Schistosome larvae stimulate macrophage cytokine production through TLR4-dependent and -independent pathways

Stephen John Jenkins¹,², James Philip Hewitson¹, Stephanie Ferret-Bernard¹ and Adrian Paul Mountford¹

¹Department of Biology, University of York, York, UK
²Institute of Immunology and Infection Research, 101 Ashworth Laboratories, King’s Buildings, University of Edinburgh, West Mains Road, Edinburgh EH9 3JT, UK

Keywords: IL-6, IL-10, IL-12p40, innate, MyD88

Abstract
Exposure of the mammalian host to infective larvae of Schistosoma mansoni causes an acute inflammatory response in the skin and the activation of several cell types of the innate immune response including macrophages. Using an in vitro model of macrophage activation, we show that schistosome larvae possess molecules that directly stimulate both thioglycollate-elicited macrophages (tMφ) and IFNγ-activated tMφ in vitro to produce several cytokines including IL-6, IL-12p40 and IL-10. The parasite-derived molecules are enriched within the material released by the parasite following transformation [0- to 3-h released larval preparation (0-3hRP)] but not within soluble preparations of whole larvae. Cytokine production was maintained in the presence of polymyxin B, confirming that contaminating endotoxin was not responsible. IL-12p40 and IL-10 production was much lower by cells from C3H/HeJ mice, which have defective Toll-like receptor 4 (TLR4), but IL-6 production was unaffected. Experiments using TLR4−/− mice confirmed that IL-12p40 production by tMφ in response to 0-3hRP was partly dependent upon functional TLR4, whereas IL-6 production was entirely independent. In contrast, tMφ from MyD88−/− mice failed to secrete either IL-12p40 or IL-6, underlining a pivotal role of TLR signalling in cytokine production by macrophages in response to stimulation with 0-3hRP. Finally, we show that glycan components of 0-3hRP are required for optimal cytokine production since protease treatment of 0-3hRP had no effect on IL-12p40 production and only a slight effect on IL-6, while sodium meta-periodate treatment almost completely abolished production of both cytokines.

Introduction
The parasitic helminth Schistosoma is the causative agent of the disease schistosomiasis suffered by ~200 million people worldwide (1). Infective cercariae gain entry into the mammalian host by active penetration of the skin, whereupon they transform into schistosomula that migrate through the dermis and, after several days, exit via the vasculature, or lymphatics (2, 3). Immune-associated events in the epidermis and dermis immediately following parasite exposure are characterized by oedema, the infiltration of mononuclear and polymorphonuclear cells (particularly, macrophages and neutrophils) and the local production of chemokines and cytokines, including IL-6, IL-10 and IL-12p40 (4–7). It is widely believed that this innate response is important in directing the subsequent generation of adaptive responses in the local draining lymph nodes (3, 7). However, the identity of the molecular interactions that occur between parasite larvae, or their released products, and the host’s innate accessory cells are at present unclear.

Immunocompetent accessory cells are some of the first to encounter invading pathogens, such as schistosomes, and recognize a restricted repertoire of microbial molecules, common to many pathogens, through cellular receptors termed pattern recognition receptors (PRRs). Ligation of these PRRs is the trigger for inflammatory cellular events that regulate innate and adaptive immunity, such as cytokine and chemokine production (8). There is also evidence that accessory cells are activated by host ‘danger molecules’ released upon tissue damage (9), which may also bind PRRs (10). Toll-like receptors (TLRs) are archetypal PRRs and play...
central roles in the induction of the signalling pathways that lead to pro-inflammatory cellular responses (11). These signalling events involve a variety of adapter proteins, with MyD88 being important for signal transduction from a variety of TLRs, including TLR2, TLR4, TLR9, and the IL-1R superfamily (11). Distinct cellular responses can be elicited by the engagement of different TLRs, or by interactions between multiple TLRs, allowing a certain degree of specificity against different pathogens (12). A number of different TLR ligands have been identified and are primarily of viral, bacterial, lower eukaryotic or host origin (11). However, recent observations have extended this identification to include components of higher eukaryotes, such as helminths (13–16).

In this paper, we wished to identify the stage of larval maturation that acts as a source of the molecules responsible for initiating the innate immune response. Consequently, we compared the capacity of live schistosome larvae of different maturation states, and soluble preparations derived from whole larvae, to stimulate cytokine production by macrophages obtained from wild-type (WT) mice and cohorts with genetic deficiencies in TLR2, TLR4, TLR9, and the IL-1R superfamily (11). We demonstrate that the material released by infective larvae upon transformation is rich in molecules that stimulate immune accessory cells via MyD88-dependent TLR4- and non-TLR4-mediated pathways.

Methods

Animals

Female C57Bl/6 strain mice (bred in-house) and C3H/HeN and C3H/HeJ mice (Harlan Seralab, Loughborough, UK) were housed at the University of York. For some experiments TLR4+/− (17) and MyD88+/− (18) mice, on a C57Bl/6 background, (kindly provided by Prof. Akira and Takeda, Osaka University, Japan), were bred and maintained under specific pathogen free (SPF) conditions in the animal unit at the University of Manchester. All animal work was carried out in accordance with the guidelines of the United Kingdom Animals (Scientific Procedures) Act 1986.

Parasites and parasite material

Cercariae of Schistosoma mansoni were shed from Biomphalaria glabrata snails harbouring patent infections, and following concentration by sedimentation on ice for 1 h were washed three times with sterile water. Cercariae were then centrifuged (1000 × g, 5 min) and the larval pellet was frozen at −20°C. Alternatively, cercariae were mechanically transformed using the method described by Ramalho-Pinto et al. (19) to generate larvae for subsequent in vitro culture. In brief, cercariae were mechanically transformed by vortexing for 90 s in RPMI-1640 containing 200 U ml−1 penicillin and 100 μg ml−1 streptomycin (Invitrogen, Paisley, UK) (RPMI-0) and cultured in vitro in RPMI-0 for 3 h at 37°C and 5% CO2 in a humidified incubator (20). The culture supernatant was then removed and the remaining larvae were washed to recover further released material. Pooled supernatants were concentrated 50-fold using centrifugal filter units (Ultrafree-MC with 5-kDa cut-off; Millipore, Watford, UK). As a control, an equivalent volume of RPMI-0 medium containing no parasite material was concentrated using the same method. The 3-h larval heads were isolated from their tails by centrifugation on a 40/70% discontinuous Percoll gradient (21), washed seven times in RPMI-0 and then frozen or cultured for a further 18 h in M169 media (20) supplemented with 200 U ml−1 penicillin, 100 μg ml−1 streptomycin and 5% heat-inactivated low-endotoxin FCS (Harlan Seralab). After the requisite culture period, the larvae were washed four times with RPMI-0 to remove all traces of serum proteins and then frozen at −20°C.

The thawed larvae of different maturation stages were sonicated (21 kHz at 6.5 μm amplitude) for 3 min and then centrifuged (100,000 × g) for 1 h to yield a soluble preparation from cercariae (soluble cercarial preparation (SCP)), and larvae at 3 h [3-h soluble schistosomula preparation (3hSSP)] and 18 h [18-h soluble schistosomulum preparation (18hSSP)]. The concentrated supernatant released by larvae between 0 and 3 h (0-3hRP) and the RPMI-0 control (RPMIc) were treated in a similar way. Protein and endotoxin concentrations of the preparations were determined using the Coomasie Plus-200 assay (Perbio Science UK Ltd, Tattenhall, UK) and Pyrogen Plus® limulus amoebocyte lysate test kit (BioWhittaker, Wokingham, UK), respectively.

Cell culture reagents

LPS (Escherichia coli strain 0111:B4), yeast Zymosan A (Saccharomyces cerevisiae) and polymyxin B (PMB) were obtained from Sigma-Aldrich (Poole, UK). Purified recombinant IFNγ from 211A CHO cell line supernatant was produced in-house.

Isolation of thioglycollate-elicited peritoneal macrophages

Adherent thioglycollate-elicited macrophages (tMφ) were used to directly test the stimulatory activities of different schistosome preparations in vitro since it was not possible to obtain an abundant supply of dermal macrophages ex vivo. Five days following intraperitoneal injection of 0.5 ml sterile 3% Brewer thioglycollate medium (Sigma-Aldrich), peritoneal exudate cells were extracted from sacrificed mice by peritoneal lavage with RPMI-1640 containing 200 U ml−1 penicillin, 100 μg ml−1 streptomycin, 2 mM L-glutamine (Invitrogen) and 10% FCS (Harlan Seralab) (RPMI-10). At this stage, cells from three to four mice were pooled to minimize individual variation. The cells were then washed, re-suspended in RPMI-10 and plated in 96-well tissue culture plates (1–2 × 105 cells per well; Nalge Nunc). After culture for 2 h, non- and semi-adherent cells were removed and discarded from adherent cells by gently washing the monolayer three times with pre-warmed RPMI-10. Resultant adherent tMφ were typically 80–90% F4/80+ as judged by flow cytometry. Adherent tMφ from the different strains of mice were phenotypically similar and recoverable in similar quantities.

Stimulation of tMφ with microbial products and live schistosome larvae

The tMφ monolayer was cultured for 24 h with soluble parasite preparations (SCP, 3hSSP, 18hSSP and 0-3hRP; 0.05–50 μg ml−1), LPS (0.0001–1 μg ml−1) or Zymosan A (0.05–50 μg ml−1) in the presence or absence of PMB (3–10 μg ml−1). Where indicated, tMφ were simultaneously activated with IFNγ (5 U ml−1) to increase sensitivity, particularly in the case of tMφ...
from C3H/HeN mice. The stimulatory properties of live larvae of different maturation stages were determined by culture of tMΦ in the presence of PMB (3 µg ml⁻¹) for 24 h with (i) whole cercariae, (ii) cercariae that had been mechanically transformed immediately prior to culture or (iii) mechanically transformed cercariae that had been cultured in vitro for 3 h followed by removal of the culture supernatant. This un-concentrated 0- to 3-h larval culture supernatant, termed 0-3 h supernatant, was also assayed for its stimulatory properties.

**Treatment regimes for 0-3hRP**

In experiments to test the contribution of different molecular structures on the stimulatory capacity of parasite material, 0-3hRP was heated treated, digested with proteases or subjected to sodium meta-periodate treatment as described subsequently. First, 0-3hRP was heated to temperatures of 37, 50, 60, 70, 80, 90 or 100°C for 45 min and the protein concentration re-checked. Second, in order to test for protease-sensitive moieties, 0-3hRP was dialysed for 4 h against PBS, pH 2.2 (Slide-A-Lyzer Mini Dialysis Unit with 3.5-kDa cut-off; Perbio Science), and then digested with pepsin (10 µg ml⁻¹) in PBS, pH 2.2; Sigma-Aldrich) overnight at 37°C. Control 0-3hRP was treated similarly but without the addition of proteases. Digested and control samples were then heated at 55°C for 30 min to inactivate the protease, and then dialysed against PBS, pH 7.2, for 4 h. Digestion of 0-3hRP was confirmed by one-dimensional SDS-PAGE of treated and control samples. Third, sodium meta-periodate treatment (22) was used to modify the carbohydrate present in 0-3hRP. Samples of 0-3hRP were dialysed against 50 mM sodium acetate buffer (pH 4.5) for 4 h and treated with sodium meta-periodate (10 mM; Sigma-Aldrich) for 45 min at 25°C in the dark, and the reaction was then stopped with sodium borohydride (50 mM; 30 min, 25°C). Treated 0-3hRP and mock-treated 0-3hRP (as above but without sodium meta-periodate) were then dialysed against several changes of PBS, pH 7.2, overnight. The protein concentrations were determined and adjusted as required. Treated and control 0-3hRP samples (50 µg ml⁻¹) from each of the above three treatment regimes were then used as before to stimulate tMΦ, or IFNγ-tMΦ where parasite material was limited (i.e. after periodate treatment).

**Cytokine detection by ELISA**

Supernatants from tMΦ cultures were stored at −20°C prior to detection by ELISA of IL-6, IL-10 and IL-12p40 as previously described (6, 7).

### Table 1. Summary of schistosome preparations analysed in this study

<table>
<thead>
<tr>
<th>Preparation acronym</th>
<th>Source material: larval developmental stage in vitro</th>
<th>Equivalent larval position in vivo</th>
</tr>
</thead>
<tbody>
<tr>
<td>SCP</td>
<td>Infective cercariae</td>
<td>Infective cercariae</td>
</tr>
<tr>
<td>3hSSP</td>
<td>3-h larvae</td>
<td>Epidermis</td>
</tr>
<tr>
<td>18hSSP</td>
<td>18-h larvae</td>
<td>Epidermal/dermal interface</td>
</tr>
<tr>
<td>0-3hRP</td>
<td>Molecules released during first 3 h of in vitro culture (×50 concentration)</td>
<td>Molecules released at the onset of infection</td>
</tr>
<tr>
<td>RPMic</td>
<td>Medium control (×50 concentration)</td>
<td>N/A</td>
</tr>
</tbody>
</table>

*aSource material in italics are schistosomula or their products following mechanical transformation and culture in vitro for the stated period of time. bApproximate location of larvae if they had matured in vivo. cEquivalent medium control to 0-3hRP. N/A, not applicable.*

**Results**

**Products released by schistosome larvae are potent inducers of cytokine production by tMΦ**

To establish whether skin-stage schistosomes possess innate stimulatory properties, soluble preparations of infective cercariae, in vitro-cultured larvae and the soluble products of mechanically transformed larvae (Table 1) were assayed for their ability to induce cytokine production by tMΦ from C3H/HeN mice. High levels of IL-10 and IL-6 were detected in the supernatants of tMΦ following 24 h culture with 0-3hRP, whereas little IL-12p40 was detected (Fig. 1A). In order to increase sensitivity, tMΦ were also activated with IFNγ (IFNγ-tMΦ) prior to co-culture with the different parasite products. Accordingly, IFNγ-tMΦ stimulated with 0-3hRP produced much greater levels of IL-12p40 and increased IL-6, but greatly reduced levels of IL-10 (Fig. 1B). In contrast to 0-3hRP, the soluble preparations of whole larvae (i.e. SCP, 3hSSP and 18hSSP) had very limited stimulatory potential, with SCP stimulating only an increase in IL-6 production by tMΦ (Fig. 1A), although low levels of both IL-6 and IL-12p40 were induced by SCP in cultures of IFNγ-tMΦ (Fig. 1B). Neither 3hSSP nor 18hSSP from skin-stage larvae induced production of IL-10, IL-6 and IL-12p40 by tMΦ, or by IFNγ-tMΦ (Fig. 1A and B). Similar results were obtained with cells from C57Bl/6 mice, except that greater levels of IL-12p40 were produced by both tMΦ and IFNγ-tMΦ but again only in response to 0-3hRP (data not shown). Differences in the stimulatory properties of the various parasite preparations were not due to contrasting kinetics of cytokine production, since the level of all cytokines detected from the supernatants of 0-3hRP- and SCP-stimulated cells had reached a peak by 24 h (data not shown).

**Live schistosome larvae have innate stimulatory activity**

Since soluble preparations of whole cercariae and in vitro-cultured larvae stimulated little, or no, cytokine production the interaction between tMΦ from C3H/HeN mice and live schistosome larvae was further explored. Cercariae stimulated a dose-dependent increase in IL-6 production by tMΦ with as few as 100 parasites per well causing a >3-fold increase in...
IL-6 (Table 2). Cercariae that had been mechanically transformed immediately prior to culture with tMφ stimulated a much greater fold increase in IL-6 compared with non-transformed cercariae when cultured at 50 or 100 larvae per well, although the difference was not apparent when parasites were cultured at a concentration of 200 per well (Table 2). Whereas mechanically transformed cercariae acquired characteristics of skin-stage larvae, such as the loss of tails, shedding of cercarial glycocalyx and release of head gland material, the non-transformed parasites remained intact throughout culture in vitro (data not shown).

The stimulatory properties of un-concentrated material released by the transformed larvae during the first 3 h of culture post-transformation (0-3 h supernatant) and the 3-h cultured live larvae devoid of this released material were compared. The 0-3 h supernatant stimulated increased IL-6 production over background levels when cultured at amounts equivalent to that released by 100 and 200 larvae per well (Table 2), and although the actual levels of IL-6 detected were much lower than those stimulated by the concentrated preparation 0-3hRP (cf. Fig. 1), and no IL-10 or IL-12p40 was detected (data not shown), these data demonstrate that the released material was stimulatory at the numbers of parasites conventionally used to infect experimental mice. It was also noted that live 3-h larvae stimulated increased IL-6 production over background levels when cultured at concentrations equivalent to 100 and 200 larvae per well (Table 2), which contrasts with the apparent non-stimulatory nature of 3hSSP (cf. Fig. 1).

### Table 2. Innate stimulatory properties of different developmental stages of live schistosome larvae and larval released products

<table>
<thead>
<tr>
<th>Larval stage/product</th>
<th>Fold increase* in IL-6 production over untreated control by tMφ cultured with larvae (number of parasites per well)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td>Cercariae**</td>
<td>1.48 ± 0.67</td>
</tr>
<tr>
<td>Transformed cercariae</td>
<td>3.71 ± 0.22</td>
</tr>
<tr>
<td>0-3 h supernatantc</td>
<td>0.65 ± 0.49</td>
</tr>
<tr>
<td>3-h schistosomulac</td>
<td>0.65 ± 0.49</td>
</tr>
</tbody>
</table>

*Values are means ± SD of fold increase in IL-6 production by tMφ (2 x 10^3 per well) cultured for 24 h compared with untreated cells. All cells were cultured in the presence of PMB (3 μg ml⁻¹). **Cercariae obtained from a single shed were left intact or mechanically transformed by vortexing immediately prior to addition to tMφ cultures. cCercariae were mechanically transformed and cultured in vitro for 3 h. The resulting transformed larval heads and tails were subsequently isolated from the culture supernatant and added to the tMφ cultures. The un-concentrated culture supernatant (0-3 h supernatant) was added to the tMφ cultures in quantities equivalent to that released by the different stated numbers of larvae.

Naturally occurring endotoxin of bacterial origin is not a major cause of cytokine production by larval released products

An important consideration of this study was the possible presence in the schistosome preparations of endotoxin (the major active component of which is LPS) that is likely to be a natural contaminant because infective cercariae emerge from the snail intermediate host in a non-sterile environment. Analysis of all parasite preparations using the limulus amoebocyte lysate assay demonstrated that the released larval preparation (0-3hRP) contained low levels of naturally occurring endotoxin contributing the equivalent of 0.0015 ± 0.0003 μg ml⁻¹ LPS to the tMφ culture, whereas only trace levels were detected in the other soluble larval preparations. Therefore, a 2-fold approach was used to determine the stimulatory properties of 0-3hRP in the absence of possible LPS contaminants. This involved, first, using the antibiotic PMB, which blocks stimulation by LPS (23), and second, obtaining macrophages from C3H/HeJ mice that have a natural mutation in the intracellular signalling Toll/IL-1 receptor (TIR) domain of TLR4 (24), through which the majority and most potent types of LPS signal (25).

The presence of PMB had little or no effect upon the levels of IL-10 and IL-6 produced by C3H/HeN tMφ when stimulated with 0-3hRP (Fig. 2A). These concentrations of PMB also had no effect on cell viability, or upon the stimulatory properties of other known microbial stimuli, such as Zymosan A (data not shown). In contrast, PMB dramatically inhibited stimulation by LPS, with a 90-fold decrease in IL-6 production and complete abrogation of IL-10 production (Fig. 2A). The serotype of LPS was selected for its highly stimulatory properties (26; data not shown) and was used at a concentration that had at least twice the endotoxin activity calculated to be contributed to the culture by 0-3hRP, to ensure that its potency was equal to, or
higher than the possible endotoxin contamination within this parasite preparation. In some experiments, tMφ were also stimulated with IFNγ to increase sensitivity and again PMB had no effect upon the levels of IL-12p40 and IL-6 produced by cells from C3H/HeN or C3H/HeJ mice stimulated with 0-3hRP but completely ablated the stimulatory properties of LPS (Fig. 2B). Similar data were obtained with both tMφ and IFNγ-tMφ from C57B/6 mice (data not shown). Combined, the data above provide strong evidence that the stimulatory properties of 0-3hRP are due to schistosome factors and not bacterial LPS contamination. However, as a precaution, all further assays of schistosome products were performed in the presence of PMB (3 μg ml−1), which was sufficient to inhibit up to 0.01 μg ml−1 LPS (data not shown).

As expected, tMφ from C3H/HeJ mice failed to produce detectable levels of IL-10 upon exposure to LPS, and produced 23-fold less IL-6 than their C3H/HeN counterparts (Fig. 2C). Furthermore, IFNγ-tMφ from C3H/HeJ mice produced 80-fold less IL-12p40 and 60-fold less IL-6 than their C3H/HeN counterparts upon exposure to LPS (Fig. 2D). In contrast, 0-3hRP stimulated IL-10 and IL-6 production by tMφ (Fig. 2C), and IL-12p40 and IL-6 production by IFNγ-tMφ (Fig. 2D) from C3H/HeJ mice, even in the presence of PMB. However, there was diversity in the requirement for functional TLR4 in the cytokine response to 0-3hRP. In this respect, C3H/HeJ tMφ produced 4-fold less IL-10 than C3H/HeN cells, but similar levels of IL-6 when stimulated with 0-3hRP in the presence of PMB (Fig. 2A and C). Furthermore, C3H/HeJ IFNγ-tMφ produced 2.7-fold less IL-12p40 than C3H/HeN cells after culture with 0-3hRP, but only 1.8-fold less IL-6 (Fig. 2B and D). That 0-3hRP stimulated less IL-10 and IL-12p40 but similar amounts of IL-6 by both tMφ and IFNγ-tMφ from C3H/HeJ compared with C3H/HeN counterparts suggests that 0-3hRP acts upon Mφ through both TLR4-dependent and -independent pathways, and that these pathways have a differential role in the resulting cytokine profile.

**0-3hRP stimulates a different cytokine profile to classic microbe-derived stimuli**

0-3hRP is unique among our soluble parasite preparations in its ability to stimulate Mφ cytokine production. To characterize the response to 0-3hRP and the pathways through which it may signal, its stimulatory properties were compared with LPS (E. coli) and Zymosan A (yeast) known to signal predominantly through TLR4 (24) and TLR2 (27), respectively. It is important to note that direct comparisons of potency of these stimuli are difficult due to the molecular heterogeneity of 0-3hRP and Zymosan, and our present lack of knowledge of the molar concentrations of their active components. Nevertheless, the overall profile of cytokine production by tMφ differed qualitatively in response to Zymosan A and 0-3hRP, such that the ratio of IL-12p40 to IL-6 was markedly different (Fig. 3). In this respect, titration of these microbial stimuli demonstrated that 0-3hRP was a weaker inducer of IL-12p40 production relative to Zymosan A, but was a more potent inducer of IL-6. Conversely, LPS stimulated high levels of both these cytokines but relatively little IL-10 (Fig. 3).

**0-3hRP contains ligands for multiple PRRs that are dependent upon signalling via MyD88**

Having shown that macrophages from C3H/HeJ mice produce less IL-12p40 and IL-10 than cells from C3H/HeN mice when cultured with 0-3hRP (Fig. 2), we used tMφ from TLR4−/− mice (17) to clarify the role of this receptor in the recognition of the larval released molecules. To control for differences in basal states of activation by tMφ from WT and TLR4−/− mice, cytokine production was expressed as a percentage of that stimulated by TLR4-independent Zymosan A (27). The 2-fold greater cytokine production by WT over TLR4−/− tMφ in response to Zymosan A was not due to endotoxin contamination because addition of PMB had no effect (data not shown), illustrating that tMφ from TLR4−/− mice most likely have a lower activation state. TLR4−/− tMφ produced little, or no, increase in IL-12p40 or IL-6 upon stimulation with LPS, whereas abundant cytokine production by WT tMφ was detected, confirming that TLR4 is essential for the recognition of LPS (Fig. 4A). In contrast, there was a dichotomy in the requirement for TLR4 in
the production of IL-6 and IL-12p40 in response to 0-3hRP. As such, TLR4/−/− tMΦ stimulated with 0-3hRP produced significantly (P < 0.001) less IL-12p40 than WT cells relative to Zymosan A; yet, there was no difference in the amounts of IL-6 (Fig. 2), reinforcing our observations made using C3H/HeN and C3H/HeJ cells (Fig. 2).

Since MyD88 is a key adapter molecule in the downstream signalling of many TLRs (28, 29) and is essential for the induction of most TLR4-dependent cellular responses, such as the production of IL-6 (18), we wanted to determine its role in the activation of tMΦ by 0-3hRP. LPS and Zymosan A are known to induce MyD88-dependent and -independent cellular responses (18, 30, 31), and in our assay, both stimulated significantly (P < 0.01 and P < 0.001, respectively) increased IL-12p40 production by tMΦ from MyD88+/− mice compared with cells cultured with media alone, although the levels were much lower than those of WT cohorts (P < 0.001) (Fig. 4A). In contrast, IL-12p40 production by tMΦ in response to 0-3hRP was totally dependent upon MyD88, with cells from MyD88−/− mice failing to produce any increase in IL-12p40 over cells cultured with media alone (Fig. 4B). IL-6 production in response to all three microbe-derived stimuli was entirely MyD88-dependent (Fig. 4B).

The cytokine-stimulating activity of 0-3hRP is heat labile but dependent upon the presence of glycans

To make an initial biochemical analysis of the stimulatory components within 0-3hRP, we first assessed their sensitivity to heat inactivation. The elements in 0-3hRP responsible for the production of IL-6 and IL-12p40 in response to 0-3hRP. As such, TLR4−/− tMΦ stimulated with 0-3hRP produced significantly (P < 0.001) less IL-12p40 than WT cells relative to Zymosan A; yet, there was no difference in the amounts of IL-6 (Fig. 2), reinforcing our observations made using C3H/HeN and C3H/HeJ cells (Fig. 2).

Since MyD88 is a key adapter molecule in the downstream signalling of many TLRs (28, 29) and is essential for the induction of most TLR4-dependent cellular responses, such as the production of IL-6 (18), we wanted to determine its role in the activation of tMΦ by 0-3hRP. LPS and Zymosan A are known to induce MyD88-dependent and -independent cellular responses (18, 30, 31), and in our assay, both stimulated significantly (P < 0.01 and P < 0.001, respectively) increased IL-12p40 production by tMΦ from MyD88+/− mice compared with cells cultured with media alone, although the levels were much lower than those of WT cohorts (P < 0.001) (Fig. 4A). In contrast, IL-12p40 production by tMΦ in response to 0-3hRP was totally dependent upon MyD88, with cells from MyD88−/− mice failing to produce any increase in IL-12p40 over cells cultured with media alone (Fig. 4B). IL-6 production in response to all three microbe-derived stimuli was entirely MyD88-dependent (Fig. 4B).

The cytokine-stimulating activity of 0-3hRP is heat labile but dependent upon the presence of glycans

To make an initial biochemical analysis of the stimulatory components within 0-3hRP, we first assessed their sensitivity to heat inactivation. The elements in 0-3hRP responsible for
stimulating tMφ production of IL-12p40 and IL-6 were heat stable at temperatures up to 50–60°C (Fig. 5A). However, exposure of 0-3hRP to higher temperatures dramatically decreased its ability to stimulate production of these cytokines, with >90% reduction following heat inactivation at 100°C (Fig. 5A). In separate experiments, treatment of 0-3hRP with either pepsin (Fig. 5B) or trypsin (data not shown) had no significant effect on the ability of stimulated tMφ to produce IL-12p40, although the production of IL-6 was reduced by up to 40% (P < 0.01). Finally, treatment of 0-3hRP with sodium metaperiodate significantly reduced by >90% the secretion of IL-12p40 and IL-6 (both P < 0.001) by IFNγ-tMφ (Fig. 5C). Although ‘mock’ periodate treatment of 0-3hRP also slightly reduced production of both IL-6 and IL-12p40, sodium metaperiodate treatment further reduced the production of IL-6 (P < 0.001) and IL-12p40, by almost 80 and 70% (P < 0.01), respectively, compared with this control (Fig. 5C).

**Discussion**

The acute cutaneous inflammatory response induced by penetrating larvae of *S. mansoni* is well documented (e.g. 6, 7), although the parasites have also developed several strategies for regulating the extent of inflammation (reviewed in 3, 32). Nevertheless, the molecular interactions between parasite and host that initiate the inflammatory response remain poorly characterized.

In this study, we show that molecules released from schistosome larvae are recognized directly by the innate immune system resulting in both pro- and anti-inflammatory cytokine production by macrophages. The stimulatory molecules are most abundant within the material released by larvae upon transformation. It is likely that the highly stimulatory properties of 0-3hRP and the apparent lack of equivalent components in the soluble preparations of whole larvae (i.e. 3hSSP and 18hSSP) reflect the physiological context in which the innate immune system has evolved to recognize molecules released by invading schistosomes. In this respect, molecules released by the parasite as it penetrates the skin are readily visible to innate accessory cells (5, 7), and contrast with the somatic antigens that form the dominant constituents of the soluble schistosomula preparations (33). The molecular composition of 0-3hRP is currently being defined using a proteomic approach (R. Curwen and R. A. Wilson, in preparation) but it comprises a heterogeneous mixture of protein constituents, probably originating from the post- and pre-acetabular glands (34), rich in proteases that aid parasite penetration of the skin (35, 36). It is unlikely to contain more than just a small number of membrane molecules (released via membrane turnover during *in vitro* culture, and subsequently solubilized by sonication) but may contain a quantity of glycans originating from the protective glycocalyx that is shed by the transforming larvae (37). Although 0-3hRP is a subset of the molecules contained in SCP (soluble cercariae), it is diluted by abundant somatic components that are unlikely to be good PRR ligands, thus accounting for the limited stimulatory nature of SCP. The accessibility of molecules expressed upon the surface of skin-stage larvae (i.e. transmembrane molecules) may provide another way for parasites to interact with innate accessory cells and could explain the stimulatory capacity of live 3-h larvae compared with the equivalent soluble preparation (3hSSP), in which membrane-bound material would likely be diluted by somatic constituents as previously noted.

One problem experienced by studies attempting to characterize pathogen-associated molecules with innate stimulatory properties is the potential interference by low levels of naturally occurring endotoxin (38). However, we provide definitive evidence that the stimulatory properties of 0-3hRP are due to molecules of schistosome origin and not LPS-like contaminants. For example, the greatly increased stimulatory capacity of transformed compared with non-transformed cercariae demonstrates that the material released by transforming larvae directly stimulates macrophages and is not of microbial origin since both groups of parasites were derived from the same pool and would contain equal quantities of any possible microbial contaminants. 0-3hRP also retained the majority of its stimulatory properties in the presence of endotoxin-neutralizing PMB, stimulated cytokine production by tMφ from C3H/HeJ mice that lack the functional receptor for LPS endotoxin and was heat labile. Further indirect evidence of the existence of schistosome PRR ligands is provided by our recent observation that 0-3hRP induces a novel maturation...
phenotype of dendritic cells compared with classic microbial products such as endotoxin and yeast glycans (39).

The initial trigger for cytokine production by accessory cells is thought to be the direct or indirect ligation of specific TLRs or other PRRs by pathogen products, resulting in translocation of transcription factors such as nuclear factor-κB to the nucleus. Comparing responses of tMφ from C3H/HeJ mice with those of C3H/HeN mice in the presence of PMB, we show that parasite-derived factors released by schistosomae larvae act partly through a TLR4-dependent pathway to induce production of a limited repertoire of cytokines. The mutation in TLR4 of C3H/HeJ mice occurs in the intracellular signalling TIR domain, allowing the receptor to potentially retain a function through its extracellular domain (24). We verified this observation using tMφ from mice with a targeted disruption of the TLR4 gene (17). The similarity of the response in C3H/HeJ and TLR4−/−tMφ versus their TLR4+/+ controls suggests that the function of this receptor in the recognition of 0-3hRP is solely dependent upon events that follow signalling through the intracellular TIR domain, unlike its role in the recognition of the helminth molecule ES-62 (16). Our observations affirm the promiscuity of TLR4, adding to the growing list of ligands identified for this receptor. Indeed, a schistosome egg glycan containing the Lewis X moiety was recently shown to be a TLR4 ligand with potent immunoregulatory function (14, 40). Since glycans containing the Lewis moiety are found within material released by cercariae (41–43), they represent likely candidates for the object of intense study, and are believed to be mediated by the different TLRs (11, 46) and the adapter proteins they recruit (12). While our observations of LPS and Zymosan A support the view that TLR4 agonists preferentially induce production of IL-6 compared with TLR2 ligands (47), 0-3hRP also preferentially stimulates IL-6 production compared with the TLR2 ligand Zymosan A (Fig. 3), but in a TLR4-independent manner (Figs 2 and 4A), demonstrating that signalling via PRRs other than TLR4 can also preferentially induce IL-6 production. Furthermore, we show that diverse signalling pathways can be activated downstream of an individual TLR depending upon the ligand, since the schistosome ligands of TLR4 contributed little to IL-6 production and more to the production of IL-12p40 and IL-10, whereas high IL-6, IL-12p40 and IL-10 production resulting from stimulation with LPS was completely dependent upon TLR4. One explanation for the dichotomy in TLR4 function in our study is that the signalling pathway controlling preferential production of IL-6 is shared by multiple receptors involved in the recognition of 0-3hRP, and represents a semi-redundant system, whereas control of IL-12p40 and IL-10 production by TLR4 is non-redundant. However, we favour the hypothesis that diverse signalling pathways may result from an individual TLR through its association with different accessory PRRs and the signalling adapter molecules they recruit (12, 14). One candidate accessory PRR that may be involved is the macrophage mannose receptor, since we have preliminary data showing ligands of this receptor within 0-3hRP (S. J. Jenkins and A. P. Mountford unpublished results). Differential expression of PRRs by accessory cells also presents an attractive explanation for the contrasting response of tMφ to 0-3hRP compared with the limited response of DC (39).

The data presented in this study contribute greatly to our understanding of the host–parasite interactions that follow exposure to infective schistosome larvae. We clearly demonstrate the existence of schistosome-derived factors that stimulate innate accessory cells through MyD88-dependent receptors, including TLR4, to produce a repertoire of pro- and anti-inflammatory cytokines similar to that stimulated by classical microbial products. As an innate organ, the skin is the major site for immune recognition of the invading parasite, and our findings warrant further investigation into the potentially parasite-beneficial or host-protective effects exerted by the recognition of these parasite factors.

Acknowledgements

We thank the staff of the University of York Animal Unit, Ann Bamford (University of York) for maintenance of the parasite life cycle and Prof. R. Grencis and Dr A. Bancroft for allowing the use of mice and facilities at the University of Manchester, UK. We also thank Prof. S. Akira...
Macrophage activation by schistosome larvae

1417


34. Stirewalt, M. A. 1974. Schistosoma mansoni: cercaria to schisto-


37. Samuelson, J. C. and Caulfield, J. P. 1982. Loss of covalently labeled glycoproteins and glycolipids from the surface of newly...


