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DESCRIPTION

PRODUCTION PROCESS FOR MUTANT

TECHNICAL FIELD

5 The present invention relates to a novel production process for a mutant comprising introducing a improved DNA construct into animal cells, the DNA construct, and a kit for producing a gene mutant or expressing a mutant gene.

10

BACKGROUND ART

Living things now in existence have evolved over a long period of time through the mutagenesis and the selection of mutants by the environment. The general evolution has very slow speed and passes through many generations to advance. On the other hand, in the case of antibody-producing cells of the immune system, the mutagenesis and the selection by antigens are completed in one generation, and the next generation will not inherit the acquired function. Such a quick rate of evolution in the immune system is interpreted as being set against the mutation of an environment-dependent pathogenic microorganism.

25 The present inventor previously produced transgenic mice to which genes of bacteria-derived enzyme

chloramphenicol acetyltransferase are introduced, and demonstrated that the mutation of an antibody gene is controlled by its promoter and enhancer and the mutation of any genes can be induced by controlling with a promoter and enhancer of an antibody gene (Azuma, T., et al., Int. Immunology, 5(2) 121-130(1992)).

The currently used production process for a mutant is one of the mutant-producing methods in which deletion, insertion and/or addition mutation is suitably introduced into a desired DNA sequence, that is, site-specific mutagenesis to site-specifically replace a certain length of DNA sequences (Site-specific mutagenesis, Zoller, et al., Nucleic Acid Res., 10, 6487-6500(1982); Zoller, et al., Methods in Enzymol., 100, 468-500(1983)).

In the general mismatch mutation using an artificially synthesized oligonucleotide as a primer, a complementary oligonucleotide consisting of around 20 bases is synthesized with an optional mutation in a base sequence nearby a site where mutation is to be induced, the oligonucleotide is hybridized to a target DNA, and then a DNA complementary to the remaining of target DNA can be produced using DNA polymerase. Thus it is possible to introduce a desired mutation into a desired site. However, this method cannot induce many mutants at once.

In the other mutagenesis systems, some examines

the disappearance or the manifestation of a specific function in the presence of a compound that damages a gene (Myers, et al., Science, 232, 613-618(1986)), and other uses bacteria etc. However, these methods are
5 fundamentally different from the above one to induce mutation by the present inventor.

Variation (diversity) in the Ig V region of mice and humans is generated by the combinatorial joining of V, D, and J gene segments existing separately in germline,
10 the deletion and addition of nucleotides at the junction of these segments during joining, and somatic hypermutation of joined V-(D)-J genes. The somatic hypermutation is related to the affinity maturation of antibody and has been frequently observed after
15 stimulation of T cell-dependent antigen (TD) (Bothwell, A.L.M., et al., Cell, 24,625(1981): Gearhart, P.J., et al., Nature, 291, 29 (1981): Griffiths, G.M., et al., Nature, 312,271(1984): Maizels, N., et al., Cell, 43, 715(1985): Wysocki, L.T., et al., Proc. Natl. Acad. Sci. USA, 83,
20 1847(1986): Cumano,A., et al., EMBO. J., 5, 2459 (1986): Berek,C., et al., Cell, 67, 1121(1991): Taketani, M., et al., Mol. Immunol., 32, 983 (1995): Furukawa,k., et al., Immunity, 11, 329 (1999): 2-10). The cis-acting elements responsible for the induction of somatic hypermutation
25 have been identified using κ -chain (O'Brien, R.L. et al.,

Nature, 326, 405(1987): Sharpe, M.J., et al., Eur. J. Immunol., 20, 1379(1990): Sharpe, M.J., et al., EMBO J., 10, 2139 (1991): Betz, A.G., et al., Cell, 77, 239(1994): Yelamos, J., et al., Nature, 376, 225(1995): Peters, A., et
5 al., Immunity, 4, 57(1996): 11-16), λ -chain (Klotz, E., et al., J. Immunol., 157, 4458(1996):17) and H-chain transgenic mice (Durdik, J., et al., Proc. Natl. Acad. Sci. USA, 86, 2346(1989): Sohn, J., et al., J. Exp. Med., 177, 493(1993): Tumas-Brundage, K.M. and Manser,
10 T., J. Exp. Med., 185, 239(1997) :18-20).

As stated above, the present inventor prepared transgenic mice carrying chloramphenicol acetyltransferase (CAT) gene that is driven by V_H 17.2.25 (Loh, D. Y., et al., Cell, 33, 85 (1983): Grosschedl, R. and Baltimore, D.,
15 Cell, 41, 885 (1985): 22, 23) and J-C intron enhancer (hereafter abbreviated to Ep) (Gillies, S.D., et al., Cell, 33, 715(1983): Banerji, J., et al., Cell, 33, 729(1983):24, 25)/ matrix attachment region (hereafter abbreviated to MAR) (Forrester, W.C., et al., Science, 265,
20 1221(1994):26). As a result, somatic hypermutation was detected in CAT but not in V_H promoter or Ep/MAR flanking. However, the frequency of mutation was approximately 1/10 that observed in endogenous V_H -D- J_H , suggesting that these cis-acting elements are critical or important for the
25 induction of hypermutation and that other components such

as $C\gamma$, $C\alpha$ of 3' or the enhancer flanking of $C\alpha$ (3' enhancer) (Pettersson, S., et al., Nature, 344,165(1990): Dariavach, P., et al., Eur. J. Immunol., 21, 1499(1991): Lieberson, R., et al., EMBO. J., 14, 6229(1995):27-29) might
5 be responsible for high frequent somatic hypermutation (Sohn, J., et al., J.Exp.Med., 177,493(1993):Tumas-Brundage, K.M. and Manser, T., J. Exp. Med., 185, 239 (1997): Giusti, A. M. and Manser, T., J. Exp. Med., 177, 793 (1997):19,20,30).

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DISCLOSURE OF THE INVENTION

In order to identify the component(s) important in raising the frequency of hypermutation in IgH gene, the present inventor employed a RAG-2-deficient (Recombination
15 Activating Gene: gene rearrangement protein-2 blastocyst complementation system developed by Chen et al. (Chen, J., et al., Proc. Natl. Acad. Sci. USA, 90, 4528(1993):31). A series of transgene constructs differed only in the 3'-flanking region were microinjected into RAG-2-deficient
20 blastocysts to transfect embryonic stem cells (hereafter abbreviated to ES cells). The chimeric mice obtained in this system were immunized with a T cell-dependent antigen.

As a result, the frequency of somatic hypermutation in V_H -D- J_H of the transgene revealed that
25 the insertion of DNaseI sensitive region 3b (hereafter

HS3b) and/or HS4 (Madisen, L. and Groudine, M., Genes Dev.8,2212(1994): Michaelson, J.S. et al., Nucleic Acids Res., 23, 975(1995): Arulampalam, V. et al., Immunol.Today, 18, 549(1997):32-34) induces random somatic hypermutation.

5 An object of the present invention is to provide a production process for a random mutant, where a DNA construct in which a modified eukaryotic promoter, an external DNA sequence, an intron enhancer and a 3' HS3/4 enhancer are linked is introduced into an animal cell.

10 Another object of the present invention is to provide a method for obtaining a mutant gene by expressing the DNA construct in an animal cell. The inventor found a DNA construct for producing such mutant and a kit for producing a gene mutant or expressing the mutant, thus to

15 accomplish the present invention. Further, a mutant expression product obtained from the expression system using animal cells is to mean that it has subjected to the screening for toxicity against animal cells.

 The present invention provides the following

20 Items 1 to 20.

Item 1. A production process for an exogenous gene random mutants, wherein a DNA construct comprising at least the following (a)-(d) is introduced into an animal cell and the exogenous gene mutant is produced in the animal cell:

25 (a) a promoter;

- (b) an exogenous gene;
- (c) an intron enhancer; and
- (d) an enhancer comprising HS3b and/or HS4 in a DNaseI sensitive region.

5

Item 2. The production process according to Item 1, wherein the (d) enhancer further comprises HS1 and HS2.

Item 3. The production process according to Item 1 or 2,
10 wherein the (d) enhancer has a DNA sequence comprising at least a DNA sequence of SEQ ID NO: 1 or 2.

Item 4. The production process according to any one of
Items 1 to 3, wherein DNA sequences of (a), (c) and (d)
15 are derived from a DNA sequence of an immunoglobulin

Item 5. The production process according to Item 2, wherein the (d) enhancer comprises HS1, HS2, HS3b and HS4.

20 Item 6. The production process according to any one of Items 1 to 5, wherein
the (a) promoter is a V_H promoter; and
the (d) enhancer has a DNA sequence comprising DNA sequences of SEQ ID NOs: 1 and 2.

25

Item 7. The production process according to Item 5 or 6, wherein the DNA construct is pvehc3'EHS3b/4.

Item 8. The production process according to any one of
5 Items 1 to 7, wherein the animal cell is a B cell line animal cell.

Item 9. The production process according to Item 8, wherein the B cell line animal cell is derived from a pre-
10 B lymphocyte line cell.

Item 10. A method for obtaining an expression product of a mutant gene by expressing the exogenous gene mutant obtained by the method according to any one of Items 1 to
15 9 in the animal cell.

Item 11. A DNA construct for producing an exogenous gene mutant, comprising at least:

- (a) a promoter;
- 20 (b) an exogenous gene;
- (c) an intron enhancer; and
- (d) an enhancer comprising HS3b and/or HS4 in a DNaseI sensitive region.

25 Item 12. The DNA construct according to Item 11, wherein

the (d) enhancer further comprises HS1 and HS2.

Item 13. The DNA construct according to Item 11 or 12,
wherein the (d) enhancer has a DNA sequence comprising at
5 least a DNA sequence of SEQ ID NO: 1 or 2.

Item 14. The DNA construct according to any one of Items
11 to 13, wherein DNA sequences of (a), (c) and (d) are
derived from a DNA sequence of immunoglobulin.

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Item 15. The DNA construct according to any one of Items
11 to 14, wherein the (d) enhancer comprises HS1, HS2,
HS3b and HS4.

15 Item 16. The DNA construct according to any one of Items
11 to 15, wherein
the (a) promoter is a V_H promoter; and
the (d) enhancer has a DNA sequence comprising DNA
sequences of SEQ ID NOs: 1 and 2.

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Item 17. The DNA construct according to Item 11 or 12,
wherein the DNA construct is pvehc3'EHS3b/4.

Item 18. A kit for production process of an exogenous
25 gene mutant or producing an expression product of a mutant

gene, comprising a vector having:

- (a) a promoter;
- (b) an exogenous gene-insertion site;
- (c) an intron enhancer; and

- 5 (d) an enhancer comprising HS3b and/or HS4 in a DNaseI sensitive region;

for achieving an expression of a mutated DNA sequence which produces an exogenous gene mutant.

- 10 Item 19. The kit according to Item 18, wherein the (d) enhancer further comprises HS1 and HS2.

Item 20. V_H 17.2.25 promoter having a sequence of SEQ ID NO: 14.

- 15 According to the present invention, an exogenous gene is introduced into the system controlled by the promoter and enhancer of an antibody gene, and therefore a gene cluster having frequent and random mutations can be obtained and a production process for an animal cell line
- 20 mutant in which a mutation is widely introduced into a peptide or protein is provided, thus it is possible to try to improve and change a function of naturally occurring protein. For example, it becomes possible to develop new type of antibiotics having an improved function against
- 25 antibiotic resistant bacteria, agricultural chemicals, or

improved peptide pharmaceuticals having a better function such as interferon and growth hormone. Moreover, by using animal cells, it is possible to carry out a toxicity screening on animal cells simultaneously, and therefore
5 the primary toxicity can be examined and the quick creation of various peptide products having a functionally improved action is attained.

Hereafter in this specification, the representation of amino acids, peptides, base sequences,
10 nucleic acids and the like by abbreviation shall follow the provision of IUPAC-IUB (IUPAC-IUB Communication on Biological Nomenclature, Eur. J. Biochem., 138: 9 (1984)), "Guideline for the preparation of specification and others containing a base sequence or an amino acid sequence"
15 (edited by the Japanese Patent Office) and the conventional symbols in the art.

The synthesis of DNA, the construction of DNA construct containing an exogenous gene (e.g., an expression vector), the preparation method of a host cell
20 transformed by the DNA construct and an expressed protein secreted by a host cell, etc. can be easily prepared or obtained in accordance with general genetic engineering techniques (Molecular Cloning 2d Ed, Cold Spring Harbor Lab. Press (1989); Continuation of Biochemical Experiment
25 Course, Genetic Research Method I, II and III, Japanese

Biochemical Society (1986)) or gene recombination techniques (refer to e.g., Science, 224, 1431 (1984); Biochem. Biophys. Res. Comm., 130, 692 (1985); Proc. Natl. Acad. Sci. USA., 80, 5990 (1983)).

5 In the present invention, a mutant refers to a gene comprising a random mutated DNA sequence of an exogenous gene or an expression product having a mutated amino acid sequence encoded by a random mutated gene.

 The mutant production in an animal cell
10 according to the invention can be achieved by preparing a DNA construct provided by the invention for inducing mutant (recombinant DNA) that can be expressed in a host cell containing a gene encoding a desired protein, and introducing the construct into the cell.

15 In the specification "introduce a DNA construct into an animal cell" means to incorporate an exogenous gene into a genome of animal cell.

 The expression product of mutant is obtained by cultivating a transformant containing the above mutant of
20 exogenous gene and then collecting the product from the resulting culture.

 The DNA construct for inducing mutant provided by the present invention has a structure comprising at least (a) a promoter, (b) an exogenous gene, (c) an intron
25 enhancer, and (d) an enhancer containing HS1, HS2 and HS3b

and/or HS4 in a DNaseI sensitive region and can be easily produced or obtained by general gene engineering technique.

For the preparation of each gene (DNA) in the invention, the gene may be derived from a suitable origin
5 or chemically synthesized.

Specifically, the DNA synthesis may be chemical synthesis based on the phosphoramidite method or the triester method, or may be carried out on the commercially available automatic oligonucleotide synthesizer. A double
10 strand fragment can also be prepared from a chemically synthesized single strand product by: synthesizing complementary strands and annealing these strands together under suitable conditions; or using a DNA polymerase together with a suitable primer sequence to add a
15 complementary strand.

In addition, each gene can be obtained by: either DNA synthesis as described above; or preparing a cDNA library in accordance with an ordinary method from a vector containing the gene or from a suitable origin in
20 which the gene is expressed, and selecting a desired clone using suitable probe or antibody specific to the gene (Proc. Natl. Acad. Sci. USA, 78, 6613 (1981); Science, 222, 778 (1983) etc).

Illustrated as the origin of cDNA in the above
25 are various kinds of cells and tissues where each gene is

expressed, cultured cells derived therefrom and the like. In addition, separation of total RNA from these origins, separation and purification of mRNA, preparation and cloning of cDNA, etc. can be all performed in accordance
5 with an ordinary method. Commercially available cDNA libraries such as cDNA library from Clontech Lab. Inc. can be used in this invention. A method of screening the gene from cDNA library is not limited and can be performed in accordance with an ordinary method.

10 Specifically, a method of immunoscreening the proteins produced by cDNA using specific antibodies against the proteins and selecting the corresponding cDNA clone, plaque hybridization and colony hybridization with the use of a probe selectively combined with a target DNA
15 sequence, a combination of these methods, etc. can be used.

As the probe used herein, a DNA chemically synthesized based on the information of the base sequence of each gene can be generally used, but a gene already obtained according to the invention or a fragment thereof
20 can also be used satisfactorily. Sense primer or antisense primer designed based on the base sequence information of exogenous gene can be used as a probe for screening.

A partial nucleotide sequence corresponding to
25 the DNA sequence of each gene is used as the above-

mentioned probe. The nucleotide sequence may contain at least 15 serial bases, preferably 20 serial bases, more preferably 30 serial bases, and most preferably 50 serial bases. Alternatively, a positive clone containing said
5 sequence itself may be used as a probe.

The DNA/RNA amplification by the PCR method (Science, 230, 1350 (1985)) can be preferably employed for obtaining a gene. Especially, when it is difficult to obtain full length of cDNA from the library, RACE method
10 (Rapid amplification of cDNA ends; Journal of Experimental Medicine, 12(6), 35 (1994)), especially 5'-RACE method (M.A. Frohman et al., Proc. Natl. Acad. Sci. USA., 8, 8998 (1988)), etc. can be preferably adopted.

When the PCR method is adopted, the primer used
15 may be designed based on the gene sequence information and synthesized in accordance with an ordinary method. The amplified DNA/RNA fragment may be separated and purified in accordance with an ordinary method as described above, for example gel electrophoresis.

20 The sequences of the genes or various kinds of DNA fragments obtained above may be determined in accordance with an ordinary method, for example the dideoxy procedure (Proc. Natl. Acad. Sci. USA., 74, 5463 (1977)) and the Maxam-Gilbert Method (Methods in
25 Enzymology, 65, 499 (1980)), or more easily using a

commercial sequencing kit and the like.

(b) Desired exogenous gene

In the constituents of the above-described DNA
5 construct, the DNA used as (b) exogenous gene may be
either an exogenous DNA sequence derived from a desired
gene or an expressed protein which is a basis for inducing
a mutant. Or the DNA used as exogenous gene may be
synthesized. The length of the exogenous gene used for
10 the present production process for a mutant may be
generally 4Kb or less, preferably 3Kb or less, more
preferably around 2Kb, but an exogenous DNA sequence
having the length more than 4Kb can be also used.

The exogenous gene is not especially limited
15 insofar as it exhibits a biological activity when
expressed. For example, growth hormone (GH), insulin,
interferon, erythropoietin, etc. can be illustrated.

For the easier checking of the insertion of
exogenous gene into a genome of host cell, the DNA
20 sequence of a gene to be used as a so-called marker may be
linked to the DNA sequence of the above (b) exogenous gene.

By using a part or the entire base sequences of
thus obtained exogenous gene, the expression of exogenous
gene and its mutant in an individual or various tissues
25 can be detected with specificity.

Such detection can be carried out in accordance with an ordinary method, for example, RNA amplification by RT-PCR (Reverse transcribed-Polymerase chain reaction; E.S. Kawasaki, et al., Amplification of RNA. In PCR Protocol, A
5 Guide to methods and applications, Academic Press, Inc., San Diego, 21-27 (1991)), Northern Blotting analysis (Molecular Cloning, Cold Spring Harbor Lab. (1989)), detection at cell-level using *in situ* RT-PCR (Nucl. Acids Res., 21, 3159-3166 (1993)) and *in situ* hybridization,
10 NASBA method (Nucleic acid sequence-based amplification, Nature, 350, 91-92 (1991)) and the other various method. The detection by RT-PCR is preferably illustrated.

The DNA fragment of the exogenous gene is provided in the form of a restriction fragment cleaved
15 with a suitable restriction enzyme, that is (b) exogenous gene of the DNA construct.

(a) Promoter

Examples of (a) promoter used as a constituent
20 of the above-mentioned DNA construct include an immunoglobulin heavy chain variable region promoter (V_H promoter), a promoter derived from SV40, a retrovirus promoter, an adenovirus promoter, a cytomegalovirus promoter, a promoter derived from viruses such as polyoma
25 virus or hepatitis B virus, a metallothionein promoter, a

heat shock promoter, a SR α promoter, other promoters effective for the mammalian cell, etc. V_H promoter, specifically V_H17.2.25 promoter (Gillies, S.D., et al., Cell, 33,715(1983); Banerji, J., et al., Cell, 33, 5 729(1983)) are preferably used in this invention.

The above promoters may include, for example, a promoter that can induce a neomycin resistance gene or a tetramycin resistance gene.

The above promoters may be in the linked form of
10 two or more.

These various DNA sequences of the above promotera are known and the information thereof is available.

15 (c) Intron enhancer

An intron enhancer is a DNA sequence that can adjust the transcription and contains the sequence of unexpressed region intervening between the exons of heavy chain JH gene and C μ gene. Heavy chain enhancer (C μ enhancer) and kappa chain enhancer (κ enhancer) of
20 immunoglobulin can be illustrated as (c) intron enhancer.

(d) Deoxyribonuclease I (DNaseI) sensitive region

A DNaseI sensitive region is a region that
25 undergoes fragmentation when the nucleus of an isolated

cell is treated with a low concentration of DNaseI. In the case of immunoglobulin heavy chain gene, HS1, HS2, HS3b and HS4 are known. HS3b/4 is both or either of HS3b and HS4. This invention is characterized in that HS3b
5 region and/or HS4 region are/is contained. In a preferable embodiment, HS1 and HS2 (these are referred to as 3'E) in addition to HS3b and/or HS4 are included. Illustrated as a 3'EHS3b/4 is a sequence without intron sequences, comprising XbaI fragment (4.0kb) from 3'
10 enhancer disclosed by Lieberon, R. et al. (EMBO. J., 14, 6229(1995)) and HS3 (1182bp) and HS4 (1381bp) from MP11 disclosed by Madisen, L. et al. (Genes Dev., 8, 2212 (1994)).

15 Connection of each gene (preparation of DNA construct)

The DNA construct of the invention can be prepared for example by inserting each of the above-described genes into a suitable expression vector in accordance with an ordinary method, for example by
20 inserting each element into a vector having suitable antibiotic resistance gene. Specifically, the DNA construct can be constructed by inserting the promoter, the exogenous gene, the intron enhancer, the DNaseI sensitive region (including DNA sequences of HS3b and/or
25 HS4 and if necessary further including an enhancer

comprising 3'E), which are cut out from a vector clone, a genomic DNA or a plasmid by an optional restriction enzyme or synthesized, into a vector used for animal cell line having an optional antibiotic resistance gene.

5 Specifically, the insertion can be carried out in accordance with an ordinary method using a restriction enzyme, ligase etc.

 In the desired recombinant expression vector, an intron enhancer and a 3'EHS3b/4 enhancer may be
10 constructed in the same direction from 5' end toward 3' end, in the same direction from 3' end toward 5' end, or in the opposite directions from 5' end toward 3' end and from 3' end toward 5' end.

 In the above DNA construct, the interval between
15 the promoter and the 3'EHS3b/4 enhancer intervening an exogenous DNA may be 0.1Kb to 100Kb, preferably 1Kb to 10Kb, more preferably 2Kb to 5Kb.

 An example of preferable expression vector is retrovirus vector. More specifically, pIND (Invitrogen),
20 pcDNA3.1/His (Invitrogen), pEGFP-N, pEGFP-C (Clontech), etc. can be illustrated.

 An example of the DNA construct containing (a) promoter and (c) intron enhancer is an expression vector containing V_H promoter and intron enhancer according to
25 the present inventor and others (Azuma, T. et al., Int.

Immunology, 5(2) 121-130 (1993)). It is preferably used for preparing the DNA construct.

In the above expression vector containing a promoter and an intron enhancer, the promoter is positioned 5' side upstream of the exogenous gene ordinary induced, and the exogenous gene, an intron enhancer follow in this order.

When the above vector for preparing the present DNA construct containing an exogenous gene is modified to contain a promoter, a cloning site for the insertion of desired exogenous gene, one or more enhancer necessary for inducing mutation (especially immunoglobulin enhancer as well as intron enhancer) and DNaseI sensitive region, it is effective as a DNA construct in a production process for an exogenous gene mutant or in a kit for producing an expression product of mutant gene.

The present DNA construct may be a circular DNA or a linear DNA.

The checking of whether the present DNA construct is attained can be conducted by an ordinary method, for example, by cleaving the construct with a restriction enzyme and subjecting it to the electrophoresis, or by combining PCR, Southern hybridization and sequencing, etc.

Production process for mutant

In a production process for an exogenous gene, first, the DNA construct resulting from the above method is introduced into a suitable host cell to obtain a
5 homologous recombinant.

Examples of the host cell include eukaryotic organisms comprising cells of mammalian, yeast, plant and the like, but animal cells of mammalian are preferable. Preferably used as such cells are established cell lines
10 to be transformed together with a determinant that can make the mutation more effective, COS cells from monkey (Cell, 23,175 (1981)), Chinese hamster oocyte, Hela cells, RatII fibroblast, alimentary canal epithelium cells and its dihydrofolate reductase defectives (Proc. Natl. Acad.
15 Sci., USA. 77: 4216 (1980)). In the case of an established cell line, it is preferable to use but not limited to, for example T cell or B cell line animal cells, especially pre-B cell line animal cells, more specifically J558 cells and J558L cells. In the case of yeast cells,
20 yeast cells belonging to the genus Saccharomyces etc. are used. In addition, cells from animal living body can be also used as host cells.

As a result of introducing an exogenous gene into the genome of a host cell, a transformant transformed
25 by the DNA construct of the present invention can be

obtained.

Such introduction of the present DNA construct into a cell can be carried out in accordance with various methods known in the art of introducing DNA into a cell, for example, electroporation method, calcium phosphate coprecipitation method, liposome method, DEAE dextran method, microinjection method, virus transduction, etc. The electroporation method is preferably employed.

Examples of the cells to which the present DNA construct is introduced are fertilised ovum, embryo, germ cells and the like. The introduction into cells can be performed in accordance with an ordinary method.

Whether a mutant is obtained or not can be checked by known method, e.g. by random sequence method or by determining the activity of the product expressed by an exogenous gene and selecting the one whose activity is different from that of the exogenous gene. Alternatively, it can be selected based on the standard activity and its degree, or on whether the activity is high or low.

The isolated form of the cell itself containing a random mutated gene transformed with the object exogenous DNA can be improved peptides having a better function related to human exogenous gene. Therefore, it can be utilized as a pharmaceutical for improving the state of disease and as a model system for the therapeutic

study.

The resulting transformant can be cultivated in accordance with an ordinary method. As a result of the cultivation, the object protein of the invention encoded
5 by a mutant of exogenous gene is expressed and produced (accumulated, secreted) intracellularly, extracellularly or on the cell membrane of the transformant.

Various common cultures can be appropriately selected depending on the adopted host cells and used for
10 the cultivation, and the cultivation can be carried out under the suitable condition for the growth of the host cell.

The recombinant protein of the invention thus obtained can be separated and purified if desired, by
15 various separation procedures utilizing physical or chemical properties of the recombinant protein (Refer to e.g. Biochemistry Data Book II, 1175-1259, Edition 1, Issue 1, published by Tokyo Kagaku Dojin (Jun 23, 1980); Biochemistry, 25(25), 8274 (1986); Eur. J. Biochem., 163,
20 313 (1987), etc).

Specifically illustrated as the above procedure are common reconstitution processing, processing by protein precipitant (salting out method), centrifugal separation, osmotic shock method, ultrasonication,
25 ultrafiltration, molecular sieve chromatography (gel

filtration), adsorption chromatography, ion exchange chromatography, affinity chromatography, various liquid chromatographies such as high performance liquid chromatography (HPLC), dialysis, and a combination thereof.

5 For example, affinity chromatography using a column bonded with antibodies specific to the protein of the invention is particularly preferable.

Thus the present recombinant expression vector is expressed in an animal cell to give an exogenous DNA,
10 resulting in random mutants encoded by said DNA. The frequency of random mutants by the recombinant expression vector of the invention is at least about 1×10^{-4} to about 1×10^{-3} /bp/generation.

The present invention provides a production
15 process for a mutant comprising an eukaryote, especially an animal cell containing the above recombinant vector having an insertable site of an exogenous DNA or an exogenous gene to achieve the expression of an introduced DNA sequence or a gene mutant.

20 According to the present production process for a mutant, a peptide having an antimicrobial activity but unavailable because of its toxicity to animal cells can be improved to provide an antimicrobial peptide having low toxicity to animal cells and higher antimicrobial activity.
25 Similarly, it is possible to make an attempt to improve or

change the functions of proteins such as naturally occurring interferon, growth hormone and the like. Therefore, a useful method for developing agrochemicals or improving the function of pharmaceuticals or the like can
5 be provided.

Moreover, the invention provides: a kit for a production process of an exogenous gene mutant; or producing an expression product of mutant gene, comprising a DNA construct having an insertable site of an exogenous
10 DNA or an exogenous gene. Specifically, the DNA construct comprises:

- (a) a promoter;
- (b) an exogenous gene-insertion site;
- (c) an intron enhancer; and
- 15 (d) an enhancer comprising HS3b and/or HS4, if necessary further HS1 and HS2, in DNaseI sensitive region.

According to the present kit for producing a mutant, an improved antimicrobial peptide with low toxicity to animal cell and higher antimicrobial activity
20 and protein pharmaceuticals with improved functions such as naturally occurring interferon, growth hormone and the like can be provided.

BRIEF DESCRIPTION OF THE DRAWINGS

25 FIG. 1A shows a schematic view of IgH mutant

transgene constructs used to generate chimeric mice.

In FIG. 1B, locations and directions of primers used to amplify the cDNA are shown by arrows.

FIG. 2 indicates the production of anti-NP
5 antibodies producing human Cp from J558L cells transfected with the IgH mutant genes shown in Fig. 1. The supernatants of transfected J558L cells (1×10^6 cells/ml) cultivated for 12 hrs were subjected to ELISA using plates coated with NP₉-BSA for estimating the antibody production.

10 FIG. 3A shows a reconstitution of RAG-2^{-/-} mouse immune systems by ES cells in chimeric mice A4-3, 2-1 and 2-3. Levels of mouse IgM in pre-immune mice were measured by ELISA using plates coated with rat anti-mouse IgM monoclonal antibody and POD-labeled goat anti-mouse IgM
15 antibody.

FIG. 3B shows a measurement of levels of human IgM in preimmune or immune sera using plates coated with mouse anti-human IgM antibody and POD-labeled goat IgM antibody. Immune sera are referred as TD.

20 Antisera from Balb/c mouse and RAG-2^{-/-} mouse were used as controls in A, and a culture supernatant of J558L in addition to these sera were used as controls in B.

FIG. 3C shows an analysis of bindings of anti-NP antibodies to plates coated with anti NP₉-BSA antibody.
25 Bound antibodies were detected using POD-labeled anti-

mouse IgG.

FIG. 3D shows binding ratios of antisera from chimeric mice to plates coated with NP₁-BSA or NP₁₂-BSA.

FIG. 4 shows a specific amplification of transgenes by RT-PCR. cDNA prepared from mRNA of IgM-B220⁺ spleen cells from the chimeric mice immunized with NP₃₄-CGG was amplified by PCR. Only chimeric mice gave bands corresponding to the transgenes (lanes 1-6), and C57BL/6 mice did not (lane 7). J558L cell transfectant (lane 8) and non-transfectant (lane 9) are also shown as positive or negative controls.

FIG. 5 shows a comparison of distribution and frequency of somatic hypermutation in transgene constructs from spleen IgM-B220⁺ cells (A) and peritoneal B1 cells (B).

The distribution and frequency of somatic hypermutation in transgene constructs from two chimeric mice are shown separately in A. The scale at the bottom indicates the amino acid position numbered according to the method of Kabat et al.

BEST MODE FOR CARRYING OUT THE INVENTION

Examples are given below to illustrate the present invention in further detail, but they are not intended to limit the scope of the invention.

Example 1

1) Construction of IgH transgenes

H-chain of Ig (IgH) carrying human C_H with
5 different 3' flanking regions were constructed.

Acquisition of fragment containing V_H promoter and
fragment containing E_μ/MAR

A fragment containing V_H17.2.25 promoter
10 (0.55kb: SEQ.No.14) with restriction enzymes KpnI and ApaI
sites and another fragment containing E_μ/MAR (1kb) with
restriction enzymes XhoI and SalI sites were cloned by PCR
from the construct [plasmid having chloramphenicol acetyl
transferase (CAT) gene controlled by V_H promoter derived
15 from V_H17.2.25 and by intron enhancer (J-C intron
enhancer/ matrix attachment region (abbreviated to
E_μ/MAR)) between restriction enzymes XbaI and XhoI sites
of pBluescript SK(+)], which the present inventor
previously prepared for generating CAT transgenic mice
20 (Azuma, T., et al., Int. Immunology, 5(2) 121-130 (1993)).

Acquisition of V_H-D-J_H gene fragment

The rearranged V_H-D-J_H gene fragment (2.0kb)
containing restriction enzymes ApaI and XhoI sites was
25 also cloned by PCR using a genomic DNA from A6, a

hybridoma producing anti-(4-hydroxy-3-nitrophenyl)acetyl
monoclonal antibody (hereafter referred to as anti-NP
monoclonal antibody) that is obtained from C57BL/6
mouse(Tokyo Animal Center)(Taketani, M., et al., Mol.
5 Immunity, 32, 983(1995):Furukawa,k., et al., Immunity, 11,
329 (1999)).

These were cloned into Bluescript II SK (from
Stratagene) in order of V_H promoter, V_H-D-J_H gene and
E_μ/MAR.

10

Acquisition of human C_μ gene

A 6.9 kb XbaI fragment of human C_μ gene was
obtained from a phage clone, CH.H.Ig_μ-24 (Takahashi, N.,
et al., Nucleic Acids Res., 8, 5983 (1980); Health Science
15 Research Resources Bank).

Acquisition of 3' enhancer

A plasmid (Lieberson, R., et al., EMBO. J,
14,6229(1995)) containing XbaI fragment (4.0kb) of 3'
20 enhancer (hereafter abbreviated to 3'E) was obtained from
Drs. J. Manis and F. Alt (Harvard Medical School).
Fragments containing HS3b (1.2kb) indicated as SEQ No.1
and HS4 (1.4kb) indicated as SEQ No.2 were cloned by PCR
using primers indicated as SEQ No.3 to 6 and genomic DNA
25 from a B cell lymphoma, MPC11 (Madisen, L.and Groudine, M.,

Genes Dev.8,2212(1994)). In each primer sequence, recognition site for SpeI or XbaI restriction enzyme is underlined.

SEQ NO.3: HS3-S: 5'-TCTAGAACCACATGCGATCTAAGGGATATTGGGG-3'

5 SEQ NO.4: HS3-anti SpeI: 5'-CAGGACTAGTGATCATTTGAGCTCCGGCTCTAAC-3'

SEQ NO.5: HS4-S XbaI: 5'-CTAGTCTAGACTGCAGACTCACTGTTCCACCATG-3'

10 SEQ NO.6: HS4-anti SpeI: 5'-GTGGACTAGTAAGCTTGGAGTTAGGTGGGTAGG-3'

These fragments were linked to the 3' end of 3'E, in order of HS3b and HS4. 3'E, HS3b and HS4 were linked in tandem without any intervening Ig intron sequences inserted.

15 Using each fragment above, three kinds of constructs pvehcΔ3'E (12kbp), pvehc3'E (16kbp) and pvehc3'EHS3b/4 (19kbp) were obtained by means of gene recombination technique.

20 2) Structural features of transgenes

The transgene constructs obtained in this invention contain V region encoded by mouse V_H186.2, a dominant V_H involved in the response to (4-hydroxy-3-nitrophenyl)acetyl haptens (Bothwell, A.L.M., et al., Cell, 25 24,625(1981)).

The mouse V_H 186.2-DFL16.1- J_H 2 gene rearranged from A6, a hybridoma producing monoclonal antibody against (4-hydroxy-3-nitrophenyl)acetyl (Taketani, M., et al., Mol. Immunity, 32, 983 (1995); Furukawa, k., et al., Immunity, 11, 329 (1999)), was linked to human C_μ , thereby facilitating a discrimination between encoded transgene and endogenous mouse H.

In addition, as the expression of transgenes on B cell surface might change the development of the B cell, transmembrane exon was removed from C_μ to prevent the expression of transgenes on the cell surface. The transgenes contained only C_μ but not gene regions encoding other H chain isotypes.

All three kinds of transgene construct contained V_H promoter and E_μ /MAR from mouse J- C_μ intron as cis-acting elements, which had been used for CAT transgene construction (Azuma, T., et al., Int. Immunol., 5, 121 (1993)). Therefore, differences in the structure of the transgenes were restricted to the 3' flanking region (Fig.1).

One of the constructs, pvehc Δ 3'E, lacked 3' enhancer region and was driven by V_H promoter and E_μ /MAR. Another construct, pvehc3'E, contained 3' enhancer (Liebersohn, R., et al., EMBO. J., 14, 6229 (1995)) in addition to V_H promoter and E_μ /MAR. The restriction enzyme BamHI

fragment (4.0kb) containing 3' enhancer (3'E) was used for the gene construction. The addition of HS3b and HS4 to pvehc3'E gave rise to a construct referred to as pvehc3'EHS3b/4.

5 Endogenous IgH genes contained an additional cis-acting element, referred to as HS3a (Matthias, P. and Baltimore, D., Mol. Cell Biol., 13, 1547 (1993)), which has an identical nucleotide sequences to those of HS3b but was in an inverted form (Arulampalam, V. et al., Immunol.
10 Today, 18, 549(1997)). The transgene constructs of the present invention did not contain this cis-acting element.

Example 2

1) Transfection of IgH mutant genes (Lundblad, A. et al.
15 Immunochemistry 9, 535-544 (1972)) into J558L cells and antibody production

To estimate the transcriptional activity of these three constructs, the constructs were tranfected into a mouse plasmacytoma cell line, J558L cells.
20 Linearized IgH gene constructs (10µg) having a restriction enzyme NotI recognition site were subjected to electroporotation (Sahagaw, B.G. et al. 1986, J.Immunol., 137; 1066-1074) into J558L cells together with a plasmid pSV2-gpt (1.6µg) (Mulligan, R.C. & Berg P. Science 209;
25 1422-1427 (1980)). Specific condition of the

electroporation is as follows.

Instrument: Gene Pulser (BIO-RAD laboratories)

Conditions:

Buffer: PBS

5 Cells: J558L 2×10^7 cells/ml in 0.5 ml PBS

Linearized DNA: IgH transgene (10 μ g) and pSV2-gpt
(1.6 μ g)

Reaction condition: capacitance (960 μ F), voltage
(350V)

10 The transfected cells were cultivated at 37°C in
the presence of CO₂ (5%) and a mixture of hypoxanthine,
xanthine and mycophenolic acid (Sigma). Limiting dilution
of transformants of a drug resistant IgH gene was carried
out, and clones were selected based on antibody production.
15 Specifically, the production of anti-NP chimeric antibody
was measured by ELISA (FIG.2).

For the quantitative analysis of antibody
production, some clones of transformant induced from each
transgene construct were cultivated for 12 hrs in a flat-
20 bottomed 96-hole plate with RPMI1640 medium (total volume:
200 μ l; Nissui Pharmaceutical Co., Ltd.) containing 10%
fetal bovine serum (FCS), at the concentration of 2×10^6
cells/ml. The culture supernatant diluted to various
concentration with phosphate buffer saline (PBS)
25 containing 1mg/ml bovine serum albumin (BSA) was analyzed

by ELISA using a polyvinyl plate coated with NP₉-BSA. Peroxidase (POD)-labeled anti-mouse λ_1 chain antibody (Southern Biotech) or anti-human μ chain antibody (ZYMED) was used for detecting bound antibodies.

5 The result is shown in FIG. 2.

Transgene constructs were designed to encode chimeric H-chains, possessing the V_H region from a mouse anti-NP monoclonal antibody and the C region from human Ig, which were expected to assemble with mouse λ_1 chains and
10 give rise to anti-NP chimeric antibody.

All constructs were actively transfected in J558L cells and produced anti-NP antibodies by pairing with endogenous λ_1 -chains. Among these constructs, pvehc Δ 3'E showed rather weaker production compared with
15 the others. The addition of 3'E (pvehc3'E) resulted in a significant increase in antibody production, however, further addition of HS3b and HS4 seemed to have no effect on antibody production. Although little information was available concerning the dependence of transcription on
20 the copy number of each transfected DNA, the inventor assumed that the amount of antibody production reflected the transcriptional activity of the constructs, which did not differ significantly between pvehc3'E and pvehc3'EHS3b/4.

Example 3

1) Generation of chimeric mice

To generate chimeric mice, the present inventor employed at least two independent transfected ES clones
5 for the microinjection into RAG-2^{-/-} mouse blastocysts. The inventor also used at least two chimeric mice/transgene constructs for the analysis of somatic hypermutation. The transfected ES cell lines and the chimeric mice used in this experiment are shown in Table 1.

10 The EcoRI fragment of pGKneo (1.5kb, 1µg) and the three kinds of linear-chain IgH transgene constructs (30µg) in the same manner as in Example 2 were electroporated into 1×10^7 of E14-1 cells (Kuhn, R. et al., Science, 254, 707 (1991)) using a Bio-Rad Gene Pulser.

15 The transfected cells were then selected with G418 (150µg/ml, Gibco). The selected colonies were screened by: PCR using two primers indicated as SEQ.No.7 and 8 that can be hybridized to DNA sequences located in the V_H17.2.25 promoter and J_H2; and Southern blotting
20 using human Cµ gene as a probe, respectively.

SEQ.No.7: 5'-GACTCAGGAGGACTCTAGTT-3'

SEQ.No.8: 5'-GGTGTCCCTAGTCCTTCATG-3'

PCR amplification was performed for 30 cycles of 95°C for 30 sec, 65°C for 30 sec and 72°C for 1 min. The
25 copy number of integrated pvehc3'E was calculated to be 2

to 3 by Southern blotting. The copy numbers of other transgenes pvehcΔ3'E and pvehc3'EHS3b/4 were estimated by the PCR analysis using DNA from pvehc3'E-transfected ES cells as a standard.

5

2) Antibody production by chimeric mice

ES cell clones containing the IgH transgene were injected into blastocysts from RAG-2^{-/-} mice (Takahashi, N. et al., Nucleic Acids Res., 8, 5983(1980)) and
10 transplanted into uteri of ICR foster mothers (8 to 12 week old, CLEA Japan Inc.). Complementarity of the immune system by B cells and T cells originated from ES cells in chimeric mice was tested by flow cytometry after staining the cells with PE anti-CD3 antibody or biotin anti-
15 B220/SA-FITC.

POD-labeled sheep anti-mouse μ -chain antibody was used for the detection of bound antibodies. For analyzing antibodies having human C μ , plates were coated with sheep anti-human IgM antibody (Southern Biotech) and
20 POD-labeled anti-human IgM monoclonal antibody (ZYMED) was used for detecting bound antibodies. The culture supernatant of J558L transformant, human Waldenstrom IgM, and anti-NP IgM monoclonal antibody B4-3 were used as control antibodies.

25 Table 1. Used ES cells and chimeric mice

Construct	Cell clone	Copy number	Chimeric mouse
pvehc Δ 3'E	3-89	<3	3-2
	3-88	<3	3-4
pvehc3'E	No.6	<3	1-1,2/3
	A-4	<3	A4-3
pvehc3'EHS3b/4	25-2-5	<3	2-1
	25-2-2	<3	2-3

A backcross with C57BL/6 mouse was carried out to examine the germinal transmission of the transgene, and the DNA was extracted from the tail of the resulting mouse to analyze the generation of chimeric mouse based on whether the transgene was contained or not.

3) Immune and antibody production

In order to examine whether the immune system was reconstituted with the lymphocytes derived from ES cells, the amounts of IgM in the pre-immune sera of chimeric mice A4-3, 2-1, and 2-3 were measured by ELISA.

The result is shown in FIG. 3A. Irrespective of the kind of transgene, detected amounts of mouse IgM in all the sera from these mice were similar, but smaller than that in a normal BALB/c mouse.

Next, NP₃₄-chicken γ globulin (NP₃₄-CGG) and NP-BSA having a different NP value were prepared in the same manner as in Azuma et al. (Azuma, T. et al., Molec. Immunol., 24, 287(1987)).

CFA (complete Freund's adjuvant: Difco,

100 μ g/mouse) including (4-hydroxy-3-nitrophenyl)acetyl
chicken γ globulin (hereafter referred to as NP₃₄-CGG) was
administered to the chimeric mice, and the same amount of
additional immune was administered using IFA (incomplete
5 Freund's adjuvant). Three days after final administration
of PBS including NP₃₄-CGG, antisera were collected from
the immunized mice. The mice were then sacrificed and
tissue samples were obtained.

Plates coated with NP₁-BSA or NP₁₆-BSA were used
10 for measuring the production of anti-NP antibodies and the
affinity maturation of these antibodies. POD-labeled
sheep anti-mouse IgG (ZYMED) was used for detecting the
antibodies.

Anti-NP monoclonal antibody, F8, was used as a
15 control for an immature antibody, and C6 as a control for
a matured monoclonal antibody (Taketani, M. et al., Mol.
Immunol., 32, 983(1995) ; Furukawa, K., et al., Immunity, 11,
329(1999)).

The result is shown in FIG. 3B. The antibody
20 expressing human Cp encoded by the transgenes in the pre-
immune or immune antisera from the mice immunized with
NP₃₄-CGG was not detected.

Hybridomas producing anti-NP monoclonal antibody
(γ 1 λ 1) were prepared from pvehc3'EHS3b/4 mice immunized
25 with NP₃₄-CGG. None of these secreted antibodies expressed

human Cp, however, some of them synthesized intracellular H-chain having human Cp.

The result suggests that the transgenes were transcribed and translated in the B cells of chimeric mice. However, the secreted H-chain products level was lower than the detectable level.

Next, antibody reaction of chimeric mice against TD antigen NP₃₄-CGG was examined by ELISA. The result is shown in FIG. 3C. The antibody value changed, but all the mice produced anti-IgG antibodies indicating that the immune systems respond to the anti-NP antibody production.

Furthermore, the binding ratio of these antibodies to NP₁-BSA, which relates to the binding to NP₁₆-BSA, was examined as the measurement of affinity maturation. The result is shown in FIG. 3D.

As shown in FIG. 3D, a control monoclonal antibody F8 showed no somatic hypermutation and had an association constant (K_a) of $2 \times 10^5/\text{M}$ with NP-Cap, and the ratio was 0.29. On the other hand, well matured monoclonal antibody C6 having $2 \times 10^7/\text{M}$ of K_a (Furukawa, K. et al., Immunity, 11, 329(1999)) had the ratio of around 1. All the sera from chimeric mice showed similar ratios to that of C6, therefore the affinity maturation of anti-NP antibodies advanced to the similar extent in all chimeric mice after immunized with NP₃₄-CGG.

Example 4. Cloning and Sequencing of V_H Gene

Although the immune system was reconstituted in the chimeric mouse, it was rather difficult to obtain
5 sufficient amount of $PNA^{hi}IgG^+B$ cells, which are popular as germinal center cells derived from a single chimeric mouse. Accordingly, after immunization with NP_{34} -CGG, spleen IgM^- $B220^+$ cells, which are expected to select isotype-switching memory B cells, XX were sorted by flow
10 cytometry using a FACS Vantage.

The single-cell suspension from spleen of chimeric mice immunized with NP_{34} -CGG were treated with 0.83% of ammonium chloride to induce hemolysis. In some experiments, T cells were also treated with anti-Thy1
15 antibody (T24/40 and HO13.4) to induce hemolysis, and then treated with a rabbit complement. The cells were stained with phycoerythrin (PE)-labeled anti-CD45R/B220 (cell surface molecule, pherMingen) or fluorescein isothiocyanate (FITC)-labeled anti-mouse IgM (ZYMED).

20 The peritoneal cells were stained with biotin-labeled anti-CD5/FITC-labeled streptavidin (pherMingen) and PE-labeled anti-CD45R(B220).

The $CD45R(B220)^+IgM^-$, $CD45R(B220)^+IgM^+$, or $CD5^+CD45R(B220)^+$ cells were fractioned by flow cytometry on
25 a fluorescence activated cell sortor (FACS) Vantage

(Becton, Dickinson and Company).

In addition, as a control experiment, CD4+ cells and/or CD8+ cells were obtained from the thymus after staining and sorting.

5 Total RNA was prepared from spleen or peritoneal cells fractionated by the above cytometry, using TRIzol (RNA preparation reagent, GIBCO BRL) according to the manufacturer's instructions.

10 The cDNA was prepared from the total RNA using oligo(dT) as a primer and Superscript II reverse transcriptase (GIBCO BRL). The cDNA samples were each PCR-amplified using: V_H186.2 primer of SEQ.No.9 and human C_μ primer of SEQ.No.10 for the transgenes; and V_H186.2 primer of SEQ.No.9 and mouse C_γ1 primer of SEQ.No.11 for
15 endogenous mouse Ig genes, respectively.

SEQ.No.9: 5'-CATGCTCTTCTTGGCAGCAAC-3'

SEQ.No.10: 5'-GCAGCCAACGGCCACGCTGC-3'

SEQ.No.11: 5'-GGCCGAATTCCATGGAGTTAGTTTGGG-3'

20 The PCR reaction was carried out under the condition of 95°C for 1 min, 62°C for 1 min and 72°C for 1 min for 30 cycles. PCR products of the transgenes and endogenous mouse IgH genes were analyzed by agarose gel electrophoresis or ligated into pCR-2.1 (Invitrogen Corporation) using TA cloning kit (Invitrogen Corporation)
25 and sequenced using M13 primer shown as SEQ.No.12 (-20,

Takara Bio Inc.), M13 reverse primer shown as SEQ.No.13 (Takara Bio Inc.) and Hitachi DNA sequencer model 5500 (Hitachi, Ltd.).

As shown in FIG. 4, the expression of antibody fragments including transgenes was identified.

2) Somatic hypermutation of transgenes in spleen or peritoneal B cells

The frequency of mutation was calculated by dividing the total of mutation by the total of sequence-determined base. The result is shown in Table 2.

Table 2. Mutations in V region sequences of spleen IgM⁺ B cells of chimeric mice

Construct	pvehcΔ3'E		pvehc3'E		pvehc3'EHS3b/4	
ES clone	3-89	3-88	No.6	A4	25-2-5	25-2-2
Number of sequence analyzed	30	18	51 ^c	30	21	21
Number of mutation observed	16	11	42	25	161	132
Percent of mutation in CDR ^a (%)	56	36	38	75	57	55
Mutation frequency ^b (%)	0.15	0.17	0.23	0.23	2.14	1.76

^a Percent of mutation in CDR = 100 x (total number of point mutations in CDR) / (total number of point mutations)

^b Mutation frequency = 100 x (total number of point mutations) / (total number of base pairs)

^c Two chimeric mice were analyzed.

For the somatic hypermutation in transgenes, primers to hybridize with either V_H186.2 or human C_μ were used. Cloning and sequencing of the cDNA prepared by RT-PCR were conducted. The result is shown in Table 3. The distribution and frequency of somatic hypermutation among the transgene constructs in spleen B cells are shown in FIG. 5.

Specific amplification of the transgenes due to the combined use of the primers was observed only in the spleen cells from chimeric mice, but not in those from C57BL/6 mice immunized with NP₃₄-CGG. In addition, it was observed that all the sequences have an identical junctional diversity in CDR3 that is a characteristic of A6DNA used to construct transgenes.

As to the gene construct pvehΔ3'E, nucleotide changes resulting from somatic hypermutation were observed in V_H186.2-DFL16.1-J_H2, but the frequency was low. The frequency of mutation was estimated to be 0.17 % (Table 2).

The gene construct pvehc3'E showed an essentially similar distribution of hypermutation, although the frequency increased slightly compared to that of pvehcΔ3'E (Table 2).

In the case of pvehc3'EHS3b/4, where HS3b/4 is

further added to the gene construct pvehc3'E of the invention, a dramatic increase of the frequency of somatic hypermutation was shown as a result in V_H186.2-DFL16.1-J_H2 gene (FIG. 5). All the sequenced PCR clones contained
5 nucleotide changes in spite of the fact that they were selected randomly. These nucleotide changes were found with high frequency in the area around CDR2 (hypervariable region 2) and CDR3 (FIG. 5 and Table 2).

The data were obtained from two chimeric mice
10 corresponding to each transgene construct shown in FIG.5, and were in good agreement with each other. A high frequency of hypermutation was apparent in pvehc3'EHS3b/4. However, mutation was clearly absent in peritoneal B1 cells from the same chimeric mouse (FIG. 5).

15 The following Table 3 shows characteristics of substitutions identified in the transgenes.

Table 3. Mutations in IgH gene

Substitution	Construct		
	pvehcΔ3'E	pvehc3'E	pvehc3'EHS3b/4
Transition			
A→G	5	12	25
G→A	3	13	47
C→T	1	5	8
T→C	3	5	34
Total	12	35	114
Transversion			
C→A	2	0	8

G→T	1	2	20
A→C	2	2	32
A→T	0	5	12
G→C	4	7	43
C→G	1	4	11
T→A	4	7	26
T→G	1	3	27
Total	15	32	179

It was clear from the comparison between IgH transgenes that the base transition and base transversion arise with almost the same frequency.

5 In addition, in the locus sequence named as RGYW motif (A/G, G, C/T, A/T) (Kabat, E. A. et al., US National Institutes of Health, Bethesda, M.D., U.S.A., p1394, (1991): Rogozin, I.B. et al., Biochim. Biophys. Acta., 1171, 11 (1992)), the emergence of somatic hypermutation
10 in V_H186.2 transgene was analyzed. The germline gene V_H186.2 (294bp) contained RGYW/WRCY motif (93bp) corresponding to 32 % of the total bases, and the frequency of somatic hypermutation in this motif appeared to be 34 %, which is not significantly higher than those
15 in the other V_H regions.

INDUSTRIAL APPLICABILITY

In the above Examples, the present inventor employed a RAG-2^{-/-} blastocyst system to prepare chimeric

mice carrying IgH transgenes with different cis-acting elements.

The producing method of the invention has some advantages over conventional transgene techniques and is superior especially when ES cells are transformed with low copy numbers of transgene. In fact, the copy number of transgene in transformant ES cells was calculated to be 3 or less in most cases by Southern blotting and PCR (Table 1).

Only a marginal numbers of hypermutation was found in pvehcΔ3'E and the finding was confirmed that the cis-acting elements, V_H promoter and E_μ/MAR were not sufficient for inducing a high frequency of hypermutation. Addition of a 3' enhancer fragment (4kb) to pvehcΔ3'E (pvehc3'E) resulted in an approximately 30 % increase in the frequency of hypermutation.

In the present invention, it was found that the expression frequency of induced hypermutation was greatly enhanced by introducing HS3b and HS4 to the 3' end of 3' enhancer of pvehc3'E, which indicates that the construct pvehc3'EHS3b/4 contains all necessary elements for somatic hypermutation. The Experiments by the inventor clarified that the motifs responsible for high frequency hypermutation are within an HS3b/4 region of 2.6 kb.

Thus it is possible to provide a recombinant

expression vector for inducing a random mutant of desired transgene. Moreover, it is possible to provide a production process for a high frequent random mutant obtained by expressing the recombinant expression vector
5 in animal cells. Furthermore, it is possible to provide a method for expressing a random mutant or a kit for producing a random mutant comprising the recombinant expression vector and animal cells.

CLAIMS

1. A production process for an exogenous gene
random mutants, wherein a DNA construct comprising at
5 least the following (a)-(d) is introduced into an animal
cell and the exogenous gene mutant is produced in the
animal cell:

- (a) a promoter;
- (b) an exogenous gene;
- 10 (c) an intron enhancer; and
- (d) an enhancer comprising HS3b and/or HS4 in a DNaseI
sensitive region.

2. The production process according to claim 1,
15 wherein the (d) enhancer further comprises HS1 and HS2.

3. The production process according to claim 1
or 2, wherein the (d) enhancer has a DNA sequence
comprising at least a DNA sequence of SEQ ID NO: 1 or 2.

20

4. The production process according to any one
of claims 1 to 3, wherein DNA sequences of (a), (c) and
(d) are derived from a DNA sequence of an immunoglobulin

25

5. The production process according to claim 2,

wherein the (d) enhancer comprises HS1, HS2, HS3b and HS4.

6. The production process according to any one of claims 1 to 5, wherein

5 the (a) promoter is a V_H promoter; and
the (d) enhancer has a DNA sequence comprising DNA sequences of SEQ ID NOs: 1 and 2.

7. The production process according to claim 5
10 or 6, wherein the DNA construct is pvehc3'EHS3b/4.

8. The production process according to any one of claims 1 to 7, wherein the animal cell is a B cell line animal cell.

15

9. The production process according to claim 8, wherein the B cell line animal cell is derived from a pre-B lymphocyte line cell.

20 10. A method for obtaining an expression product of a mutant gene by expressing the exogenous gene mutant obtained by the method according to any one of claims 1 to 9 in the animal cell.

25 11. A DNA construct for producing an exogenous

gene mutant, comprising at least:

- (a) a promoter;
- (b) an exogenous gene;
- (c) an intron enhancer; and
- 5 (d) an enhancer comprising HS3b and/or HS4 in a DNaseI sensitive region.

12. The DNA construct according to claim 11, wherein the (d) enhancer further comprises HS1 and HS2.

10

13. The DNA construct according to claim 11 or 12, wherein the (d) enhancer has a DNA sequence comprising at least a DNA sequence of SEQ ID NO: 1 or 2.

15

14. The DNA construct according to any one of claims 11 to 13, wherein DNA sequences of (a), (c) and (d) are derived from a DNA sequence of immunoglobulin.

15. The DNA construct according to any one of
20 claims 11 to 14, wherein the (d) enhancer comprises HS1, HS2, HS3b and HS4.

16. The DNA construct according to any one of
claims 11 to 15, wherein
25 the (a) promoter is a V_H promoter; and

the (d) enhancer has a DNA sequence comprising DNA sequences of SEQ ID NOs: 1 and 2.

17. The DNA construct according to claim 11 or 12, wherein the DNA construct is pvehc3'EHS3b/4.

18. A kit for production process of an exogenous gene mutant or producing an expression product of a mutant gene, comprising a vector having:

- 10 (a) a promoter;
- (b) an exogenous gene-insertion site;
- (c) an intron enhancer; and
- (d) an enhancer comprising HS3b and/or HS4 in a DNaseI sensitive region;
- 15 for achieving an expression of a mutated DNA sequence which produces an exogenous gene mutant.

19. The kit according to claim 18, wherein the (d) enhancer further comprises HS1 and HS2.

20

20. V_H17.2.25 promoter having a sequence of SEQ ID NO: 14.