Research

Phylogenetic variation and polymorphism at the Toll-like receptor 4 locus (TLR4)

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Abstract

Background: Differences in responses to bacterial surface lipopolysaccharides (LPSs) are apparent between and within mammalian species. It has been shown in mice that resistance to LPS is caused by defects in the Toll-like receptor 4 gene (Tlr4), the product of which is thought to bind LPS and mediate LPS signal transduction in immune system cells.

Results: We have sequenced the Toll-like receptor 4 gene of humans (TLR4; 19.0 kilobases, kb) and mice (Tlr4; 91.7 kb), as well as the coding region and splice junctions of Tlr4 from 35 mouse (Mus musculus) strains, from the chimpanzee and from the baboon. No other discernible genes or regions of interspecies conservation lie close to Tlr4 and, in both humans and mice, flanking sequences and introns are rich in repeats of retroviral origin. Interstrain analyses reveal that Tlr4 is a polymorphic protein and that the extracellular domain is far more variable than the cytoplasmic domain, both among strains and among species. The cytoplasmic domain of the Tlr4 protein is highly variable at the carboxy-terminal end.

Conclusions: We suggest that selective evolutionary pressure exerted by microbes expressing structurally distinguishable LPS molecules has produced the high level of variability in the Tlr4 extracellular domain. The highly variable carboxy-terminal region of the cytoplasmic domain is likely to determine the magnitude of the response to LPS within a species.

Background

Cells of the innate immune system sense host invasion by detecting structural determinants that are broadly conserved among pathogens of a given phylogenetic group [1]. The lipopolysaccharides (LPS or endotoxin) that decorate the outer membrane of Gram-negative bacteria are excellent examples of such determinants. In response to minute concentrations of LPS derived from certain Gram-negative bacteria, macrophages secrete cytokines such as tumor necrosis factor (TNF), interleukin-1 (IL-1) and IL-6, which contribute to the containment of infection and help to initiate a specific immune response. On the other hand, overstimulation of the innate immune system through this channel can lead to acute systemic inflammation and shock [2,3].

Dramatic differences in LPS responses are apparent between closely related species [4], and there are substantial differences even among genetically heterogeneous members of the
same species [5]. For example, whereas humans and chim-panzees are generally considered to be very sensitive to LPS [4], baboons and most other primates are highly resistant. It is likely that part of the difference in sensitivity may be explained at a very proximal level, although differences in responses to cytokines (for example TNF) may also have a role. Neither interspecific differences nor interindividial differences in LPS responses have, until recently, been accessible to systematic genetic analysis. Advances in understanding LPS signal transduction now permit these issues to be examined.

Although LPS was once thought to exert its effects through intercalation into biological membranes, or to bind to many different receptors on the cell surface, it has been clear for nearly three decades that there is, in fact, a single biochemical pathway for LPS detection. This was indicated by the observation that allelic [6] mutations of a single gene (Lps) in mice could entirely abrogate the response to LPS, and did so with great specificity. Mice of the strains C3H/HeJ [7] and C57BL/10ScCr [8] are highly resistant to LPS, showing none of the usual biological effects, yet respond normally to other bacterial products and to most cytokines induced by LPS [9].

We recently identified the Lps locus through positional cloning [10] and showed that the LPS-resistance phenotype was caused by defects in the Toll-like receptor 4 gene (Tlr4) [11]. In C3H/HeJ mice, a point mutation (P712H; single-letter amino-acid notation) modifies the protein within the cytoplasmic domain, creating a codominant inhibitory effect on LPS signal transduction. In C57BL/10ScCr mice the gene is deleted, yielding a recessive abolition of the LPS response. Subsequently, overexpression of the wild-type Tlr4 protein was found to enhance LPS signal transduction in wild-type macrophages, lowering the effective concentration (EC_{50}) for LPS by a factor of 30, whereas overexpression of the Tlr4P712D isoform represented in C3H/HeJ mice almost completely suppresses signaling [12]. Furthermore, genetic complementation studies have demonstrated that LPS and Tlr4 enter into close physical proximity in the course of signal transduction - Tlr4 appears to bind directly to LPS. Hence, the species origin of Tlr4 is the sole determinant of species preference for a given LPS structure [13].

Mice of the C3H/HeJ and C57BL/10ScCr strains are abnormally susceptible to infection by certain Gram-negative bacteria, suggesting that timely recognition of LPS is essential for successful containment of infection [14,15]. Because deleterious mutations of Tlr4 have become fixed spontaneously in two strains of mice, we considered it possible that other functionally important mutations might be identified in mice and humans. Moreover, information on the degree of Tlr4 polymorphism in these and other species might allow inferences about the importance of different Tlr4 domains. Accordingly, we decided to sequence the entire Tlr4 gene of both humans and mice, and to survey genetic variation at the locus in each species. We also examined the TLR4 sequence from two species of subhuman primates that have dramatically different responses to LPS.

Results

Overall structure of TLR4 and Tlr4

The mouse Tlr4 gene is somewhat longer than its human counterpart TLR4, owing to the greater length of intronic sequence - 15,337 base pairs (bp) from beginning to end of the transcribed sequence in the mouse, compared with 11,467 bp in the human. There are three exons in Tlr4, and each corresponds to a homologous sequence in the human gene. Rock et al. [16] reported a human cDNA sequence (GenBank accession number U88880) that includes a fourth exon, positioned between the ‘normal’ first and second introns. When included in the processed transcript, however, this exon specifies early termination of the polypeptide chain. Although it is possible that translation is initiated distal to the added stop codon, and that a shorter product results in the human than in the mouse, this would be unusual, given the length of the 5’ untranslated region (UTR) that would then exist and the presence of multiple upstream initiation codons. Moreover, there is no murine sequence homologous to the alternative second exon of the human gene. The biological significance of this exon is therefore unclear and, in all likelihood, its inclusion in the mRNA leads to the formation of a nonfunctional protein.

Neither the human nor the mouse gene has a TATA element or CAAT box in the proximal promoter region. A number of conserved promoter and enhancer motifs are apparent on alignment of the murine and human 5’ flanking sequences, and are described in detail elsewhere [17]. Both Tlr4 and TLR4 lie amid repetitive sequences of retroviral origin, and no other genes are detected close to either of them using homology searches or the gene prediction algorithm GRAIL. In Figure 1, the grayscale images of the human and mouse genes call attention to the repetitive elements in the region and illustrate the relationship between exons and spacing in the two species.

Genetic variation at the mouse Tlr4 locus

Among 35 strains of Mus musculus, 10 different alleles were identified on the basis of mutations at 22 sites with respect to the reference sequence. Of these, 13 create amino-acid substitutions (Table 1, Figure 2). The most common murine allele was represented at a frequency of only 69%. The ancestry of different Tlr4 alleles can be traced by haplotype analysis, as many deviations from the reference allele occur in conjunction with one another. A plausible arrangement of strain relationships is presented in Figure 3. Some strains have accumulated many more mutations than others. For example, the P/J strain Tlr4 gene exhibits 11 mutations that distinguish it from the most common haplotype, six of them specifying changes in the Tlr4 amino-acid sequence; the SEA/GnJ strain differs by nine mutations, and the strains NZW/J and VM/Dk, which are identical to one another,
differ from the most common haplotype at six sites. Shared mutations suggest that introgression took place after mutational separation had occurred, leading to the introduction of groups of mutations by genetic recombination. Hence, mice of the P/J, NZW/J, and VM/Dk strains have several mutations that are observed in the A/J and BALB/c strains, but also lack some of the mutations of the latter strains and have unique mutations of their own.

Most of the murine mutations reside within exon 3, and only two substitutions are noted that modify the cytoplasmic domain (Figure 2). Of these, however, one mutation (R761H) is fairly common among the strains surveyed, and the corresponding residue has been reported as an H in the hamster. A single conservative substitution (V637I) was noted within the transmembrane domain of the P/J strain.

Anthropoid ape and lower primate TLR4 sequences, and their relationship to the human and rodent sequences

The human and chimpanzee amino-acid sequences are nearly identical over the interval studied, distinguished by only three substitutions. The baboon sequence is 93.5% identical to the human in the extracellular domain, differs in the transmembrane domain by one substitution out of 30 residues, and differs in the proximal cytoplasmic domain by only one residue in 155. At the carboxyl terminus, however, homology is badly disrupted, so that 16 of the last 21 human residues are not replicated in the baboon protein, which is 13 amino acids shorter than the human protein. Similarly, among rodents, the carboxyl terminus of the protein is the least conserved. Overall, the order of conservation with respect to domain is: proximal cytoplasmic domain > transmembrane domain > extracellular domain > distal cytoplasmic domain (Table 2, Figure 4).
The pathway by which LPS activates host innate immune defenses has been illuminated by the positional cloning of Lps and establishment of its identity with Tlr4 [11], a mammalian representative of an ancient family of receptors [16] that serve both developmental and defensive functions. The function of Tlr4 as the LPS signal transducer became clear when mapping [10] and sequencing [11] data revealed Tlr4 as the only gene in a critical 2.6 megabase (Mb) region, and, furthermore, showed that two strains of LPS-resistant mice are homozygous for mutations of Tlr4 that are absent in closely related LPS-sensitive strains.

In Drosophila, the prototypic homolog of the mammalian Toll-like receptors (Toll) defends against fungal infection [18], whereas the protein 18-wheeler defends against bacterial infection [19]. In the case of Toll, there is no evidence of direct contact between the receptor and the microbial pathogen or its components. Rather, infection activates a proteolytic cascade that leads to the production of an endogenous ligand (Spätzle), which in turn engages Toll. The situation is apparently different in mammals, in which Tlr4 is clearly a direct interface with the microbial world. As such, the primary structure of a Tlr4 molecule determines ligand specificity [13], accounting for the well-known observation that mouse cells can recognize tetra-acyl lipid A as an agonist [20] whereas human cells recognize it as an antagonist [20-24].

As mice of the C3H/HeJ strain are highly susceptible to infection by Gram-negative bacteria, it would seem plausible

| Mouse strain* | Genomic nucleotide affected | Exon/intron affected | Altered amino acid | Affected receptor domain† | Conserved‡ |
|---------------|-----------------------------|----------------------|-------------------|---------------------------|------------|
| 10            | 26,400 A → G                | Exon 1               | -                 | -                         |            |
| 4, 8, 20, 21, 22 | 37,685: (T)10              | Intron 2             | -                 | -                         |            |
| 34, 35        | 37,685: (T)12              | Intron 2             | -                 | -                         |            |
| 23            | 37,754 G → A               | Exon 3               | D94N              | Ecto                      | Yes        |
| 4, 8, 10, 20, 21, 34, 35 | 38,101 G → A              | Exon 3               | M209I             | Ecto                      | Yes        |
| 21            | 38,130 A → G               | Exon 3               | D219G             | Ecto                      | No         |
| 4, 8, 21      | 38,234 G → A               | Exon 3               | V254I             | Ecto                      | Yes        |
| 10            | 38,584 A → G               | Exon 3               | -                 | -                         |            |
| 21            | 38,742 A → T               | Exon 3               | Q423L             | Ecto                      | Yes        |
| 10            | 38,794 G → A               | Exon 3               | -                 | -                         |            |
| 10            | 38,903 G → T               | Exon 3               | A477S             | Ecto                      | Yes        |
| 18            | 39,020 A → G               | Exon 3               | T516A             | Ecto                      | Yes        |
| 4, 8, 10, 20, 21, 22, 34, 35 | 39,199 C → T       | Exon 3               | -                 | -                         |            |
| 4, 8, 10, 20, 21, 34, 35 | 39,253 A → C              | Exon 3               | E593D             | Ecto                      | Yes        |
| 23            | 39,273 A → T               | Exon 3               | N600I             | Ecto                      | No         |
| 10            | 39,294 C → T               | Exon 3               | A607V             | Ecto                      | Yes        |
| 10            | 39,383 G → A               | Exon 3               | V637I             | TM                        | No         |
| 10            | 39,544 G → A               | Exon 3               | -                 | -                         |            |
| 19            | 39,604 T → C               | Exon 3               | -                 | -                         |            |
| 4, 8, 20, 21, 34, 35 | 39,631 C → T            | Exon 3               | -                 | -                         |            |
| 4, 8, 20, 21, 34, 35 | 39,756 G → A            | Exon 3               | R761H             | Cyto                      | Yes        |
| 10            | 39,907 T → G               | Exon 3               | N811K             | Cyto                      | Yes        |

*Strains are as follows: 1, NZO/HILt; 2, SJ/Cal; 3, DBA/2J; 4, A/J; 5, FL/Su; 6, CBA/J; 7, AKR/J; 8, BALB/c; 9, DDD/1c; 10, P/J; 11, MRL/MpJ; 12, SJL/J; 13, NOD/LtJ; 14, 129Sv; 15, FL/J; 16, MA/MgJ; 17, SW RJ; 18, LP/J; 19, PRO/ReJ; 20, SD/Ei; 21, SEA/GnJ; 22, SM/J; 23, KK/HkJ; 24, ST/ReJ; 25, WBR/Ei; 26, YBR/Ei; 27, FVB/NJ; 28, PL/J; 29, LT/CHRe5v; 30, RIII/J; 31, RF/J; 32, NZB/BIN; 33, AU/SsJ; 34, NZW/Laxc; 35, VM/DXB.
†Exon 1, 26041-26424; Exon 2, 32397-32563; Exon 3, 37732-41297. A TG at 26335; TGA at 39982. Ecto, extracellular domain; Cyto, cytoplasmic domain; TM, transmembrane domain.
‡Yes implies one or two forms; No implies three to five forms among the six mammalian species examined.
that among human patients with Gram-negative sepsis, some individuals may have been at risk by virtue of mutations in \textit{TLR4}. The first step in determining whether different isoforms of \textit{Tlr4} confer added risk of, or protection against, sepsis is the assessment of genetic variability at this locus in the normal population.

Whereas ablative mutations of \textit{Tlr4} abolish the capacity to detect LPS, the question arises as to whether less severe mutation might alter either the specificity of LPS detection, as discussed above, or the magnitude of the LPS signal. Our present knowledge of \textit{Tlr4} structural variation in mice may permit an answer to this question, insofar as the mutations might easily be recreated, and measurements of signal-transducing activity through the modified receptors carried out in immortalized C3H/HeJ macrophages [13]. Animals of the A/J and P/J strains have defective tumoricidal capacity [25-29], although in neither case is the \textit{Lps} locus implicated in the defect.

Deleterious mutations of \textit{TLR4} might reasonably be sought in individuals who have developed serious Gram-negative infections, on the premise that mice with deleterious mutations of \textit{Tlr4} are rendered susceptible to such infections. Similarly, in birds, a polymorphism at the \textit{tenascin} locus (which lies a few megabases proximal to \textit{Tlr4} in mice) predicts susceptibility to Gram-negative infection [30], suggesting that it may lie in linkage disequilibrium with a particular form of the avian \textit{Tlr4} gene. Beyond this, it may be assumed, given the powerful pro-inflammatory activity of the receptor, that germline or

\begin{figure}
\centering
\includegraphics[width=\textwidth]{figure2.png}
\caption{Distribution of coding mutations found in \textit{Tlr4} of 35 \textit{Mus musculus} strains. All coding mutations reside within exon 3. Most occupy portions of the gene corresponding to the extracellular domain. The transmembrane domain is denoted by a blue-green bar. Mutations occurring at sites that are relatively conserved among species (only one or two forms among six species) are shown in blue; those occurring at sites that are less conserved (three to five forms among six species) are shown in black.}
\end{figure}

\begin{figure}
\centering
\includegraphics[width=\textwidth]{figure3.png}
\caption{Genetic distance and probable ancestral relationships among \textit{Tlr4} genes of 35 \textit{Mus musculus} strains. Numbers within circles denote strains, in accordance with the legend to Table 1. Numbers adjacent to arrows indicate the mutational distance (number of mutations separating each strain from its presumed ancestor), with reference to both coding and noncoding substitutions listed in Table 1. Arrows point in the direction of descent, and their lengths are proportional to distance. Dotted lines suggest introgression, given the similarity of the haplotypes observed. The symbol '?' denotes the likelihood of an intermediate form before interbreeding of strains.}
\end{figure}
somatic mutations of TLR4 could in some instances cause constitutive signal transduction (as observed with the Drosophila Toll locus [31]). Such mutations might, in principle, account for certain inflammatory diseases. In fact, in the Lpr model of systemic lupus erythematosus, the Lps locus was believed a likely modulator of phenotype [32]. This would seem still more likely at present, given the paucity of other candidate genes in the immediate vicinity of Tlr4 [10].

The evolutionary conservation of Tlr4 is of particular interest, in that different species show preferential responses to some LPS forms and not others [20-24], and have particular ‘set points’ for responses to toxic LPS molecules. An assessment of variability may be made through comparison of different species, but is complemented by the study of a large number of individuals within species. Both approaches reveal that the extracellular domain of Tlr4 is highly variable compared with the transmembrane domain and proximal cytoplasmic domain of the protein. Pooling the number of mutable sites in the extracellular domain and transmembrane domain of humans and mice, 17 coding changes are observed, compared with two in the proximal cytoplasmic domain. Moreover, variability does not seem confined to any particular region of the extracellular domain, but is spread uniformly across its length.

The extracellular domain of any receptor is concerned principally with ligand recognition. Given that the ligand is an endogenous protein, extracellular domain conservation tends to be strict, insofar as mutations affecting extracellular domain structure are likely to diminish specificity or affinity of binding. Hence, protein-binding receptors tend to be minimally polymorphic. The presumed ligand for Tlr4 is LPS itself, presented alone or in conjunction with another protein (for example CD14) [13], and the relatively high frequency of polymorphism observed in Tlr4 may be viewed as a consequence of the protective effect rendered by LPS recognition and the variability of LPS structure.

The cytoplasmic domain of Tlr4 is far more stringently conserved than the extracellular domain. This is probably due to the fact that the cytoplasmic domain of Tlr4 is not required to cope with a ligand of variable structure. Rather, when called upon to signal, it must do so, utilizing transducing
molecules with conserved structures. On the other hand, the intensity of the LPS signal has apparently been optimized for each species [33], perhaps ensuring that the response to LPS is appropriately integrated into the immune response as a whole. Humans, ungulates and rabbits, for example, exhibit an intense response to low concentrations of LPS, whereas the lower primates and most rodents are relatively more resistant [4]. The highly variable carboxy-terminal end of the Tlr4 cytoplasmic domain may be seen as the embodiment of interspecific differences in LPS sensitivity, although the poor conservation of this portion of the molecule might alternatively be taken to indicate a neutral effect of mutation. It is also possible that this region of the molecule is subject to a higher rate of mutation than that applying to the rest of the protein. Although most such mutations might be removed by selection, some might be discovered in populations defined by the occurrence of Gram-negative sepsis.

Materials and methods
Determination of the complete mouse (Tlr4) and human (TLR4) genomic sequences
The mouse bacterial artificial chromosome (BAC) 152C16 (from the 129/J strain, Research Genetics) was previously shown by us to contain the Tlr4 gene in its entirety [10], and a small fraction of Tlr4 was also found in the overlapping BAC 309I17 [10]. Human TLR4 was identified in BAC 110P15 (Genome Systems) by hybridization screening using a PCR-amplified human TLR4 cDNA sequence as a probe. All three BACs were fragmented by ultrasound, shotgun cloned into the vector pBluescript-KS, and extensively sequenced using ABI model 373 and 377 sequencers, using Big Dye terminators; 959 reads were obtained from 309I17, 1503 reads from 110P15, and 2731 reads from 152C16. The average read length was approximately 700 nucleotides. To concentrate data acquisition efforts on the Tlr4 and TLR4 genes themselves, PCR primers were fashioned to match regions flanking each gene. A 16 kb fragment was amplified from the mouse BAC 152C16, and a 12 kb fragment was amplified from the human BAC 110P15, each containing all exons of the respective gene. These fragments were also shotgun cloned and extensively sequenced, so that the depth of sequence reached an average of 12 reads over the area of greatest interest. Sequence assembly used the programs phred and Phrap (obtained from Brent Ewing and Phil Green, University of Washington Genome Center). Interpretation of repetitive elements was achieved with the program RepeatMasker (obtained from Arian Smit, University of Washington Genome Center). A contiguous high-quality sequence 18,974 bp in length, containing TLR4, was obtained from the human BAC, and a contig 91,748 bp in length, containing Tlr4, was obtained from the mouse BAC. Over these intervals, the error rate was estimated at <1 per 10^4 bp. The sequences have been posted to GenBank in annotated form (accession number AF177767 for the murine sequence and AF177765 for the human sequence). All data related to mutations are presented with reference to these sequences.

Figure 4
Spline curve illustrating interspecific sequence variation across the Tlr4 protein. A multiple alignment of Tlr4 sequences from three rodent species (mouse, rat and hamster) and three primate species (human, chimpanzee and baboon) was generated using the GCG program Pileup. The number of amino acids observed at each residue was plotted using the program Prism 3.0 (a value of 1 was assigned if a single amino acid was observed in the six species; a value of 5 was assigned if five different amino acids were observed among the six species, and so on). The points were then connected using a cubic spline curve. TM, transmembrane domain. Numbering refers to the human sequence. Where a deletion was introduced by Pileup, a single mismatch was assumed. Where the sequence was truncated, each missing residue was tabulated as a separate mismatch.
Sequencing DNA from mouse, chimpanzee and baboon samples

Mouse DNA, obtained from animals of 35 Mus musculus strains, was ordered from the Jackson Laboratories. Chimpanzee and baboon DNA were obtained from Kurt Benirschke (University of California, San Diego) and Gregory Delzoppo (Scripps Research Institute), respectively.

The three principal exons of Tlr4 were amplified independently from mouse genomic DNA samples, leaving a margin of approximately 50 bp to each side of the exons so as to identify intronic mutations that might alter splicing. All exons of the chimpanzee were amplified and sequenced using the same primers used to amplify and sequence the human exons. For the baboon, the first two exons were also amplified using these same primers; however, the third exon of the baboon was amplified with a substituted primer at the 5' end.

The PCR products were isolated by agarose gel electrophoresis. Exons 1 and 2 were sequenced using the same primers that were used for amplification. Exon 3 was sequenced using the flanking primers as well as a collection of eight internal primers. In this manner, the entire coding region and all splice junctions of Tlr4 could be covered with a total of 14 sequencing reads. All primers used for amplification and sequencing are presented in Table 3; separate sets were used to amplify and sequence mouse and the primate samples.

Independent assembly of each sample was required as a condition for further analysis, and if such assembly failed, additional reads were executed using a secondary collection of primers. Thereafter, mutations were identified en masse, by pooling all of the reads from 25 to 30 samples at a time and reassembling with the program polypHred, using the phredPhrapPoly script (obtained from Natalie Kolker, University of Washington Genome Center). Consed alpha (obtained from David Gordon, University of Washington Genome Center) was used to visualize reads and mutations.

### Table 3

| Oligonucleotide primers used to amplify and sequence mouse and human Tlr4 genes |
|-----------------------------------|-----------------------------------|
| **Mouse**                          | **Primate**                       |

#### Amplification

| Exon 1 | Exon 2 | Exon 3 |
|--------|--------|--------|
| ↑ CAGTCGGTCAGCAAGACGCTTCTTCTC | ↑ GCTCGGTAAAACGGTGATAG | ↓ TGGAAATCTCTGSSGCAGAG |
| ↓ CAAGCGAGCCTGACGAAAGGATTG | ↓ TTAATGCTTTGGAGAGG | ↓ CAGTCGCAATAGAAAGTTG |
| ↓ AAGGGAAGTATGAGAACACCGTCTG | ↓ AAATCTGGGGCTGTGTCCTT | ↓ CACATACATCGATGAGAGG |
| ↑ TCTCGCTGACACCATCATCACCTG | ↑ CATGTGTTCTTATGCTGACG | ↓ TGACTTCTTTGCTCAGGTTG |
| ↓ ATGCACTGCTTTGCTCTT | ↓ TGCGCGTCAATCAGGTCGT | ↓ GCTCGGTAAAACGGTGATAG |

#### Sequencing

| Exon 1 same as above | Exon 2 same as above | Exon 3 primary |
|----------------------|----------------------|----------------|
| 1 ↓ ATGCACTGCTTTGCTCTT | 31 ↑ GAGCTGGTAAATGATATTAG | 31 ↓ TCAAAATGCGAGCCTCTCTAG |
| 2 ↓ TTTATATCCTCACAAG | 32 ↑ CAAATCTCCTCTAGACCTGTCCT | 32 ↓ AATACTTTGCTGGTTCCTCCC |
| 3 ↓ CAGCTCCTCTCACG | 33 ↑ TTTCCTGACACTCTGTCAGA | 33 ↓ GGAATCTTATTGATTGAGAC |
| 4 ↓ TGGAAAGACATGAAATCCTT | 34 ↑ TCTCCGGTCACTCTGTCAGACC | 34 ↓ GAAAGCGAGCTGAAATGAGAG |
| 5 ↑ TGATGATGCAACAGAT | 35 ↑ TCCAGATGAAATAGCAGCATTGTCG | 35 ↓ AACAAGGTGGAGCCAG |
| 6 ↑ TGGAAAGACATGAAATCCTT | 36 ↑ CAATCTCCTGACACTCTGTCAGA | 36 ↓ AATACTTTGCTGGTTCCTCCC |
| 7 ↑ CGCTCCTGCTCTCAGGTCG | 37 ↑ TCCATCGTCTGACACTCTGTCAGA | 37 ↓ GGAATCTTATTGATTGAGAC |
| 8 ↑ TGGAAAGACATGAAATCCTT | 38 ↑ AAATCTGGGGCTGTGTCCTT | 38 ↓ GAGCTGGTAAATGATATTAG |
| 9 ↑ AAGGAGACTGCACTCAC | 39 ↑ CTCCGCTGGCTCTTCCTCCTGTG | 39 ↓ TCAAAATGCGAGCCTCTCTAG |
| 10 ↓ TCCAGATGAAATAGCAGCATTGTCG | 40 ↑ TCCGCTCTGCTGACACTCTGTCAGA | 40 ↓ GGAATCTTATTGATTGAGAC |

#### Exon 3 secondary

| 5 ↑ TGATGATGCAACAGAT | 1 ↑ GAAATTTTGGAGCAGT | ↓ TTAATTTTTTGGAGCAGT |
| 7 ↑ CTGCAACACTGCTAGCATTG | 2 ↑ TTCAATGATCCTTACATCTCAG | 2 ↓ TTCAATGATCCTTACATCTCAG |
| 9 ↓ ATGCACTGCTTTGCTCTT | 3 ↑ TTAATTTTTTGGAGCAGT | 3 ↓ TTAATTTTTTGGAGCAGT |
| 10 ↓ TCCAGATGAAATAGCAGCATTGTCG | 4 ↑ TCCGCTCTGCTGACACTCTGTCAGA | 4 ↓ TCAAAATGCGAGCCTCTCTAG |
| 5 ↑ TGATGATGCAACAGAT | 5 ↓ GTGTCCTGACACTCTGTCAGA | 5 ↓ GTGTCCTGACACTCTGTCAGA |
| 6 ↑ TCCGCTCTGCTGACACTCTGTCAGA | 6 ↓ AACAAGGTGGAGCCAG |
| 7 ↓ GTTTCGCTCTTATGAGAGG | 7 ↓ GTTTCGCTCTTATGAGAGG |
| 8 ↓ AAGGAGACTGCACTCAC | 8 ↓ AAGGAGACTGCACTCAC | 8 ↓ AAGGAGACTGCACTCAC |
| 9 ↓ AAGGAGACTGCACTCAC | 9 ↓ AAGGAGACTGCACTCAC | 9 ↓ AAGGAGACTGCACTCAC |
| 10 ↓ TCCAGATGAAATAGCAGCATTGTCG | 10 ↓ TCCAGATGAAATAGCAGCATTGTCG | 10 ↓ TCCAGATGAAATAGCAGCATTGTCG |
| 0 ↑ TTTATATCCTCACAAGAT | 0 ↑ TTTATATCCTCACAAGAT | 0 ↑ TTTATATCCTCACAAGAT |
| 0 ↓ GCTCAGATGAAATAGCATTAG | 0 ↓ GCTCAGATGAAATAGCATTAG | 0 ↓ GCTCAGATGAAATAGCATTAG |

↑ Primer matches + strand; ↓ primer matches - strand.
The annotated chimpanzee exon sequences have been submitted to GenBank with the accession numbers AF179218, AF179219 and AF179220. The baboon sequences have been submitted with the accession numbers AF180962, AF180963 and AF180964. For genetic comparisons, rat (GenBank AF057025) and hamster (AF153676) Tlr4 sequences were also used.

**Genetic computation**

A 500 MHz DEC-alpha system equipped with 256 Mbytes of memory was used for direct analysis of sequence data as described above. In addition to the programs already mentioned, the GCG software (version 9.0) was used for alignment analysis, with the program Pileup used in multiple alignments of protein sequences. The windows-based program Generunner 3.0 (Hastings Software) was used for the design of oligonucleotide primers. A spline curve describing heterogeneity of the Tlr4 polypeptide sequence from different species was produced using the program Prism 3.0 (Graphpad Software). Sequences were prepared for submission with the use of the program Sequin 2.90 (obtained from the National Center for Biotechnological Information).

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