A human homologue of the Drosophila Toll protein signals activation of adaptive immunity

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Induction of the adaptive immune response depends on the expression of co-stimulatory molecules and cytokines by antigen-presenting cells. The mechanisms that control the initial induction of these signals upon infection are poorly understood. It has been proposed that their expression is controlled by the non-clonal, or innate, component of immunity that preceded evolution of the development of an adaptive immune system in vertebrates. We report here the cloning and characterization of a human homologue of the Drosophila toll protein (Toll) which has been shown to induce the innate immune response in adult Drosophila-like Drosophila Toll, human Toll is a type I transmembrane protein with an extracellular domain consisting of a leucine-rich repeat (LRR) domain, and a cytoplasmic domain homologous to the cytoplasmic domain of the human interleukin (IL)-1 receptor. Both Drosophila Toll and the IL-1 receptor are known to signal through the NF-kB pathway. We show that a constitutively active mutant of human Toll transfected into human cell lines can induce the activation of NF-kB and the expression of NF-kB-controlled genes for the inflammatory cytokines IL-1, IL-6 and IL-8, as well as the expression of the co-stimulatory molecule B7.1, which is required for the activation of naive T cells.

The Toll protein controls dorsal–ventral patterning in Drosophila embryos and activates the transcription factor Dorsal upon binding to its ligand Spatzle. In adult Drosophila, the Toll/Dorsal signalling pathway participates in an anti-fungal immune response. Signal-ling through Toll parallels the signalling pathway induced by the IL-1 receptor (IL-1R) in mammalian cells: IL-1R signals through the NF-kB pathway, and Dorsal and its inhibitor Cactus are homologous to NF-kB and I-kB proteins, respectively. Moreover, the cytoplasmic domain of Drosophila Toll is homologous to the cytoplasmic domain of IL-1R (ref. 9). Remarkably, the tobacco virus-resistance gene that encodes N-protein is also similar to Toll in that it contains both a Toll signalling domain and an LRR domain. It thus appears that the immune-response system mediated by Toll represents an ancient host defence mechanism (Fig. 1). To investigate the possibility that this pathway has been retained in the immune system of vertebrates, we used sequence and pattern searches of the expressed-sequence tag (EST) database at the National Center for Biotechnology Information (NCBI). A search with a sequence profile of the Toll/IL-1R signalling domain identified a matching sequence in the EST database derived from human fetal liver/spleen library (Genebank accession number H48602, corresponding to clone 202057 from the IMAGE consortium). Sequencing of this clone showed that it corresponds to the 3′ untranslated region (UTR) and the C terminus of the coding region of the messenger RNA. A fragment of the clone amplified by using the polymerase chain reaction (PCR) was used to screen a human spleen complementary DNA library by hybridization, and the 5′ end of the cDNA was cloned using the 5′-RACE technique as described. The full-length 4,874-base-pair (bp) cDNA clone contained an open reading frame of 2,523 bp which encoded an 841-amino-acid protein chain (Fig. 2a), as well as a LINE-1 reverse transcriptase pseudogene in the 3′-UTR. Alignment of the sequences of the human and Drosophila Toll proteins shows that there is homology over the entire length of the protein chains (Fig. 2b). Notably, the similarity between the cytoplasmic domains is higher than between the human proteins Toll and IL-1R (not shown). The extracellular domain of human Toll (hToll) contains 21 tandemly repeated leucine-rich motifs separated by a non-LRR region, similar to Drosophila Toll (dToll). At the N-terminal end of the LRR domain there is a 31-amino-acid long N-flanking region that is also present in several other LRR-containing proteins, for example RP105, Decorin and Biglycan. The C-terminal end of the LRR domain is flanked by a cysteine-rich domain which is also present in dToll and some other transmembrane proteins.

To examine the expression pattern of hToll, a 720-bp cDNA fragment was used to probe northern blots containing poly(A)+ RNA from several organs. Most organs expressed two mRNA species: one of ~5 kilobases (kb) was predominant in most tissues except peripheral blood leukocytes (PBL), and corresponded to the length of the cDNA that we cloned. The lower band was ~4 kb long and this band was predominant in the PBL. The 4-kb band was not detectable in kidney, and liver did not contain any mRNA at all (Fig. 3). We also tested different mouse and human cell lines for expression of hToll mRNA by using PCR with reverse transcription (RT-PCR). We found mRNA for hToll in monocytes, macrophages, dendritic cells, γ/δ T cells, Th1 and Th2 α/β T cells, a small intestinal epithelial cell line, and a B-cell line (data not shown). The hToll gene is expressed most strongly in spleen and PBL (Fig. 3); its expression in other tissues may be due to the presence of macrophages and dendritic cells, in which it could act as an early-warning system for infection. Alternatively, hToll may be widely expressed because hToll signals through the conserved NF-kB pathway (see below) and NF-kB is a ubiquitous transcription factor.

To characterize hToll functions and see whether it can induce transcription of immune response genes like dToll, we generated a dominant-positive mutant of hToll because the natural ligand of hToll is unknown. To produce a constitutively active mutant of hToll, we made use of genetic information from dToll: analysis of ventralizing mutants in Drosophila embryos had identified the function of the ectodomain C-flanking cysteine-rich region in dToll as controlling the activity of dToll in signal transduction. In three dominant
mutations, one of four conserved cysteine residues was changed to a tyrosine, rendering $dT_{oll}$ constitutively active\textsuperscript{16}. In another dominant mutant allele of $dT_{oll}$, this region is deleted altogether, together with the rest of the extracellular part of $dT_{oll}$, indicating that a constitutively active dominant mutant of $dT_{oll}$ can be generated by disrupting the disulphide bonds in the C-terminal region flanking the ectodomain or deleting these with the rest of the extracellular part of $dT_{oll}$\textsuperscript{18}. As this region is conserved in $hT_{oll}$ (Fig. 2a) and most of the C-terminal flanking region, including three of the four conserved cysteine residues, were removed, $mCD_{4}$ acted as an epitope tag to monitor the expression of the construct in human cells by using monoclonal antibodies. This chimeraic construct was cloned into an expression vector and transfected into the human monocytic cell line THP-1 which endogenously expresses $hT_{oll}$ and should thus contain the downstream signalling components. Seven stably transfected clones were selected for further analysis. Expression of this CD4/$hT_{oll}$ chimera in these clones was identified by surface $mCD_{4}$ expression detected by FACS analysis (Fig. 4b). We used RT-PCR on seven independent clones to test whether the dominant-positive mutant of $hT_{oll}$ can induce expression of NF-κB-controlled immune-response genes; untransfected or mock-transfected THP-1 cells acted as controls. Each experiment was repeated 8–12 times and the results from two representative CD4/$hT_{oll}$ transfected clones and untransfected THP-1 cells are shown in Fig. 4c. Expression of a constitutively active mutant of $hT_{oll}$ caused induction of the IL-1, IL-8 and B7.1 genes. In the presence of interferon (IFN)-γ, IL-6 mRNA was also strongly induced, which was not due to induction of IL-1 by IFN-γ because during the time of the experiments (8–20 hours) IL-1 was induced in control cells but IL-6 was not. Other genes that might be controlled by signalling through $hT_{oll}$, for example inducible nitric oxide synthase (iNOS) and B7.2, although expressed in the transfectants, could not be tested in this system because of constitutive expression (B7.2) or clonal heterogeneity in the bulk population of control cells (iNOS).

As $dT_{oll}$ signals through the NF-κB pathway, we tested whether $hT_{oll}$ could induce activation of NF-κB. The CD4/$hT_{oll}$ chimaeric construct was transiently transfected into Jurkat cells together with an NF-κB-promoter-driven luciferase reporter gene. As shown in Fig. 4d, expression of CD4/$hT_{oll}$ causes induction of NF-κB, confirming that $T_{oll}$/NF-κB signalling is conserved in $Drosophila$ and humans.
These findings indicate that Toll functions in vertebrates as a non-clonal receptor of the immune system. Pro-inflammatory cytokines such as IL-1, IL-6 and IL-8 are induced during infection but, apart from CD14, the cellular receptors that are activated during this initial phase have not yet been identified. It has been argued that the innate immune response has been preserved in vertebrates not only to resist infection before adaptive immunity is induced, but also to induce signals that inform the adaptive system of the presence of pathogens within the internal milieu. We have shown that the Toll/NF-κB host defence pathway is conserved from Drosophila to humans and that the human homologue of the Drosophila Toll protein can induce signals for activating both an innate and an adaptive immune response in vertebrates.

Methods

Cloning and sequence analysis. BLAST searches of the NCBI sequence database were made using sequence profiles of the dToll/IL-1R signalling domains. Matching sequence (GenBank accession number H48602) corresponded to the IMAGE consortium clone number 202057. PCR primers were designed based on the EST sequence and used to amplify a 220-bp fragment from a human spleen cDNA library (Clontech). The PCR-amplified fragment was used to screen a human spleen cDNA library as described. The 5' end of the cDNA was cloned by 5'-RACE protocol as described. Cloned products were subjected to automated sequencing (ABI).

RNA expression. Tissue northern blots were purchased from Clontech and analysed according to the manufacturer’s instructions. The presence of equal amounts of RNA on the blots was confirmed by hybridization with a β-actin probe.

RT-PCR analysis. Total RNA was isolated using TRIzol Reagent (Gibco BRL) and 0.5 μg was used for reverse transcription using AMV reverse transcriptase (Seikagaku America) or Superscript reverse transcriptase (Gibco BRL). One-tenth to one-twentieth of this reaction was used as a template for PCR amplification with Taq polymerase (Gibco) for 25–35 cycles at 94°C for 1 min, 55°C for 1 min, and 72°C for 1 min. Sequences of the primers were taken from ref. 21.

Construction of mCD4/toll chimaera. An SphI site was introduced between V365 and N366 in the mouse CD4 gene by site-directed mutagenesis. An internal SphI site in hToll at the position between M620 and P621 was used for the construction of the chimaera.

Monoclonal antibodies and FACS analysis. Anti-mouse CD4 antibody (L3T4) (Pharmingen) was used for staining. Flow cytometry analysis was done on a FACScan (Becton–Dickinson).

NF-κB assay. Jurkat cells were transiently transfected using Lipofectamine reagent (Gibco BRL) according to the manufacturer’s instructions. 1 μg pB2XLuc plasmid, containing NF-κB-driven luciferase gene was cotransfected with 1 μg pSRαN expression vector with or without (control sample) the insert encoding CD4/Toll chimaera. For a positive control, cells were treated with PMA (phorbol myristate acetate; 20 ng ml⁻¹) and PHA (phytohaemagglutinin; 2 μg ml⁻¹) for 3–4 h. Cells were lysed 2 or 3 days later for measurement of luciferase activity using reagents from Promega.

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A protein related to splicing factor U2AF\textsuperscript{35} that interacts with U2AF\textsuperscript{65} and SR proteins in splicing of pre-mRNA

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Recognition of a functional 3' splice site in pre-mRNA splicing requires a heterodimer of the proteins U2AF\textsuperscript{65}/U2AF\textsuperscript{35}. U2AF\textsuperscript{65} binds to RNA at the polypyrimidine tract\textsuperscript{1,2}, whereas U2AF\textsuperscript{35} is thought to interact through its arginine/serine-rich (RS) domain with other RS-domain-containing factors bound at the 5' splice site, assembled in splicing enhancer complexes, or associated with the U4/U6.U5 small nuclear ribonucleoprotein complex\textsuperscript{3-7}. It is unclear, however, how such network interactions can all be established through the small RS domain in U2AF\textsuperscript{35}. Here we describe the function of a U2AF\textsuperscript{35}-related protein (Urp), which is the human homologue of a mouse imprinted gene. Nuclear extracts depleted of Urp are defective in splicing, but activity can be restored by addition of recombinant Urp. U2AF\textsuperscript{35} could not replace Urp in complementation, indicating that their functions do not overlap. Co-immunodepletion showed that Urp is associated with the U2AF\textsuperscript{65}/U2AF\textsuperscript{35} heterodimer. Binding studies revealed that Urp specifically interacts with U2AF\textsuperscript{65} through a U2AF\textsuperscript{35}-homologous region and with SR proteins (a large family of RS-domain-containing proteins) through its RS domain. Therefore, Urp and U2AF\textsuperscript{35} may independently position RS-domain-containing factors within spliceosomes.

We previously purified and cloned a mammalian serine kinase, SRPK1, which is highly specific for the RS-rich superfamily of splicing factors\textsuperscript{8-10}. This stringent substrate specificity enabled us to do a yeast two-hybrid screen to search for new RS-domain-containing proteins. We isolated a positive clone from a HeLa complementary DNA library that contains a typical RS domain at the carboxy terminus (Fig. 1a). The recombinant protein was efficiently phosphorylated in the RS domain, like other RS-domain proteins, by SRPK1 \textit{in vitro} (data not shown). This protein of 483 amino acids does not contain any RNA-recognition motif characteristic of the SR family of splicing factors\textsuperscript{11-12}; instead, it contains two stretches of sequences, H1 and H2 (boxed in Fig. 1b), homologous to the human splicing factor U2AF\textsuperscript{35}. We therefore named it Urp, for U2AF\textsuperscript{35}-related protein. Northern blotting analysis indicates that both U2AF\textsuperscript{35} and Urp are ubiquitously expressed in all human tissues and cell lines examined (data not shown). Homology searches revealed that Urp had been independently cloned from humans as U2AF1-RS2 (ref. 13), which is nearly identical (94% amino-acid identity) to another human gene, U2AF1-RS1 (ref. 13) (Fig. 1b). These two human genes are equally homologous (~80% identical) to two mouse genes\textsuperscript{14}, one of which is imprinted\textsuperscript{15}, and the other has a high transmission distortion in interspecific backcross progeny\textsuperscript{16}. Thus, several U2AF\textsuperscript{35}-related proteins are expressed in mammalian cells and may be members of the family of RS-domain-containing splicing factors\textsuperscript{17}.

To determine the function of Urp, we raised rabbit polyclonal antibodies against recombinant Urp expressed in bacteria. Immunoblotting of HeLa cell nuclear extracts demonstrated that anti-Urp antibodies specifically detect a single band of relative molecular mass 66K (Fig. 2a), which corresponds to endogeneous Urp as it matches the size of Urp translated in reticulocyte lysate (data not shown). To see whether Urp is essential for splicing, we constructed affinity columns with anti-Urp antibodies or preimmune serum. Control depletion with preimmune serum had no effect, but passing nuclear extract through an anti-Urp affinity column efficiently depleted it of Urp, as evidenced by western blotting and complete inhibition of splicing (data not shown). Although this result is consistent with Urp being a splicing factor, this Urp-depleted extract could not be complemented by recombinant Urp, probably because of co-depletion of other essential splicing factors (see below). We therefore did the immunodepletion in the presence of high salt (0.5 M KCl) to minimize co-depletion. As shown in Fig. 2c, human β-globin pre-mRNA splicing in Urp-depleted nuclear extract was significantly decreased (compare lanes 1 and 2). The second step of the splicing reaction, which generates released lariat intermediate and...