Note: Within nine months of the publication of the mention of the grant of the European patent in the European Patent Bulletin, any person may give notice to the European Patent Office of opposition to that patent, in accordance with the Implementing Regulations. Notice of opposition shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).
Description

Technical Field

[0001] The present invention relates to a Toll-like receptor (TLR) 7-knockout mouse unresponsive to synthetic compounds such as imidazoquinoline group compound R-848. The present invention also relates to a screening method for identifying an immunopotentiating synthetic compound using such a TLR7 knockout mouse.

Background Art

[0002] A Toll gene is known to be necessary for patterning dorsoventral axis while an embryo is developing in Drosophila (Cell 52, 269-279, 1998; Annu. Rev. Cell Dev. Biol. 12, 393-416, 1996), and for an antifungal immune response in an adult body (Cell 86, 973-983, 1996). The Toll is a type-I transmembrane receptor with an extracellular domain including leucine-rich repeats (LRRs) and this intracellular domain has been elucidated that it demonstrates high homology with that of mammalian interleukin-1 receptor (IL-1R) (Nature 351, 355-356, 1991; Annu. Rev. Cell Dev. Biol. 12, 393-416, 1996; J. Leukoc. Biol. 63, 650-657, 1998).

[0003] Recently, mammalian homologs of Toll, called Toll-like receptors (TLR) were identified, and heretofore, 10 families such as TLR2, TLR3, TLR4, TLR6, TLR7, TLR8, TLR9, and TLR10 have been reported (Nature 388, 394-397, 1997; Proc. Natl. Acad. Sci. USA 95, 588-593, 1998; Blood 91, 4020-4027, 1998; Gene 231, 59-65, 1999). This TLR family is known that it recruits IL-1R-associated kinase (IRAK) through MyD88 which is an adapter protein as well as aforementioned IL-1R, and then activates downstream mitogen-activated protein (MAP) kinase and NF-κB which is a nuclear factor (J. Exp. Med. 187, 2097-2101, 1998; Mol. Cell 2, 253-258, 1998; Immunity 11, 115-122, 1999). In addition, it is also believed that the role of TLR family in mammals is related to innate immune recognition as a pattern recognition receptor (PRR) that recognizes the consensus structure of bacteria (Cell 91, 295-298, 1997).

[0004] One of pathogen-associated molecular patterns (PAMPs) recognized by the aforementioned PRR is a lipopolysaccharide (LPS) which is a principal component of outer-membrane of Gram-negative bacteria (Cell 91, 295-298, 1997). It is known that host cells are stimulated by the LPS to generate various kinds of inflammatory cytokines such as TNF-α, IL-1 and IL-6 and the like on the host cells (Adv. Immunol. 28, 293-450, 1979, Annu. Rev. Immunol. 13, 437-457, 1995), and LPS captured by LPS-binding protein (LBP) is delivered to CD14 on cell surface (Science 249, 1431-1433, 1990; Annu Rev. Immunol. 13, 437-457, 1995). The present inventors generated knockout mice of TLR4 and reported that TLR4 knockout mice are unresponsive to LPS which is a principal component of the aforementioned outer-membrane of Gram-negative bacteria (J. Immunol. 162, 3749-3752, 1999). They generated TLR2 knockout mice and reported that macrophages of themselves lower the reactivity to Gram-positive bacteria cell walls or peptidoglycan which is a component thereof (Immunity, 11, 443-451, 1999), and that biological reaction is induced via TLR2 and MyD88 signalling pathway (J. Immunol. 164, 554-557, 2000).

[0005] Further, the present inventors compared/analyzed TLR6 knockout mice, wild-type mice and TLR2 knockout mice to clarify that TLR6 is a receptor protein specifically recognizing mycoplasma-derived lipoprotein/lipopeptide. They also found full length cDNA (GenBank Accession No. AF245704) for TLR9 and clarified that TLR9 is a receptor protein specifically recognizing bacterial DNA including unmetylated CpG sequence. Moreover, two new members of TLR7 and TLR8 are registered to GenBank (Accession No. AF240467 and AF246971).

[0006] On the other hand, imiquimod, which is an immune response-modulator that has been found when screening antitherpesvirus activity, shows antiviruss activity and antitumor activity in animal model. It is clarified this agent shows antiviruss activity and antiproliferative activity by inducing cytokines such as IFN-γ, IL-6 and IL-12, similar to a secondary induction of IFN-γ in various types of cells (J. Leukoc. Biol. 58, 365-372, 1995; J. Interferon Res, 1989; S2115, Antimicrob. Agents Chemother. 38, 2059-2064, 1994; Am. J. Clin. Pathol. 102, 768-774, 1994), and that imiquimod stimulates NF-κB and MAP kinase (J. Immunol. 165, 5552-5557, 2000; Mol. Cell. Biol. 15, 2207-2218, 1995), as well. In addition, imiquimod and its related compounds are known to inhibit the replication of type 2 herpes simplex virus and cytomegalo virus (J. Infect. Dis. 183, 844-849, 2001; Antimicrob. Agents Chemother. 32, 678-83, 1988). In actual treatment, it is reported that imiquimod is effective for treating pudendal wart (condyoma acuminatum) caused by human papilloma virus (Sex. Transm. Infec. 76, 162-8, 2000), and that resiquimod which is a derivative of imiquimod, abbreviated as R-848, is also effective for treating genital herpes (JAMA 285, 2182-2183, 2001). The structural formulae of these imiquimod and R-848 that are imidazoquinoline group compounds are shown as follows:
TLR families are known to be receptors, involved in recognizing the components of pathogens, and it has been clarified by the present inventors that the components derived from various kinds of bacteria and mycoses by TLR family are recognized by TLR family, as mentioned above. In other words, the present inventors have already generated TLR2-, TLR4-, TLR6-, and TLR9-deficient mice, and revealed that TLR4, TLR2, TLR6, and TLR9 are receptors which respond to lipopolysaccharide (LPS), to peptidoglycan and lipopeptide derived from Gram-positive bacteria, to polypeptide of mycoplasma similarly to TLR2, and to bacterium DNA including non-methylate CpG sequence, respectively. However, it is still unknown which substances the other TLR family member such as TLR3, TLR7, TLR8 or TLR10 and the like recognize. In addition, proteins, which can recognize immunopotentiating synthetic compounds, have been unknown.

Chuang and Ulevitch (Eur. Cytokine Netw., vol 11, no. 3, 2000, p. 372-378) show that TLR7, TLR8, and TLR9 signal via NF-κB. WO 03/043572 relates to a method of eliciting a TLR6-mediated cellular response in a cell that expresses TLR6 comprising: selecting a compound identified as a TLR5 agonist; and administering to the cell the compound in an amount that affects at least one TLR6-mediated cellular signalling pathway. It is suggested to create a TLR7 knock out mouse, and use it in screening methods. No phenotype is disclosed for said mice.

The object of the present inventors is to provide a relationship between each member of TLR family to signalling by stimulation with an immunopotentiating synthetic compound in vivo. In particular, they aim to provide a non-human animal model unresponsive to synthetic compounds, said compound being useful to elucidate the relationship of each TLR family member toward signalling by stimulation of immunopotentiating synthetic compounds in vivo. In particular, they aim to provide a Toll-like receptor (TLR) 7-knockout mouse.

The present inventors generated MyD88-deficient mice, and have already reported that the macrophages derived from MyD88-deficient mice do not respond to any of TLR ligands when producing cytokines. Since it is not possible to confirm the response in cells of MyD88-deficient mice, various kinds of compounds which might activate cells via TLR were examined. Large number of compounds were screened, and through the process of inducing cytokines from mononuclear leukocytes or macrophages, imidazoquinoline group compounds which might induce humoral immunity and cellular immunity, i.e. imidazoquinoline group compounds such as imiquimod, R-848 and the like which act on various cells and which are known as an immunopotentiator inducing generation of inflammatory cytokines such as IFN-α, IL-6 and IL-12 and the like were treated as target compounds. Meanwhile, the primary structure of the protein of TLR7 is quite similar to that of TLR9 which recognizes microbial DNA. However, the lack of reactivity to microbial DNA is not observed in TLR7-deficient mice as in TLR9-deficient mice. Then, TLR7-deficient mice of TLR family were generated, and the substances recognized by TLR7 were screened subsequently having a TLR7 as a target.

Then, the present inventors stimulated TLR7-deficient mice with R-848 and found that induction of generation of inflammatory cytokines from macrophages, induction of proliferation of B cells, and induction of maturation of dendritic cells were not observed at all. Moreover, in macrophages derived from TLR7-deficient mice, the activation of intracellular signals such as NF-κB, JNK, IRAK by stimulation with R-848 was not observed at all. This suggests that TLR7 is involved not only in recognition of components of pathogen but also of a synthetic compound. Considering that the reactivity to R-848 was completely defective also in MyD88-deficient mice, wherein MyD88 is an adapter molecule that plays an essential role to TLR family signalling pathway, it was thought that TLR7 was an essential receptor for recognition of imidazoquinoline group compounds R-848, and imidazoquinoline group compounds R-848 exerted adjuvancy by signalling pathway via TLR7-MyD88. As imidazoquinoline group compounds R-848 has an intensive antiviral action and immunopotentiating effect for immune cells, and used for treating actually human pudendal wart caused by papilloma virus, it was thought that TLR7-deficient mice could be not only a very useful model mice for elucidating the action mechanism of imidazoquinoline group compounds, but also to be a very useful model mice for clinical application of synthetic compound agents such as a therapeutic agent for viral infectious diseases, with TLR7 as a target. The present invention has been completed based on the above-mentioned knowledge.
Disclosure of the Invention

[0012] The present invention relates to a homozygous Toll-like receptor (TLR) 7-knock out mouse, characterized in that the 1.8kb fragment at the 5' end of the TLR7 exon, encoding part of the leucine-rich repeat region of TLR7, is replaced with a neomycin resistance gene cassette ("1"); a homozygous Toll-like receptor (TLR) 7-knock out mouse according to "1", wherein the mouse is unresponsive to an imidazoquinoline compound ("2"); a homozygous Toll-like receptor (TLR) 7-knock out mouse according to "2", wherein the imidazoquinoline compound is resiquimod ("3"); and a homozygous Toll-like receptor (TLR) 7-knock out mouse according to any one of "1" to "3", wherein the neomycin resistance gene is derived from pMC1neo ("4").

[0013] The present invention also relates to a method for obtaining a homozygous Toll-like receptor (TLR) 7-knock out mouse according to any one of "1" to "4", comprising the steps of:

(a) constructing a targeting vector by replacing the 1.8kb fragment of the 5' end of the TLR7 exon, encoding part of the leucine-rich repeat region of TLR7, with a neomycin resistance gene cassette;
(b) introducing the linearized targeting vector into a embryonic stem cell;
(c) microinjecting the targeted embryonic stem cell comprising a mutated TLR7 allele into a blastocyst of the mouse to generate a chimeric mouse;
(d) intercrossing the chimeric mouse with a wild-type mouse to generate a mouse comprising a heterozygous deletion of the TLR7 gene; and
(e) intercrossing the heterozygous mice to generate a mouse comprising a homozygous deletion of the TLR7 gene ("5"); the method according to "5", wherein the targeting vector further comprises a negative selection marker in its 3' terminal region ("6"); the method according to "6", wherein the negative selection marker is a diphtheria toxin A fragment (DT-A) gene or a herpes simplex virus thymidine kinase (HSV-tk) gene ("7").

[0014] The present invention also provides a screening method for identifying an immunopotentiating synthetic compound, wherein an immunopotentiating effect in immune cells derived from the homozygous Toll-like receptor (TLR) 7-knockout mouse according to any one of "1" to "4" is measured/evaluated with the use of said immune cells derived from said knockout mouse, and a test synthetic compound ("8"); and the screening method according to "8", which comprises a step of comparing the immunopotentiating effect in immune cells derived from the homozygous Toll-like receptor (TLR) 7-knockout mouse according to any one of "1" to "4", with that in immune cells derived from a wild-type mouse ("10").

[0015] The present invention also provides a screening method for identifying an immunopotentiating synthetic compound, wherein the immunopotentiating effect in the homozygous Toll-like receptor (TLR) 7-knockout mouse according to any one of "1" to "4" is measured/evaluated with the use of said knockout mouse and a test synthetic compound ("9"); and the screening method according to "9", which comprises a step of comparing the immunopotentiating effect in the homozygous Toll-like receptor (TLR) 7-knockout mouse according to any one of "1" to "4", with that in a wild-type mouse ("11"); and the screening method according to any one of "8" to "11", wherein imiquimod or resiquimod is a lead compound ("12").

Brief Description of Drawings

[0016] Fig. 1 is a figure that shows gene maps indicating mouse TLR7 gene locus, targeting vector and mutated allele by mutation.
Fig. 2 is a photograph that shows the result of Southern blotting analysis of TLR7 knockout mouse of the present invention.
Fig. 3 are photographs that show the result by Northern blotting analysis of TLR7 knockout mouse of the present invention.
Fig. 4 is a figure that shows the result of generation of IFN-α, TNF-α and IL-12 by administrating orally imiquimod and R-848 in BALB/c mouse (wild-type).
Fig. 5 is a figure that shows the result of generation of TNF-α and IL-12 by stimulation of imiquimod and R-848 in the presence or absence of IFN-γ in peritoneal macrophages derived from wild-type mice and MyD88 knockout mice.
Fig. 6 is a figure that shows the result of generation of IL-12p40 by various kinds of ODN stimulation in peritoneal macrophages derived from TLR7 knockout mice and TLR9 knockout mice of the present invention.
Fig. 7 is a figure that shows the result of generation of TNF-α by stimulation of chemical complex R-848, LPS, and ODN1668 in the presence or absence of IFN-γ in peritoneal macrophages derived from TLR7 knockout mice of the present invention.
Fig. 8 is a figure that shows the result of generation of IL-12p40 by stimulation of chemical complex R-848, LPS, and ODN1688 in the presence or absence of IFN-γ in peritoneal macrophages from TLR7 knockout mice of the present invention.

Fig. 9 is a figure that shows the result of generation of TNF-α, IL-6 and IL-12p40 by stimulation of R-848, LPS in the presence or absence of IFN-γ in peritoneal macrophages derived from TLR7 knockout mice of the present invention.

Fig. 10 is a figure that shows the result of the amount of [3H] thymidine incorporation by stimulation of R-848 and CpGDNA in dendritic cells from born marrow derived from TLR7 knockout mice of the present invention.

Fig. 11 is a figure that shows the result of the expression on the cell surface of CD40, CD80 and CD86 molecules by stimulation of R-848 and CpGDNA in dendritic cells from TLR7 knockout mice of the present invention.

Fig. 12 are photographs that show the result of the activation of NF-κB by stimulation of R-848 and CpGDNA stimulation in macrophages derived from TLR7 knockout mice of the present invention.

Fig. 13 are photographs that show the result of the transactivation of NF-κB reporter genes induced by R-848 in macrophages derived from TLR7 knockout mice of the present invention.

Fig. 14 is a figure that shows the result of the response in vivo to R-848 in TLR7 knockout mice of the present invention.

**Best Mode for Carrying Out the Invention**

[0017] As for the TLR7 knockout mouse of the present invention, its reactivity to stimulation by immunopotentiating synthetic compound of the living body, or the reactivity of the cells, tissues or organs constituting the living body is specifically decreased or deleted compared to a wild-type mouse. Stimulation by an immunopotentiating synthetic compound may either be in vivo, where an immunopotentiating synthetic compound is to be administered to a living body, or in vitro wherein cells separated from a living body are contacted with an immunopotentiating synthetic compound.

[0018] The present invention also provides a method for generating a TLR7 knockout mouse unresponsive to an immunopotentiating synthetic compound. With a gene fragment obtained from a mouse gene library by PCR method or the like, a gene encoding TLR7 is screened, and the screened gene which encodes TLR7 is subcloned with the use of a viral vector or the like and identified by DNA sequencing. A targeting vector is constructed by replacing the 1.8 kb fragment at the 5’ end of this gene encoding TLR7 with a pMC1neo gene cassette or the like, and introducing a gene such as a diphtheria toxin A fragment (DT-A) gene or a herpes simplex virus thymidine kinase gene (HSV-tk) into 3’-terminal side.

[0019] This constructed targeting vector is linearized and introduced into ES cells by a method such as electroporation for homologous recombination, and then selecting the ES cells which are homologously recombined by an antibiotic such as G418 or ganciclovir (GANC) among those recombinants. It is preferable to confirm by Southern blotting analysis etc. whether this selected ES cells are the targeted recombinants. The clone of the confirmed ES cell is microinjected into blastocyst of mouse, and the blastocyst is transplanted into a uterus of a recipient mouse to generate a chimeric mouse. A heterozygous mouse (F1 mouse: +/-) can be obtained by intercrossing this chimeric mouse with a wild-type mouse. In addition, as a confirming method for whether TLR7 is produced in a TLR7 knockout mouse, an examining method by Northern blotting or the like by isolating RNA from the mouse obtained by the above-mentioned method, or an examining method for the expression of TLR7 in this mouse by Western blotting or the like can be exemplified.

[0020] Moreover, it can be confirmed that generated TLR7 knockout mouse was unresponsive to an immunopotentiating synthetic compound, for example by contacting an immunopotentiating synthetic compound with immune cells such as macrophages, mononuclear cells, or dendritic cells of TLR7 knockout mouse in vitro or in vivo, to measure the production level of TNF-α, IL-6, IL-12, IFN-γ and the like in the cells, the proliferative response of splenic B cells; expression level of antigens such as CD40, CD80, CD86, and MHC class II on the surface of splenic B cells, and activation of a molecule in signal transduction pathway of TLR7 such as NF-κB, JNK, and IRAK. Thus, the TLR7 knockout mouse of the present invention can be a useful model on elucidating action mechanism of an immunopotentiating synthetic compound, and devising a treatment strategy for viral infection.

[0021] Meanwhile, homozygotic mice born at the expected Mendelian ratios include TLR7-deficient types recognizing an immunopotentiating synthetic compound and the wild-types of their littermates. Precise comparative experiments can be conducted at an individual level by using the deficient types and their littermates of wild-types in these homozygotic mice simultaneously, therefore it is preferable to combine a wild-type mouse and its littermate, for example in screening substances for inhibiting or promoting a response to an immunopotentiating synthetic compound, as described below.

[0022] The TLR7 knockout mouse of the present invention, and immune cells such as macrophages, splenic cells and dendritic cells derived from the TLR7 knockout mouse, can be used for screening various kinds of viral infectious diseases by various kinds of viral infection such as herpesvirus, cytomegalovirus, and human papillomavirus, or for screening substances for inhibiting or promoting a response to an immunopotentiating synthetic compound such as an agonist or an antagonist to TLR7, besides for elucidating action mechanism of an immunopotentiating synthetic com-
pound. The screening method for substances for inhibiting or promoting a response to viral infectious diseases such as pudendal wart, or substances for inhibiting or promoting a response to an immunopotentiating synthetic compound such as an agonist or an antagonist to TLR7 will be described with examples in the following.

[0023] The TLR7 knockout mouse may be used in a screening method for substances for inhibiting or promoting a response to an immunopotentiating synthetic compound. It may also be used in a method for measuring/evaluating the response to an immunopotentiating synthetic compound using immune cells such as macrophages, splenic cells, and dendritic cells derived from the TLR 7 knockout mouse.

[0024] The screening method using immune cells derived from the TLR7 knockout mouse may comprise the following steps:

(a) contacting immune cells obtained from the TLR7 knockout mouse with a test substance in vitro;
(b) culturing the immune cells in the presence of an immunopotentiating synthetic compound; and
(c) measuring/evaluating the response to an immunopotentiating synthetic compound in the immune cells.

[0025] Alternatively the method may comprise the following steps:

(a) contacting immune cells obtained from the TLR7 knockout mouse with an immunopotentiating synthetic compound in vitro;
(b) culturing the immune cells in the presence of the test substance; and
(c) measuring/evaluating the response to an immunopotentiating synthetic compound in the immune cells.

[0026] Alternatively the method may comprise the following steps:

(a) administering the test substance to the TLR7 knockout mouse;
(b) culturing the immune cells obtained from the non-human animal in the presence of an immunopotentiating synthetic compound; and
(c) measuring/evaluating the response to the immunopotentiating synthetic compound in the immune cells.

[0027] Alternatively the method may comprise the following steps:

(a) administering the test substance to the TLR7 knockout mouse of the present invention;
(b) administering the immunopotentiating synthetic compound to the knockout mouse; and
(c) measuring/evaluating the response to the immunopotentiating synthetic compound in the immune cells obtained from the knockout mouse.

[0028] Alternatively the method may comprise the following steps:

(a) administering the immunopotentiating synthetic compound to the TLR7 knockout mouse of the present invention;
(b) culturing the immune cells obtained from the knockout mouse in the presence of the test substance; and
(c) measuring/evaluating the response to the immunopotentiating synthetic compound in the immune cells.

[0029] Alternatively the method may comprise the following steps:

(a) administering the immunopotentiating synthetic compound to the TLR7 knockout mouse of the present invention;
(b) administering the test substance to the knockout mouse; and
(c) measuring/evaluating the response to the immunopotentiating synthetic compound in the immune cells obtained from the knockout mouse.

[0030] Alternatively, the method may comprise the following steps:

(a) administring the test substance to the TLR7 knockout mouse;
(b) administring the immunopotentiating synthetic compound to the knockout mouse; and
(c) measuring/evaluating the response to the immunopotentiating synthetic compound in the knockout mouse.

[0031] Alternatively the method may comprise the following steps:

(a) infecting the TLR7 knockout mouse with the immunopotentiating synthetic compound;
(b) administring the test substance to the knockout mouse; and
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(c) measuring/evaluating the response to the immunopotentiating synthetic compound in the knockout mouse.

[0032] The measurement/evaluation of the response to the immunopotentiating synthetic compound in the present invention relates to a measurement/evaluation of the function of a specific response to the immunopotentiating synthetic compound, and signal transduction into the cells. As for a function of signal transduction, a function for producing cytokine such as TNF-α, IL-6, IL-12 and IFN-γ, a function of generating nitrite ion, a function of proliferating cells, a function of expressing antigen such as CD40, CD80, CD86 and MHC class II on the cell surface, a function of activating molecule in the signal transduction pathway of TLR7 such as NF-κB, JNK and IRAK, and the like can be specifically exemplified but it is not limited to these. Moreover, as mentioned above, in measuring/evaluating the response to the immunopotentiating synthetic compound, it is preferable to measure/evaluate with the measured value of a wild-type mouse, especially that of a littermate as a control, because dispersion by individual difference can be reduced.

[0033] By using the TLR7 knockout mouse of the present invention, or the immune cells derived from the knockout mouse, it is possible to screen for an immunopotentiating synthetic compound.

[0034] The screening method may comprise the following steps:

(a) contacting immune cells such as macrophages, splenic cells and dendritic cells obtained from the TLR7 knockout mouse with the test synthetic compound in vitro; and
(b) measuring/evaluating immunopotentiating effect in the immune cells,

[0035] Alternatively the method may comprise:

(a) administering the test synthetic compound to the knockout mouse; and
(b) measuring/evaluating immunopotentiating effect in the immune cells obtained from the knockout mouse.

[0036] The screening method may comprise the following steps:

(a) administering the test synthetic compound to the knockout mouse; and
(b) measuring/evaluating immunopotentiating effect on the knockout mouse.

[0037] The measurement/evaluation of immunopotentiating effect in the screening method relates to measurement/evaluation of the level of function, for example, a function for producing cytokine such as IFN-α, TNF-α, IL-6, IL-12 and IFN-γ, a function of producing nitrite oxide, a function of proliferating cells, a function of expressing antigen such as CD40, CD80, CD86 and MHC class II on the cell surface, and a function for activating molecules in signalling pathway of TLR7 such as NF-κB, JNK and IRAK, in the immune cells derived from the TLR7 knockout mouse. For instance, by using the method of combinatorial chemistry or the like, using imiquimod or R-848 as a lead compound, when the screening of the above-mentioned immunopotentiating synthetic compound is performed and the behavior in the TLR7 knockout mouse or in the immune cells derived from the knockout mouse is similar to that obtained by using the lead compound, the test synthetic compound has a high possibility to be a candidate substance of the immunopotentiating synthetic compound.

[0038] Further, by using the TLR7 knockout mouse of the present invention and the immune cells derived from the knockout mouse, and a wild-type mouse and the immune cells derived from the wild-type mouse as control, it is possible to screen the immunopotentiating synthetic compound. The method may comprise measuring/comparing the immunopotentiating effect in a TLR7 knockout mouse of the present invention and that from a wild-type mouse. The method may comprise measuring/comparing the immunopotentiating effect in the immune cells derived from the TLR7 knockout mouse of the present invention, and the immune cells derived from the wild-type mouse as a control.

[0039] The screening method may comprise:

(a) contacting the immune cells such as macrophages, splenic cells and dendritic cells obtained from the knockout mouse and the wild-type mouse with the test synthetic compound in vitro; and
(b) measuring/comparing and evaluating the immunopotentiating effect in the immune cells.

[0040] Alternatively the method may comprise:

(a) administering the test synthetic compound to the knockout mouse and the wild-type mouse; and
(b) measuring/comparing and evaluating the immunopotentiating effect in the immune cells obtained from the knockout mouse and the wild-type mouse.

[0041] The method may comprise:
(a) administering the test synthetic compound to the knockout mouse and the wild-type mouse; and
(b) measuring/comparing and evaluating the immunopotentiating effect in the knockout mouse and the wild-type mouse.

[0042] The measuring/comparing and evaluating of immunopotentiating effect in the screening method for immunopotentiating synthetic compound relates to measure the level of for example, a function for producing cytokine such as IFN-α, TNF-α, IL-6, IL-12 and IFN-γ, a function for producing nitrite ion, a function for proliferating cells, a function of expressing antigen such as CD40, CD80, CD86 and MHC class II on the cell surface, and a function for activating molecules in signalling pathway of TLR7 such as NF-κB, JNK and IRAK, in TLR7 knockout mouse or the immune cells derived from the knockout mouse and the wild-type mouse or the immune cells derived from the wild-type mouse as a control, and to compare/evaluate the level of the function. For instance, the test synthetic compound which induces the generation of cytokine in the wild-type mouse, while does not induce the generation of cytokine in the knockout mouse, has a high possibility to be a candidate substance for the immunopotentiating synthetic compound. Further, by using the method of combinatorial chemistry or the like, using imiquimod or R-848 as a lead compound, when the screening of the aforementioned immunopotentiating synthetic compound results in the similarity between the behavior in the knockout mouse or the immune cells derived from the knockout mouse, and that in the immune cells derived from the wild-type mouse or the wild type mouse as a control, the test synthetic compound has a high possibility to be a candidate substance for the immunopotentiating synthetic compound.

[0043] As it was revealed that TLR7 is involved in the recognition of the immunopotentiating synthetic compound, by the TLR7 knockout mouse of the present invention, it is thought that the knockout mouse will become an extremely useful animal model in devising a treatment strategy for various kinds of viral infectious diseases by various kinds of viral infections such as herpesvirus, cytomegalovirus, and human papillomavirus, as well as in elucidating the action mechanism of the immunopotentiating synthetic compound. Moreover, there is a possibility for an agonist and an antagonist of TLR7 to be substances for inhibiting or promoting the aforementioned various kinds of viral infectious diseases, and useful substances for diagnose/treatment for diseases and the like caused by deletion or abnormality of TLR7 activity.

[0044] The present invention will be explained more specifically with examples and reference example in the following.

[0045] The followings were used as reagents and the like in the examples.

[0046] DNA derived from Escherichia coli and DNA derived from Micrococcus deikticus were purchased from Sigma, and further purified with phenol/chloroform extract. LPS derived from Salmonella minnesota Re-595 was purchased from Sigma. Mycoplasma polypeptide MALP-2 used was provided by Dr. Peter F. Muhlradt (Institute of Immunology, Philipps University, Germany) was used. All of the used ODNs (oligodeoxynucleotide) were purchased from Hokkaido System Science Co., Ltd. In addition, the sequence and backbone of CpGDNA are as follows:

**Phosphorothioate 1688:** tccatgacgtt cctgatgct (Seq. ID No. 1)
**D19:** ggTGCATCGAT GCAagggggg (Seq. ID No. 2)
**ACC-30:** ACCGATAACGTT GCCGGTGACGGCACCACG (Seq. ID No. 3)
**Phosphodiester 1688:** TTCATGACGTT CCTGATGCT (Seq. ID No. 4)

[0047] The underlined portions show typical CpG motifs. Capital and lower-case letters indicate the phosphodieter backbone (natural backbone) and phosphothioate modification, respectively.

Reference Example 1 (Generation of MyD 88 knockout mice)

[0048] MyD 88 gene was screened from 129/SvJ mouse gene library (Stratagene), subcloned in pBluescript vector (Stratagene), and identified by restriction mapping and DNA sequencing. Targeting vector was contructed by replacing 1.0-kb gene fragment with a neomycin resistance gene from pMC1neo (Stratagene). The replaced gene fragment included two exons encoding domain similar to the cytoplasmic domain of IL-1RacP (receptor accessory protein). The neomycin resistance gene comprised a 1.1-kb 5' gene fragment as flanking sequences. Next, HSV-tk cassette was introduced to 3' terminal of the gene fragment. ES cell E 14.1 was transfected with linealized marker vector, and selected with G418 and ganciclovir. The targeted ES clone comprising mutated MyD88 alleles was microinjected into the blastocyst of C57BL/6 mice. The obtained chimeric mice were crossed with C57BL/6 female mice in order to generate heterozygote mice. The heterozygote mice were intercrossed to obtain homozygotic mice, and thus MyD88 deficient mice were generated. MyD88 knockout mice grew healthy, not showing abnormality by the age of 20 weeks.

Reference Example 2 (Generation of TLR9 knockout mice)

[0049] TLR9 genome DNA was isolated from 129/SvJ mouse gene library (Stratagene) with a probe derived from
TLR7 genome DNA was screened with a probe (Seq. ID No. 5) derived from human genome DNA sequence (reg. AC003046) comprising a human TLR7 gene from 129/SvJ mouse gene library (Stratagene), subcloned in pBluescript II SK (+) vector (Stratagene), and identified by restriction mapping and DNA sequencing. Targeting vector was constructed by replacing 1.8-kb TLR7 gene fragment encoding a part of leucine-rich repeat with a neomycin resistance gene cassette (pMC1-neo; Stratagene), and inserting herpes simplex thymidine kinase (HSV-TK) as a negative selection marker. The gene loci of mouse TLR7, the targeting vector, and mutated allele by mutation are shown in Fig. 1. In Fig. 1, "I" shows coding exons. The restriction enzymes used were ScaI (Sc) and BamHI (B). This targeting vector was linealized, electroporated into E 14.1 embryonic stem (ES) cells, 340 clones showing resistance to G418 and ganciclovir were selected, and six clones were screened by PCR method and Southern blotting.

Three targeted ES clones comprising mutated TLR7 alleles were microinjected into the blastocyst of C57BL/6 mice to generate chimeric mice. These chimeric male mice were crossed with C57BL/6 female mice to generate heterozygote F1 mice, and homozygotic mice (TLR7 knockout mice: TLR7-/-) were obtained by intercrossing the heterozygote F1 mice. TLR7 knockout mice grew healthy, not showing abnormality by the age of 20 weeks.

Example 1 (Generation of TLR7 knockout mice)

Example 2 (Preparation of peritoneal macrophage)

2 ml of 4% thioglicolic acid medium (DIFCO) was injected into the intraperitoneum of the wild-type mice, TLR7 knockout mice (TLR7KO), MyD88 knockout mice (MyD88KO) and TLR9 knockout mice (TLR9KO), respectively, three days later, peritoneal exudate cells were isolated from the intraperitoneal cavity of each mouse, these cells were cultured in PRMI 1640 medium (GIBCO) added with 10% fetal bovine serum (GIBCO) at 37°C for two hours, nonadherent cells were recovered by washing with Hank’s buffered salt solution (HBBS; GIBCO) at iced temperature, and the adherent cells were used as peritoneal macrophages in following experiment. Total RNA (10 μg) obtained from macrophages derived from the wild-type male mice (+/Y), hemizygous male mice, and hemizygous female mice, respectively. The TLR7 knockout mice of the present invention (TLR7-/-) were generated according to Mendelian rate, not showing significant abnormality by 25th week.

Example 3 (Activating immune cells by imidazoquinoline group compound)

Before examining the reactivity of TLR7 knockout mice to imiquimod and R-848 which are imidazoquinoline group compounds, firstly imiquimod or R-848 was administrated to BALB/c mice (wild-type) orally at the predetermined concentration, respectively, the concentration of IFN-α, TNF-α and IL-12 in serum were measured after two hours. The results are shown in Fig. 4. As a result, the concentration of IFN-α, TNF-α and IL-12 in serum increased depending on applied dose. The data is shown as a mean value +/- S.D. (N=3). Similarly, the ability of these compounds to activate immune cells via TLR was examined using imiquimod or R-848 respectively. The peritoneal macrophages (5 x 10^4 cells)
from wild-type mice and MyD88 knockout mice prepared in example 2, were stimulated by imiquimod or R-848, in the presence or absence of 30 U/ml of IFN-γ, and the concentration of TNF-α and IL-12 produced by peritoneal macrophages were measured by ELISA. The results are shown in Fig. 5. As a result, the macrophages derived from wild-type mice produced cytokines such as TNF-α and IL-12 in large amount, while the macrophages derived from MyD88 knockout mice hardly produced any cytokines, which led to elucidation that the cytokine produced from the peritoneal macrophages responding to imidazoquinoline group compounds, completely depends on MyD88. These results also revealed that imiquimod and R-848 activate immune cells via TLR. The data is shown as a mean value +/- S.D. (N=3).

Example 4 (ODN stimulation in macrophages derived from TLR7KO mice)

Subsequently, the peritoneal macrophages (5 x 10^4 cells) derived from TLR7 knockout mice (TLR7KO) and wild-type mice, were stimulated with imidazoquinoline group compound R-848, LPS and ODN 1688 (phosphothioate 1688) at a predetermined concentration in the presence (+) or absence (-) of 30 U/ml of IFN-γ, and the concentration of TNF-α produced by peritoneal macrophages contained in cultured supernatant was measured by ELISA after 24h of culture. The results are shown in Fig. 7 (in the figure, marks of * show detection limits and below). As shown in Fig. 7, the production of TNF-α by LPS and ODN 1688 stimulation in the peritoneal macrophages from TLR7 knockout mice was confirmed as in normal mice, while induction of production of TNF-α by imidazoquinoline group compound R-848 was not confirmed. Similarly, the concentration of IL-12p40 produced by peritoneal macrophages contained in cultured supernatant after 24h of culture was measured by ELISA. The result are shown in Fig. 8 (in the figure, marks of * show detection limits and below). As shown in Fig. 8, the production of IL-12p40 by LPS and ODN 1688 stimulation in the peritoneal macrophages derived from TLR7 knockout mice were confirmed as in normal mice, while induction for production of IL-12p40 by imidazoquinoline group compound R-848 was not confirmed.

Example 5 (Imidazoquinoline group compounds stimulation in the macrophages derived from TLR7KO mice)

As TLR2 knockout mice, TLR4 knockout mice, TLR6 knockout mice or TLR9 knockout mice which had already been generated by the present inventors, showed reactivity to imidazoquinoline group compounds, it was thought that TLR members except TLR2, TLR4, TLR6 and TLR9 were involved in recognizing the imidazoquinoline group compounds. Therefore, the reactivity of TLR7 knockout mice to R-848 was examined to confirm whether TLR7 recognized imidazoquinoline group compound R-848 specifically. The peritoneal macrophages (5 x 10^4 cells) derived from TLR7 knockout mice (TLR7KO) and wild-type mice were stimulated with R-848 and LPS at a predetermined concentration in the presence (+) or absence (-) of 30 U/ml of IFN-γ, and the concentration of TNF-α, IL-6 and IL-12p40 produced by peritoneal macrophages contained in cultured supernatant was measured by ELISA after 24h of culture. The results are shown in Fig. 9 (in the figure, marks of * show detection limits and below). As shown in Fig. 9, the production of TNF-α, IL-6 and IL-12p40 by LPS stimulation in the peritoneal macrophages from TLR7 knockout mice was confirmed as in the wild-type mice, while induction for production of TNF-α, IL-6 and IL-12p40 was not confirmed by R-848 stimulation.

Example 6 (Stimulation by imidazoquinoline group compounds in the macrophages derived from TLR7KO mice)

As activated phenotypes were proliferated and expressed by stimulating B-lymphocytes with imidazoquinoline group compounds, the proliferation of splenic cells was examined (Cell. Immunol. 203, 55-62, 2000; J. Immunol. 165, 5552-5557, 2000). Splenic cells (5 x 10^4 cells) derived from TLR7 knockout mice (TLR7KO) and wild-type mice were stimulated with R-848 and CpGDNA at a predetermined concentration, cultured for 48h (cultured by adding [3H] thymidine for last eight hours), and measured the level of thymidine incorporation using a β scintillation counter. The results are shown in Fig. 10. A mean value +/- S.D., obtained from representative experiment among triplicate experiments is shown.
As shown in Fig. 10, splenic cells derived from wild-type mice showed proliferation depending on concentration, by R-848 and stimulation. On the contrary, the splenic cells derived from TLR7 knockout mice proliferated depending on concentration by CpGDNA stimulation, while they did not show proliferation by stimulation of R-848.

Example 8 (Imidazoquinoline group compounds stimulation in dendritic cells derived from TLR7 KO mice)

[0059] The reactivity to R-848 in dendritic cells derived from bone marrow (BMDC) derived from TLR7 knockout mice (TLR7KO) and the wild-type mice was examined. The bone marrow cells obtained from TLR7 knockout mice and wild-type mice were cultured for six days in RPMI-1640 medium supplemented with 10% fetal bovine serum and 10 ng/ml of GM-CSF (mouse granulocyte/macrophage colony stimulating factor). The obtained mouse undifferentiated dendritic cells were stimulated with R-848 or CpGDNA for 24h, the cells were stained with biotinylated antibody to CD40, CD80 and CD86, and phycoerythrin (PE)-associated streptavidin was detected, which was analyzed by automated-cell analysing system (FACSCalibur). The results of examination of the expression of the molecules of CD40, CD80 and CD86 onto the surface of cells by flow cytometry are shown in Fig. 11. Similar to the case with CpGDNA, when BMDC derived from wild-type mice was stimulated with R-848, upregulation of CD40, CD80 and CD86 expression was induced. On the contrary, when BMDC derived from TLR7 knockout mice (TLR7KO) was stimulated with R-848, the expression of these molecules on cell surface was not enhanced.

Example 9 (Induction of NF-κB complexes by imidazoquinoline group compounds stimulation.)

[0060] Next, the activation of intracellular signalling cascade to an antiviral immune-modulator was examined. Peritoneal macrophages (2 x 10^6 cells) from TLR7 knockout mice (TLR7KO) and wild-type mice respectively stimulated with thiglycollic acid, were stimulated with 100 nM of R-848 or 1.0 μM of CpGDNA 1668 (CpG) for a certain period of time, nucleoprotein was extracted, the extract obtained with the specific probe including NF-κB associated-site was incubated, electrophoresed, visualized by autoradiography, and NF-κB activation was examined as DNA-associated protein by electrophoretic mobility shift assay (EMSA). The results are shown in Fig. 12. In Fig. 12, an arrow indicates induced NF-κB complexes, and an arrow head indicates a free probe. As shown in Fig. 12, in the macrophages derived from wild-type mice, enhanced DNA avidity of NF-κB transcription, activation of c-Jun N terminal kinase (JNK), and phosphorylation of IRAK were induced when they were stimulated with imidazoquinoline group compound R-848 or CpGDNA. On the contrary, similarly to the macrophages derived from MyD88 knockout mice, all these signal event was completely destructed in macrophages derived from TLR7 knockout mice, as well.

Example 10 (Transactivation of NF-κB reporter gene by stimulation of imidazoquinoline group compounds.)

[0061] The functional role of TLR7 in transactivation of NF-κB reporter gene induced by R-848 was examined. HEK293 cells were transiently cotransfected with TLR7 expression plasmids derived from human, and luciferase activity was measured before and after R-848 stimulation. The peritoneal macrophages derived from TLR7 knockout mice (TLR7KO) and wild-type mice respectively stimulated with thiglycollic acid were stimulated with 100 nm of R-848 or 1.0 μM of phosphothioate modified CpGDNA 1668 (CpG), all cell lysate were prepared, immunoprecipitated with anti-JNK antibody, and JNK activity was examined by In Vitro Kinase Assay using GST-c-Jun fusion protein as a substrate. The results are shown in Fig. 13 (top). The same lysate was blotted with anti-JNK antibody. Moreover, the lysate was immunoprecipitated with anti-IRAK, and kinase activity of IRAK was examined by In Vitro Kinase Assay. The results are shown in Fig. 13 (bottom). The same lysate was blotted with anti-IRAK antibody. As shown in Fig. 13, NF-κB-dependent promoter to R-848 was promoted by transfection with TLR7. It is important to activate TLR7-MyD88-dependent signal pathway for the effect of imidazoquinoline in vivo.

Example 11 (Involvement of TLR7 in vivo)

[0062] It was examined in vivo whether TLR7 was involved in a response to R-848. The wild-type mice, MyD88 knockout mice or TLR7 knockout mice were injected intraperitoneal with 50 nmol of R-848, the sera were harvested after a predetermined period of time, and the concentration of IFN-α, TNF-α and IL12p40 in sera was determined by ELISA. The results are shown in Fig. 14. When wild-type mice were injected with R-848 intraperitoneally, the concentration of IFN-α, TNF-α and IL12p40 in sera increased significantly. On the contrary, the increment of these cytokines was not observed in TLR7 knockout mice. Similarly, MyD88 knockout mice did not show any reactivity to R-848.

Industrial Applicability

[0063] As the TLR7 knockout mouse of the present invention is unresponsive to immunopotentiating synthetic com-
pounds, it can not only be a very useful mouse model to elucidate action mechanism of imidazoquinoline group compounds, but also be a very useful mouse model for applying agents made by synthetic compounds such as therapeutical agents for viral infectious diseases targeting TLR7 to clinical use.

SEQUENCE LISTING

[0064]

<110> JAPAN SCIENCE AND TECHNOLOGY CORPORATION
Akira, Shizuo
Sumitomo Pharmaceuticals Corporation

<120> Model non-human animal unresponsive to synthetic compounds having immunostimulating activity

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<150> JP P2001-358295
<151> 2001-11-22

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Claims

1. A homozygous Toll-like receptor (TLR) 7-knock out mouse, characterized in that the 1.8kb fragment at the 5’ end of the TLR7 exon, encoding part of the leucine-rich repeat region of TLR7, is replaced with a neomycin resistance gene cassette.

2. A homozygous Toll-like receptor (TLR) 7-knock out mouse according to claim 1, wherein the mouse is unresponsive to an imidazoquinoline compound.

3. A homozygous Toll-like receptor (TLR) 7-knock out mouse according to claim 2, wherein the imidazoquinoline compound is resiquimod.

4. A homozygous Toll-like receptor (TLR) 7-knock out mouse according to any one of claims 1 to 3, wherein the neomycin resistance gene is derived from pMC1neo.

5. A method for obtaining a homozygous Toll-like receptor (TLR) 7-knock out mouse according to any one of claims 1 to 4, comprising the steps of:

   (a) constructing a targeting vector by replacing the 1.8kb fragment of the 5’ end of the TLR7 exon, encoding part of the leucine-rich repeat region of TLR7, with a neomycin resistance gene cassette;

   (b) introducing the linearized targeting vector into an embryonic stem cell;
(c) microinjecting the targeted embryonic stem cell comprising a mutated TLR7 allele into a blastocyst of the mouse to generate a chimeric mouse;
(d) intercrossing the chimeric mouse with a wild-type mouse to generate a mouse comprising a heterozygous deletion of the TLR7 gene; and
(e) intercrossing the heterozygous mice to generate a mouse comprising a homozygous deletion of the TLR7 gene.

6. The method according to claim 5, wherein the targeting vector further comprises a negative selection marker in its 3' terminal region.

7. The method according to claim 6, wherein the negative selection marker is a diphtheria toxin A fragment (DT-A) gene or a herpes simplex virus thymidine kinase (HSV-tk) gene.

8. A screening method for identifying an immunopotentiating synthetic compound, wherein an immunopotentiating effect in immune cells derived from the homozygous Toll-like receptor (TLR) 7-knockout mouse according to any one of claims 1 to 4 is measured/evaluated with the use of said immune cells derived from said knockout mouse, and a test synthetic compound.

9. A screening method for identifying an immunopotentiating synthetic compound, wherein the immunopotentiating effect in the homozygous Toll-like receptor (TLR) 7-knockout mouse according to any one of claims 1 to 4 is measured/evaluated with the use of said knockout mouse and a test synthetic compound.

10. The screening method according to claim 8, which comprises a step of comparing the immunopotentiating effect in immune cells derived from the homozygous Toll-like receptor (TLR) 7-knockout mouse according to any one of claims 1 to 4, with that in immune cells derived from a wild-type mouse.

11. The screening method according to claim 9, which comprises a step of comparing the immunopotentiating effect in the homozygous Toll-like receptor (TLR) 7-knockout mouse according to any one of claims 1 to 4, with that in a wild-type mouse.

12. The screening method according to any one of claims 8 to 11, wherein imiquimod or resiquimod is a lead compound.

**Patentansprüche**

1. Homozygote Toll-like-Rezeptor (TLR) 7-Knockout-Maus, **dadurch gekennzeichnet, dass** das 1,8 kb-Fragment am 5'-Ende des TLR7-Exons, welches einen Teil der an Leucin reichen Wiederholungsregion von TLR7 codiert, durch eine Neomycinresistenzgen-Kassette ersetzt ist.

2. Homozygote Toll-like-Rezeptor (TLR) 7-Knockout-Maus nach Anspruch 1, wobei die Maus nicht auf eine Imidazochinolin-Verbindung anspricht.

3. Homozygote Toll-like-Rezeptor (TLR) 7-Knockout-Maus nach Anspruch 2, wobei die Imidazochinolin-Verbindung Resiquimod ist.

4. Homozygote Toll-like-Rezeptor (TLR) 7-Knockout-Maus nach einem der Ansprüche 1 bis 3, wobei das Neomycinresistenzgen von pMC1 neo abgeleitet ist.

5. Verfahren zum Erhalten einer homozygoten Toll-like-Rezeptor (TLR) 7-Knockout-Maus nach einem der Ansprüche 1 bis 4, welches die folgenden Stufen umfasst:

   (a) Konstruieren eines zielenden Vektors durch Ersetzen des 1,8 kb-Fragments des 5'-Endes des TLR7-Exons, welches einen Teil der an Leucin reichen Wiederholungsregion von TLR7 codiert, durch eine Neomycinresistenzgen-Kassette,
   (b) Einbringen des linearisierten zielenden Vektors in eine embryonale Stammzelle,
   (c) Mikroinjizieren der embryonalen Stammzelle, auf die gezielt wurde und welche ein mutiertes TLR7-Allel umfasst, in eine Keimblase der Maus, um eine chimäre Maus zu erzeugen,
   (d) Kreuzen der chimären Maus mit einer Wildtyp-Maus, um eine Maus zu erzeugen, welche eine homozygote
Deletion des TLR7-Gens aufweist, und
(e) Kreuzen der heterozygoten Mäuse, um eine Maus zu erzeugen, welche eine homozygote Deletion des
TLR7-Gens aufweist.

6. Verfahren nach Anspruch 5, wobei der zieliende Vektor weiterhin einen negativen Selektionsmarker in seiner 3'-
terminalen Region umfasst.

7. Verfahren nach Anspruch 6, wobei der negative Selektionsmarker ein Diptherietoxin A-Fragment (DT-A) -Gen oder
ein Herpes simplex-Virus-Thymidinkinase (HSV-tk) -Gen ist.

8. Screeningverfahren zum Identifizieren einer immunpotenzierenden synthetischen Verbindung, wobei eine immu-
npotenzierende Wirkung in Immunzellen, abgeleitet von der homozygoten Toll-like-Rezeptor (TLR) 7-Knockout-Maus
nach einem der Ansprüche 1 bis 4, unter Verwendung von der Knockout-Maus abgeleiteten Immunzellen und
einer synthetischen Testverbindung gemessen/beurteilt wird.

9. Screeningverfahren zum Identifizieren einer immunpotenzierenden synthetischen Verbindung, wobei die immu-
npotenzierende Wirkung in der homozygoten Toll-like-Rezeptor (TLR) 7-Knockout-Maus nach einem der Ans-
sprüche 1 bis 4 unter Verwendung der Knockout-Maus und einer synthetischen Testverbindung gemessen/beurteilt wird.

10. Screeningverfahren nach Anspruch 8, welches eine Stufe umfasst, bei der man die immunpotenzierende Wirkung
in Immunzellen, abgeleitet von der homozygoten Toll-like-Rezeptor (TLR) 7-Knockout-Maus nach einem der An-
sprüche 1 bis 4, mit derjenigen in Immunzellen, abgeleitet von einer Wildtyp-Maus, vergleicht.

11. Screeningverfahren nach Anspruch 9, welches eine Stufe umfasst, bei der man die immunpotenzierende Wirkung
in der homozygoten Toll-like-Rezeptor (TLR) 7-Knockout-Maus nach einem der Ansprüche 1 bis 4 mit derjenigen
in einer Wildtyp-Maus vergleicht.

12. Screeningverfahren nach einem der Ansprüche 8 bis 11, wobei Imiquimod oder Resiquimod eine Leitverbindung ist.

Revendications

1. Souris homozygote knock-out pour le récepteur de type Toll-7 (TLR), caractérisée en ce que le fragment de 1,8
kb à l’extrémité 5’ de l’exon de TLR7, codant pour une partie de la région de répétition riche en leucine de TLR7,
est remplacé par une cassette contenant un gène de résistance à la néomycine.

2. Souris homozygote knock-out pour le récepteur de type Toll-7 (TLR) selon la revendication 1, où la souris est
insensible à un composé imidazoquinoline.

3. Souris homozygote knock-out pour le récepteur de type Toll-7 (TLR) selon la revendication 2, où le composé
imidazoquinoline est le resiquimod.

4. Souris homozygote knock-out pour le récepteur de type Toll-7 (TLR) selon l’une quelconque des revendications 1
to 3, où le gène de résistance à la néomycine est dérivé de pMC1neo.

5. Procédé pour obtenir une souris homozygote knock-out pour le récepteur de type Toll-7 (TLR) selon l’une quelconque
des revendications 1 à 4, comprenant les étapes consistant à :
   (a) construire un vecteur de ciblage en remplaçant le fragment de 1,8 kb de l’extrémité 5’ de l’exon de TLR7,
codant pour une partie de la région de répétition riche en leucine de TLR7, par une cassette contenant un gène
de résistance à la néomycine ;
   (b) introduire le vecteur de ciblage linéarisé dans une cellule souche embryonnaire ;
   (c) micro-injecter la cellule souche embryonnaire ciblée, comprenant un allèle de TLR7 muté, dans un blastocyte
de la souris afin de générer une souris chimérique ;
   (d) croiser la souris chimérique avec une souris de type sauvage afin de générer une souris comprenant une
délétion hétérozygote du gène TLR7 ; et
   (e) croiser les souris hétérozygotes afin de générer une souris comprenant une délétion homozygote du gène
TLR7.
6. Procédé selon la revendication 5, dans lequel le vecteur de ciblage comprend en outre un marqueur de sélection négative dans sa région terminale 3’.

7. Procédé selon la revendication 6, dans lequel le marqueur de sélection négative est un gène du fragment A de la toxine diphtérique (DT-A) ou un gène de la thymidine kinase du virus herpès simplex (HSV-tk).

8. Procédé de criblage pour identifier un composé synthétique d’immunopotentialisation, dans lequel un effet d’immunopotentialisation dans des cellules immunitaires dérivées de la souris homozygote knock-out pour le récepteur de type Toll-7 (TLR) selon l’une quelconque des revendications 1 à 4 est mesuré/évalué en utilisant lesdites cellules immunitaires dérivées de ladite souris knock-out, et un composé synthétique de test.

9. Procédé de criblage pour identifier un composé synthétique d’immunopotentialisation, dans lequel l’effet d’immunopotentialisation chez la souris homozygote knock-out pour le récepteur de type Toll-7 (TLR) selon l’une quelconque des revendications 1 à 4 est mesuré/évalué en utilisant ladite souris knock-out et un composé synthétique de test.

10. Procédé de criblage selon la revendication 8, qui comprend une étape consistant à comparer l’effet d’immunopotentialisation dans des cellules immunitaires issues de la souris homozygote knock-out et un composé de test avec celui observé dans des cellules immunitaires dérivées d’une souris de type sauvage.

11. Procédé de criblage selon la revendication 9, qui comprend une étape consistant à comparer l’effet d’immunopotentialisation chez la souris homozygote knock-out et un composé de test avec celui observé chez une souris de type sauvage.

12. Procédé de criblage selon l’une quelconque des revendications 8 à 11, dans lequel l’imiquimod ou le resiquimod est un composé tête de série.
Figure 1

Wild-type allele

Targeting vector

Mutated allele

6 kbp

5 kbp

2 kbp

B: BamHI
E: EcoRI
Sc: ScaI

P-bluescript II SK (+) vector
Figure 5

![Graph showing TNF-α and IL-12 levels in response to different concentrations of Imiquimod and R-848.](image)
Figure 8

[Graph showing IL-12 p40 levels under different conditions]
Figure 10

[Graph showing the incorporation of [H]thymidine in wild-type and TLR7 KO cells in response to increasing concentrations of R-848 and CpG DNA.]

Figure 11

[Histograms showing the expression levels of CD40, CD80, and CD86 in wild-type and TLR7 KO cells treated with medium, CpG DNA, or R-848.]
REFERENCES CITED IN THE DESCRIPTION

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