Corynebacterium macginleyi: a conjunctiva specific pathogen

Antonia M Joussen, Guido Funke, Frank Joussen, Georg Herbertz

Abstract

Background—Although non-diphtherial corynebacteria are ubiquitous in nature and commonly colonise the skin and mucous membranes of humans, they rarely account for clinical infection. Methods and results—10 patients with unilateral conjunctivitis are described in which Corynebacterium macginleyi was isolated. This species has only recently been reported to be exclusively isolated form ocular surfaces. C macginleyi was uniformly susceptible to topical antibiotics commonly used in ophthalmology. Conclusion—Despite the fact that the pathogenicity of C macginleyi is not yet assured, this micro-organism should be recognised as a potential cause of bacterial superinfections. Appropriate antibiotic therapy leads to its elimination and resolution of the conjunctivitis.

Patients and microbiological analysis

We examined the conjunctival smears of 181 patients with clinically diagnosed bacterial conjunctivitis. Part of the conjunctival smear material was used for light microscopy. For enrichment of the micro-organisms patients' material was cultured on Columbia agar supplemented with 5% sheep blood, on chocolate agar, both in air with 3% carbon dioxide. For detection of Gram negative rods we used Endo agar cultured in ambient air. Anaerobic cultures were on Schaedler's blood-agar in oxygen-free, carbon dioxide enriched atmosphere. Thioglycolate broth was used as fluid enrichment medium. The aerobic cultures were inspected for the first time after overnight incubation. Total incubation time for all media was 48 hours. Biochemical identification was performed using the commercial API-Coryne system (bioMerieux Marcy l'Etoile France). A prolonged incubation time was not necessary. For evaluation of results we used the data given by Funke et al.8 and the API-Coryne database 2.0. Susceptibility tests were performed according to the standards for agar diffusion tests published by DIN, on Müller-Hinton agar supplemented with 5% sheep blood, incubated in air.

Results

A total of 181 patients with the clinical diagnosis of bacterial infection of the conjunctiva were examined. In 107 cases cultures yielded one or more micro-organisms. We found pathogenic bacteria in following frequency: 18.7% Staphylococcus aureus, 12.1% C macginleyi, 10.3% Streptococcus pneumoniae, 8.4% Haemophilus influenzae.

We found the pathogenic species Arcanobacterium haemolyticum, Branhamella catharrhalis,
Neisseria meningitidis, Pseudomonas aeruginosa, cultivated in one case each.

Bacteria of apparently low pathogenic characteristics had been isolated as follows: 40.1% Staphylococcus epidermidis, 13.1% Staphylococcus epidermidis group. Besides these, Acinetobacter spp and Enterobacteriaceae have been found and, rarely, Micrococcus species, non-haemolytic Streptococcus species, and Aeromonas hydrophila. C. macginleyi was isolated in a total of 10 patients from the conjunctival smears in 13 separate cases. Three patients appeared with bilateral infection, two others had recurrent disease 2–4 months after primary detection of C. macginleyi. The age of these patients ranged from 33 to 86 with an average of 64.5. Five patients were females, five were males. All appeared with unilateral conjunctivitis worsening over a few days, without any history of infection. Visual acuity was unchanged as was intraocular pressure. Most patients showed conjunctival hyperaemia with moderate follicular reaction (Fig 1A). Seven of the patients had a whitish discharge. There were no signs of keratitis or intraocular infection or inflammation in any patient. C. macginleyi was the only bacterial micro-organism isolated in five out of 13 swabs. From these patients one had a bacterial superinfection of a viral aetiology. In eight of the 13 swabs with C. macginleyi, other bacterial micro-organisms were co-cultivated. Even C. pseudodiphthericum was isolated in one case.

Tests for Chlamydia trachomatis or cultures for fungi were performed only if clinical suspicion existed. There were no positive results.

Light microscopy of the conjunctival swab before cultivation sometimes showed Gram positive rods and only in few cases a significant increase in leucocytes. In cultures on Columbia agar supplemented with 5% sheep blood almost no growth after 1 night of incubation was visible. Under anaerobic conditions there is no growth to be seen after 48 hours of cultivation. Even carbon dioxide enriched air retards the growth of C. macginleyi. Colony size after 48 hours of incubation in air remained below 0.2 mm in diameter. These colonies were stained by Gram reagents and analysed microscopically. They consisted of coryneform rods with a distinctly striped pattern, forming irregular arrangements, similar to Chinese letters (Fig 1B).

Susceptibility tests were performed for antibiotics, which are usually applied topically for external infections (Table 1). All patients were sensitive to gentamicin. Under treatment with gentamicin 0.5% eye drops four times daily and gentamicin ointment at night, eight out of 10 patients recovered uneventfully after 8 days. The remaining conjunctival irritation was successfully treated with zinc sulphate-containing eye drops (0.2%) over 2–3 weeks.

Discussion

More specific detection methods in recent years have allowed further investigation of the coryneform bacteria. C. macginleyi was first identified in 1995 by Riegel et al during investigations on lipophilic corynebacteria. It has been uniquely isolated from ocular surfaces. The first 18 cases of C. macginleyi conjunctivitis have been detected in Switzerland. Within the recent past we found in 10 patients 13 cases of C. macginleyi conjunctivitis in Germany, indicating that the presence of this micro-organism is not geographically limited.

Thiel et al report on increasing percentage of patients positive for corynebacteria. We found in our patients 18.7% Staphylococcus.

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S = sensitive, I = intermediate, R = resistant, — = not tested.

Table 1

Corynebacterium macginleyi: a conjunctiva specific pathogen

Figure 1 (A) Corynebacterium macginleyi conjunctivitis—clinical appearance. (B) Corynebacterium macginleyi—microscopic appearance (original magnification ×1000, Gram stain).

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Weiss et al. described by Funke almost all of the topical antibiotic agents. C. macginleyi. In our study group we found C. macginleyi predominantly in middle aged patients with no preference regarding sex. This is in accordance with the data described by Funke et al. Younger children have not been analysed; however, infection with C. macginleyi can be expected according to Weiss et al.

Many systems fail to identify C. macginleyi strains correctly. The identification of C. macginleyi is described in the review by Funke et al. Positive nitrate reduction, negative pyrazinamidase, positive alkaline phosphatase, acid production from glucose and sucrose but not from maltose. It is important to use a system for biochemical identification that is able to detect C. macginleyi and that the database employed contains information about C. macginleyi.

C. macginleyi is reported to be sensitive to almost all of the topical antibiotic agents. This might be one reason for only rare reports in the literature, as most patients are successfully treated. However, with the availability of more sensitive detection systems diagnosis is likely to become more frequent. Here we show for the first time more resistant species of this bacterium, especially to erythromycin and kanamycin, making susceptibility testing advisable in these patients.

Till now reason for exclusive infection of the ocular surface is unknown. From our data, as well as from Funke et al., C. macginleyi seems to predominantly affect already injured conjunctivae. In our patients only three cases showed C. macginleyi as only infectious agent. All patients had obvious signs of infection. Further investigations should analyse the normal healthy conjunctival flora for C. macginleyi.

The role of C. macginleyi in causing severe ocular infections is also unknown. It has not yet been shown in corneal ulcers, but other low pathogenic agents such as Mycobacterium chelonae can mimic corynebacterium keratitis, so that C. macginleyi is also likely to be able to cause serious damage to the cornea. All cases suspicious for C. diphtheriae should therefore also be PCR-detected for C. macginleyi.

C. minutissimum, as well as Propionibacterium acnes, both belonging to the low pathogenic coryneform bacteria, may cause intraocular infections such as endophthalmitis following penetrating ocular trauma, keratoconjunctivitis, and cataract extraction. Similar events could potentially cause C. macginleyi endophthalmitis.


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