

Induction of protective immunity to *Theileria annulata* using two major merozoite surface antigens presented by different delivery systems

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Allelic forms (Tams1-1 and Tams1-2) of the major merozoite surface antigen gene of Theileria annulata have recently been expressed in Escherichia coli and in Salmonella typhimurium aroA vaccine strain SL3261. To test the potential of subunit vaccines against T. annulata infection, we immunized four groups of three calves with either recombinant (re-) (Tams1-1 and Tams1-2) proteins or naked DNA encoding these antigens. Group I was immunized intramuscularly with both re-proteins incorporated into immunostimulating complexes (ISCOMs). Group II was inoculated intramuscularly with naked plasmid DNA encoding Tams1-1 and Tams1-2. Groups III and IV received S. typhimurium SL3261 [pSTams1-1][pIP5] and SL3261[pSTams1-2][pIP5] subcutaneously and orally, respectively. A final group of three animals (Group V) served as an unimmunized control group. Four weeks after the last immunization all calves were challenged with a T. annulata stabilate generated from blood of an infected animal with 30% piroplasm parasitaemia. All calves vaccinated with ISCOMs proved to be protected from T. annulata infection and had generated antibodies against both re-(Tams1-1 and Tams1-2) at the time of challenge. In two of these animals the antibody had a surface binding profile by IFAT. Two of three calves immunized with naked DNA also proved to be protected, but none of the animals had generated any detectable antibodies against the recombinants. Salmonella-based delivery of the recombinants did not induce any protection; two of six animals died of theileriosis and there was no difference between subcutaneous or oral administration. These preliminary results show that re-(Tams1-1 and/or Tams1-2) may elicit protective immune responses in cattle, depending on the antigen delivery system. © 1997 Elsevier Science Ltd.

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The protozoan parasite *Theileria annulata* infects cattle and domestic buffalo (*Bubalus bubalis*) across an area stretching from southern Europe and northern Africa, through the Middle East and southern Russia into India and southern China¹. *T. annulata* is transmitted by ticks of the genus *Hyalomma* and more than 250

million domestic cattle have been estimated to be at risk from tropical theileriosis². Infection in the bovine is initiated by inoculation of sporozoites from infected ticks during feeding. These invade mononuclear leucocytes (thought primarily to be monocytes and B-cells expressing MHC class II antigens^{3,4}), where they develop into multinucleated macroschizonts and induce host cells to divide in synchrony with the macroschizont. These parasites develop further from microschantos into free merozoites, which upon release from the leucocyte invade erythrocytes to develop into piroplasms.

Tropical theileriosis is currently controlled by chemotherapy, tick control using acaricides and vaccination using attenuated macroschizont-infected lymphoid cell lines. Although these live vaccines are being used with success in endemic areas^{5–10}, their

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application has a number of drawbacks. The vaccine lines need to be evaluated for their ability to protect against challenge, require prolonged *in vitro* passage to become attenuated and it is not clear whether attenuated cells can revert to virulence *in vivo*. Such prolonged passage can also lead to lines losing their ability to immunize recipient animals as a result of the inability of the macroschizonts to transfer into the cells of the vaccinated animal. Furthermore, although cattle are protected by the attenuated cell line vaccine, it does not prevent the development of the erythrocytic stages when animals are challenged¹¹. In addition, once thawed the vaccine has a short half life and there is the possibility of transferring other pathogens by the use of such live vaccines. Many of these limitations could be overcome by the production of immunogenic antigens by re-DNA techniques and development of a subunit vaccine against tropical theileriosis.

One antigen (SPAG-1) has been identified on the surface of the sporozoite by a mAb which blocks sporozoite penetration of bovine peripheral blood mononuclear cells *in vitro*¹². A C-terminal fragment (SR1) of SPAG-1 has been expressed as a fusion protein in the e1 loop of hepatitis B core antigen (HBcAg) and used to immunize calves in a small scale vaccination trial. High antibody titres were generated in all four animals immunized with HBcAg-SR1 and IFAT confirmed that there was recognition of the sporozoite surface. Clinical symptoms were delayed on sporozoite challenge; however, all the animals eventually developed disease¹³.

The merozoite stage is, like the sporozoite, invasive and also a potential target for a protective immune response. Recently, allelic forms of *Tams1* (*Tams1-1* and *Tams1-2*) which encode the 30 kDa and 32 kDa major merozoite surface antigens of *T. annulata*, respectively, have been expressed in a *Salmonella typhimurium aroA* vaccine strain and *Escherichia coli*¹⁴. Immunostimulating complexes (ISCOMs) were prepared from both recombinants purified from *E. coli*. As a third route to deliver re-(*Tams1-1* and *Tams1-2*) to the bovine host, both genes were subcloned into an eukaryotic expression vector and used as a DNA vaccine. In this paper we evaluate the potential of these three systems as a component of a multistage subunit vaccine against tropical theileriosis in cattle.

MATERIALS AND METHODS

Bacterial strains and culture

Escherichia coli M15[pREP4]¹⁵ and *Salmonella typhimurium* SL3261¹⁶ producing re-(*Tams1-1* or *Tams1-2*) were grown as described previously¹⁴. *S. typhimurium* SL3261 [pSTams1-1][pIP5] and SL3261[pSTams1-2][pIP5] used for animal experiments were washed in PBS and each was resuspended at 1×10^9 ml⁻¹ and 2×10^{10} ml⁻¹ for subcutaneous and oral administration, respectively. The bacteria were checked for the correct *aro*, *his*, Ap^R and Km^R phenotypes by selection on appropriate minimal plates¹⁷ with or without 10 µg ml⁻¹ para-aminobenzoic acid, 10 µg ml⁻¹ 2,3-dihydroxybenzoic acid, 40 µg ml⁻¹ phenylalanine, 40 µg ml⁻¹ tyrosine, 40 µg ml⁻¹ histidine, 40 µg ml⁻¹ tryptophan, and by culture on LB plates supplemented with 100 µg ml⁻¹ ampicillin (Ap) and 25 µg ml⁻¹ kanamycin

(Km). To distinguish *S. typhimurium* from *E. coli*, growth was tested on brilliant green agar (BGA) plates. Smooth LPS phenotype was determined by examining LPS profiles on silver-stained gels¹⁸. Briefly, whole cell lysates were boiled for 10 min, incubated with 125 µg proteinase K at 60°C for 1 h and subsequently examined by sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE) followed by silver staining. *S. typhimurium*-Smooth and -Rough (ATCC 14028), kindly provided by Dr J.G. Kusters, and *E. coli* were used as controls.

Preparation of ISCOMs containing re-(*Tams1-1* or *Tams1-2*)

Production and isolation from *E. coli* M15[pETams1-1+][pREP4] and M15[pETams1-2+][pREP4] of re-(*Tams1-1* and *Tams1-2*), respectively, were performed as described previously¹⁴. For the preparation of ISCOMs, recombinants were mixed separately with cholesterol, phosphatidylcholine (Sigma Chemical Co., St. Louis, MO, USA) and Quil A (Spikoside, Iscotec, Lulea, Sweden) at a ratio of 1:1:5 (w/w/w). After ultrasonic treatment for 10 min the mixture was incubated for 1 h at room temperature. Subsequently, the mixture was extensively dialysed against 10 mM Tris-HCl (pH 7.4), 0.3 M NaCl, layered over a linear (10–60%) sucrose gradient and centrifuged for 18 h at 25 K (Beckmann SW28 rotor). Fractions containing peak levels of the recombinants (coinciding with the presence of ISCOM-like structures, as judged by negative contrast electron microscopy) were pooled. For SDS-PAGE and subsequent Western blotting, ISCOMs were pelleted by ultracentrifugation for 4 h at 40 K and resuspended in PBS (1/10 of the initial volume).

Cloning of *Tams1-1* and *Tams1-2* in pSLRSV.Nul

Primers for PCR were chosen which were complementary to conserved sequences flanking *Tams1-1* and *Tams1-2*. Primer 1: 5'-GGAATTAGATCTATGTTGTCCAGGACCACC-3' and primer 2: 5'-AATTCACGATCTGGGTTTTAAAGGAAGTAAAGG-3'. *Bgl*II restriction sites (presented in bold) were incorporated to facilitate subsequent cloning. The positions of the start and stop codons are underlined in primer 1 and primer 2, respectively. PCR was performed in a final reaction volume of 100 µl containing 50 mM KCl, 10 mM Tris-HCl (pH 8.3), 0.1% Triton X-100, 1.5 mM MgCl₂, 200 µM (each) deoxynucleoside triphosphate, 80 pmol of each primer, 2.5 U *Taq* polymerase (Promega, Madison, WI, USA) and 10 ng of plasmid DNA containing *Tams1-1* or *Tams1-2*. Reactions were performed in an automatic DNA thermal cycler (Perkin-Elmer, Foster City, CA, USA) for 30 cycles comprising denaturation at 94°C for 1 min, annealing at 55°C for 1 min and extension at 72°C for 1 min. The resulting 851-bp fragments were separated by electrophoresis on a 1.2% low-melting point agarose gel and isolated using QiaexII (Qiagen, Chatsworth, CA, USA). These fragments containing the complete coding sequence of *Tams1-1* and *Tams1-2*, were ligated into expression vector pSLRSV.Nul, digested with *Bgl*II and treated with Calf-Intestinal-Phosphatase¹⁷. Plasmid pSLRSV.Nul/*Bgl*II-version was kindly provided by Prof. L. Babiuk and modified by Dr J.

Lewis (personal communication)¹⁹⁻²¹. The resulting plasmids pBgIII30 and pBgIII32 contained the 30 kDa and 32 kDa merozoite surface antigen coding sequences, respectively, both under control of the Rous Sarcoma Virus-Long Terminal Repeat promoter and the SV-40 polyadenylation signal. The plasmid backbone was derived from pSL301TM (Invitrogen, San Diego, CA, USA). Both plasmids were purified from *E. coli* DH5 α using Qiagen columns (Qiagen) and verified by nucleotide sequence analysis²².

Immunizations

Experimental calves (*Bos taurus*), 5–6 months of age and kept in experimental tick-proof pens were used in the experiments. Fifteen calves were allocated randomly into five groups of three animals each and immunized according to the schedule summarized in Table 1. Each immunization consisted of two preparations: one contained re-(Tams1-1), given on the right-hand side in the neck of the animal, and the other contained re-(Tams1-2), given on the left-hand side in the neck.

- (1) Group I: 1 ml ISCOM vaccine given intramuscularly, containing 20 μ g re-(Tams1-1 or Tams1-2);
- (2) Group II: 1 ml DNA vaccine given intramuscularly, containing 500 μ g of pBgIII30 or pBgIII32 dissolved in TE [10 mM Tris-HCl (pH 8), 1 mM EDTA];
- (3) Group III: 1 ml *Salmonella* vaccine given subcutaneously, containing 1×10^9 *S. typhimurium* SL3261[pSTams1-1][pIP5] or SL3261[pSTams1-2][pIP5];
- (4) Group IV: 5 ml *Salmonella* vaccine given orally, containing 1×10^{11} *S. typhimurium* SL3261-[pSTams1-1][pIP5] or SL3261[pSTams1-2][pIP5];
- (5) Group V: three unimmunized control calves were infected with *T. annulata* only.

Rectal temperatures were taken daily and blood was collected weekly for sera analysis. Additional blood samples were collected three times weekly in citrate buffer for the preparation of blood smears and determination of packed cell volume (PCV). The remainder of the blood sample was stored at -20°C for PCR analysis. *T. annulata* piroplasms were counted in Giemsa-stained blood smears. When very scanty, their presence was confirmed by PCR and subsequently Southern hybridization using *Tams1-1* cDNA²³ as a probe, as described previously²⁴. Blood samples for PCR were collected from Groups I and II prior to challenge and in weeks 2, 3, 4, 8 and 12 postchallenge. Saponin lysis mix (0.22% NaCl, 0.015% saponin, 1 mM EDTA) was added to 50 μ l of blood and the mixture

was centrifuged for 1 min. Pellets were washed three times with saponin lysis mix, resuspended in 100 μ l PCR buffer (50 mM KCl, 10 mM Tris-HCl (pH 8), 0.5% Tween 20, 100 μ g proteinase K ml⁻¹) and incubated at 56°C for 1 h. After 10 min at 95°C , 5 μ l were used for PCR, performed as described above.

Challenge

Calf 184 was experimentally infected with *T. annulata* (Ankara, Turkey)²⁵ administered as a ground-up tick supernate, kindly provided by Prof. C.G.D. Brown. When the parasitaemia (piroplasms) reached 30%, blood was frozen with 10% DMSO in 2 ml aliquots at -80°C , representing approximately 6×10^9 piroplasms per aliquot, plus an unknown number of macroschizont-infected leucocytes. Control and immunized animals were challenged with this stablate administered subcutaneously (1 ml) and intravenously (1 ml) in the neck of the animal (Table 1).

Western blotting

Column-purified re-(Tams1-1 and Tams1-2) and proteinase K-digested whole cell lysates of *S. typhimurium* SL3261 were separated by 0.1% SDS–12% PAGE¹⁷. Purified *S. typhimurium* LPS (Sigma) was used as a control. Subsequently, Western blot analysis was performed using preimmune, prechallenge and postchallenge sera from immunized calves at a 1:100 dilution and, as a control, rabbit anti-LPS serum, kindly provided by Edwin J. Tijhaar. Pelleted ISCOMs were separated and the Western blot analysed as described above, using immune serum from calf 184. Ab–Ag complexes were visualized using ECL immunodetection (Amersham, Buckinghamshire, UK).

Indirect immunofluorescent antibody assay

T. annulata (Ankara, Turkey)²⁵ piroplasm infected erythrocytes were prepared by centrifugation of blood at 800g for 10 min. Pelleted cells were washed twice in PBS, resuspended to a density of approximately 3×10^7 ml⁻¹ and fixed in acetone at -20°C for 20 min. The indirect antibody test was carried out essentially as described by Minami *et al.*²⁶. Briefly, sera isolated at the time of challenge from ISCOM immunized animals (Group I) and control calf 476 (Group V) and at weeks 2, 3 and 9 postchallenge were assayed by incubating 20 μ l of a 1:50 dilution (in PBS/20% horse serum) on each well at room temperature for 30 min, followed by washing three times with PBS. Fluorescein-conjugated rabbit anti-bovine IgG (Sigma) was added to each well and incubated for a further 30 min at room tempera-

Table 1 Vaccination schedule for four groups of three calves each immunized with re-(Tams1-1 and Tams1-2) either prepared as ISCOMs, encoded on plasmid pBgIII30 and pBgIII32, respectively, or produced in *S. typhimurium* vaccine strain SL3261

Animal group	Delivery system	Route	Week ^a					
			0	1	4	5	8	12
I	ISCOMs	Intramuscular	+	–	+	–	C	
II	pBgIII30 and -32	Intramuscular	+	–	+	–	+	C
III	<i>Salmonella</i> SL3261	Subcutaneous	+	+	–	C		
IV	<i>Salmonella</i> SL3261	Oral	+	+	–	C		
V	Unimmunized control	Intravenous and subcutaneous	–	–	–	–	C	

^aC, challenged with *T. annulata*

ture. After washing as above, slides were stained with 0.1% Evans blue, mounted with 50% glycerol in PBS and examined by microscopy under UV illumination at $\times 500$ magnification²⁷.

RESULTS

ISCOM immunized calves

The characteristic cage-like structure of the ISCOMs was confirmed by electron microscopy after negative staining (data not shown). After performing SDS-PAGE of the pelleted ISCOMs, Western blot analysis using *T. annulata* immune calf serum revealed bands at approximately 32 kDa and 33 kDa (Figure 1(A), lanes 1 and 2), confirming the presence of purified re-(Tams1-1) and re-(Tams1-2), respectively (Figure 1(A), lanes 3 and 4).

Group I calves were immunized twice with ISCOMs intramuscularly, with an interval of 4 weeks (Table 1) and proved to be protected against *T. annulata* challenge (Table 2). Western blot analysis revealed that all three animals had generated antibodies against

re-(Tams1-1 and Tams1-2) at the time of challenge [Figure 2(A), lanes 1, 2, 5, 6, 9 and 10]. Although there were equal amounts of recombinant proteins loaded onto the gel [Figure 1(A), lanes 3 and 4], the sera preferentially detected re-(Tams1-2) [Figure 2(A), lanes 2, 6 and 10], indicating that it was more immunogenic than the ISCOMs prepared from re-(Tams1-1) [Figure 2(A), lanes 1, 5 and 9]. Specific antibody levels were maintained in the sera of all three animals at least until week 23 postchallenge when the experiment was terminated [Figure 2(A), lanes 3, 4, 7, 8, 11 and 12], whereas the levels in control calf 476 (Group V) were almost non-detectable [Figure 2(A), lanes 15 and 16]. Furthermore, sera taken at the time of challenge from calves 473 and 474 reacted with piroplasm antigen in IFAT, whereas serum from calf 475 showed a weak recognition (Figure 3). Sera taken from the three immunized and one control animal at week 9 postchallenge recognized the piroplasms and preimmune sera did not react with either the recombinants or piroplasms (data not shown).

The protected animals (Nos. 473, 474 and 475) were confirmed to be *T. annulata* carriers by PCR (Figure 4 and Southern hybridization [Figure 4(B)]. The expected 778-bp fragment was amplified in all three animals at week 12 postchallenge [Figure 4(B), lanes 9 (after longer exposure), 15 and 22]. Calf 475, which developed a maximum piroplasm parasitaemia of 1% was already positive by PCR at week 3 postchallenge (Figure 4, lane 19) and showed a profile similar to control calf 476 (Figure 4, lanes 23–28). Calf 474 developed a 25% piroplasm parasitaemia in week 23 postchallenge (Figure 4, lane 16) and was treated with ButalexTM (Mallinckrodt Veterinary, Uxbridge, UK) at a dose rate of 2.5 mg kg⁻¹ body weight.

Naked DNA immunized calves

Group II was immunized three times intramuscularly at 4-week intervals using 500 µg pBgIII30 and 500 µg pBgIII32 in each immunization (Table 1). SDS-PAGE and Western blot analysis with sera collected just before every boost or challenge did not reveal any detectable antibodies against the recombinants. Sera used at a 1:10 dilution instead of 1:100 was also

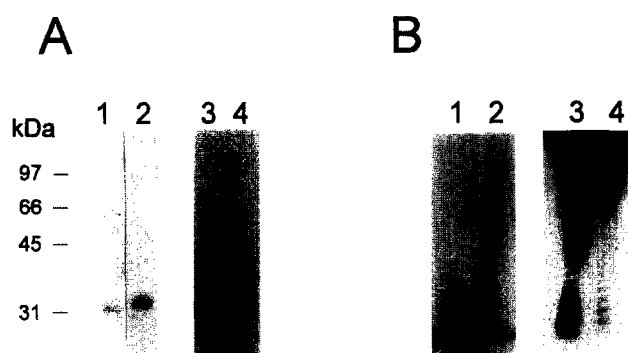


Figure 1 (A) Western blot analysis of re-(Tams1-1 and Tams1-2) ISCOMs (lanes 1 and 2, respectively) using *T. annulata* immune serum, and Coomassie brilliant blue-stained re-Tams1-1 and re-Tams1-2 proteins run in parallel (lanes 3 and 4, respectively). (B) SDS-PAGE of *S. typhimurium* LPS stained with silver (lanes 1 and 2) and Western blot analysed using rabbit anti-LPS serum (lanes 3 and 4). Lanes: 1, *S. typhimurium* SL3261; 3, purified *S. typhimurium* LPS (Sigma)

Table 2 Induction of protective immunity to *T. annulata* in different groups of calves immunized with re-(Tams1-1 and Tams1-2) either prepared as ISCOMs, encoded on plasmid pBgIII30 and pBgIII32, or produced in *S. typhimurium* vaccine strain SL3261

Group no.	Calf no.	Delivery system	Temperature (°C)	Maximum values		Final outcome
				Parasitaemia (%)	PCV reduction (%)	
I	473	ISCOMs	39.2	<0.1	14	Protected
	474		39.1	<0.1	18	Protected ^a
	475		39.2	1	19	Protected
	519		41.6	35	52	Theileriosis ^b
II	520	Plasmid DNA	40.8	1	18	Protected
	521		39.0	0.1	14	Protected
	3033		41.0	8	56	Theileriosis ^b
III	7643	<i>Salmonella</i> s.c.	41.5	8	49	Theileriosis ^b
	9263		41.9	50	78	Fatal theileriosis
	7775		41.2	3	44	Theileriosis ^b
IV	1045	<i>Salmonella</i> oral	41.9	50	68	Fatal theileriosis
	4063		41.9	3	58	Theileriosis ^b
	397		41.7	20	66	Theileriosis ^b
V	403	None	41.8	18	57	Theileriosis ^b
	476		41.2	10	39	Theileriosis ^b

^aAnimal developed tropical theileriosis at week 23 postchallenge, and was treated with ButalexTM, dose rate 2.5 mg ml body weight

^bAnimal recovered without treatment

negative (data not shown). However, calves 520 and 521 were protected, whereas calf 519 was not (Table 2). PCR analysis confirmed that all animals were carriers of *T. annulata* (data not shown).

Recombinant *Salmonella* immunized calves

SL3261[pSTam1-1 or pSTam1-2][pIP5] showed the characteristic pattern of smooth LPS after SDS-PAGE required by *Salmonella* for infecting host macrophages (data not shown). The *aroA* and *his* deletions of SL3261[pSTam1-1 or pSTam1-2][pIP5], required for the avirulent *Salmonella* to be useful as vaccine, were confirmed by growth limited to minimal plates supplemented with both the aromatic compounds and histidine. Growth of transformed SL3261 on LB containing both Ap and Km confirmed the presence of both pSTam1-1 or pSTam1-2 and pIP5. As an additional control, *S. typhimurium* and *E. coli* could be distinguished on BGA plates.

Groups III and IV were immunized twice subcutaneously and orally, respectively, with *Salmonella* producing re-(Tams1-1 and Tams1-2) (Table 1). All animals developed fever accompanied by piroplasm parasitaemia. Calves 9263 (Group III) and 1045 (Group IV) died of acute theileriosis. The other two animals from Group III developed a parasitaemia of 8%, whereas the two remaining animals from Group IV exhibited a maximum parasitaemia of 3%. Western blot analysis showed that all immunized animals had been infected with *Salmonella* at the time of challenge (Figures 2(B) and 2(C)). Sera from calves 9263 and 1045 (both died subsequently) contained fewer antibodies against *Salmonella* LPS [Figure 2(B), lanes 11 and 12, Figure 2(C), lanes 7 and 8, respectively] in comparison with sera from animals that survived challenge [Figure 2(B), lanes 3, 4, 7 and 8, Figure 2(C), lanes 3, 4, 11 and 12]. Furthermore, only calves 3033 and 9263, both immunized subcutaneously, had generated antibodies against re-Tams1-1 and to a lesser

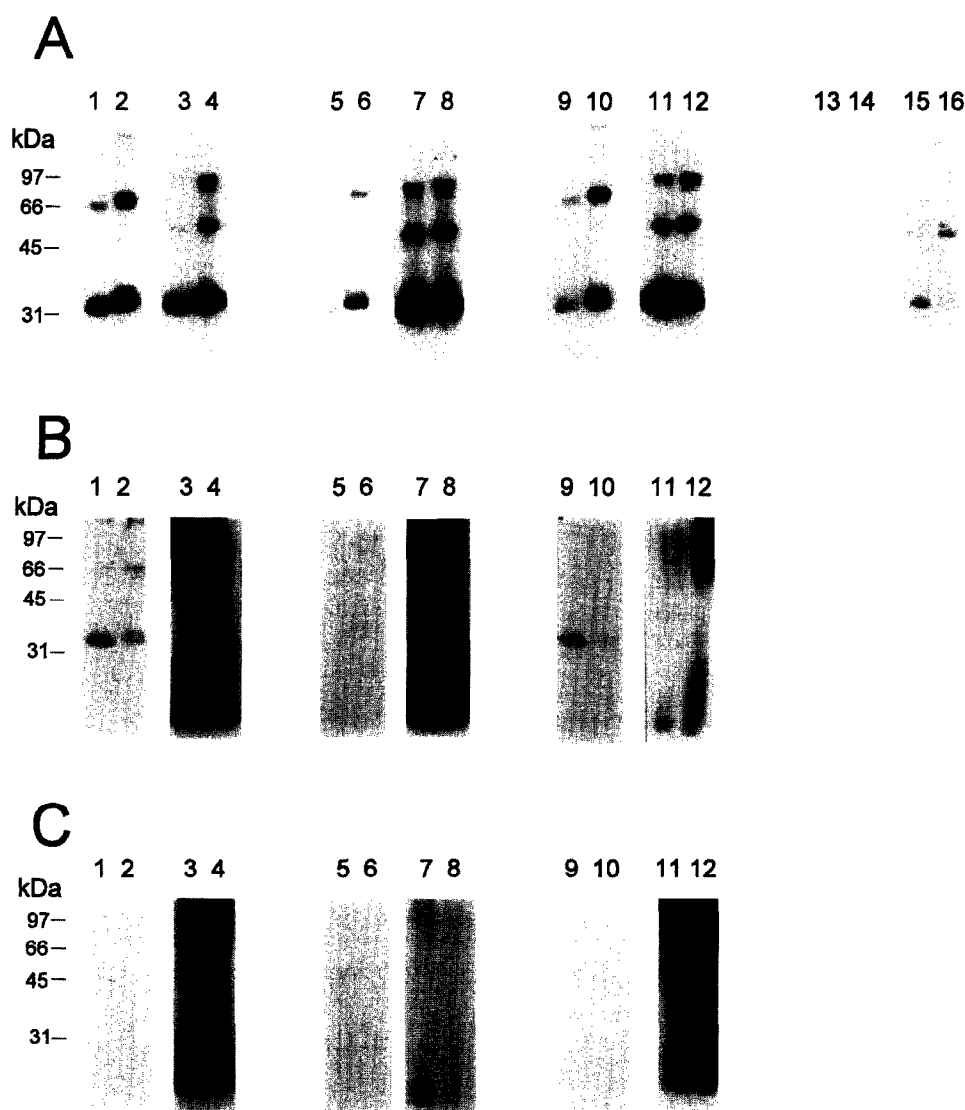


Figure 2 (A) Western blot analysis of re-Tams1-1 and re-Tams1-2, respectively, using sera from ISCOM immunized calves (Group I) and control animal 476 (Group V). Lanes 1 and 2, serum from calf 473 at the time of challenge (t.c); lanes 3 and 4, week 23 postchallenge (p.c); lanes 5 and 6, calf 474 t.c; lanes 7 and 8, week 23 p.c; lanes 9 and 10, calf 475 t.c; lanes 11 and 12, week 23 p.c; lanes 13 and 14, calf 476 preimmune serum; lanes 15 and 16, week 23 p.c. (B) Western blot analysis of re-(Tams1-1 and Tams1-2) (lanes 1, 2, 5, 6, 9 and 10) and *S. typhimurium* LPS (lanes 3, 4, 7, 8, 11 and 12) using sera from recombinant *Salmonella* immunized calves (Group III, immunized subcutaneously). Lanes 1–4, serum from calf 3033; lanes 5–8, calf 7643; lanes 9–12, calf 9263. (C) As (B) using sera from recombinant *Salmonella* immunized calves (Group IV, immunized orally). Lanes 1–4, calf 7775; lanes 5–8, calf 1045; lanes 9–12, calf 4063

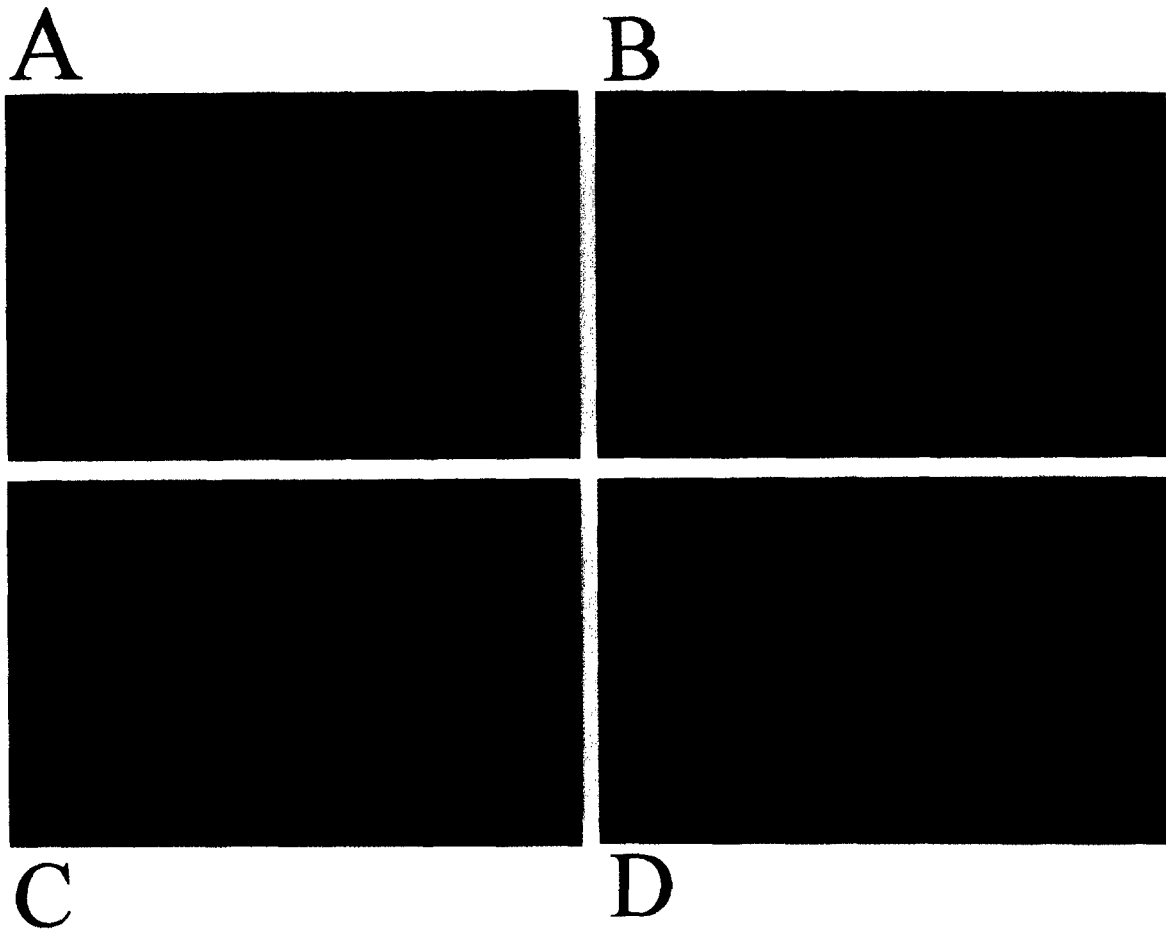


Figure 3 Immunofluorescence reactivity of antibodies generated at the time of challenge in ISCOM-immunized animals against acetone-fixed slide preparations of *T. annulata* (Ankara) piroplasms ($\times 500$). (A) Control mAb 5E1; (B) calf 473; (C) calf 474; (D) calf 475

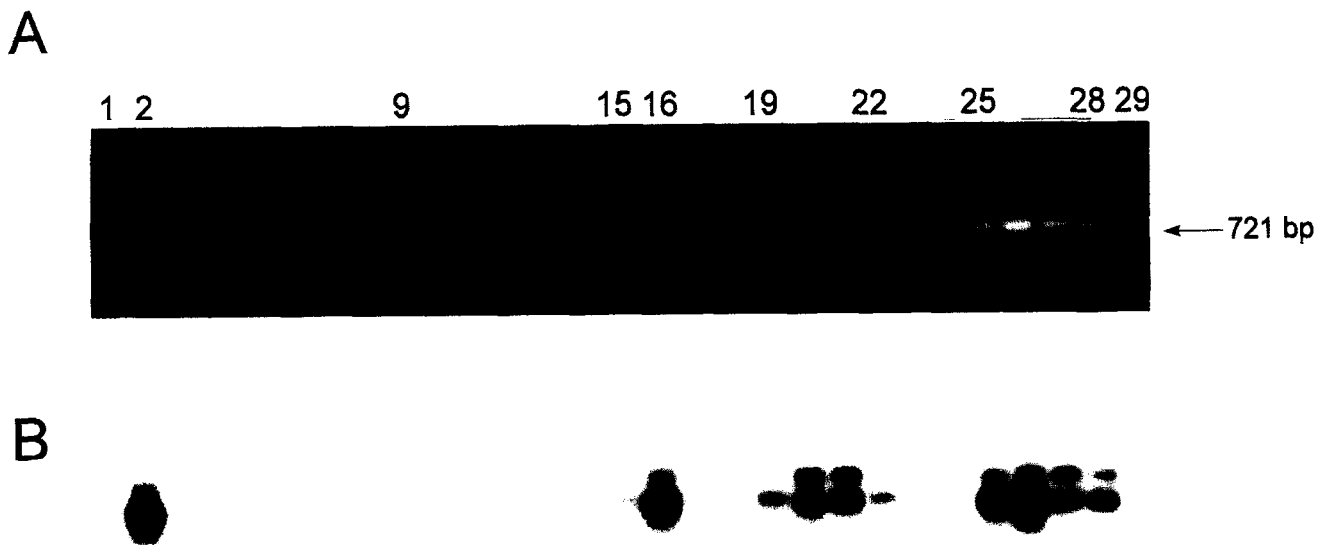


Figure 4 Agarose gel electrophoresis (A) and Southern blot hybridization using the c-DNA probe (B) of amplified DNA from calves immunized with iscoms (Group I) and control calf #476 (Group V) using primers 1 and 2. Lanes: 1, distilled water; 2, *T. annulata* positive control; 3 and 29, molecular size markers (pBluescript[Stratagene, La Jolla, CA, USA] digested with *Hinf*I); 4 to 9, calf #473: 4, prior to challenge; 5, week 2 post-challenge; 6, week 3 post-challenge; 7, week 4 post-challenge; 8, week 8 post-challenge; 9, week 12 post-challenge; 10 to 16, calf #474: same as above, except an additional lane 16, week 23 post-challenge; 17 to 22, calf #475: same as above; 23 to 28, calf #476: same as above.

extent against re-Tams1-2 at the time of challenge [Figure 2(B), lanes 1, 2, 9 and 10]. Preimmune sera taken from the same animals did not react with either recombinant antigen or LPS (data not shown). As a control, anti-*Salmonella* LPS serum raised in rabbit reacted with *Salmonella*-LPS [Figure 1(B), lanes 3 and 4].

DISCUSSION

In the present paper we have shown the ability of re-(Tams1-1 and/or Tams1-2) to induce protective immunity in cattle against tropical theileriosis, using ISCOMs and naked DNA to deliver these antigens, whereas all three unimmunized control animals developed disease. *Salmonella*-based delivery did not induce any protection; in contrast, two of six animals died of tropical theileriosis.

ISCOM immunized calves

At the time of challenge, all three ISCOM immunized calves had generated antibodies which recognized the piroplasm by IFAT. The pattern of staining was identical to that detected by monoclonal 5E1, which has been shown to bind to the surface of the merozoite²⁸, suggesting that antibody generated against the ISCOMs results against the merozoite surface. Therefore, protection may have been due to high amounts of circulating antibodies to re-(Tams1-1 and Tams1-2), which bind to the merozoite surface and prevent invasion of erythrocytes or opsonize the parasite for clearance in the spleen by macrophages. Serum from calf 475 showed a weak reaction in IFAT compared with the other two calves, which might be the reason why the animal developed a higher piroplasm parasitaemia (1%). To investigate the possibility of a Th-cell mediated immunity, a lymphocyte-stimulation assay will be used in further studies to characterize the phenotype of Th cells elicited by inoculation of re-(Tams1-1 and/or Tams1-2).

Since ISCOMs were described by Morein *et al.*²⁹, numerous studies have demonstrated their potential as a vaccine with built-in adjuvant properties. Bacterial, parasitic, viral and other proteins and peptides have been incorporated into ISCOMs to enhance the immune response in various ways, including activation of helper³⁰ and cytotoxic³¹ T cells and generation of potent long-lasting antibody responses³². Protection against lethal *Toxoplasma gondii*³³, *Trypanosoma cruzi*³⁴ and *Eimeria falciformis*³⁵ infections has been induced in mice by immunization with ISCOMs containing one or several isolated parasite antigens.

Naked DNA immunized calves

Two of three naked DNA immunized calves were protected, although one had fever for 1 day accompanied by a maximum piroplasm parasitaemia of 1%. There were no detectable antibodies generated against the recombinants, which suggests that protective immunity is dependent on T cells. Sedegah *et al.*³⁶ have shown 68% protection in mice against *Plasmodium yoelii* infection using plasmid DNA encoding the *P. yoelii* circumsporozoite protein, and that protection was dependent on CD8⁺ T cells.

Intradermal instead of intramuscular administration may improve the results³⁷. Skin-associated lymphoid tissues contain specialized cells that enhance immune responses, e.g. the dendritic cell and the macrophages of the dermis can also take up antigen and initiate immune responses. Therefore, the *in vivo* transfection of epidermal or dermal cells by DNA would be expected to provide a more efficient route for DNA immunization. A further improvement may be gene gun-based DNA immunization using a hand-held gene delivery instrument that requires >5000-fold less DNA to induce similar responses via intramuscular or intradermal inoculation³⁷.

Recombinant *Salmonella* immunized calves

None of the recombinant *Salmonella* immunized calves were protected, and two of the six animals died of tropical theileriosis. Only calves 3033 and 9263 had generated antibodies against re-(Tams1-1 or Tams1-2) at a relatively low level at the time of challenge and, in contrast to the ISCOM antisera, these were biased towards the re-Tams1-1 protein. Moreover, calf 9263 died of tropical theileriosis. This observation suggests that the recombinants were not produced at sufficient levels or in the appropriate form to induce adequate humoral and/or cell-mediated responses, possibly due to plasmid instability or toxicity of the recombinants *in vivo*. We recently reported that pSTams1-1 or pSTams1-2 and pIP5 can be maintained for at least 48 h in *S. typhimurium* SL3261 *in vitro*¹⁴. Stabilization of antigen production could be further attained by insertion of the gene into the bacterial chromosome³⁸. Also, Sadoff *et al.*³⁹ have shown that the gene encoding the circumsporozoite (CS) antigen of *P. berghei* when expressed in *S. typhimurium* can induce an antigen-specific cell-mediated immune response in mice and provided protection against sporozoite challenge in the absence of measurable antibody to CS protein.

Control calves

Group V animals received infected blood only and exhibited a very quick increase in piroplasm parasitaemia (from 5 to 20% in 24 h). Furthermore, on Giemsa-stained blood smears no Maltese crosses were detected which are typical for division of piroplasms within erythrocytes. Instead, multiple piroplasm infections were seen within individual erythrocytes, indicating the presence of macroschizont-infected leucocytes in the blood stabilate which differentiate to produce merozoites. This could be directly from infected cells within the stabilate or, more probably, following transfer of the schizont to the host cell of the immunized animal. Indeed, macroschizont-infected leucocytes could be detected in the spleen and liver of both animals that died. As it is known that *Tams1-1* is produced by macroschizonts²³ (at a low level) before differentiation to merozoites occurs, it is conceivable that peptides from these molecules are presented on the infected leucocyte together with MHC-I and recognized by cytotoxic T cells. Alternatively, the production of INF- γ by Th1 cells may activate macrophages and prevent the establishment of the macroschizont within leucocytes, as previously shown *in vitro* by Preston *et al.*⁴⁰ and Visser *et al.*⁴¹. As we did not obtain data on the T cell response in immunized animals, the

mechanism by which protection occurs is unclear at the present time. Further immunization trials are being conducted using ISCOMs with re-(Tams1-1 and Tams1-2) proteins combined with re-SPAG, and naked DNA encoding these proteins. Ultimately, it is likely that a subunit recombinant vaccine consisting of engineered antigens from sporozoites, schizonts as well as from merozoites/piroplasms will be necessary to induce protection against *T. annulata* infection, initiated through the bite of an infected tick.

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