

Identification of common epitopes between different serotypes of *Streptococcus pneumoniae* group 19

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Vaccination with polysaccharide-based conjugate vaccines (PCVs) is recognized as one of the most successful strategy to prevent morbidity and mortality from pneumococcal diseases [1]. PCV vaccines contain capsular polysaccharide fragments from the *Streptococcus pneumoniae* (Sp) serotypes causing the majority of the diseases, covalently linked to a carrier protein. The most relevant limitation of PCV vaccines is caused by the large structural diversity of capsular polysaccharides: the protection offered by vaccination is serotype-specific and serotype prevalence is dynamic. Shifts in worldwide serotype distribution constitute a major challenge for eliminating pneumococcal infections, because commercial vaccines are unable to protect against serotypes not included in the vaccine [2]. This phenomenon is stimulating the search for a new generation of vaccines. Ideal candidates should be protective against a broader range of pneumococcal serotypes, with the possibility of the addition in the vaccine formulation of emerging new clinical isolates. In this framework, to simplify vaccine composition and to elicit a broader protection, we propose the identification of saccharide fragments containing chemical structures shared by different serotypes as cross-reactive and potentially cross-protective common antigens. In particular, we will present recent data on our ongoing work on the identification of common epitopes between different serotypes of Sp group 19 [3]. A small library of saccharides containing chemical structures shared by the 19F and 19A serotypes of *S. pneumoniae* has been synthesized and tested with a glycan array. The ability of the new compounds to be recognized by antibodies in reference group 19 antisera and factor reference antisera has been evaluated. Our study has shown that a phosphorylated simple disaccharide can be considered as a common carbohydrate epitope shared among different Sp 19 serotypes, setting the stage for exploring new common synthetic epitopes as potential candidates for a new generation of carbohydrate-based vaccines.

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Bibliographic references:

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