

## Publication

A porin from *Klebsiella pneumoniae* : sequence homology, three-dimensional model, and complement binding**JournalArticle (Originalarbeit in einer wissenschaftlichen Zeitschrift)****ID** 153905**Author(s)** Albertí, S; Rodríguez-Quñones, F; Schirmer, T; Rummel, G; Tomás, J M; Rosenbusch, J P; Benedí, V J**Author(s) at UniBasel** [Schirmer, Tilman](#) ;**Year** 1995**Title** A porin from *Klebsiella pneumoniae* : sequence homology, three-dimensional model, and complement binding**Journal** Infection and immunity**Volume** 63**Number** 3**Pages / Article-Number** 903-10**Keywords** Amino Acid Sequence; \*Bacterial Proteins; Cloning; Molecular; Complement C1q/\*metabolism; Genes; Bacterial/\*genetics; *Klebsiella pneumoniae*/\*genetics; Models; Molecular Sequence Data; Porins/\*genetics/immunology; Restriction Mapping; Sequence Analysis; DNA; Sequence Homology; Amino Acid

A recombinant plasmid containing ompK36, the gene coding for the *Klebsiella pneumoniae* outer membrane protein OmpK36, was constructed by transposon mutagenesis and subcloning. Clones were identified in a cosmid library in *Escherichia coli* on the basis of their reaction with antiserum against the OmpK36 protein and by the presence in gel electrophoretic analysis of a band in *E. coli* outer membranes migrating with a mobility corresponding to 36 kDa. The ompK36-encoded protein exhibited characteristic properties of porins, such as heat modifiability and resistance to trypsin. The sequence of the gene revealed that OmpK36 is a close relative of the enterobacterial porin family, with a high degree of homology with *E. coli* OmpC, PhoE, and OmpF. On the basis of the structures of OmpF and PhoE porins, determined previously by X-ray analysis, it appears likely that the three-dimensional structure of OmpK36 also contains the motif of a 16-stranded beta-barrel, with long loops on one end and short turns on the other. Like the OmpC porin from *E. coli*, OmpK36 contains a long insertion in loop 4. The results of a binding study of complement component C1q to OmpK36 and the analysis of the OmpK36 model suggest that C1q binding sites are covered by the lipopolysaccharide core in the native porin.

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