Heterologous Expression of an Immunogenic Pneumococcal Type 3 Capsular Polysaccharide in Lactococcus lactis

CHRISTOPHE GILBERT,* KAREN ROBINSON, RICHARD W. F. LE PAGE, AND JEREMY M. WELLS
Cortecs Centre for Vaccine Discovery, Department of Pathology, University of Cambridge, Cambridge CB2 1QF, United Kingdom

Received 21 June 1999/Returned for modification 27 July 1999/Accepted 23 March 2000

In order to develop a new system for the analysis of capsular biosynthetic pathways we have explored the possibility of expressing type 3 capsular polysaccharide (CPS) from the pathogen Streptococcus pneumoniae in Lactococcus lactis, an unencapsulated lactic acid bacterium being developed as a vaccine delivery vehicle for mucosal immunization. Only three of the four type 3 CPS biosynthesis genes were found to be necessary for the abundant formation (120 mg liter

−1) of an extracellular type 3 CPS in L. lactis, implying a role for the type 3-specific synthase in the extracellular transport of the CPS or implying the existence of an alternative export system in L. lactis. The authenticity of the expressed heterologous polysaccharide was established by chemical and immunological analyses. Proton and carbon nuclear magnetic resonance spectroscopy of CPSs purified from L. lactis and S. pneumoniae showed that the two CPS structures were identical. When mice were immunized intraperitoneally with 3.5 × 10

6 CFU of live recombinant lactococci expressing a total of approximately 0.5 μg of type 3 CPS, the immune responses elicited appeared identical to those observed in mice inoculated with 0.5 μg of type 3 CPS purified from S. pneumoniae. These findings show that L. lactis is a useful host in which to study the role and function of genes involved in the production of bacterial capsules. Additionally, L. lactis shows potential as a host for the safe production of capsule antigens and as a vaccine delivery vehicle for polysaccharide antigens.

Highly invasive bacteria, regardless of whether they are gram positive or gram negative, commonly have an organized, adherent layer of polysaccharide associated with their outer surface. Such encapsulated bacteria are major causes of morbidity and mortality worldwide (7, 25, 29). The polysaccharide(s) comprising a capsule can be multichained or composed of linear homopolymers or heteropolymers. Capsules mediate important roles during certain stages of infection. Early in infection, during colonization, they facilitate bacteria-to-host adhesion (which may aid in quorum sensing); later, they enable bacteria to survive in the infected host despite attack by the complement- and phagocyte-mediated mechanisms associated with both innate and acquired immunity. Hence, virtually all bacteria capable of rapid proliferation in the bloodstream are encapsulated.

Because of this vital role in pathogenicity and because many capsular polysaccharides (CPSs) are immunogenic and used as vaccine antigens, it is important to improve our understanding of the biological properties of these bacterial polysaccharides and of the genes and enzymes involved in their biosynthesis. A number of recent studies (2, 13, 17, 18, 21, 24) have described the genetic organization of capsular biosynthetic operons. This paper describes a novel system of heterologous capsule expression which can be used to augment and refine these studies further. It also opens the way to certain unique possibilities, such as the production of CPSs in nonpathogenic bacteria and the analysis of the biological properties of such polysaccharides on bacterial surfaces in isolation from all other virulence determinants. Intriguingly, it also points to the possibility of recombining protective polysaccharide epitopes and generating chimeric vaccine antigens. Single vaccine antigens of this type could then be used to elicit protection against a range of bacterial serotypes.

To establish this system we used as an initial model the expression in Lactococcus lactis of the capsule biosynthesis genes of Streptococcus pneumoniae type 3 (strain WU2). The type 3 capsule biosynthetic genes of S. pneumoniae have a cassette-like organization, a feature common to the capsule operons of all serotypes analyzed to date. Genetic and biochemical analyses of the type 3 capsule locus have revealed four type-specific genes (Fig. 1). Phosphoglucomutase (Cps3M) and glucose-1-phosphate uridylyltransferase (Cps3U) are required for the synthesis of one of the saccharide precursors, UDP-glucose. UDP-glucose dehydrogenase (Cps3D) is the enzyme necessary for conversion of UDP-glucose to the second saccharide precursor, UDP-glucuronic acid, and the type 3 synthase (Cps3S) is required for polymerization of the precursors into a repeating unit (2, 9, 10). The region flanking the left end of the cassette is repeated in the pneumococcal chromosome and found upstream of other operons. Downstream of the last type 3 capsule-specific gene lies an internal portion of a gene (designated tnpA) that has homology with putative transposases of insertion sequences. Adjacent to tnpA is a sequence homologous to plpA, a gene encoding a permease-like protein involved in the transport of oligopeptides that has undergone a deletion at the 5’ terminus (9, 10). Both tnpA and plpA are physically linked and present in single copy in other capsule types.

**MATERIALS AND METHODS**

**Bacterial strains and growth conditions.** S. pneumoniae strain WU2 of serotype 3 (8) was grown at 37°C in Trypticase soy broth (Difco Laboratories, East Molesey, United Kingdom). The plasmid-cured L. lactis strain MG1363 (14) and recombinant L. lactis MG1363 were cultured at 30°C in M17 broth or on M17 agar plates (Difco) supplemented with 0.5% glucose (Sigma, Dorset, United Kingdom). For antibiotic selection erythromycin was used at a concentration of 5 μg ml

−1 (Sigma). To purify CPS from L. lactis UCP1619 the bacteria were
DNA techniques. Plasmid pTREP, a derivative of the pTREX vector, was purified from *L. lactis* using the QIAprep spin plasmid preparation kit (Qiagen Inc., Valencia, Calif.) according to the method suggested by the manufacture, except that the bacteria were first recovered by centrifugation and incubated in 100 μl of Tris-EDTA buffer (10 mM Tris-HCl [pH 8.0]; 1 mM EDTA) containing 5 mg of lysozyme (Sigma) per ml and 100 U of mutanolysin (Sigma) per ml for 10 min at 37°C to partially digest the cell wall. General molecular cloning was carried out by techniques described by Maniatis et al. (20). The method for the transformation of *L. lactis* strain MG1363 has been described previously (32).

Chromosomal DNA was isolated as described by Ausubel et al. (5). Purification of the CPS. The freeze-dried polysaccharide and the purified pneumococcal type 3 pneumococcal CPS (10).

Cells extracts and protein determinations. The optical density values of overnight cultures of *L. lactis* strains were recorded, and a 5-ml sample of each strain was centrifuged. The pellet was washed in NaH₂PO₄ buffer (50 mM, pH 7) and resuspended in 5 ml of the same buffer. Cells were broken with a French pressure cell and centrifuged to remove cells debris. The protein contents were determined on the supernatants by the method of Bradford (6) using bovine serum albumin (BSA) (Sigma) as the standard.

Mice. Specific pathogen-free female (BALB/c) mice (6 weeks old) were purchased from Harlan UK Ltd. (Bicester, Oxford, United Kingdom).

Immunizations. *L. lactis* strains were cultured at 30°C in the MOPS minimal medium supplemented with 0.5% glucose (Sigma) and 5 μg of erythromycin (Sigma) per ml. The cultures were diluted directly into phosphate-buffered saline (PBS) to achieve a concentration of 3.5 × 10⁸ cells in a 100-μl volume prior to immunization. Groups of 60 mice (180 mice in total) were immunized intraperitoneally with 3.5 × 10⁵ live lactococci harboring the control vector pTREP or with 3.5 × 10⁶ cells of *L. lactis* UCP1619 (expressing the equivalent of 0.5 μg of polysaccharide) or 0.5 μg of purified CPS from *S. pneumoniae* type 3 (Merck, Darmstadt, Germany). In each group, one subgroup of mice was bled daily from day 1 to 10 following the immunization to assay the immunoglobulin M (IgM) response. Subsequently, subgroups of 12 mice from each treatment group were bled weekly and tested for CPS-specific IgG responses. Moreover, a naive nonvaccinated control group was also included in the experiment.

**Antibody detection of CPS on live recombinant strains of *L. lactis*.** Overnight cultures were centrifuged, and the pellets were resuspended in a volume of medium to achieve a concentration of 10⁶ live recombinant *L. lactis* cells per ml (an optical density at 600 nm of 1 was considered to indicate a concentration of 5 × 10⁶ cells ml⁻¹). A 10-μl volume of the suspension (10⁷ live cells) was pipetted onto nitrocellulose, and the filter was blocked for 1 h at room temperature with 1% BSA in PBS (Sigma). The presence of CPS was detected by using a rabbit anti-CPS type 3 polyclonal serum (Statens Seruminstitut, Copenhagen, Denmark) as the primary antiserum, followed by anti-rabbit immunoglobulin horseradish peroxidase conjugate (Dako AAS, Glostrup, Denmark). The conjugate was detected by incubating the filter in a solution of 4-chloronaphthol (Sigma). Cell culture supernatants were applied to nitrocellulose in volumes of approximately 10-μl, with each application being allowed to dry thoroughly before subsequent applications were made.

**ELISA for the detection of antigen-specific serum antibody.** Antibodies (of IgM, IgG, and IgG subclasses) to pneumococcal type 3 CPS were measured by enzyme-linked immunosorbent assay (ELISA). ELISA plates (Nunc-Immuno plates with MaxiSorp surfaces; Gibco, Paisley, Scotland) were coated with 10 μg of type 3 CPS (Merck) per ml in PBS and incubated for 5 h at 37°C and then overnight at 4°C. Wells were blocked for 1 h at room temperature with 1% BSA (Sigma). Primary antisera were tested in duplicate wells using a twofold dilution series (in PBS containing 1% BSA and 0.05% Tween 20), including replicate wells of a diluted (1:50) preimmune serum on every plate. After 2 h, secondary anti-mouse immunoglobulin-alkaline phosphatase conjugates diluted in PBS.
containing 1% BSA and 0.05% Tween 20 (Southern Biotechnology Associates, Inc., Birmingham, Alabama) were applied for 90 min. Reactions were developed by using n-nitrophenyl phosphate (Sigma) as a substrate, and the optical density was read at 405 nm in an ELISA reader (Titertek Multiskan MCC340; ICN Biomedicals Ltd., Thame, Oxfordshire, United Kingdom). Dilution curves were drawn for each sample, and the end-point titers were calculated as the dilution of the sample producing the same optical density as the 1/50 dilution of a pooled preimmune serum. Statistical comparisons between results for different groups were made by the Mann-Whitney U test. A P value of >0.05 was considered insignificant.

RESULTS

Cloning and expression of genes for pneumococcal type 3 capsule biosynthesis in L. lactis. A schematic diagram of the pneumococcal type 3 locus and the genes required for capsule biosynthesis is shown in Fig. 2. Since UDP-glucose is necessary for many bacteria to produce essential cell constituents, including techoic acid and lipotechoic acid (4), we predicted that L. lactis would possess homologues of cps3U and cps3M. In order to investigate which of the pneumococcal type 3 enzymes would be required for capsule production in L. lactis, two different PCR fragments were generated by using oligonucleotide primers designed from the nucleic sequence of the genomic region involved in the capsule biosynthesis in S. pneumoniae type 3 (Fig. 2) (10). The PCR-amplified fragment CPS1 included the cps3D and cps3S genes and their cognate promoter but lacked the cps3U and cps3M genes. The PCR-amplified fragment CPS2 contained the cps3U gene in addition to cps3D and cps3S (shown in Fig. 2). DNA fragments CPS1 and CPS2 were ligated into the BamHI site of plasmid pTREP, a promoterless derivative of the vector pTREX (31) and then were used to transform electrocompetent L. lactis strain MG1363. Clones UCP1618 and UCP1619 harbored recombi-
nant plasmids pTREP-CPS1 and pTREP-CPS2 containing the fragments CPS1 and CPS2, respectively (Fig. 2).

Pneumococcal type 3 capsule production in *L. lactis*. A suspension of 10^8 cells of live lactococci of strains UCP1618 and UCP1619 and a control strain harboring the vector pTREP were spotted onto nitrocellulose membranes (corresponding to 12, 13.4, and 12 μg of protein content, respectively). The presence of type 3 CPS was detected by immunoblotting with rabbit type 3 specific antisera (Fig. 3). A strong signal was detected with *L. lactis* UCP1619(pTREP-CPS2), indicating extracellular production of the type 3 capsule in this strain. Compared to the control nonexpressor strain, a signal was also observed with *L. lactis* UCP1618(pTREP-CPS1) but this signal was substantially weaker than that obtained with strain UCP1619. The low level of CPS production in strain UCP1618 was confirmed by the analysis of nine other independently selected clones harboring the plasmid pTREP-CPS2. The quantity of CPS produced by each of these nine clones was identical. These findings indicated that the Cps3M function can be complemented in *L. lactis* but that expression of the pneumococcal glucose-1-phosphate uridylyltransferase (Cps3U) is required for efficient production of the saccharide precursor UDP-glucose in *L. lactis*.

In *L. lactis* the level (or activity) of cellular UDP-Glc-1-phosphate uridylyltransferase might be lower than the level (or activity) of pneumococcal cellular UDP-Glc-1-phosphate uridylyltransferase (GalU) (22). We expected the type 3 polysaccharide to be loosely associated with lactococcal cells since it is the only CPS which has not been shown to be covalently linked to the cell wall in pneumococcus (27). Interestingly, however, when *L. lactis* UCP1619 cells were mixed by vortexing and pelleted by centrifugation, the amount of CPS released in the supernatant of the culture was low compared to the amount of CPS that remained associated with cells (Fig. 3).

The CPS produced by *L. lactis* UCP1619 was purified from a minimal medium broth in order to avoid the presence of other polysaccharides in the culture (contaminant saccharides were precipitated from M17 rich medium). In minimal medium, an *L. lactis* culture harvested at stationary phase (at 30°C) yielded approximately 15 μg of polysaccharide per 10^8 CFU of lactococcal cells (corresponding to 120 mg liter^-1^). The calculated amount of polysaccharide produced by *L. lactis* may in fact be an underestimate due to inefficiencies in the extraction procedure and loss of material during the purification steps. Possible errors in our calculation of the amounts of polysaccharide present on *L. lactis* cells would not have influenced our immunogenicity studies (see Results), since similar immune responses were measured over a 10-fold range of *L. lactis* expressing type 3 CPS.

NMR spectroscopy. The proton NMR spectra (Fig. 4) and the carbon NMR spectra (Fig. 5) of CPSs purified from *S. pneumoniae* type 3 (Fig. 5I) and *L. lactis* strain UCP1619 (Fig. 5II) were very similar. On the proton spectra, the integration of the hydrogen peaks associated with the saccharides were comparable for the two polysaccharides. Moreover, two specific peaks corresponding to hydrogens involved in a β-link were observed in the proton NMR chromatograms for both *S. pneumoniae* and *L. lactis* (Fig. 5I and II) at 4.76 and 5.03 ppm (data not shown). All these results confirmed that *L. lactis* strain UCP 1619 was producing a surface CPS with a structure identical to that of the native CPS produced by *S. pneumoniae* type 3. No contaminants were evident from the proton and carbon spectra of the type 3 CPS purified from *L. lactis* UCP1619.

Immune responses to purified pneumococcal type 3 CPS and recombinant lactococci expressing type 3 CPS. Three
groups of 60 mice were inoculated with either live *L. lactis* UCP1619 (expressing in total 0.5 mg of type 3 polysaccharide), lactococci harboring the control vector pTREP, or 0.5 mg of purified pneumococcal type 3 CPS via the intraperitoneal route. Since Snippe et al. (26) had shown previously that IgM responses are short-lived and occur within the first 2 weeks of vaccination with CPS, 6 mice were selected at random on each of the first 10 days after inoculation from each of the three treatment groups, and blood samples were collected for IgM assay. In the subsequent period, 12 mice from each treatment group were bled weekly and tested for CPS-specific IgG responses.

On days 4 to 9 following the primary immunization a slightly elevated type 3 CPS-specific IgM response was detected in mice given lactococci expressing CPS or purified CPS, compared with that detected in mice given the control lactococcal strain (Fig. 6A). On day 6, the IgM response elicited by lactococci expressing CPS reached a peak (median titer, 102) and
this was significantly higher \( (P = 0.001) \) than the IgM responses of the control group whose median titer at this point was 45. The peak IgM response elicited by vaccination with pure CPS occurred a little earlier, on day 4, when the median titer reached 72. This was also significantly higher than the response of the control group \( (P = 0.032) \) at this time, with a median titer of 55. On days 4 to 8 the IgM responses of mice given pure CPS or lactococci expressing CPS were not significantly different \( (P > 0.05) \).

After 4 weeks the low-level IgG responses elicited by pure CPS or lactococci expressing CPS reached a peak (Fig. 6B). The serum IgG titers of the mice given pure CPS (median titer, 74) were significantly higher \( (P = 0.001) \) than those of the control group (median titer, 41). The responses of mice inoc-
ulated with lactococci expressing CPS (median titer, 77) were also significantly higher than those of the control group \( (P = 0.001) \). The IgG responses of mice inoculated with either CPS formulation were not significantly different from each other \( (P > 0.05) \). After the initial primary responses had declined, at 7 weeks, a booster inoculation was given to each group to determine whether any of the immunized groups of mice had acquired immunological memory. For T-cell-dependent antigens the induction of memory B- and T-cells ensures that serum antibody responses to a secondary antigenic challenge appear more quickly, attain a higher titer, and consist predominantly of IgG. In contrast, T-cell-independent antigens such as bacterial polysaccharides cannot recruit specific T-cell help and therefore a second exposure to these antigens elicits an antibody response of similar isotype, kinetics, and magnitude to the primary response.

As after the primary inoculations, mice were bled daily for 10 days postboost so that IgM responses could be detected; they were subsequently bled every week for a further 6 weeks. A rise in CPS-specific serum IgM titer was noted between days 3 and 10 postboost. The peak in the IgM responses of mice given either form of CPS occurred on day 9 (Fig. 7A), when the titers of the group given lactococci expressing CPS (median titer, 110) and those elicited by pure CPS (median titer, 155) were significantly higher than those of the control group \( (P = 0.015 \text{ and } P = 0.009 \text{ respectively}) \).

The peak in the postboost anti-CPS IgG responses occurred 2 weeks postboost (Fig. 7B), when the titers of the group given lactococci expressing CPS (median titer, 91) were significantly higher than those of the control group \( (P = 0.010) \) and remained at this level for the ensuing 3 weeks of the experiment.

The peak IgM and IgG responses elicited by either form of CPS were not significantly different from those obtained after primary immunization, indicating that the responses were characteristically T cell independent in nature. The IgG serum antibodies to CPS measured 2 weeks after the boost were primarily of the IgG1 and IgG3 subclasses. (From pTREP control, pTREP-CPS, and pure CPS groups the median IgG1 titers were 66, 122, and 80, respectively. Median IgG3 titers were 213, 360, and 480, respectively). The titer of IgG3 in sera from pTREP control mice increased to 213 compared with the preimmune level of 50. This might be due to the cross-reaction of antibodies elicited by \( L. \text{ lactis} \) with \( S. \text{ pneumoniae} \) polysaccharide. Moreover, it is generally known that anti-mouse whole-molecule IgG reagents do not recognize all IgG subclasses equally, and IgG3 may not have been detected with the same sensitivity as the other IgG subclasses.

**DISCUSSION**

We have shown that it is possible to express a heterologous type 3 CPS from \( S. \text{ pneumoniae} \) in \( L. \text{ lactis} \). This was achieved by cloning fragments of the pneumococcal type 3 CPS biosynthetic operon into the lactococcal cloning vector pTREP and transforming \( L. \text{ lactis} \). Reports of the instability of pneumococcal DNA in \( E. \text{ coli} \) are well documented (11, 19), but in \( L. \text{ lactis} \) the pneumococcal DNA could be stably manipulated and was efficiently expressed using the native pneumococcal promoters. The suitability of \( L. \text{ lactis} \) as a host for the cloning and expression of pneumococcal operons is presumed to reflect the fact that pneumococci and lactococci are more closely related phylogenetically to each other than they are to \( E. \text{ coli} \) and suggests that \( L. \text{ lactis} \) may be a useful alternative host to \( E. \text{ coli} \) for the manipulation and expression of streptococcal genes. Previous attempts to express type 3 CPS in \( E. \text{ coli} \) failed to yield an extracellular capsule; instead, the polysaccharide accumulated in the periplasm, leading to poor growth (3, 17). In contrast, the growth rates of the lactococcal CPS expressor strain were identical to those of the host strain and plasmid instability was not observed during repeated subcultures.

On the basis of antibody detection of CPS on the surface of intact recombinant lactococcal cells, it was apparent that type 3 CPS was produced in low amounts when only two (\( cps3D \) and \( cps3S \)) of the four pneumococcal biosynthetic genes were introduced into \( L. \text{ lactis} \). The two enzymes encoded by \( cps3M \) and \( cps3U \) genes are not required for the expression of type 3 CPS (9, 10), but their roles in the synthesis of the precursor requirements for CPS biosynthesis (glucose-1-phosphate, UDP-glucose) must be fulfilled by essential enzymes involved in the cell metabolism. In \( L. \text{ lactis} \), two distinct forms of phosphoglucomutase have been described and their activities may fulfill the role of \( Cps3M \) (23). Recently, a glucose-1-phosphate uridylyltransferase, \( GalU \), has been identified in \( L. \text{ lactis} \) (15), and it may fulfill the role of \( Cps3U \). However, the amount of CPS produced by \( L. \text{ lactis} \) was dramatically increased by incorporating the pneumococcal \( Cps3U \) gene into \( L. \text{ lactis} \) along with the genes encoding \( Cps3D \) and \( Cps3S \). This indicated that the normal levels of lactococcal \( GalU \) activity appear to be too low to match the activities of the \( Cps3D \) and \( Cps3S \) enzymes expressed in \( L. \text{ lactis} \) and become rate-limiting for the production of significant quantities of type 3 CPS. These results are in agreement with a recent publication reporting that the cloning and functional overexpression of the lactococcal \( galU \) gene resulted in increased levels of UDP-glucose and UDP-galactose in \( L. \text{ lactis} \) (16).

The amount of CPS produced by \( L. \text{ lactis} \) strain UCP1619 was estimated to be around 120 mg liter \(^{-1} \) in test-tube culture. It may well be possible to increase these yields still further by engineering strains that express functional phosphoglucomutase (Cps3M-like) in addition to the other biosynthetic enzymes. Proton and carbon NMR studies on purified type 3 CPS from \( S. \text{ pneumoniae} \) and recombinant \( L. \text{ lactis} \) strain UCP1619 showed that the composition and structure of the two polysaccharides were identical. By determining the minimum number of genes required for a good expression level of pneumococcal type 3 capsule in \( L. \text{ lactis} \), we have shown that specific pneumococcal genes providing transport functions are not required for type 3 polysaccharide synthesis, assembly, and transport in \( L. \text{ lactis} \). A common feature of pneumococcal capsule biosynthetic operons is the presence of conserved sequences at both ends of the operon that contain genes thought to be involved in the export and polymerization of CPS. It has been shown that a number of type 3 CPS operons contain deletions in these conserved regions, suggesting that type 3 capsule synthesis may not require the lipid intermediates and additional transport proteins involved in the export of other capsular serotypes across the cell membrane (9). Our results support the hypothesis that the type 3 synthase encoded by the gene \( cps3S \) carries out both the polymerization and export functions necessary for type 3 capsule production. However, it remains possible that an alternative transport mechanism exists in both \( L. \text{ lactis} \) and \( S. \text{ pneumoniae} \) that can export type 3 CPS. Previous work on the expression of a plasmid encoded exopolysaccharide in \( L. \text{ lactis} \) indicated that this organism may indeed possess a lipid carrier involved in the export of exopolysaccharide (28, 30).

Preliminary experiments in our laboratory showed that 5-\( \mu g \) doses of CPS in mice elicited antibody titers similar to those elicited by 0.5-\( \mu g \) doses (data not shown). Therefore, 0.5-\( \mu g \) doses of CPS (or equivalent \( L. \text{ lactis} \) cell doses) were used to study the immune responses in detail. After a single inocula-
tion, groups of mice immunized with *L. lactis* expressing in total 0.5 µg of type 3 polysaccharide or 0.5 µg of purified type 3 CPS from pneumococcus elicited similar titers of anti-CPS IgM antibody in the serum. The primary immune responses were monitored over time, and once antibody levels had declined to background levels, the groups of mice were boosted with a second dose of either CPS-expressing lactococci or of purified pneumococcal CPS. A T-cell-dependent antigen would be expected to induce memory B and T cells during the primary immunization, leading to more rapid secondary anti-

FIG. 7. Anti-type 3 CPS serum antibody responses following secondary intraperitoneal administration of recombinant *L. lactis* or purified pneumococcal CPS. Three groups of 60 mice were inoculated with either $3.5 \times 10^6$ CFU of *L. lactis* UCP1619 expressing 0.5 µg of type 3 polysaccharide (pTREP-CPS2), the same dose of lactococci harboring the control vector pTREP (pTREP control), or 0.5 µg of purified pneumococcal type 3 CPS (pure CPS) via the intraperitoneal route and boosted with a second identical injection after 7 weeks. (A) Anti-CPS IgM responses of 6 mice selected at random from each of the three treatment groups on day 9 postboost. (B) Anti-CPS IgG responses of 12 mice selected at random from each of the three treatment groups at 2 weeks postboost. The end-point titers were calculated as the dilution of serum producing the same optical density as a 1/50 dilution of a pooled preimmune serum taken at the start of the experiment. End-point titers of individual samples (open circles) and the group median values (bars) are shown.
body responses characterized by higher titers and consisting mainly of IgG. After boosting, the anti-CPS antibody responses in the groups of mice immunized with CPS-expressing lactococci or with purified CPS were not elevated above the levels observed for the initial primary antibody responses. These responses were therefore characteristic of those elicited by a classic T-cell-independent antigen such as an unconjugated bacterial polysaccharide, which cannot recruit antigen specific helper T cells. Immunization with \textit{L. lactis} UCP1619 therefore did not affect the T-cell-independent nature of the anti-CPS responses. Although IgG antibodies to CPS could be detected in the serum of mice immunized with purified CPS or lactococci expressing CPS they were either of the IgG1 or IgG3 subclass, as observed for other T-cell-independent antigens in mice (1).

The antibody titers elicited in our experiments were relatively low. Nevertheless, previous workers have shown that inoculation of BALB/c mice with the same amount (0.5 μg) of purified type 3 CPS as used here (without adjuvant) can protect against challenge with 25 U of a mean lethal dose of \textit{S. pneumoniae} type 3 (26). In our experiments, the rise in mean titer of anti-CPS antibody observed in the majority of mice was only twofold. However, following the boost with pure CPS, two of six mice showed a rise in IgM titer of four- to fivefold and a further two mice showed a three- to fourfold rise. Following boosting with strain UCP1619 only two of six mice displayed more than threefold rises in antibody titers. The increases observed here in IgM titer were not as great as those reported by Snippe et al. (26), who detected peak titers of 2-mercaptoethanol sensitive antibodies in vaccinated mice (5 μg of pure CPS) that were eightfold higher than those in the controls (26).

They used haemagglutination assays rather than ELISA to measure the antibody responses, and differences between the sensitivity of the assays may at least partly account for the lower titers recorded in our experiments. Snippe et al. showed that immunization with 0.5 μg of CPS resulted in complete protection against a lethal dose of \textit{S. pneumoniae} type 3. The work of Snippe et al. indicates that we are likely to obtain protection from challenge with virulent pneumococci in a mouse model following immunization with \textit{L. lactis} expressing between 0.5 to 5 μg of type 3 CPS. Future work is aimed at investigating the dose-dependent response to immunization with type 3 \textit{L. lactis} by using different routes of administration and testing the capacity of the immune responses elicited to protect against pneumococcal colonization and respiratory challenge.

Overall, these results clearly show the advantage of using \textit{L. lactis} rather than \textit{E. coli} as a heterologous host to clone, express, and study the genes involved in the biosynthesis of capsules in \textit{S. pneumoniae}. They also confirm that \textit{L. lactis} is a potential host for the safe production and delivery of capsule vaccine antigens. The \textit{L. lactis} model described here for pneumococcal capsule expression in \textit{L. lactis} would permit significant quantities of capsule antigen to be manufactured without the need to culture large quantities of pathogenic bacteria, thereby reducing the biological hazard involved with the growth of \textit{S. pneumoniae} strains in order to purify the polysaccharides.

The question of whether immune clearance of lactococci was inhibited by CPS expression was not addressed; however, vaccinated mice were monitored following the immunizations. No obvious adverse effects could be observed.

The encouraging results described in this paper indicate that further investigations into the delivery of recombinant and chimeric capsule antigens via systemic and mucosal routes of administration using recombinant lactococci are now warranted.

**ACKNOWLEDGMENTS**

We thank Peter Grice (University Chemical Laboratory, University of Cambridge) for performing the NMR analysis of the polysaccharides.

At the time these studies were conducted, J.M.W. held an Advanced Research Fellowship from the Biotechnology and Biological Sciences Research Council, United Kingdom, and C.G. held a Fellowship from Cortice plc, whose financial support is gratefully acknowledged.

**REFERENCES**


Editor: E. I. Tuomanen