

STUDIES ON SIGNIFICANT ASPECTS OF *NEISSERIA MUCOSA* IN CLINICAL MICROBIOLOGY

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ABSTRACT

The current advances in cellular microbiology, genomics, and immunology has opened novel horizons in the considerate of neisserial pathogenesis and in the definition of novel prophylactic involvement. It is now clear that *Neisseria mucosa* has evolved a number of surface structures to mediate communication with host cells and a numeral of mechanisms to undermine the immune system and escape complement-mediated killing. In this review, we revealed significant aspects of *Neisseria mucosa* for readers.

Keywords: *Neisseria mucosa*, review.

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Introduction

Neisseria mucosa was described by von Lingelsheim in 1906. One year later 1960s, Berger and coworkers showed that *Neisseria mucosa* may be differentiated from the *N. sicca*, *N. subflava* biovar *perflava* by a nitrate reduction test. There are no biochemical methods for differentiating between these latter species [1,4]

Berger and coworkers showed that *N. subflava* biovar *perflava* and *N. sicca* were serologically distinct [2,3,6].

Neisseria mucosa belongs to the family *Neisseriaceae*. It is clinically important pathogenic bacteria and is relatively easy to identify from other *Neisserial* species [1,2,6]. The other species of *Neisseria* such as *N. lactamica* and *N. cinerea* are commonly painstaking

commensals, but have been alarmed as causes of infection in patients who are immunocompromised. More up to date species to the genus *Neisseria* are *N. oralis*, *N. shayegani*, *N. wadsworthii*, *N. zoodegmatis* and *N. animaloris* isolated from human clinical samples [1,2,3,6].

2. Taxonomy [1,2,3,]

Phylum: - Proteobacteria

Class: - Betaproteobacteria

Order: - Neisseriales

Family: - Neisseriaceae

Genus: - *Neisseria*

Species: - *mucosa*

3. Media for Isolation [1,2,3,4,5]

Non- Selective media:

- a) Blood agar
- b) Chocolate agar
- c) Muller Hinton starch casein Hydrolysate agar

Selective Media

- a) GC selective agar
- b) Modified T-M agar

4. Incubation Conditions

5-10% CO₂ at 37±2°C, 24hrs - 48hrs.

5. Morphological appearance

Neisseria mucosa shows adherent colonial morphology, pigmented or non-pigmented and opaque [1,2,5,6]. However, they form smooth, round, moist, or uniform gray/brown/red colonies with a greenish color underneath on primary isolation medium. It is gram negative, non motile, non spore forming bacteria [3,4].

6. Biochemical appearance

Neisseria mucosa shows positive results for Oxidase, Catalase, chitinase, iodine, carbonic anhydrase test, nitrite and nitrate production, Sugar fermentation tests (Fructose, Glucose, Maltose and Sucrose) [4,5,6] while they show negative results for acid production from Lactose and Mannose, Beta- galactosidase, Gamma glutamyl transferase and tributyrin hydrolysis [4,5,6].

7. Chemistry Of *Neisseria Mucosa*

Neisseria mucosa is a part of normal human nasopharyngeal flora and infrequently causes human infections, including meningitis, cerebrospinal fluid shunt infection [1,2,3,4,5,6].

Neisseria mucosa LPS consists of lipid A, a central part containing two 2-keto-3-deoxy-octulosonic acids (KDO) and two heptoses (L-glycero-D-manno-heptopyranoside) substituted with short polysaccharide side chains. Due to the short polysaccharide side chains and lack of repeating O antigen units, meningococcal LPS is frequently referred to as lipo-oligosaccharide (LOS). The inherited basis for synthesis, structure, and function of meningococcal LPS has recently been thoroughly reviewed [3,4,6].



Purified *Neisseria mucosa* LPS forms huge complexes in aqueous solution [6]. In cells expressing CD14 (monocytes, macrophages, neutrophils), better LPS aggregates come into view to have lesser cell-activating potency than lesser sized aggregates. The capsule polysaccharide does not affect the activation of these cells [4,5]. In endothelial cells, which are dependent on soluble CD14 for LPS-induced activation, the size of the aggregates does not influence cell activation, whereas capsular polysaccharide reduces the activation [2,4,8].

8. Conclusion

One can use and aim for the clinical research on *Neisseria mucosa* by using the present review article.

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