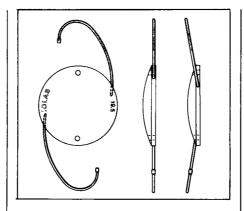


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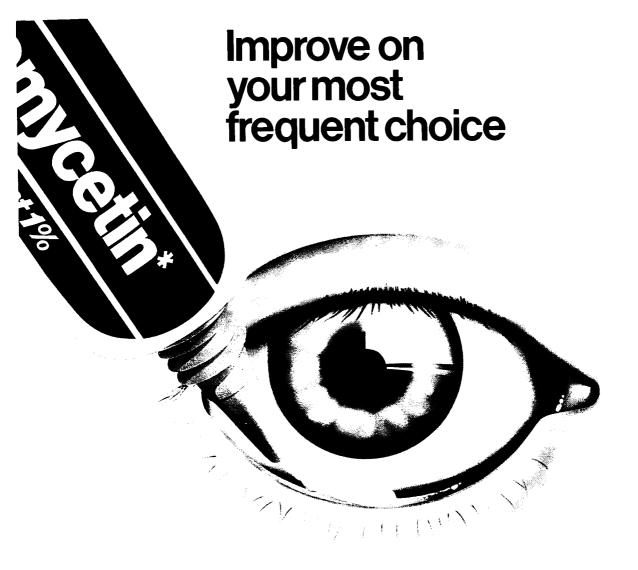


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Chloromycetin and the device showing a tube having a blue nozzle are the trade marks of Parke, Davis and Company for ophthalmic preparations containing chloramphenicol. † Blue Nozzle patent no. 8018334 pending.

P456-UK-May 81

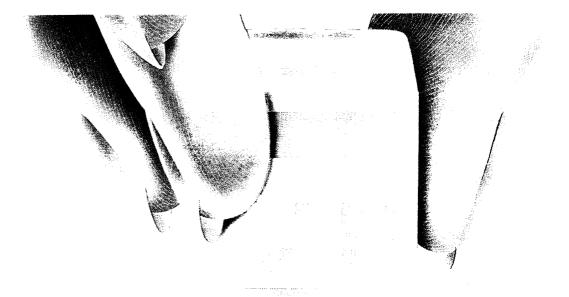
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Eumovate Eye Drops have significantly less effect on intra-ocular pressure than hydrocortisone, betamethasone, prednisolone or dexamethasone eye drops.

'It is therefore of some significance that with clobetasone butyrate it has been possible to dissociate the adverse intraocular pressure effect from the advantageous anti-inflammatory effects, and thus we may well have a "safer" steroid for use in ophthalmology.'

Ramsell TG, Bartholomew RS, Walker SR. Br J Ophthalmol 1980; 64:43-5.

EUMOVOITE EYE Drops (clobetasone butyrate)
EUMOVOITE-N EYE Drops (clobetasone butyrate and neomycin)

Prescribing information

for the treatment of non-infected

Indications Eumovate Eye Drops are indicated

inflammatory conditions of the eye. Eumovate-N Eye Drops are indicated for inflammatory conditions of the eye where secondary bacterial infection is likely to occur. **Dosage and administration**The usual dosage is one to two drops four times a day: for severe inflammatory conditions one or two drops should be instilled into the eye every one or two hours until control is achieved, when the frequency may be reduced. **Contra-indications**

Viral, fungal, tuberculous or purulent conditions of the eye, hypersensitivity to any component of the preparation. Use is contra-indicated if glaucoma is present. Eurovate Drops and Eurovate-N Drops contain benzalkonium chloride as a preservative and therefore should not be used to treat patients who wear soft contact lenses.

Precautions

Although Eumovate Eye Drops have been shown to have little effect on intra-ocular pressure in most patients, those receiving long term treatment should have their intra-ocular pressure monitored frequently.

Cataract is reported to have occurred

after unduly prolonged treatment with some topical corticosteroids and in those diseases which cause thinning of the cornea, perforation has been known to occur in general, topical steroids should not be used extensively in pregnancy, i.e., in large amounts or

for prolonged periods Side effects

Rises in intra-ocular pressure have been reported in susceptible patients but these are generally much less than with other corticosteroid eye preparations, including hydrocortisone. Product Licence numbers

 Product Licence numbers

 Eumovate Drops
 4/0260

 Eumovate-N Drops
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 Basic NHS cost (exclusive of VAT)

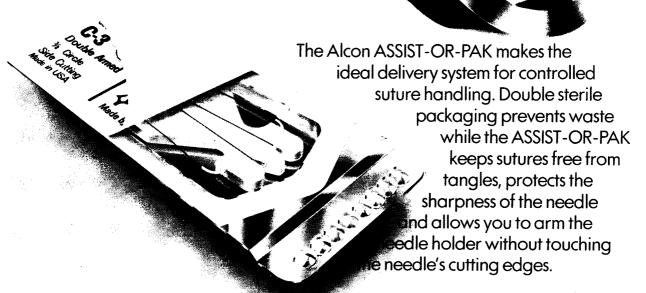
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USES

Elevated intra-ocular pressure including: ocular hypertension, chronic open-angle glaucoma (including aphakia); secondary glaucoma.

DOSAGE AND ADMINISTRATION

Usually one drop 0.25% solution in affected eye twice a day. If necessary change to one drop 0.5% solution twice a day.

CONTRA-INDICATIONS

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Not studied; weigh benefit against possible hazards. **PRECAUTIONS**

'Timoptol' may be absorbed systemically.

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Children

Not currently recommended.

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Ocular irritation, including conjunctivitis, belpharitis, and keratitis reported occasionally. Visual disturbances reported infrequently. Rash and urticaria reported rarely.

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Rare effects reported are aphakic cystoid macular oedema, headache, dry mouth, anorexia, dyspepsia, nausea, dizziness, CNS effects, palpitation, and hypertension.

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0.25% Ophthalmic Solution 'Timoptol', £5.18 per 5 ml pack. 0.5% Ophthalmic Solution 'Timoptol', £5.82 per 5 ml pack.

PRODUCT LICENCE NUMBERS

0.25% Ophthalmic Solution 'Timoptol', 0025/0134 0.5% Ophthalmic Solution 'Timoptol', 0025/0135. ®denotes registered trademark. Issued February 1982.



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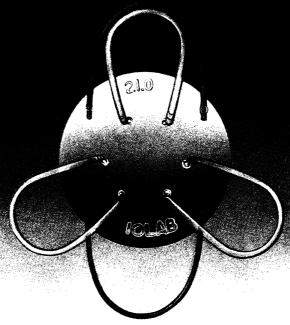
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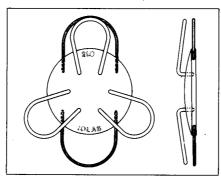
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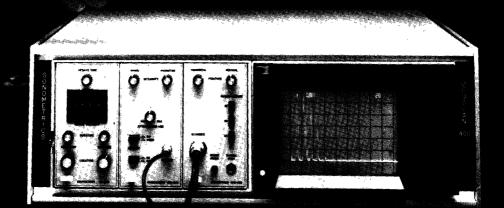
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Directors: Sanford D. Hecht, M.D., Pierre Guibor, M.D., Eugene Wiggs, M.D., and Darrell Wolfley, M.D.

Date: Thursday and Friday June 3-4, 1982

Faculty: Holger Busse, M.D., Universitata Augenklinik. West Germany; Edward Goodman, M.D., Boston University; Pierre Guibor, New York Medical College; Sanford Hecht, M.D., Boston University and Tufts University; Howard M. Leibowitz, M.D., Boston University; Eugene Wiggs, M.D., University of Colorado; Darrell Wolfley, M.D., University of Utah.

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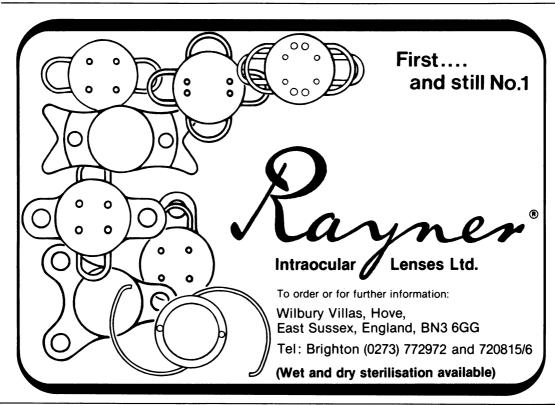
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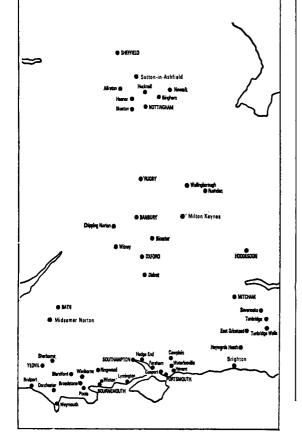
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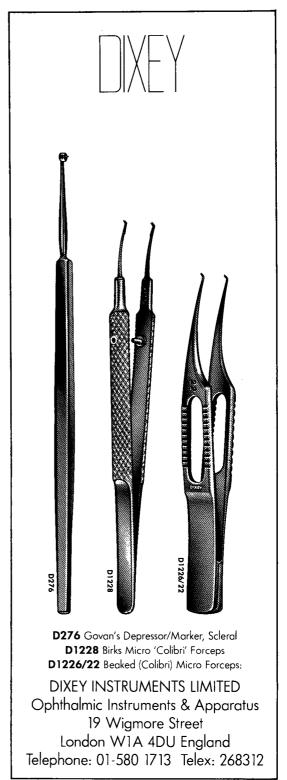
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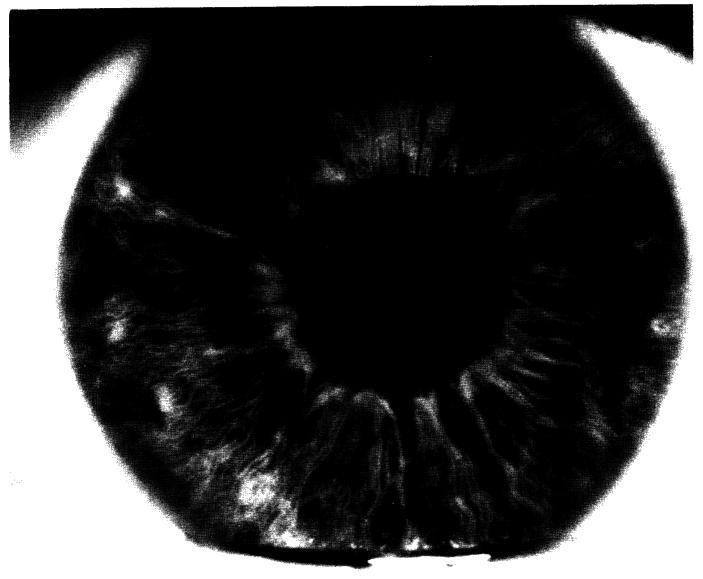
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