

SECTION C — CHEMISTRY; METALLURGY

C12 BIOCHEMISTRY; BEER; SPIRITS; WINE; VINEGAR; MICROBIOLOGY; ENZYMOLOGY; MUTATION OR GENETIC ENGINEERING

C12N MICROORGANISMS OR ENZYMES; COMPOSITIONS THEREOF; PROPAGATING, PRESERVING, OR MAINTAINING MICROORGANISMS; MUTATION OR GENETIC ENGINEERING; CULTURE MEDIA (microbiological testing media C12Q 1/00) [3]

Note(s) [3, 4, 6, 7, 2006.01]

- Attention is drawn to Notes (1) to (3) following the title of class C12.
- Biocidal, pest repellent, pest attractant or plant growth regulatory activity of compounds or preparations is further classified in subclass A01P.
- Therapeutic activity of single-cell proteins or enzymes is further classified in subclass A61P.
- When classifying in this subclass, classification is also made in group B01D 15/08 insofar as subject matter of general interest relating to chromatography is concerned.
- In this subclass, it is desirable to add the indexing codes of subclass C12R.

Subclass index

MICROORGANISMS; SPORES; UNDIFFERENTIATED CELLS; VIRUSES.....	1/00, 3/00, 5/00, 7/00, 11/00
ENZYMES.....	9/00, 11/00
TREATMENT WITH ELECTRICAL OR WAVE ENERGY.....	13/00
MUTATION OR GENETIC ENGINEERING.....	15/00

1/00 Microorganisms, e.g. protozoa; Compositions thereof (medicinal preparations containing material from protozoa, bacteria or viruses A61K 35/66, from algae A61K 36/02, from fungi A61K 36/06; preparing medicinal bacterial antigen or antibody compositions, e.g. bacterial vaccines, A61K 39/00); Processes of propagating, maintaining or preserving microorganisms or compositions thereof; Processes of preparing or isolating a composition containing a microorganism; Culture media therefor [3, 2006.01]	1/19	• • • modified by introduction of foreign genetic material [5, 2006.01]
1/02 • Separating microorganisms from their culture media [3, 2006.01]	1/20	• Bacteria; Culture media therefor [3, 2006.01]
1/04 • Preserving or maintaining viable microorganisms (immobilised microorganisms C12N 11/00) [3, 2006.01]	1/21	• • modified by introduction of foreign genetic material [5, 2006.01]
1/06 • Lysis of microorganisms [3, 2006.01]	1/22	• Processes using, or culture media containing, cellulose or hydrolysates thereof [3, 2006.01]
1/08 • Reducing the nucleic acid content [3, 2006.01]	1/24	• Processes using, or culture media containing, waste sulfite liquor [3, 2006.01]
1/10 • Protozoa; Culture media therefor [3, 2006.01]	1/26	• Processes using, or culture media containing, hydrocarbons (refining of hydrocarbon oils by using microorganisms C10G 32/00) [3, 2006.01]
1/11 • • modified by introduction of foreign genetic material [5, 2006.01]	1/28	• • aliphatic [3, 2006.01]
1/12 • Unicellular algae; Culture media therefor (as new plants A01H 13/00) [3, 2006.01]	1/30	• • • having five or less carbon atoms [3, 2006.01]
1/13 • • modified by introduction of foreign genetic material [5, 2006.01]	1/32	• Processes using, or culture media containing, lower alkanols, i.e. C ₁ to C ₆ [3, 2006.01]
1/14 • Fungi (culture of mushrooms A01G 18/00; as new plants A01H 15/00); Culture media therefor [3, 2006.01]	1/34	• Processes using foam culture [3, 2006.01]
1/15 • • modified by introduction of foreign genetic material [5, 2006.01]	1/36	• Adaptation or attenuation of cells [3, 2006.01]
1/16 • • Yeasts; Culture media therefor [3, 2006.01]	1/38	• Chemical stimulation of growth or activity by addition of chemical compounds which are not essential growth factors; Stimulation of growth by removal of a chemical compound (C12N 1/34 takes precedence) [3, 2006.01]
1/18 • • • Baker's yeast; Brewer's yeast [3, 2006.01]	3/00 Spore-forming or isolating processes [3, 2006.01]	
	5/00 Undifferentiated human, animal or plant cells, e.g. cell lines; Tissues; Cultivation or maintenance thereof; Culture media therefor (plant reproduction by tissue culture techniques A01H 4/00) [3, 5, 2006.01]	

- 5/02 • Propagation of single cells or cells in suspension; Maintenance thereof; Culture media therefor [3, 2006.01]
- 5/04 • Plant cells or tissues [5, 2006.01]
- 5/07 • Animal cells or tissues [2010.01]
- Note(s) [2010.01]**
- The last place priority rule does not apply between the subgroups of this group.
- 5/071 • • Vertebrate cells or tissues, e.g. human cells or tissues [2010.01]
- 5/073 • • • Embryonic cells or tissues; Foetal cells or tissues [2010.01]
- 5/0735 • • • • Embryonic stem cells; Embryonic germ cells [2010.01]
- 5/074 • • • Adult stem cells [2010.01]
- 5/075 • • • Oocytes; Oogonia [2010.01]
- 5/076 • • • Sperm cells; Spermatogonia [2010.01]
- 5/077 • • • Mesenchymal cells, e.g. bone cells, cartilage cells, marrow stromal cells, fat cells or muscle cells [2010.01]
- 5/0775 • • • • Mesenchymal stem cells; Adipose-tissue derived stem cells [2010.01]
- 5/078 • • • Cells from blood or from the immune system [2010.01]
- 5/0781 • • • • B cells; Progenitors thereof [2010.01]
- 5/0783 • • • • T cells; NK cells; Progenitors of T or NK cells [2010.01]
- 5/0784 • • • • Dendritic cells; Progenitors thereof [2010.01]
- 5/0786 • • • • Monocytes; Macrophages [2010.01]
- 5/0787 • • • • Granulocytes, e.g. basophils, eosinophils, neutrophils or mast cells [2010.01]
- 5/0789 • • • • Stem cells; Multipotent progenitor cells [2010.01]
- 5/079 • • • Neural cells [2010.01]
- 5/0793 • • • Neurons [2010.01]
- 5/0797 • • • • Stem cells; Progenitor cells [2010.01]
- 5/09 • Tumour cells [2010.01]
- 5/095 • • Stem cells; Progenitor cells [2010.01]
- 5/10 • Cells modified by introduction of foreign genetic material, e.g. virus-transformed cells [5, 2006.01]
- 5/12 • • Fused cells, e.g. hybridomas [5, 2006.01]
- 5/14 • • • Plant cells [5, 2006.01]
- 5/16 • • • Animal cells [5, 2006.01]
- 5/18 • • • • Murine cells, e.g. mouse cells [5, 2006.01]
- 5/20 • • • • • one of the fusion partners being a B lymphocyte [5, 2006.01]
- 5/22 • • • • Human cells [5, 2006.01]
- 5/24 • • • • • one of the fusion partners being a B lymphocyte [5, 2006.01]
- 5/26 • • • Cells resulting from interspecies fusion [5, 2006.01]
- 5/28 • • • • one of the fusion partners being a human cell [5, 2006.01]
- 7/00 Viruses, e.g. bacteriophages; Compositions thereof; Preparation or purification thereof** (medicinal preparations containing viruses A61K 35/76; preparing medicinal viral antigen or antibody compositions, e.g. virus vaccines, A61K 39/00) [3, 2006.01]
- 7/01 • Viruses, e.g. bacteriophages, modified by introduction of foreign genetic material (vectors C12N 15/00) [5, 2006.01]
- 7/02 • Recovery or purification [3, 2006.01]
- 7/04 • Inactivation or attenuation; Producing viral sub-units [3, 2006.01]
- 7/06 • • by chemical treatment [3, 2006.01]
- 7/08 • • by serial passage of virus [3, 2006.01]
- 9/00 Enzymes, e.g. ligases (6.); Proenzymes; Compositions thereof** (preparations containing enzymes for cleaning teeth A61K 8/66, A61Q 11/00; medicinal preparations containing enzymes or proenzymes A61K 38/43; enzyme containing detergent compositions C11D); **Processes for preparing, activating, inhibiting, separating, or purifying enzymes** [3, 2006.01]
- Note(s) [3, 5]**
- In this group:
- proenzymes are classified with the corresponding enzymes;
 - enzymes are generally categorised according to the "Nomenclature and Classification of Enzymes" of the International Commission on Enzymes. Where appropriate, this designation appears in the subgroups below in parenthesis.
- 9/02 • Oxidoreductases (1.), e.g. luciferase [3, 2006.01]
- 9/04 • • acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3, 2006.01]
- 9/06 • • acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3, 2006.01]
- 9/08 • • acting on hydrogen peroxide as acceptor (1.11) [3, 2006.01]
- 9/10 • Transferases (2.) (ribonucleases C12N 9/22) [3, 2006.01]
- 9/12 • • transferring phosphorus containing groups, e.g. kinases (2.7) [3, 2006.01]
- 9/14 • Hydrolases (3.) [3, 2006.01]
- 9/16 • • acting on ester bonds (3.1) [3, 2006.01]
- 9/18 • • • Carboxylic ester hydrolases [3, 2006.01]
- 9/20 • • • • Triglyceride splitting, e.g. by means of lipase [3, 2006.01]
- 9/22 • • • Ribonucleases [3, 2006.01]
- 9/24 • • acting on glycosyl compounds (3.2) [3, 2006.01]
- 9/26 • • • acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3, 2006.01]
- 9/28 • • • • Alpha-amylase from microbial source, e.g. bacterial amylase [3, 2006.01]
- 9/30 • • • • • Fungal source [3, 2006.01]
- 9/32 • • • • Alpha-amylase from plant source [3, 2006.01]
- 9/34 • • • • Glucoamylase [3, 2006.01]
- 9/36 • • • acting on beta-1, 4 bonds between N-acetylmuramic acid and 2-acetyl amino 2-deoxy-D-glucose, e.g. lysozyme [3, 2006.01]
- 9/38 • • • acting on beta-galactose-glycoside bonds, e.g. beta-galactosidase [3, 2006.01]
- 9/40 • • • acting on alpha-galactose-glycoside bonds, e.g. alpha-galactosidase [3, 2006.01]
- 9/42 • • • acting on beta-1, 4-glucosidic bonds, e.g. cellulase [3, 2006.01]
- 9/44 • • • acting on alpha-1, 6-glucosidic bonds, e.g. isoamylase, pullulanase [3, 2006.01]
- 9/46 • • • • Dextranase [3, 2006.01]
- 9/48 • • acting on peptide bonds, e.g. thromboplastin, leucine aminopeptidase (3.4) [3, 2006.01]
- 9/50 • • • Proteinases [3, 2006.01]
- 9/52 • • • • derived from bacteria [3, 2006.01]
- 9/54 • • • • • bacteria being Bacillus [3, 2006.01]

9/56	• • • • • Bacillus subtilis or Bacillus licheniformis [3, 2006.01]	13/00	Treatment of microorganisms or enzymes with electrical or wave energy, e.g. magnetism, sonic waves [3, 2006.01]
9/58	• • • • • derived from fungi [3, 2006.01]	15/00	Mutation or genetic engineering; DNA or RNA concerning genetic engineering, vectors, e.g. plasmids, or their isolation, preparation or purification; Use of hosts therefor (mutants or genetically engineered microorganisms C12N 1/00, C12N 5/00, C12N 7/00; new plants A01H; plant reproduction by tissue culture techniques A01H 4/00; new animals A01K 67/00; use of medicinal preparations containing genetic material which is inserted into cells of the living body to treat genetic diseases, gene therapy A61K 48/00; peptides in general C07K) [3, 5, 6, 2006.01]
9/60	• • • • • from yeast [3, 2006.01]		
9/62	• • • • • from Aspergillus [3, 2006.01]		
9/64	• • • • • derived from animal tissue, e.g. rennin [3, 2006.01]		
9/66	• • • Elastase [3, 2006.01]		
9/68	• • • Plasmin, i.e. fibrinolysin [3, 2006.01]		
9/70	• • • Streptokinase [3, 2006.01]		
9/72	• • • Urokinase [3, 2006.01]		
9/74	• • • Thrombin [3, 2006.01]		
9/76	• • • Trypsin; Chymotrypsin [3, 2006.01]		
9/78	• • acting on carbon to nitrogen bonds other than peptide bonds (3.5) [3, 2006.01]		
9/80	• • • acting on amide bonds in linear amides [3, 2006.01]		
9/82	• • • Asparaginase [3, 2006.01]		
9/84	• • • Penicillin amidase [3, 2006.01]		
9/86	• • • acting on amide bonds in cyclic amides, e.g. penicillinase [3, 2006.01]		
9/88	• Lyases (4.) [3, 2006.01]	15/01	• Preparation of mutants without inserting foreign genetic material therein; Screening processes therefor [5, 2006.01]
9/90	• Isomerases (5.) [3, 2006.01]	15/02	• Preparation of hybrid cells by fusion of two or more cells, e.g. protoplast fusion [5, 2006.01]
9/92	• • Glucose isomerase [3, 2006.01]	15/03	• • Bacteria [5, 2006.01]
9/94	• Pancreatin [3, 2006.01]	15/04	• • Fungi [5, 2006.01]
9/96	• Stabilising an enzyme by forming an adduct or a composition; Forming enzyme conjugates [3, 2006.01]	15/05	• • Plant cells [5, 2006.01]
9/98	• Preparation of granular or free-flowing enzyme compositions (C12N 9/96 takes precedence) [3, 2006.01]	15/06	• • Animal cells [5, 2006.01]
9/99	• Enzyme inactivation by chemical treatment [3, 2006.01]	15/07	• • Human cells [5, 2006.01]
		15/08	• • Cells resulting from interspecies fusion [5, 2006.01]
		15/09	• Recombinant DNA-technology [5, 2006.01]
11/00	Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3, 2006.01]	15/10	• • Processes for the isolation, preparation or purification of DNA or RNA (chemical preparation of DNA or RNA C07H 21/00; preparation of non-structural polynucleotides from microorganisms or with enzymes C12P 19/34) [5, 2006.01]
11/02	• Enzymes or microbial cells immobilised on or in an organic carrier [3, 2006.01]	15/11	• • DNA or RNA fragments; Modified forms thereof (DNA or RNA not used in recombinant technology C07H 21/00) [5, 2006.01]
11/04	• • entrapped within the carrier, e.g. gel or hollow fibres [3, 2006.01]	15/113	• • • Non-coding nucleic acids modulating the expression of genes, e.g. antisense oligonucleotides [2010.01]
11/06	• • attached to the carrier via a bridging agent [3, 2006.01]	15/115	• • • Aptamers, i.e. nucleic acids binding a target molecule specifically and with high affinity without hybridising therewith [2010.01]
11/08	• • the carrier being a synthetic polymer [3, 2006.01, 2020.01]	15/117	• • • Nucleic acids having immunomodulatory properties, e.g. containing CpG-motifs [2010.01]
11/082	• • • obtained by reactions only involving carbon-to-carbon unsaturated bonds [2020.01]	15/12	• • • Genes encoding animal proteins [5, 2006.01]
11/084	• • • Polymers containing vinyl alcohol units [2020.01]	15/13	• • • Immunoglobulins [5, 2006.01]
11/087	• • • Acrylic polymers [2020.01]	15/14	• • • Human serum albumins [5, 2006.01]
11/089	• • • obtained otherwise than by reactions only involving carbon-to-carbon unsaturated bonds [2020.01]	15/15	• • • Protease inhibitors, e.g. antithrombin, antitrypsin, hirudin [5, 2006.01]
11/091	• • • Phenol resins; Amino resins [2020.01]	15/16	• • • Hormones [5, 2006.01]
11/093	• • • Polyurethanes [2020.01]	15/17	• • • Insulins [5, 2006.01]
11/096	• • • Polyesters; Polyamides [2020.01]	15/18	• • • Growth hormones [5, 2006.01]
11/098	• • • formed in the presence of the enzymes or microbial cells [2020.01]	15/19	• • • Interferons; Lymphokines; Cytokines [5, 2006.01]
11/10	• • the carrier being a carbohydrate [3, 2006.01]	15/20	• • • Interferons [5, 2006.01]
11/12	• • Cellulose or derivatives thereof [3, 2006.01]	15/21	• • • Alpha-interferons [5, 2006.01]
11/14	• Enzymes or microbial cells immobilised on or in an inorganic carrier [3, 2006.01]	15/22	• • • Beta-interferons [5, 2006.01]
11/16	• Enzymes or microbial cells immobilised on or in a biological cell [3, 2006.01]		
11/18	• Multi-enzyme systems [3, 2006.01]		

Note(s) [3]

This group covers processes wherein there is a modification of the genetic material which would not normally occur in nature without intervention of man which produce a change in the gene structure which is passed on to succeeding generations.

- 15/23 • • • • • Gamma-interferons [5, 2006.01]
- 15/24 • • • • • Interleukins [5, 2006.01]
- 15/25 • • • • • Interleukin-1 [5, 2006.01]
- 15/26 • • • • • Interleukin-2 [5, 2006.01]
- 15/27 • • • • • Colony stimulating factors [5, 2006.01]
- 15/28 • • • • • Tumor necrosis factors [5, 2006.01]
- 15/29 • • • Genes encoding plant proteins, e.g. thaumatin [5, 2006.01]
- 15/30 • • • Genes encoding protozoal proteins, e.g. from Plasmodium, Trypanosoma, Eimeria [5, 2006.01]
- 15/31 • • • Genes encoding microbial proteins, e.g. enterotoxins [5, 2006.01]
- 15/32 • • • • Bacillus crystal proteins [5, 2006.01]
- 15/33 • • • • Genes encoding viral proteins [5, 2006.01]
- 15/34 • • • • Proteins from DNA viruses [5, 2006.01]
- 15/35 • • • • • Parvoviridae, e.g. feline panleukopenia virus, human parvovirus [5, 2006.01]
- 15/36 • • • • • Hepadnaviridae [5, 2006.01]
- 15/37 • • • • • Papovaviridae, e.g. papillomaviruses, polyomavirus, SV40 [5, 2006.01]
- 15/38 • • • • • Herpetoviridae, e.g. herpes simplex virus, varicella-zoster virus, Epstein-Barr virus, cytomegalovirus, pseudorabies virus [5, 2006.01]
- 15/39 • • • • • Poxviridae, e.g. vaccinia virus, variola virus [5, 2006.01]
- 15/40 • • • • • Proteins from RNA viruses, e.g. flaviviruses [5, 2006.01]
- 15/41 • • • • • Picornaviridae, e.g. rhinovirus, coxsackie viruses, echoviruses, enteroviruses [5, 2006.01]
- 15/42 • • • • • • Foot-and-mouth disease virus [5, 2006.01]
- 15/43 • • • • • • Poliovirus [5, 2006.01]
- 15/44 • • • • • Orthomyxoviridae, e.g. influenza virus [5, 2006.01]
- 15/45 • • • • • Paramyxoviridae, e.g. measles virus, mumps virus, Newcastle disease virus, canine distemper virus, rinderpest virus, respiratory syncytial viruses [5, 2006.01]
- 15/46 • • • • • Reoviridae, e.g. rotavirus, bluetongue virus, Colorado tick fever virus [5, 2006.01]
- 15/47 • • • • • Rhabdoviridae, e.g. rabies viruses, vesicular stomatitis virus [5, 2006.01]
- 15/48 • • • • • Retroviridae, e.g. bovine leukaemia virus, feline leukaemia virus [5, 2006.01]
- 15/49 • • • • • • Lentiviridae, e.g. immunodeficiency viruses such as HIV, visna-maedi virus, equine infectious anaemia virus [5, 2006.01]
- 15/50 • • • • • Coronaviridae, e.g. infectious bronchitis virus, transmissible gastroenteritis virus [5, 2006.01]
- 15/51 • • • • • Hepatitis viruses [5, 2006.01]
- 15/52 • • • Genes encoding for enzymes or proenzymes [5, 2006.01]
- 15/53 • • • • Oxidoreductases (1) [5, 2006.01]
- 15/54 • • • • Transferases (2) [5, 2006.01]
- 15/55 • • • • Hydrolases (3) [5, 2006.01]
- 15/56 • • • • • acting on glycosyl compounds (3.2), e.g. amylase, galactosidase, lysozyme [5, 2006.01]
- 15/57 • • • • • acting on peptide bonds (3.4) [5, 2006.01]
- 15/58 • • • • • Plasminogen activators, e.g. urokinase, TPA [5, 2006.01]
- 15/59 • • • • • Chymosin [5, 2006.01]
- 15/60 • • • • Lyases (4) [5, 2006.01]
- 15/61 • • • • Isomerases (5) [5, 2006.01]
- 15/62 • • • DNA sequences coding for fusion proteins [5, 2006.01]
- 15/63 • • Introduction of foreign genetic material using vectors; Vectors; Use of hosts therefor; Regulation of expression [5, 2006.01]
- 15/64 • • • General methods for preparing the vector, for introducing it into the cell or for selecting the vector-containing host [5, 2006.01]
- 15/65 • • • using markers (enzymes used as markers C12N 15/52) [5, 2006.01]
- 15/66 • • • General methods for inserting a gene into a vector to form a recombinant vector using cleavage and ligation; Use of non-functional linkers or adaptors, e.g. linkers containing the sequence for a restriction endonuclease [5, 2006.01]
- 15/67 • • • General methods for enhancing the expression [5, 2006.01]
- 15/68 • • • • Stabilisation of the vector [5, 2006.01]
- 15/69 • • • • Increasing the copy number of the vector [5, 2006.01]
- 15/70 • • • Vectors or expression systems specially adapted for E. coli [5, 2006.01]
- 15/71 • • • • Expression systems using regulatory sequences derived from the trp-operon [5, 2006.01]
- 15/72 • • • • Expression systems using regulatory sequences derived from the lac-operon [5, 2006.01]

Note(s) [5]

In this group, the following term is used with the meaning indicated:

- "fusion" means the fusion of two different proteins.

Note(s) [5]

In this group, the following expression is used with the meaning indicated:

- "non-functional linkers" means DNA sequences which are used to link DNA sequences and which have no known function of structural gene or regulating function.

Note(s) [5]

In this group:

- genes encoding for proenzymes are classified with the corresponding genes encoding enzymes;

- 15/73 • • • • Expression systems using phage lambda regulatory sequences [5, 2006.01]
- 15/74 • • • Vectors or expression systems specially adapted for prokaryotic hosts other than E. coli, e.g. Lactobacillus, Micromonospora [5, 2006.01]
- Note(s) [5]**
- This group covers the use of prokaryotes as hosts.
- 15/75 • • • • for Bacillus [5, 2006.01]
- 15/76 • • • • for Actinomyces; for Streptomyces [5, 2006.01]
- 15/77 • • • • for Corynebacterium; for Brevibacterium [5, 2006.01]
- 15/78 • • • • for Pseudomonas [5, 2006.01]
- 15/79 • • • Vectors or expression systems specially adapted for eukaryotic hosts [5, 2006.01]
- Note(s) [5]**
- This group covers the use of eukaryotes as hosts.
- 15/80 • • • • for fungi [5, 2006.01]
- 15/81 • • • • • for yeasts [5, 2006.01]
- 15/82 • • • • for plant cells [5, 2006.01]
- 15/83 • • • • • Viral vectors, e.g. cauliflower mosaic virus [5, 2006.01]
- 15/84 • • • • • Ti-plasmids [5, 2006.01]
- 15/85 • • • • for animal cells [5, 2006.01]
- 15/86 • • • • • Viral vectors [5, 2006.01]
- 15/861 • • • • • • Adenoviral vectors [7, 2006.01]
- 15/863 • • • • • • Poxviral vectors, e.g. vaccinia virus [7, 2006.01]
- 15/864 • • • • • • Parvoviral vectors [7, 2006.01]
- 15/866 • • • • • • Baculoviral vectors [7, 2006.01]
- 15/867 • • • • • • Retroviral vectors [7, 2006.01]
- 15/869 • • • • • • Herpesviral vectors [7, 2006.01]
- 15/87 • • Introduction of foreign genetic material using processes not otherwise provided for, e.g. co-transformation [5, 2006.01]
- 15/873 • • • Techniques for producing new embryos, e.g. nuclear transfer, manipulation of totipotent cells or production of chimeric embryos [2010.01]
- 15/877 • • • • Techniques for producing new mammalian cloned embryos [2010.01]
- 15/88 • • • using microencapsulation, e.g. using liposome vesicle [5, 2006.01]
- 15/89 • • • using microinjection [5, 2006.01]
- 15/90 • • • Stable introduction of foreign DNA into chromosome [5, 2006.01]