

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

Notes

- (1) In subclasses C12M to C12Q or C12S, and within each of these subclasses, in the absence of an indication to the contrary, classification is made in the last appropriate place. [3]
- (2) In this class, viruses, undifferentiated human, animal or plant cells, protozoa, tissues and unicellular algae are considered as micro-organisms. [3,5]
- (3) In this subclass, unless specifically provided for, undifferentiated human, animal or plant cells, protozoa, tissues and unicellular algae are classified together with micro-organisms. Sub-cellular parts, unless specifically provided for, are classified with the whole cell. [5]

Note

The codes of subclass C12R are only for use as indexing codes associated with subclasses C12C to C12Q or C12S, so as to provide information concerning the micro-organisms used in the processes classified in these subclasses. [3]

C12C BREWING OF BEER (cleaning of raw materials A23N; pitching or depitching machines, cellar tools C12L; propagating yeasts C12N 1/14; non-beverage ethanolic fermentation C12P 7/06)

Note

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

Subclass Index

RAW MATERIALS FOR PREPARING BEER	1/00, 3/00, 5/00	SPECIAL BEER	12/00
PREPARATION AND TREATMENT OF WORT; FERMENTATION PROCESSES FOR BEER	7/00, 11/00	BREWING DEVICES.....	13/00

1/00 Preparation of malt	5/00 Other raw materials for the preparation of beer
1/02 . Pretreatment of grains, e.g. washing, steeping	5/02 . Additives for beer
1/027 . Germinating [6]	5/04 . . Colouring additives
1/033 . . in boxes or drums [6]	7/00 Preparation of wort (malt extract C12C 1/18)
1/047 . . Influencing the germination by chemical or physical means [6]	7/01 . Pretreatment of malt, e.g. malt grinding [6]
1/053 . . . by irradiation or electric treatment [6]	7/04 . Preparation or treatment of the mash
1/067 . Drying [6]	7/047 . . part of the mash being unmalted cereal mash [6]
1/073 . . Processes or apparatus specially adapted to save or recover energy [6]	7/053 . . part of the mash being non-cereal material [6]
1/10 . . Drying on fixed supports	7/06 . . Mashing apparatus
1/12 . . Drying on moving supports	7/14 . Clarifying wort (Läuterung)
1/125 . Continuous or semi-continuous processes for steeping, germinating or drying [6]	7/16 . . by straining
1/13 . . with vertical transport of the grains [6]	7/165 . . . in mash filters [6]
1/135 . . with horizontal transport of the grains [6]	7/17 . . . in lautertuns [6]
1/15 . Grain or malt turning, charging or discharging apparatus [6]	7/175 . . by centrifuging [6]
1/16 . After-treatment of malt, e.g. malt cleaning, detachment of the germ	7/20 . . Boiling the beerwort (brew kettles C12C 13/02) [6]
1/18 . Preparation of malt extract or of special kinds of malt, e.g. caramel, black malt (malt products for use as foodstuffs A23L)	7/22 . . . Processes or apparatus specially adapted to save or recover energy [6]
3/00 Treatment of hops	7/24 . Clarifying beerwort between hop boiling and cooling [6]
3/02 . Drying	7/26 . Cooling beerwort; Clarifying beerwort during or after the cooling [6]
3/04 . Conserving; Storing; Packing	7/28 . After-treatment [6]
3/06 . . Powder or pellets from hops [6]	11/00 Fermentation processes for beer
3/08 . . Solvent extracts from hops [6]	11/02 . Pitching yeast
3/10 . . . using carbon dioxide [6]	11/06 . Acidifying the wort
3/12 . . Isomerised products from hops [6]	11/07 . Continuous fermentation [6]

C12C – C12G

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| <ul style="list-style-type: none"> 11/09 . Fermentation with immobilised yeast [6] 11/11 . Post fermentation treatments, e.g. carbonation, concentration (C12H takes precedence; containers with means specially adapted for effervescing potable liquids B65D 85/73) [6] <p>12/00 Processes specially adapted for making special kinds of beer [6]</p> <ul style="list-style-type: none"> 12/02 . Beer with low calorie content (C12C 12/04 takes precedence) [6] 12/04 . Beer with low alcohol content (removal of alcohol C12H 3/00) [6] | <ul style="list-style-type: none"> 13/00 Brewing devices, not covered by a single group of C12C 1/00 to C12C 12/04 [3,6] 13/02 . Brew kettles [3] 13/06 . . heated with fire [3] 13/08 . . with internal heating elements [6] 13/10 . Home brew equipment [6] |
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C12F RECOVERY OF BY-PRODUCTS OF FERMENTED SOLUTIONS; DENATURING OF, OR DENATURED, ALCOHOL [6]**Note**

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

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| <p>3/00 Recovery of by-products</p> <ul style="list-style-type: none"> 3/02 . of carbon dioxide 3/04 . . Recovery of volatile fermentation products from carbon dioxide 3/06 . from beer or wine (C12F 3/02 takes precedence; removal of yeast C12G 1/08) | <ul style="list-style-type: none"> 3/08 . . Recovery of alcohol from press residues or other waste material (from carbon dioxide C12F 3/04) 3/10 . from distillery slops <p>5/00 Preparation of denatured alcohol</p> |
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C12G WINE; OTHER ALCOHOLIC BEVERAGES; PREPARATION THEREOF (beer C12C)**Note**

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

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| <p>1/00 Preparation of wine or sparkling wine</p> <ul style="list-style-type: none"> 1/02 . Preparation of must from grapes; Must treatment or fermentation 1/022 . . Fermentation; Microbiological or enzymatic treatment [6] 1/024 . . . in a horizontally mounted cylindrical vessel (C12G 1/026 takes precedence) [6] 1/026 . . . in vessels with movable equipment for mixing the content [6] 1/028 . . . with thermal treatment of the grapes or the must [6] 1/032 . . . with recirculation of the must for pommage extraction [6] 1/036 . . . by use of a home wine making vessel [6] 1/04 . Sulfiting the must; Desulfiting 1/06 . Preparation of sparkling wine, e.g. champagne; Impregnating wine with carbon dioxide 1/067 . . Continuous processes [6] 1/073 . . Fermentation with immobilised yeast [6] | <ul style="list-style-type: none"> 1/08 . Removal of yeast (“degorgeage”) 1/09 . . Agitation, centrifugation or vibration of bottles [6] 1/10 . Deacidifying of wine [6] 1/12 . Processes for preventing winestone precipitation [6] <p>3/00 Preparation of other alcoholic beverages</p> <ul style="list-style-type: none"> 3/02 . by straight fermentation 3/04 . by mixing, e.g. liqueurs 3/06 . . with flavouring ingredients 3/07 . . . Flavouring with wood or wood extract; Pretreatment of the wood used therefor [6] 3/08 . by other methods for varying the composition of fermented solutions (removal of alcohol from alcoholic beverages to obtain alcohol-free or low-alcohol beverages C12H 3/00) 3/10 . . Increasing the alcohol content 3/12 . . . by distillation (distillation processes or apparatus, in general B01D 3/00) 3/14 . . . by freezing [6] |
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C12H PASTEURISATION, STERILISATION, PRESERVATION, PURIFICATION, CLARIFICATION, AGEING OF ALCOHOLIC BEVERAGES OR REMOVAL OF ALCOHOL THEREFROM (deacidifying wine C12G 1/10; preventing winestone precipitation C12G 1/12; simulation ageing by flavouring C12G 3/06) [6]

Note

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. [8]

Note

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

<p>1/00 Pasteurisation, sterilisation, preservation, purification, clarification, or ageing of alcoholic beverages</p> <p>1/02 . . combined with removal of precipitate or added materials, e.g. adsorption material</p> <p>1/04 . . with the aid of ion-exchange material or inert clarification material, e.g. adsorption material</p> <p>1/044 . . . with the aid of inorganic material [6]</p> <p>1/048 with silicon containing material [6]</p> <p>1/052 . . . with the aid of organic material [6]</p> <p>1/056 with the aid of polymers [6]</p> <p>1/06 . . Precipitation by physical means, e.g. by irradiation, vibrations</p> <p>1/065 . . . Separation by centrifugation [6]</p> <p>1/07 . . . Separation by filtration [6]</p> <p>1/075 by cross-flow filtration [6]</p> <p>1/08 . . . by heating</p> <p>1/10 . . Precipitation by chemical means</p> <p>1/12 . without precipitation</p>	<p>1/14 . . with non-precipitating compounds, e.g. sulfiting; Sequestration, e.g. with chelate-producing compounds</p> <p>1/15 . . . with enzymes [6]</p> <p>1/16 . . by physical means, e.g. irradiation</p> <p>1/18 . . . by heating</p> <p>1/20 in containers allowing for expansion of the contents</p> <p>1/22 . Ageing or ripening by storing, e.g. lagering of beer</p> <p>3/00 Removal of alcohol from alcoholic beverages to obtain alcohol-free or low-alcohol beverages (distillation or rectification of fermented solutions to obtain pure alcohol B01D 3/00; recovery of by-products of wine or beer other than low-alcohol beverages C12F 3/06; preparation of alcoholic beverages other than wine or beer by varying the composition of fermented solutions C12G 3/08) [6]</p> <p>3/02 . by evaporating [6]</p> <p>3/04 . using semi-permeable membranes [6]</p>
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C12J VINEGAR; ITS PREPARATION

Note

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

<p>1/00 Vinegar; Preparation; Purification</p> <p>1/02 . from wine</p> <p>1/04 . from alcohol</p>	<p>1/06 . from milk</p> <p>1/08 . Addition of flavouring ingredients</p> <p>1/10 . Apparatus</p>
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C12L PITCHING OR DEPITCHING MACHINES; CELLAR TOOLS (cleaning of casks B08B 9/00)

Note

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

- 3/00** Pitching or depitching machines
- 9/00** Venting devices for casks, barrels, or the like
- 11/00** Cellar tools

C12M APPARATUS FOR ENZYMOLOGY OR MICROBIOLOGY (installations for fermenting manure A01C 3/02; preservation of living parts of humans or animals A01N 1/02; physical or chemical apparatus in general B01; brewing apparatus C12C; fermentation apparatus for wine C12G; apparatus for preparing vinegar C12J 1/10) [3]

Note

Attention is drawn to Notes (1) to (3) following the title of class C12. [4]

Note

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

<p>1/00 Apparatus for enzymology or microbiology [3]</p> <p>Note</p> <p>This group covers:</p> <ul style="list-style-type: none"> – apparatus where micro-organisms or enzymes are produced or isolated; – apparatus where the characteristics of micro-organisms or enzymes are investigated, e.g. which growth factors are necessary; – apparatus specially adapted to employ micro-organisms or enzymes as “reactants” or biocatalysts; – apparatus of both the laboratory and industrial scale. [3] <p>1/02 . with agitation means; with heat exchange means [3]</p> <p>1/04 . with gas introduction means [3]</p> <p>1/06 . . with agitator, e.g. impeller [3]</p> <p>1/08 . . with draft tube [3]</p> <p>1/09 . . Flotation apparatus [5]</p> <p>1/10 . rotatably mounted [3]</p> <p>1/107 . with means for collecting fermentation gases, e.g. methane (producing methane by anaerobic treatment of sludge C02F 11/04) [5]</p> <p>1/113 . . with transport of the substrate during the fermentation [5]</p> <p>1/12 . with sterilisation, filtration, or dialysis means [3]</p> <p>1/14 . with means providing thin layers or with multi-level trays [3]</p> <p>1/16 . containing, or adapted to contain, solid media [3]</p> <p>1/18 . . Multiple fields or compartments [3]</p> <p>1/20 . . . Horizontal planar fields [3]</p>	<p>1/21 . Froth suppressors [5]</p> <p>1/22 . Petri type dish [3]</p> <p>1/24 . tube or bottle type [3]</p> <p>1/26 . Inoculator or sampler [3]</p> <p>1/28 . . being part of container [3]</p> <p>1/30 . . . Sampler being a swab [3]</p> <p>1/32 . . multiple field or continuous type [3]</p> <p>1/33 . Disintegrators [5]</p> <p>1/34 . Measuring or testing with condition measuring or sensing means, e.g. colony counters [3]</p> <p>1/36 . including condition or time responsive control, e.g. automatically controlled fermentors (controlling or regulating in general G05) [3]</p> <p>1/38 . . Temperature-responsive control [3]</p> <p>1/40 . Apparatus specially designed for the use of free, immobilised, or carrier-bound enzymes, e.g. apparatus containing a fluidised bed of immobilised enzymes [3]</p> <p>1/42 . Apparatus for the treatment of micro-organisms or enzymes with electrical or wave energy, e.g. magnetism, sonic wave [5]</p> <p>3/00 Tissue, human, animal or plant cell, or virus culture apparatus [3]</p> <p>3/02 . with means providing suspensions [3]</p> <p>3/04 . with means providing thin layers [3]</p> <p>3/06 . with filtration, ultrafiltration, inverse osmosis or dialysis means [5]</p> <p>3/08 . Apparatus for tissue disaggregation [5]</p> <p>3/10 . for culture in eggs [5]</p>
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C12N MICRO-ORGANISMS OR ENZYMES; COMPOSITIONS THEREOF (biocides, pest repellants or attractants, or plant growth regulators containing micro-organisms, viruses, microbial fungi, enzymes, fermentates, or substances produced by, or extracted from, micro-organisms or animal material A01N 63/00; food compositions A21, A23; medicinal preparations A61K; chemical aspects of, or use of materials for, bandages, dressings, absorbent pads or surgical articles A61L; fertilisers C05); **PROPAGATING, PRESERVING, OR MAINTAINING MICRO-ORGANISMS** (preservation of living parts of humans or animals A01N 1/02); **MUTATION OR GENETIC ENGINEERING; CULTURE MEDIA** (microbiological testing media C12Q) [3]

Notes

- (1) Attention is drawn to Notes (1) to (3) following the title of class C12. [3,4]
- (2) Biocidal, pest repellant, pest attractant or plant growth regulatory activity of compounds or preparations is further classified in subclass A01P. [8]
- (3) Therapeutic activity of single-cell proteins or enzymes is further classified in subclass A61P. [7]
- (4) 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. [8]

Note

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

Subclass Index

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

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| <p>1/00 Micro-organisms, 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 micro-organisms or compositions thereof; Processes of preparing or isolating a composition containing a micro-organism; Culture media therefor [3]</p> <p>1/02 . Separating micro-organisms from their culture media [3]</p> <p>1/04 . Preserving or maintaining viable micro-organisms (immobilised micro-organisms C12N 11/00) [3]</p> <p>1/06 . Lysis of micro-organisms [3]</p> <p>1/08 . Reducing the nucleic acid content [3]</p> <p>1/10 . Protozoa; Culture media therefor [3]</p> <p>1/11 . . . modified by introduction of foreign genetic material [5]</p> <p>1/12 . Unicellular algae; Culture media therefor (culture of multi-cellular plants A01G; as new plants A01H 13/00) [3]</p> <p>1/13 . . . modified by introduction of foreign genetic material [5]</p> <p>1/14 . Fungi (culture of mushrooms A01G 1/04; as new plants A01H 15/00); Culture media therefor [3]</p> <p>1/15 . . . modified by introduction of foreign genetic material [5]</p> <p>1/16 . . Yeasts; Culture media therefor [3]</p> <p>1/18 . . . Baker's yeast; Brewer's yeast [3]</p> <p>1/19 . . . modified by introduction of foreign genetic material [5]</p> <p>1/20 . Bacteria; Culture media therefor [3]</p> <p>1/21 . . . modified by introduction of foreign genetic material [5]</p> <p>1/22 . Processes using, or culture media containing, cellulose or hydrolysates thereof [3]</p> <p>1/24 . Processes using, or culture media containing, waste sulfite liquor [3]</p> <p>1/26 . Processes using, or culture media containing, hydrocarbons (refining of hydrocarbon oils by using micro-organisms C10G 32/00) [3]</p> <p>1/28 . . aliphatic [3]</p> <p>1/30 . . . having five or less carbon atoms [3]</p> <p>1/32 . Processes using, or culture media containing, lower alkanols, i.e. C₁ to C₆ [3]</p> <p>1/34 . Processes using foam culture [3]</p> <p>1/36 . Adaptation or attenuation of cells [3]</p> <p>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]</p> <p>3/00 Spore-forming or isolating processes [3]</p> | <p>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]</p> <p>5/02 . Propagation of single cells or cells in suspension; Maintenance thereof; Culture media therefor [3]</p> <p>5/04 . Plant cells or tissues [5]</p> <p>5/06 . Animal cells or tissues [5]</p> <p>5/08 . Human cells or tissues [5]</p> <p>5/10 . Cells modified by introduction of foreign genetic material, e.g. virus-transformed cells [5]</p> <p>5/12 . . Fused cells, e.g. hybridomas [5]</p> <p>5/14 . . . Plant cells [5]</p> <p>5/16 . . . Animal cells [5]</p> <p>5/18 Murine cells, e.g. mouse cells [5]</p> <p>5/20 one of the fusion partners being a B lymphocyte [5]</p> <p>5/22 . . . Human cells [5]</p> <p>5/24 one of the fusion partners being a B lymphocyte [5]</p> <p>5/26 . . . Cells resulting from interspecies fusion [5]</p> <p>5/28 one of the fusion partners being a human cell [5]</p> <p>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]</p> <p>7/01 . Viruses, e.g. bacteriophages, modified by introduction of foreign genetic material (vectors C12N 15/00) [5]</p> <p>7/02 . Recovery or purification [3]</p> <p>7/04 . Inactivation or attenuation; Producing viral sub-units [3]</p> <p>7/06 . . by chemical treatment [3]</p> <p>7/08 . . by serial passage of virus [3]</p> |
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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** (preparation of malt C12C 1/00) [3]

Note

In this group:

- proenzymes are classified with the corresponding enzymes; [5]
- 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. [3]

- 9/02 . Oxidoreductases (1.), e.g. luciferase [3]
- 9/04 . . acting on CHO groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3]
- 9/06 . . acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3]
- 9/08 . . acting on hydrogen peroxide as acceptor (1.11) [3]
- 9/10 . Transferases (2.) (ribonucleases C12N 9/22) [3]
- 9/12 . . transferring phosphorus containing groups, e.g. kinases (2.7) [3]
- 9/14 . Hydrolases (3.) [3]
- 9/16 . . acting on ester bonds (3.1) [3]
- 9/18 . . . Carboxylic ester hydrolases [3]
- 9/20 Triglyceride splitting, e.g. by means of lipase [3]
- 9/22 . . . Ribonucleases [3]
- 9/24 . . acting on glycosyl compounds (3.2) [3]
- 9/26 . . . acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3]
- 9/28 Alpha-amylase from microbial source, e.g. bacterial amylase [3]
- 9/30 Fungal source [3]
- 9/32 Alpha-amylase from plant source [3]
- 9/34 Glucoamylase [3]
- 9/36 . . . acting on beta-1, 4 bonds between N-acetylmuramic acid and 2-acetyl amino 2-deoxy-D-glucose, e.g. lysozyme [3]
- 9/38 . . . acting on beta-galactose-glycoside bonds, e.g. beta-galactosidase [3]
- 9/40 . . . acting on alpha-galactose-glycoside bonds, e.g. alpha-galactosidase [3]
- 9/42 . . . acting on beta-1, 4-glucosidic bonds, e.g. cellulase [3]
- 9/44 . . . acting on alpha-1, 6-glucosidic bonds, e.g. isoamylase, pullulanase [3]
- 9/46 Dextranase [3]
- 9/48 . . acting on peptide bonds, e.g. thromboplastin, leucine aminopeptidase (3.4) [3]
- 9/50 . . . Proteinases [3]
- 9/52 derived from bacteria [3]
- 9/54 bacteria being Bacillus [3]
- 9/56 Bacillus subtilis or Bacillus licheniformis [3]
- 9/58 derived from fungi [3]
- 9/60 from yeast [3]
- 9/62 from Aspergillus [3]
- 9/64 derived from animal tissue, e.g. rennin [3]
- 9/66 . . . Elastase [3]

- 9/68 . . . Plasmin, i.e. fibrinolysin [3]
- 9/70 . . . Streptokinase [3]
- 9/72 . . . Urokinase [3]
- 9/74 . . . Thrombin [3]
- 9/76 . . . Trypsin; Chymotrypsin [3]
- 9/78 . . acting on carbon to nitrogen bonds other than peptide bonds (3.5) [3]
- 9/80 . . . acting on amide bonds in linear amides [3]
- 9/82 Asparaginase [3]
- 9/84 Penicillin amidase [3]
- 9/86 . . . acting on amide bonds in cyclic amides, e.g. penicillinase [3]
- 9/88 . Lyases (4.) [3]
- 9/90 . Isomerases (5.) [3]
- 9/92 . . Glucose isomerase [3]
- 9/94 . Pancreatin [3]
- 9/96 . Stabilising an enzyme by forming an adduct or a composition; Forming enzyme conjugates [3]
- 9/98 . Preparation of granular or free-flowing enzyme compositions (C12N 9/96 takes precedence) [3]
- 9/99 . Enzyme inactivation by chemical treatment [3]

11/00 Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3]

- 11/02 . Enzymes or microbial cells being immobilised on or in an organic carrier [3]
- 11/04 . . entrapped within the carrier, e.g. gel, hollow fibre [3]
- 11/06 . . attached to the carrier via a bridging agent [3]
- 11/08 . . the carrier being a synthetic polymer [3]
- 11/10 . . the carrier being a carbohydrate [3]
- 11/12 . . . Cellulose or derivatives thereof [3]
- 11/14 . Enzymes or microbial cells being immobilised on or in an inorganic carrier [3]
- 11/16 . Enzymes or microbial cells being immobilised on or in a biological cell [3]
- 11/18 . Multi-enzyme systems [3]

13/00 Treatment of micro-organisms or enzymes with electrical or wave energy, e.g. magnetism, sonic waves [3]

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 micro-organisms 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]

Note

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. [3]

- 15/01 . Preparation of mutants without inserting foreign genetic material therein; Screening processes therefor [5]
- 15/02 . Preparation of hybrid cells by fusion of two or more cells, e.g. protoplast fusion [5]

- 15/03 . . . Bacteria [5]
 15/04 . . . Fungi [5]
 15/05 . . . Plant cells [5]
 15/06 . . . Animal cells [5]
 15/07 . . . Human cells [5]
 15/08 . . . Cells resulting from interspecies fusion [5]
 15/09 . . . Recombinant DNA-technology [5]
 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 micro-organisms or with enzymes C12P 19/34) [5]
 15/11 . . . DNA or RNA fragments; Modified forms thereof (DNA or RNA not used in recombinant technology C07H 21/00) [5]
 15/12 . . . Genes encoding animal proteins [5]
 15/13 . . . Immunoglobulins [5]
 15/14 . . . Human serum albumins [5]
 15/15 . . . Protease inhibitors, e.g. antithrombin, antitrypsin, hirudin [5]
 15/16 . . . Hormones [5]
 15/17 . . . Insulins [5]
 15/18 . . . Growth hormones [5]
 15/19 . . . Interferons; Lymphokines; Cytokines [5]
 15/20 . . . Interferons [5]
 15/21 . . . Alpha-interferons [5]
 15/22 . . . Beta-interferons [5]
 15/23 . . . Gamma-interferons [5]
 15/24 . . . Interleukins [5]
 15/25 . . . Interleukin-1 [5]
 15/26 . . . Interleukin-2 [5]
 15/27 . . . Colony stimulating factors [5]
 15/28 . . . Tumor necrosis factors [5]
 15/29 . . . Genes encoding plant proteins, e.g. thaumatin [5]
 15/30 . . . Genes encoding protozoal proteins, e.g. from Plasmodium, Trypanosoma, Eimeria [5]
 15/31 . . . Genes encoding microbial proteins, e.g. enterotoxins [5]
 15/32 . . . Bacillus crystal proteins [5]
 15/33 . . . Genes encoding viral proteins [5]
 15/34 . . . Proteins from DNA viruses [5]
 15/35 . . . Parvoviridae, e.g. feline panleukopenia virus, human parvovirus [5]
 15/36 . . . Hepadnaviridae [5]
 15/37 . . . Papovaviridae, e.g. papillomaviruses, polyomavirus, SV40 [5]
 15/38 . . . Herpetoviridae, e.g. herpes simplex virus, varicella-zoster virus, Epstein-Barr virus, cytomegalovirus, pseudorabies virus [5]
 15/39 . . . Poxviridae, e.g. vaccinia virus, variola virus [5]
 15/40 . . . Proteins from RNA viruses, e.g. flaviviruses [5]
 15/41 . . . Picornaviridae, e.g. rhinovirus, coxsackie viruses, echoviruses, enteroviruses [5]
 15/42 . . . Foot-and-mouth disease virus [5]
 15/43 . . . Poliovirus [5]
 15/44 . . . Orthomyxoviridae, e.g. influenza virus [5]
 15/45 . . . Paramyxoviridae, e.g. measles virus, mumps virus, Newcastle disease virus, canine distemper virus, rinderpest virus, respiratory syncytial viruses [5]
 15/46 . . . Reoviridae, e.g. rotavirus, bluetongue virus, Colorado tick fever virus [5]
 15/47 . . . Rhabdoviridae, e.g. rabies viruses, vesicular stomatitis virus [5]
 15/48 . . . Retroviridae, e.g. bovine leukaemia virus, feline leukaemia virus [5]
 15/49 . . . Lentiviridae, e.g. immunodeficiency viruses such as HIV, visna-maedi virus, equine infectious anaemia virus [5]
 15/50 . . . Coronaviridae, e.g. infectious bronchitis virus, transmissible gastroenteritis virus [5]
 15/51 . . . Hepatitis viruses [5]
 15/52 . . . Genes encoding for enzymes or proenzymes [5]
- Note**
- In this group:
- genes encoding for proenzymes are classified with the corresponding genes encoding 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 groups below in parenthesis. [5]
- 15/53 . . . Oxidoreductases (1) [5]
 15/54 . . . Transferases (2) [5]
 15/55 . . . Hydrolases (3) [5]
 15/56 . . . acting on glycosyl compounds (3.2), e.g. amylase, galactosidase, lysozyme [5]
 15/57 . . . acting on peptide bonds (3.4) [5]
 15/58 . . . Plasminogen activators, e.g. urokinase, TPA [5]
 15/59 . . . Chymosin [5]
 15/60 . . . Lyases (4) [5]
 15/61 . . . Isomerases (5) [5]
 15/62 . . . DNA sequences coding for fusion proteins [5]
- Note**
- In this group, the following term is used with the meaning indicated:
- “fusion” means the fusion of two different proteins. [5]
- 15/63 . . . Introduction of foreign genetic material using vectors; Vectors; Use of hosts therefor; Regulation of expression [5]
 15/64 . . . General methods for preparing the vector, for introducing it into the cell or for selecting the vector-containing host [5]
 15/65 . . . using markers (enzymes used as markers C12N 15/52) [5]

- 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]

Note

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. [5]

- 15/67 . . . General methods for enhancing the expression [5]
 15/68 Stabilisation of the vector [5]
 15/69 Increasing the copy number of the vector [5]
 15/70 . . . Vectors or expression systems specially adapted for E. coli [5]

Notes

- (1) This group covers the use of E. coli as host. [5]
 (2) Shuttle vectors also replicating in E. coli are classified according to the other host. [5]
- 15/71 Expression systems using regulatory sequences derived from the trp-operon [5]
 15/72 Expression systems using regulatory sequences derived from the lac-operon [5]
 15/73 Expression systems using phage lambda regulatory sequences [5]
 15/74 . . . Vectors or expression systems specially adapted for prokaryotic hosts other than E. coli, e.g. Lactobacillus, Micromonospora [5]

Note

This group covers the use of prokaryotes as hosts. [5]

- 15/75 for Bacillus [5]
 15/76 for Actinomyces; for Streptomyces [5]
 15/77 for Corynebacterium; for Brevibacterium [5]
 15/78 for Pseudomonas [5]
 15/79 . . . Vectors or expression systems specially adapted for eukaryotic hosts [5]

Note

This group covers the use of eukaryotes as hosts. [5]

- 15/80 for fungi [5]
 15/81 for yeasts [5]
 15/82 for plant cells [5]
 15/83 Viral vectors, e.g. cauliflower mosaic virus [5]
 15/84 Ti-plasmids [5]
 15/85 for animal cells [5]
 15/86 Viral vectors [5]
 15/861 Adenoviral vectors [7]
 15/863 Poxviral vectors, e.g. vaccinia virus [7]
 15/864 Parvoviral vectors [7]
 15/866 Baculoviral vectors [7]
 15/867 Retroviral vectors [7]
 15/869 Herpesviral vectors [7]
 15/87 . . Introduction of foreign genetic material using processes not otherwise provided for, e.g. co-transformation [5]
 15/88 . . . using micro-encapsulation, e.g. using liposome vesicle [5]
 15/89 . . . using micro-injection [5]
 15/90 . . . Stable introduction of foreign DNA into chromosome [5]

C12P FERMENTATION OR ENZYME-USING PROCESSES TO SYNTHESISE A DESIRED CHEMICAL COMPOUND OR COMPOSITION OR TO SEPARATE OPTICAL ISOMERS FROM A RACEMIC MIXTURE (fermentation processes to form a food composition A21, A23; compounds in general, see the relevant compound class, e.g. C01, C07; brewing of beer C12C; producing vinegar C12J; processes for producing enzymes C12N 9/00; DNA or RNA concerning genetic engineering, vectors, e.g. plasmids, or their isolation, preparation or purification C12N 15/00) [3]

Notes

- (1) This subclass covers both major and minor chemical modifications. [3]
 (2) Group C12P 1/00 covers processes for producing organic compounds not sufficiently identified to be classified in groups C12P 3/00 to C12P 37/00. Compounds identified only by their empirical formulae are not considered to be sufficiently identified. [3]
 (3) Attention is drawn to Notes (1) to (3) following the title of class C12. [4]
 (4) If a particular reaction is considered of interest, it is also classified in the relevant chemical compound class, e.g. C07, C08. [3]
 (5) In this subclass:
 – metal or ammonium salts of a compound are classified as that compound.
 – compositions are classified in the relevant compound groups. [3]

Note

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

Subclass Index

BIOSYNTHESIS OF CHEMICAL SUBSTANCES

Inorganic compounds3/00

Acyclic or carbocyclic organic compounds5/00 to 15/00
 peptides or proteins 21/00
 Carotenes..... 23/00

Tetracyclines	29/00
Prostaglandins	31/00
Steroids	33/00
Heterocyclic organic compounds	17/00
containing saccharide radicals	19/00
Riboflavin	25/00

Giberellin	27/00
Cephalosporin; penicillin	35/00; 37/00
SEPARATION OF OPTICAL ISOMERS	41/00
OTHER PROCESSES FOR BIOSYNTHESIS	
PREPARATIONS	1/00, 39/00

1/00 Preparation of compounds or compositions, not provided for in groups C12P 3/00 to C12P 39/00, by using micro-organisms or enzymes; General processes for the preparation of compounds or compositions by using micro-organisms or enzymes [3]

- 1/02 . by using fungi [3]
- 1/04 . by using bacteria [3]
- 1/06 . by using actinomycetales [3]

3/00 Preparation of elements or inorganic compounds except carbon dioxide [3]

5/00 Preparation of hydrocarbons [3]

- 5/02 . acyclic (producing methane by anaerobic treatment of sludge C02F 11/04) [3]

7/00 Preparation of oxygen-containing organic compounds [3]

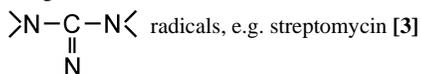
- 7/02 . containing a hydroxy group [3]
- 7/04 . . acyclic [3]
- 7/06 . . . Ethanol, i.e. non-beverage [3]
- 7/08 produced as by-product or from waste or cellulosic material substrate [3]
- 7/10 substrate containing cellulosic material [3]
- 7/12 substrate containing sulfite waste liquor or citrus waste [3]
- 7/14 Multiple stages of fermentation; Multiple types of micro-organisms or reuse for micro-organisms [3]
- 7/16 Butanols [3]
- 7/18 polyhydric [3]
- 7/20 Glycerol [3]
- 7/22 aromatic [3]
- 7/24 containing a carbonyl group [3]
- 7/26 Ketones [3]
- 7/28 Acetone-containing products [3]
- 7/30 produced from substrate containing inorganic compounds other than water [3]
- 7/32 produced from substrate containing inorganic nitrogen source [3]
- 7/34 produced from substrate containing protein as nitrogen source [3]
- 7/36 produced from substrate containing grain or cereal material [3]
- 7/38 Cyclopentanone- or cyclopentadione-containing products [3]
- 7/40 containing a carboxyl group [3]
- 7/42 Hydroxy carboxylic acids [3]
- 7/44 Polycarboxylic acids [3]
- 7/46 Dicarboxylic acids having four or less carbon atoms, e.g. fumaric acid, maleic acid [3]
- 7/48 Tricarboxylic acids, e.g. citric acid [3]
- 7/50 having keto groups, e.g. 2-ketoglutaric acid [3]
- 7/52 Propionic acid; Butyric acids [3]
- 7/54 Acetic acid (vinegar C12J) [3]
- 7/56 Lactic acid [3]

- 7/58 . . . Aldonic, ketoaldonic or saccharic acids (uronic acids C12P 19/00) [3]
- 7/60 2-Ketogulonic acid [3]
- 7/62 Carboxylic acid esters [3]
- 7/64 Fats; Fatty oils; Ester-type waxes; Higher fatty acids, i.e. having at least seven carbon atoms in an unbroken chain bound to a carboxyl group; Oxidised oils or fats [3]
- 7/66 containing the quinoid structure [3]
- 9/00 Preparation of organic compounds containing a metal or atom other than H, N, C, O, S, or halogen [3]
- 11/00 Preparation of sulfur-containing organic compounds [3]
- 13/00 Preparation of nitrogen-containing organic compounds [3]
- 13/02 . . Amides, e.g. chloramphenicol [3]
- 13/04 . . Alpha- or beta-amino acids [3]
- 13/06 . . . Alanine; Leucine; Isoleucine; Serine; Homoserine [3]
- 13/08 . . . Lysine; Diaminopimelic acid; Threonine; Valine [3]
- 13/10 . . . Citrulline; Arginine; Ornithine [3]
- 13/12 . . . Methionine; Cysteine; Cystine [3]
- 13/14 . . . Glutamic acid; Glutamine [3]
- 13/16 using surfactants, fatty acids or fatty acid esters, i.e. having at least seven carbon atoms in an unbroken chain bound to a carboxyl group or a carboxyl ester group [3]
- 13/18 using biotin or its derivatives [3]
- 13/20 . . . Aspartic acid; Asparagine [3]
- 13/22 . . . Tryptophan; Tyrosine; Phenylalanine; 3,4-Dihydroxyphenylalanine [3]
- 13/24 . . . Proline; Hydroxyproline; Histidine [3]
- 15/00 Preparation of compounds containing at least three condensed carbocyclic rings [3]
- 17/00 Preparation of heterocyclic carbon compounds with only O, N, S, Se, or Te as ring hetero atoms (C12P 13/04 to C12P 13/24 take precedence) [3]
- 17/02 . . Oxygen as only ring hetero atoms [3]
- 17/04 . . . containing a five-membered hetero ring, e.g. griseofulvin [3]
- 17/06 . . . containing a six-membered hetero ring, e.g. fluorescein [3]
- 17/08 . . . containing a hetero ring of at least seven ring members, e.g. zearalenone, macrolide aglycons [3]
- 17/10 . . Nitrogen as only ring hetero atom [3]
- 17/12 . . . containing a six-membered hetero ring [3]
- 17/14 . . Nitrogen or oxygen as hetero atom and at least one other diverse hetero ring atom in the same ring [3]
- 17/16 . . containing two or more hetero rings [3]
- 17/18 . . containing at least two hetero rings condensed among themselves or condensed with a common carbocyclic ring system, e.g. rifamycin [3]

19/00 Preparation of compounds containing saccharide radicals (ketoaldonic acids C12P 7/58) [3]**Note**

Attention is drawn to Note (3) following the title of subclass C07H, which defines the expression "saccharide radical". [3]

- 19/02 . Monosaccharides (2-ketogulonic acid C12P 7/60) [3]
- 19/04 . Polysaccharides, i.e. compounds containing more than five saccharide radicals attached to each other by glycosidic bonds [3]
- 19/06 . . Xanthan, i.e. Xanthomonas-type heteropolysaccharides [3]
- 19/08 . . Dextran [3]
- 19/10 . . Pullulan [3]
- 19/12 . Disaccharides [3]
- 19/14 . produced by the action of a carbohydrase, e.g. by alpha-amylase [3]
- 19/16 . produced by the action of an alpha-1, 6-glucosidase, e.g. amylose, debranched amylopectin (non-biological hydrolysis of starch C08B 30/00) [3]
- 19/18 . produced by the action of a glycosyl transferase, e.g. alpha-, beta- or gamma-cyclodextrins [3]
- 19/20 . produced by the action of an exo-1, 4 alpha-glucosidase, e.g. dextrose [3]
- 19/22 . produced by the action of a beta-amylase, e.g. maltose [3]
- 19/24 . produced by the action of an isomerase, e.g. fructose [3]
- 19/26 . Preparation of nitrogen-containing carbohydrates [3]
- 19/28 . . N-glycosides [3]
- 19/30 . . . Nucleotides [3]
- 19/32 having a condensed ring system containing a six-membered ring having two nitrogen atoms in the same-ring, e.g. purine nucleotides, nicotineamide-adenine dinucleotide [3]
- 19/34 Polynucleotides, e.g. nucleic acids, oligoribonucleotides [3]
- 19/36 Dinucleotides, e.g. nicotineamide-adenine dinucleotide phosphate [3]
- 19/38 . . . Nucleosides [3]
- 19/40 having a condensed ring system containing a six-membered ring having two nitrogen atoms in the same ring, e.g. purine nucleosides [3]
- 19/42 . . . Cobalamins, i.e. vitamin B₁₂, LLD factor [3]
- 19/44 . Preparation of O-glycosides, e.g. glucosides [3]
- 19/46 . . having an oxygen atom of the saccharide radical bound to a cyclohexyl radical, e.g. kasugamycin [3]
- 19/48 . . . the cyclohexyl radical being substituted by two or more nitrogen atoms, e.g. destomycin, neamin [3]
- 19/50 having two saccharide radicals bound through only oxygen to adjacent ring carbon atoms of the cyclohexyl radical, e.g.ambutyrosin, ribostamycin [3]
- 19/52 containing three or more saccharide radicals, e.g. neomycin, lividomycin [3]
- 19/54 . . . the cyclohexyl radical being bound directly to a nitrogen atom of two or more



- 19/56 . . having an oxygen atom of the saccharide radical directly bound to a condensed ring system having three or more carbocyclic rings, e.g. daunomycin, adriamycin [3]
- 19/58 . . having an oxygen atom of the saccharide radical directly bound through only acyclic carbon atoms to a non-saccharide heterocyclic ring, e.g. bleomycin, phleomycin [3]
- 19/60 . . having an oxygen of the saccharide radical directly bound to a non-saccharide heterocyclic ring or a condensed ring system containing a non-saccharide heterocyclic ring, e.g. coumermycin, novobiocin [3]
- 19/62 . . . the hetero ring having eight or more ring members and only oxygen as ring hetero atoms, e.g. erythromycin, spiramycin, nystatin [3]
- 19/64 . Preparation of S-glycosides, e.g. lincomycin [3]
- 21/00 **Preparation of peptides or proteins** (single-cell protein C12N 1/00) [3]
- 21/02 . having a known sequence of two or more amino acids, e.g. glutathione [3]
- 21/04 . . Cyclic or bridged peptides or polypeptides, e.g. bacitracin (cyclised by -S-S-bonds only C12P 21/02) [3]
- 21/06 . produced by the hydrolysis of a peptide bond, e.g. hydrolysate products (preparing foodstuffs by protein hydrolysis A23J 3/00) [3]
- 21/08 . Monoclonal antibodies [5]

- 23/00 **Preparation of compounds containing a cyclohexene ring having an unsaturated side chain containing at least ten carbon atoms bound by conjugated double bonds, e.g. carotenes** (containing hetero-rings C12P 17/00) [3]
- 25/00 **Preparation of compounds containing alloxazine or isoalloxazine nucleus, e.g. riboflavin** [3]
- 27/00 **Preparation of compounds containing a gibbane ring system, e.g. gibberellin** [3]
- 29/00 **Preparation of compounds containing a naphthacene ring system, e.g. tetracycline** (C12P 19/00 takes precedence) [3]
- 31/00 **Preparation of compounds containing a five-membered ring having two side-chains in ortho position to each other, and having at least one oxygen atom directly bound to the ring in ortho position to one of the side-chains, one side-chain containing, not directly bound to the ring, a carbon atom having three bonds to hetero atoms with at the most one bond to halogen, and the other side-chain having at least one oxygen atom bound in gamma-position to the ring, e.g. prostaglandins** [3]
- 33/00 **Preparation of steroids** [3]

Note

Attention is drawn to Note (1) following the title of subclass C07J, which explains what is covered by the term "steroids". [3]

Note

In groups C12P 33/02 to C12P 33/20, the following terms are used with the meaning indicated:

- “acting”, “forming”, “hydroxylating”, “dehydroxylating” or “dehydrogenating” means the action of a micro-organism or enzyme rather than other chemical action. [3]

33/02	. Dehydrogenating; Dehydroxylating [3]
33/04	. . Forming an aryl ring from A ring [3]
33/06	. Hydroxylating [3]
33/08	. . at 11 position [3]
33/10	. . . at 11alpha-position [3]
33/12	. Acting on D ring [3]
33/14	. . Hydroxylating at 16 position [3]
33/16	. . Acting at 17 position [3]
33/18	. . . Hydroxylating at 17 position [3]
33/20	. containing heterocyclic rings [3]

35/00 Preparation of compounds having a 5-thia-1-azabicyclo [4.2.0] octane ring system, e.g. cephalosporin [3]

35/02	. by desacylation of the substituent in the 7 position [3]
35/04	. by acylation of the substituent in the 7 position [3]
35/06	. Cephalosporin C; Derivatives thereof [3]
35/08	. disubstituted in the 7 position [3]

37/00 Preparation of compounds having a 4-thia-1-azabicyclo [3.2.0] heptane ring system, e.g. penicillin [3]

37/02	. in presence of phenylacetic acid or phenylacetamide or their derivatives [3]
37/04	. by acylation of the substituent in the 6 position [3]
37/06	. by desacylation of the substituent in the 6 position [3]

39/00 Processes involving micro-organisms of different genera in the same process, simultaneously [3]**41/00 Processes using enzymes or micro-organisms to separate optical isomers from a racemic mixture [4]****C12Q MEASURING OR TESTING PROCESSES INVOLVING ENZYMES OR MICRO-ORGANISMS (immunoassay G01N 33/53); COMPOSITIONS OR TEST PAPERS THEREFOR; PROCESSES OF PREPARING SUCH COMPOSITIONS; CONDITION-RESPONSIVE CONTROL IN MICROBIOLOGICAL OR ENZYMOLOGICAL PROCESSES [3]****Notes**

- (1) This subclass does not cover the observation of the progress or of the result of processes specified in this subclass by any of the methods specified in groups G01N 3/00 to G01N 29/00, which is covered by subclass G01N. [3]
- (2) In this subclass, the following expression is used with the meaning indicated:
 - “involving”, when used in relation to a substance, includes the testing for the substance as well as employing the substance as a determinant or reactant in a test for a different substance. [3]
- (3) Attention is drawn to Notes (1) to (3) following the title of class C12. [4]
- (4) In this subclass, test media are classified in the appropriate group for the relevant test process. [3]

Note

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

1/00	Measuring or testing processes involving enzymes or micro-organisms (measuring or testing apparatus with condition measuring or sensing means, e.g. colony counters, C12M 1/34); Compositions therefor; Processes of preparing such compositions [3]	1/26	. involving oxidoreductase [3]
1/02	. involving viable micro-organisms [3]	1/28	. . involving peroxidase [3]
1/04	. . Determining presence or kind of micro-organism; Use of selective media for testing antibiotics or bacteriocides; Compositions containing a chemical indicator therefor [3]	1/30	. . involving catalase [3]
1/06	. . . Quantitative determination [3]	1/32	. . involving dehydrogenase [3]
1/08	. . . using multifield media [3]	1/34	. involving hydrolase [3]
1/10	. . . Enterobacteria [3]	1/37	. . involving peptidase or proteinase [5]
1/12	. . . Nitrate to nitrite reducing bacteria [3]	1/40	. . involving amylase [3]
1/14	. . . Streptococcus; Staphylococcus [3]	1/42	. . involving phosphatase [3]
1/16	. . . using radioactive material [3]	1/44	. . involving esterase [3]
1/18	. . Testing for antimicrobial activity of a material [3]	1/46	. . . involving cholinesterase [3]
1/20	. . . using multifield media [3]	1/48	. involving transferase [3]
1/22	. . Testing for sterility conditions [3]	1/50	. . involving creatine phosphokinase [3]
1/24	. . Methods of sampling, or inoculating or spreading a sample; Methods of physically isolating an intact micro-organism [3]	1/52	. . involving transaminase [3]
1/25	. involving enzymes not classifiable in groups C12Q 1/26 to C12Q 1/70 [5]	1/527	. involving lyase [5]
		1/533	. involving isomerase [5]
		1/54	. involving glucose or galactose [3]
		1/56	. involving blood clotting factors, e.g. involving thrombin, thromboplastin, fibrinogen [3]
		1/58	. involving urea or urease [3]
		1/60	. involving cholesterol [3]
		1/61	. involving triglycerides [5]
		1/62	. involving uric acid [3]
		1/64	. Geomicrobiological testing, e.g. for petroleum [3]

1/66 . involving luciferase [3]

1/68 . involving nucleic acids [3]

1/70 . involving virus or bacteriophage [3]

3/00 Condition-responsive control processes (apparatus therefor C12M 1/36; controlling or regulating in general G05) [3]

C12R INDEXING SCHEME ASSOCIATED WITH SUBCLASSES C12C TO C12Q OR C12S, RELATING TO MICRO-ORGANISMS [3]

Notes

- (1) This subclass constitutes an indexing scheme associated with the other subclasses of class C12, relating to micro-organisms used in the processes classified in subclasses C12C to C12Q or C12S. [3]
- (2) The bacteria terminology is based on “Bergey’s Manual of Determinative Bacteriology”, Eighth Edition, 1975. [3]

1/00	Micro-organisms [3]	1/325	. . . Mycobacterium avium [3]
1/01	. Bacteria or actinomycetales [3]	1/33	. . . Mycobacterium fortuitum [3]
1/02	. . Acetobacter [3]	1/34	. . . Mycobacterium smegmatis [3]
1/025	. . Achromobacter [3]	1/35	. . Mycoplasma [3]
1/03	. . Actinomadura [3]	1/36	. . Neisseria [3]
1/04	. . Actinomyces [3]	1/365	. . Nocardia [3]
1/045	. . Actinoplanes [3]	1/37	. . Proteus [3]
1/05	. . Alcaligenes [3]	1/38	. . Pseudomonas [3]
1/06	. . Arthrobacter [3]	1/385	. . . Pseudomonas aeruginosa [3]
1/065	. . Azotobacter [3]	1/39	. . . Pseudomonas fluorescens [3]
1/07	. . Bacillus [3]	1/40	. . . Pseudomonas putida [3]
1/08	. . . Bacillus brevis [3]	1/41	. . Rhizobium [3]
1/085	. . . Bacillus cereus [3]	1/42	. . Salmonella [3]
1/09	. . . Bacillus circulans [3]	1/425	. . Serratia [3]
1/10	. . . Bacillus licheniformis [3]	1/43	. . . Serratia marcescens [3]
1/11	. . . Bacillus megaterium [3]	1/44	. . Staphylococcus [3]
1/12	. . . Bacillus polymyxa [3]	1/445	. . . Staphylococcus aureus [3]
1/125	. . . Bacillus subtilis [3]	1/45	. . . Staphylococcus epidermidis [3]
1/13	. . Brevibacterium [3]	1/46	. . Streptococcus [3]
1/14	. . Chainia [3]	1/465	. . Streptomyces [3]
1/145	. . Clostridium [3]	1/47	. . . Streptomyces albus [3]
1/15	. . Corynebacterium [3]	1/48	. . . Streptomyces antibioticus [3]
1/16	. . . Corynebacterium diphtheriae [3]	1/485	. . . Streptomyces aureofaciens [3]
1/165	. . . Corynebacterium poinsettiae [3]	1/49	. . . Streptomyces aureus [3]
1/17	. . . Corynebacterium pyogenes [3]	1/50	. . . Streptomyces bikiniensis [3]
1/18	. . Erwinia [3]	1/51	. . . Streptomyces candidus [3]
1/185	. . Escherichia [3]	1/52	. . . Streptomyces chartreusis [3]
1/19	. . . Escherichia coli [3]	1/525	. . . Streptomyces diastatochromogenes [3]
1/20	. . Flavobacterium [3]	1/53	. . . Streptomyces filipinensis [3]
1/21	. . Haemophilus [3]	1/54	. . . Streptomyces fradiae [3]
1/22	. . Klebsiella [3]	1/545	. . . Streptomyces griseus [3]
1/225	. . Lactobacillus [3]	1/55	. . . Streptomyces hygroscopicus [3]
1/23	. . . Lactobacillus acidophilus [3]	1/56	. . . Streptomyces lavendulae [3]
1/24	. . . Lactobacillus brevis [3]	1/565	. . . Streptomyces lincolnensis [3]
1/245	. . . Lactobacillus casei [3]	1/57	. . . Streptomyces noursei [3]
1/25	. . . Lactobacillus plantarum [3]	1/58	. . . Streptomyces olivaceus [3]
1/26	. . Methylomonas [3]	1/585	. . . Streptomyces platensis [3]
1/265	. . Micrococcus [3]	1/59	. . . Streptomyces rimosus [3]
1/27	. . . Micrococcus flavus [3]	1/60	. . . Streptomyces sparsogenes [3]
1/28	. . . Micrococcus glutamicus [3]	1/61	. . . Streptomyces venezuelae [3]
1/285	. . . Micrococcus lysodeikticus [3]	1/62	. . Streptosporangium [3]
1/29	. . Micromonospora [3]	1/625	. . Streptoverticillium [3]
1/30	. . . Micromonospora chalcone [3]	1/63	. . Vibrio [3]
1/31	. . . Micromonospora purpurea [3]	1/64	. . Xanthomonas [3]
1/32	. . Mycobacterium [3]	1/645	. Fungi [3]

1/65	. . Absidia [3]	1/79	. . Paecilomyces [3]
1/66	. . Aspergillus [3]	1/80	. . Penicillium [3]
1/665	. . . Aspergillus awamori [3]	1/81	. . . Penicillium brevis [3]
1/67	. . . Aspergillus flavus [3]	1/82	. . . Penicillium chrysogenum [3]
1/68	. . . Aspergillus fumigatus [3]	1/825	. . . Penicillium notatum [3]
1/685	. . . Aspergillus niger [3]	1/83	. . . Penicillium patulum [3]
1/69	. . . Aspergillus oryzae [3]	1/84	. . Pichia [3]
1/70	. . . Aspergillus ustus [3]	1/845	. . Rhizopus [3]
1/71	. . . Aspergillus wentii [3]	1/85	. . Saccharomyces [3]
1/72	. . Candida [3]	1/86	. . . Saccharomyces carlsbergensis [3]
1/725	. . . Candida albicans [3]	1/865	. . . Saccharomyces cerevisiae [3]
1/73	. . . Candida lipolytica [3]	1/87	. . . Saccharomyces lactis [3]
1/74	. . . Candida tropicalis [3]	1/88	. . Torulopsis [3]
1/745	. . Cephalosporium [3]	1/885	. . Trichoderma [3]
1/75	. . . Cephalosporium acremonium [3]	1/89	. Algae [3]
1/76	. . . Cephalosporium coeruleum [3]	1/90	. Protozoa [3]
1/765	. . . Cephalosporium crotocinigenum [3]	1/91	. Cell lines [3,7]
1/77	. . Fusarium [3]	1/92	. Viruses [5,7]
1/78	. . Hansenula [3]	1/93	. . Animal viruses [7]
1/785	. . Mucor [3]	1/94	. . Plant viruses [7]

C12S PROCESSES USING ENZYMES OR MICRO-ORGANISMS TO LIBERATE, SEPARATE OR PURIFY A PRE-EXISTING COMPOUND OR COMPOSITION (biological treatment of water, waste water, or sewage C02F 3/00, of sludge C02F 11/02; processes using enzymes or micro-organisms to separate optical isomers from a racemic mixture C12P 41/00); **PROCESSES USING ENZYMES OR MICRO-ORGANISMS TO TREAT TEXTILES OR TO CLEAN SOLID SURFACES OF MATERIALS** [5]

Notes

- (1) This subclass covers processes already provided for in:
- Section A: A21, A23, A61L, A62D;
 - Section B: B01D, B08B, B09C;
 - Section C: C01, C05F, C08, C09B, C09H, C10G, C13, C14C, C21B, C22B, C23F, C23G;
 - Section D: D01C, D01F, D06L, D06M, D06P, D21C, D21H;
 - Section E: E21B;
 - Section F: F24F, F24J, F26B;
 - Section H: H01M.
- This subclass is intended to provide a basis for a complete search to be made with respect to the subject matter defined by the subclass title and, therefore, all relevant information is classified in this subclass, even if classified elsewhere. [5]
- (2) Attention is drawn to Notes (1) to (3) following the title of class C12. [5]
- (3) The classification symbols of this subclass are not listed first when printed on the patent documents. [5]

Note

In this subclass, it is desirable to add the indexing codes of subclass C12R. [6]

1/00	Treatment of petroleum oils, shale oils or sand oils [5]	3/18	. Recovery or purification of glyceridic oils, fats, ester-type waxes or fatty acids [5]
1/02	. Desulfurising [5]	3/20	. Removal of nucleic acids from intact or disrupted cells [5]
3/00	Treatment of animal or plant materials or micro-organisms [5]	3/22	. Treatment of blood fractions [5]
3/02	. Recovery or purification of carbohydrate material [5]	3/24	. Treatment of animal secretions or organs [5]
3/04	. . Cellulose, e.g. plant fibres [5]	5/00	Treatment of emulsions, gases or foams [5]
3/06	. . . Treatment of hemp or flax [5]	7/00	Treatment of hides, e.g. depilating, bating [5]
3/08	. . . in the production of paper pulp [5]	9/00	Cleaning solid surfaces of materials [5]
3/10	. . Treatment of sugar or molasses [5]	11/00	Treatment of textiles, e.g. cleaning [5]
3/12	. . Treatment of pectin or starch [5]	13/00	Processes not provided for in groups C12S 1/00 to C12S 11/00 [5]
3/14	. Recovery or purification of proteinaceous material [5]		
3/16	. . Collagen or gelatin [5]		