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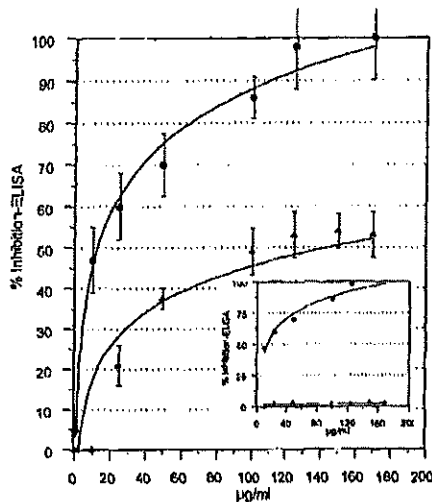
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(54) Title: BROAD-SPECTRUM LPS BASED VACCINES OF UNENCAPSULATED STRAINS OF HAEMOPHILUS INFLUENZAE AND OTHER PATHOGENIC SPECIES OF GRAM-NEGATIVE BACTERIA

Specificity of NTHi LPS-induced IgG PAb for LPS in supramolecular, micelle-like structure, as compared to the same LPS in non-micellar, monomeric structure



● Inhibition by NTHi LPS in supramolecular, micelle like, structure
▲ inhibition by NTHi LPS in monomeric structure

The window inside the graph represents the binding curve of the MAH1-3 monoclonal antibody for LPS in micellar and non-micellar structure (p<0.01).

(57) Abstract: Vaccines prepared using purified LPS of unencapsulated (non-typeable) Haemophilus influenzae detoxified by a strategy which leaves structurally intact the native Lipid A moiety, preserving broadly cross-reactive epitopes expressed by an antigenic supramolecular structure composed of at least 50 LPS monomers. The detoxification strategy preferentially involves the formation of the equimolar complex between by the Lipid A moiety of LPS and the cyclic synthetic peptide with the sequence NH₂-Lys-Thr-Lys-Cys-Lys-Phe-Ile-Ile-Ile-Ile-Cys-COOH, CysA-Cys10 disulfide. The same strategy of detoxification resulting in the preservation of the supramolecular, micelle-like, structure of native NTHi LPS is extended to LPS antigens purified from other pathogenic species of bacteria, in order to generate the homologous endotoxin vaccines.

WO 2004/052394 A1



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