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

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

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

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

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11/06

. Acidifying the wort

11/07 11/09 11/11	 Continuous fermentation [6] Fermentation with immobilised yeast [6] Post fermentation treatments, e.g. carbonation, concentration (C12H takes precedence; containers with means specially adapted for effervescing potable liquids B65D 85/73) [6] 	13/00 13/02 13/06 13/08 13/10	Brewing devices, not covered by a single group of C12C 1/00 to C12C 12/04 [3,6] Brew kettles [3] heated with fire [3] with internal heating elements [6] Home brew equipment [6]
12/00	Processes specially adapted for making special kinds of beer [6]		11 .,
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]		

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]

3/00 3/02 3/04 3/06	 Recovery of by-products of carbon dioxide Recovery of volatile fermentation products from carbon dioxide from beer or wine (C12F 3/02 takes precedence; removal of yeast C12G 1/08) 	3/08 3/10 5/00	Recovery of alcohol from press residues or other waste material (from carbon dioxide C12F 3/04) from distillery slops Preparation of denatured alcohol
3/06	. from beer or wine (C12F 3/02 takes precedence;	5/00	Preparation of denatured alcohol

C12G WINE; OTHER ALCOHOLIC BEVERAGES; PREPARATION THEREOF (beer C12C)

<u>Note</u>

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

1/00	Preparation of wine or sparkling wine	1/08	. Removal of yeast ("degorgeage")
1/02	 Preparation of must from grapes; Must treatment or 	1/09	Agitation, centrifugation or vibration of bottles [6]
	fermentation	1/10	. Deacidifying of wine [6]
1/022	Fermentation; Microbiological or enzymatic treatment [6]	1/12	. Processes for preventing winestone precipitation [6]
1/024	in a horizontally mounted cylindrical vessel	3/00	Preparation of other alcoholic beverages
	(C12G 1/026 takes precedence) [6]	3/02	 by straight fermentation
1/026	in vessels with movable equipment for mixing	3/04	by mixing, e.g. liqueurs
	the content [6]	3/06	with flavouring ingredients
1/028	with thermal treatment of the grapes or the	3/07	Flavouring with wood or wood extract;
	must [6]		Pretreatment of the wood used therefor [6]
1/032	with recirculation of the must for pompage	3/08	. by other methods for varying the composition of
	extraction [6]		fermented solutions (removal of alcohol from
1/036	by use of a home wine making vessel [6]		alcoholic beverages to obtain alcohol-free or low-
1/04	Sulfiting the must; Desulfiting		alcohol beverages C12H 3/00)
1/06	 Preparation of sparkling wine, e.g. champagne; 	3/10	Increasing the alcohol content
	Impregnating wine with carbon dioxide	3/12	by distillation (distillation processes or
1/067	Continuous processes [6]		apparatus, in general B01D 3/00)
1/073	Fermentation with immobilised yeast [6]	3/14	by freezing [6]

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]

1/00 Pasteurisation, sterilisation, pre- purification, clarification, or age beverages	*	 with non-precipitating compounds, e.g. sulfiting; Sequestration, e.g. with chelate-producing compounds
1/02 . combined with removal of prec materials, e.g. adsorption mater 1/04 . with the aid of ion-exchange clarification material, e.g. ad 1/044 with the aid of inorganic 1/048 with silicon containing 1/052 with the aid of organic materials of the aid of organic materials of the aid of polyme 1/06 with the aid of polyme 1/06 with the aid of polyme 1/06 Separation by physical materials of the aid of polyme 1/065 Separation by centrifugat 1/07 Separation by filtration [6] 1/075 by cross-flow filtration 1/08 by heating	rial 1/16 e material or inert 1/18 dsorption material 1/20 material [6] g material [6] 1/22 aterial [6] 3/00 eans, e.g. by ion [6]	 with enzymes [6] by physical means, e.g. irradiation by heating in containers allowing for expansion of the contents . Ageing or ripening by storing, e.g. lagering of beer 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]
1/10 . Precipitation by chemical model 1/12 . without precipitation	ans 3/02 3/04	 by evaporating [6] using semi-permeable membranes [6]

C12J VINEGAR; ITS PREPARATION

Note

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

1/00	Vinegar; Preparation; Purification	1/06 • from milk
1/02	. from wine	1/08 . Addition of flavouring ingredients
1/04	. from alcohol	1/10 . Apparatus

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]

1/00	Apparatus for enzymology or microbiology [3]	1/21	. Froth suppressors [5]
		1/22	. Petri type dish [3]
<u>Note</u>		1/24	. tube or bottle type [3]
	This group covers:	1/26	. Inoculator or sampler [3]
	apparatus where micro-organisms or enzymes are	1/28	being part of container [3]
	produced or isolated;	1/30	Sampler being a swab [3]
	 apparatus where the characteristics of micro- 	1/32	multiple field or continuous type [3]
	organisms or enzymes are investigated, e.g. which	1/33	. Disintegrators [5]
	growth factors are necessary; apparatus specially adapted to employ micro-	1/34	. Measuring or testing with condition measuring or
	 apparatus specially adapted to employ micro- organisms or enzymes as "reactants" or biocatalysts; 		sensing means, e.g. colony counters [3]
	 apparatus of both the laboratory and industrial scale. 	1/36	. including condition or time responsive control,
	[3]		e.g. automatically controlled fermentors (controlling or regulating in general G05) [3]
		1/38	. Temperature-responsive control [3]
1/02	. with agitation means; with heat exchange means [3]	1/40	. Apparatus specially designed for the use of free,
1/04	. with gas introduction means [3]	1740	immobilised, or carrier-bound enzymes,
1/06	with agitator, e.g. impeller [3]		e.g. apparatus containing a fluidised bed of
1/08	with draft tube [3]		immobilised enzymes [3]
1/09	Flotation apparatus [5]	1/42	. Apparatus for the treatment of micro-organisms or
1/10	. rotatably mounted [3]		enzymes with electrical or wave energy,
1/107	. with means for collecting fermentation gases,		e.g. magnetism, sonic wave [5]
	e.g. methane (producing methane by anaerobic treatment of sludge C02F 11/04) [5]	3/00	Tissue, human, animal or plant cell, or virus culture apparatus [3]
1/113	with transport of the substrate during the	3/02	with means providing suspensions [3]
	fermentation [5]	3/04	with means providing thin layers [3]
1/12	with sterilisation, filtration, or dialysis means [3]	3/06	with filtration, ultrafiltration, inverse osmosis or
1/14	with means providing thin layers or with multi-level		dialysis means [5]
1/16	trays [3]	3/08	. Apparatus for tissue disaggregation [5]
1/16	. containing, or adapted to contain, solid media [3]	3/10	. for culture in eggs [5]
1/18	Multiple fields or compartments [3]		
1/20	Horizontal planar fields [3]		

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]

- (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]

<u>Note</u>

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

Subclass index

	ORGANISMS; SPORES; ERENTIATED CELLS; VIRUSES1/00; 3/00; 5/00; 7/00; 11/00	TREATMENT WITH ELECTRICAL OR WAVE ENERGY13/00 MUTATION OR GENETIC ENGINEERING15/00
ENZYM	ES	
1/00	Micro-organisms, e.g. protozoa; Compositions thereof (medicinal preparations containing material from protozoa, bacteria or viruses A61K 35/66, from place A61K 36/02, from funci A61K 36/06, proporting	5/00 Undifferentiated human, animal or plant cells, e.g. cell lines; Tissues; Cultivation or maintenance thereof; Culture media therefor (plant reproduction by tissue outly to techniques AOLH 4/00) [3-5]
	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	tissue culture techniques A01H 4/00) [3,5] 5/02 • Propagation of single cells or cells in suspension; Maintenance thereof; Culture media therefor [3]
	propagating, maintaining or preserving micro-	5/04 . Plant cells or tissues [5]
	organisms or compositions thereof; Processes of	5/07 . Animal cells or tissues [2010.01]
	preparing or isolating a composition containing a micro-organism; Culture media therefor [3]	
1/02	Separating micro-organisms from their culture media [3]	Note The last place priority rule does not apply between the
1/04	Preserving or maintaining viable micro-organisms (immobilised micro-organisms C12N 11/00) [3]	subgroups of this group. [2010.01]
1/06	. Lysis of micro-organisms [3]	5/071 Vertebrate cells or tissues, e.g. human cells or
1/08	. Reducing the nucleic acid content [3]	tissues [2010.01]
1/10	 Protozoa; Culture media therefor [3] 	5/073 Embryonic cells or tissues; Foetal cells or
1/11	 modified by introduction of foreign genetic material [5] 	tissues [2010.01] 5/0735 Embryonic stem cells; Embryonic germ
1/12	 Unicellular algae; Culture media therefor (culture of multi-cellular plants A01G; as new plants A01H 13/00) [3] 	cells [2010.01] 5/074 Adult stem cells [2010.01]
1/13	 . modified by introduction of foreign genetic material [5] 	5/075 Oocytes; Oogonia [2010.01] 5/076 Sperm cells; Spermatogonia [2010.01] 5/077 Mesenchymal cells, e.g. bone cells, cartilage
1/14	• Fungi (culture of mushrooms A01G 1/04; as new plants A01H 15/00); Culture media therefor [3]	cells, marrow stromal cells, fat cells or muscle cells [2010.01]
1/15	 modified by introduction of foreign genetic material [5] 	5/0775 Mesenchymal stem cells; Adipose-tissue derived stem cells [2010.01]
1/16	. Yeasts; Culture media therefor [3]	5/078 Cells from blood or from the immune
1/18	Baker's yeast; Brewer's yeast [3]	system [2010.01]
1/19	modified by introduction of foreign genetic material [5]	5/0781 B cells; Progenitors thereof [2010.01] 5/0783 T cells; NK cells; Progenitors of T or NK
1/20 1/21	Bacteria; Culture media therefor [3]modified by introduction of foreign genetic	cells [2010.01] 5/0784 Dendritic cells; Progenitors
1/21	material [5] Processes using, or culture media containing,	thereof [2010.01] 5/0786 Monocytes; Macrophages [2010.01]
1/22	cellulose or hydrolysates thereof [3]	5/0787 Granulocytes, e.g. basophils, eosinophils,
1/24	Processes using, or culture media containing, waste sulfite liquor [3]	neutrophils or mast cells [2010.01] 5/0789 Stem cells; Multipotent progenitor
1/26	Processes using, or culture media containing,	cells [2010.01]
	hydrocarbons (refining of hydrocarbon oils by using	5/079 Neural cells [2010.01]
	micro-organisms C10G 32/00) [3]	5/0793 Neurons [2010.01]
1/28	aliphatic [3]	5/0797 Stem cells; Progenitor cells [2010.01]
1/30	having five or less carbon atoms [3]	5/09 . Tumour cells [2010.01]
1/32	. Processes using, or culture media containing, lower	5/095 Stem cells; Progenitor cells [2010.01]
	alkanols, i.e. C_1 to C_6 [3]	5/10 . Cells modified by introduction of foreign genetic
1/34	. Processes using foam culture [3]	material, e.g. virus-transformed cells [5]
1/36	. Adaptation or attenuation of cells [3]	5/12 Fused cells, e.g. hybridomas [5]
1/38	. Chemical stimulation of growth or activity by	5/14 Plant cells [5]
	addition of chemical compounds which are not	5/16 Animal cells [5]
	essential growth factors; Stimulation of growth by	5/18 Murine cells, e.g. mouse cells [5]
	removal of a chemical compound (C12N 1/34 takes precedence) [3]	5/20 one of the fusion partners being a B lymphocyte [5]
3/00	Spore-forming or isolating processes [3]	5/22 Human cells [5]

5/24			
	one of the fusion partners being a B lymphocyte [5]	9/40	acting on alpha-galactose-glycoside bonds, e.g. alpha-galactosidase [3]
5/26	Cells resulting from interspecies fusion [5]	9/42	acting on beta-1, 4-glucosidic bonds,
5/28	one of the fusion partners being a human cell [5]	9/44	e.g. cellulase [3] acting on alpha-1, 6-glucosidic bonds,
	con [e]	<i>,,</i>	e.g. isoamylase, pullulanase [3]
7/00	Viruses, e.g. bacteriophages; Compositions thereof;	9/46	Dextranase [3]
	Preparation or purification thereof (medicinal	9/48	. acting on peptide bonds, e.g. thromboplastin,
	preparations containing viruses A61K 35/76; preparing	2740	leucine aminopeptidase (3.4) [3]
	medicinal viral antigen or antibody compositions,	9/50	Proteinases [3]
7/01	e.g. virus vaccines, A61K 39/00) [3]	9/52	derived from bacteria [3]
7/01	 Viruses, e.g. bacteriophages, modified by introduction of foreign genetic material (vectors 	9/54	bacteria being Bacillus [3]
	C12N 15/00) [5]	9/56	Bacillus subtilis or Bacillus
7/02	Recovery or purification [3]	2720	licheniformis [3]
7/04	Inactivation or attenuation; Producing viral sub-	9/58	derived from fungi [3]
7704	units [3]	9/60	from yeast [3]
7/06	by chemical treatment [3]	9/62	from Aspergillus [3]
7/08	by serial passage of virus [3]	9/64	derived from animal tissue, e.g. rennin [3]
		9/66	Elastase [3]
9/00	Enzymes, e.g. ligases (6.); Proenzymes; Compositions	9/68	Plasmin, i.e. fibrinolysin [3]
	thereof (preparations containing enzymes for cleaning	9/70	Streptokinase [3]
	teeth A61K 8/66, A61Q 11/00; medicinal preparations containing enzymes or proenzymes A61K 38/43;	9/72	Urokinase [3]
	enzyme containing detergent compositions C11D);	9/74	Thrombin [3]
	Processes for preparing, activating, inhibiting,	9/76	Trypsin; Chymotrypsin [3]
	separating, or purifying enzymes (preparation of malt C12C 1/00) [3]	9/78	acting on carbon to nitrogen bonds other than
	C12C 1/00) [3]	0.400	peptide bonds (3.5) [3]
Note		9/80	acting on amide bonds in linear amides [3]
		9/82	Asparaginase [3]
	In this group:	9/84	Penicillin amidase [3]
	 proenzymes are classified with the corresponding enzymes; [5] 	9/86	acting on amide bonds in cyclic amides, e.g. penicillinase [3]
	 enzymes are generally categorised according to the 	9/88	. Lyases (4.) [3]
	"Nomenclature and Classification of Enzymes" of	9/90	. Isomerases (5.) [3]
	the International Commission on Enzymes. Where	9/92	Glucose isomerase [3]
	appropriate, this designation appears in the subgroups below in parenthesis. [3]	9/94	. Pancreatin [3]
	subgroups below in parentilesis. [5]	9/96	. Stabilising an enzyme by forming an adduct or a
9/02			composition; Forming enzyme conjugates [3]
	Oxidoreductases (1) e g luciferase [3]		
	. Oxidoreductases (1.), e.g. luciferase [3]	9/98	Preparation of granular or free-flowing enzyme
9/04	acting on CHOH groups as donors, e.g. glucose		compositions (C12N 9/96 takes precedence) [3]
9/04	• acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3]	9/98 9/99	
	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as 	9/99	compositions (C12N 9/96 takes precedence) [3]
9/04	• acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3]		compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3]
9/04 9/06	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] 	9/99 11/00	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3]
9/04 9/06 9/08	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, 	9/99	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or
9/04 9/06 9/08 9/10 9/12	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] 	9/99 11/00 11/02	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3]
9/04 9/06 9/08 9/10 9/12	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] 	9/99 11/00	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow
9/04 9/06 9/08 9/10 9/12 9/14 9/16	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] 	9/99 11/00 11/02 11/04	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3]
9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] 	9/99 11/00 11/02 11/04 11/06	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3]
9/04 9/06 9/08 9/10 9/12 9/14 9/16	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of 	9/99 11/00 11/02 11/04 11/06 11/08	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3]
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9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18 9/20	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of lipase [3] Ribonucleases [3] acting on glycosyl compounds (3.2) [3] acting on alpha-1, 4-glucosidic bonds, 	9/99 11/00 11/02 11/04 11/06 11/08 11/10 11/12	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3] . the carrier being a carbohydrate [3] . Cellulose or derivatives thereof [3] . Enzymes or microbial cells being immobilised on or
9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18 9/20 9/22 9/24 9/26	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of lipase [3] Ribonucleases [3] acting on glycosyl compounds (3.2) [3] acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3] 	9/99 11/00 11/02 11/04 11/06 11/08 11/10 11/12 11/14	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3] . the carrier being a carbohydrate [3] . Cellulose or derivatives thereof [3] . Enzymes or microbial cells being immobilised on or in an inorganic carrier [3]
9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18 9/20 9/22 9/24 9/26	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of lipase [3] Ribonucleases [3] acting on glycosyl compounds (3.2) [3] acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3] Alpha-amylase from microbial source, e.g. bacterial amylase [3] 	9/99 11/00 11/02 11/04 11/06 11/08 11/10 11/12 11/14	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3] . the carrier being a carbohydrate [3] . Cellulose or derivatives thereof [3] . Enzymes or microbial cells being immobilised on or in an inorganic carrier [3] . Enzymes or microbial cells being immobilised on or
9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18 9/20 9/22 9/24 9/26 9/28	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of lipase [3] Ribonucleases [3] acting on glycosyl compounds (3.2) [3] acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3] Alpha-amylase from microbial source, e.g. bacterial amylase [3] Fungal source [3] 	9/99 11/00 11/02 11/04 11/06 11/08 11/10 11/12 11/14 11/16	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3] . the carrier being a carbohydrate [3] . Cellulose or derivatives thereof [3] . Enzymes or microbial cells being immobilised on or in an inorganic carrier [3] . Enzymes or microbial cells being immobilised on or in a biological cell [3] . Multi-enzyme systems [3]
9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18 9/20 9/22 9/24 9/26 9/28	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of lipase [3] Ribonucleases [3] acting on glycosyl compounds (3.2) [3] acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3] Alpha-amylase from microbial source, e.g. bacterial amylase [3] Fungal source [3] Alpha-amylase from plant source [3] 	9/99 11/00 11/02 11/04 11/06 11/08 11/10 11/12 11/14 11/16	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3] . the carrier being a carbohydrate [3] . Cellulose or derivatives thereof [3] . Enzymes or microbial cells being immobilised on or in an inorganic carrier [3] . Enzymes or microbial cells being immobilised on or in a biological cell [3] . Multi-enzyme systems [3] Treatment of micro-organisms or enzymes with electrical or wave energy, e.g. magnetism, sonic
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9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18 9/20 9/22 9/24 9/26 9/28	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of lipase [3] Ribonucleases [3] acting on glycosyl compounds (3.2) [3] acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3] Alpha-amylase from microbial source, e.g. bacterial amylase [3] Fungal source [3] Alpha-amylase from plant source [3] Glucoamylase [3] acting on beta-1, 4 bonds between N- 	9/99 11/00 11/02 11/04 11/06 11/08 11/10 11/12 11/14 11/16	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3] . the carrier being a carbohydrate [3] . Cellulose or derivatives thereof [3] . Enzymes or microbial cells being immobilised on or in an inorganic carrier [3] . Enzymes or microbial cells being immobilised on or in a biological cell [3] . Multi-enzyme systems [3] Treatment of micro-organisms or enzymes with electrical or wave energy, e.g. magnetism, sonic
9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18 9/20 9/22 9/24 9/26 9/28 9/30 9/32 9/34	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of lipase [3] Ribonucleases [3] acting on glycosyl compounds (3.2) [3] acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3] Alpha-amylase from microbial source, e.g. bacterial amylase [3] Fungal source [3] Alpha-amylase from plant source [3] Glucoamylase [3] acting on beta-1, 4 bonds between N-acetylmuramic acid and 2-acetylamino 	9/99 11/00 11/02 11/04 11/06 11/08 11/10 11/12 11/14 11/16	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3] . the carrier being a carbohydrate [3] . Cellulose or derivatives thereof [3] . Enzymes or microbial cells being immobilised on or in an inorganic carrier [3] . Enzymes or microbial cells being immobilised on or in a biological cell [3] . Multi-enzyme systems [3] Treatment of micro-organisms or enzymes with electrical or wave energy, e.g. magnetism, sonic
9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18 9/20 9/22 9/24 9/26 9/28 9/30 9/32 9/34 9/36	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of lipase [3] Ribonucleases [3] acting on glycosyl compounds (3.2) [3] acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3] Alpha-amylase from microbial source, e.g. bacterial amylase [3] Fungal source [3] Glucoamylase [3] Glucoamylase [3] acting on beta-1, 4 bonds between N-acetylmuramic acid and 2-acetylamino 2-deoxy-D-glucose, e.g. lysozyme [3] 	9/99 11/00 11/02 11/04 11/06 11/08 11/10 11/12 11/14 11/16	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3] . the carrier being a carbohydrate [3] . Cellulose or derivatives thereof [3] . Enzymes or microbial cells being immobilised on or in an inorganic carrier [3] . Enzymes or microbial cells being immobilised on or in a biological cell [3] . Multi-enzyme systems [3] Treatment of micro-organisms or enzymes with electrical or wave energy, e.g. magnetism, sonic
9/04 9/06 9/08 9/10 9/12 9/14 9/16 9/18 9/20 9/22 9/24 9/26 9/28 9/30 9/32 9/34	 acting on CHOH groups as donors, e.g. glucose oxidase, lactate dehydrogenase (1.1) [3] acting on nitrogen containing compounds as donors (1.4, 1.5, 1.7) [3] acting on hydrogen peroxide as acceptor (1.11) [3] Transferases (2.) (ribonucleases C12N 9/22) [3] transferring phosphorus containing groups, e.g. kinases (2.7) [3] Hydrolases (3.) [3] acting on ester bonds (3.1) [3] Carboxylic ester hydrolases [3] Triglyceride splitting, e.g. by means of lipase [3] Ribonucleases [3] acting on glycosyl compounds (3.2) [3] acting on alpha-1, 4-glucosidic bonds, e.g. hyaluronidase, invertase, amylase [3] Alpha-amylase from microbial source, e.g. bacterial amylase [3] Fungal source [3] Alpha-amylase from plant source [3] Glucoamylase [3] acting on beta-1, 4 bonds between N-acetylmuramic acid and 2-acetylamino 	9/99 11/00 11/02 11/04 11/06 11/08 11/10 11/12 11/14 11/16	compositions (C12N 9/96 takes precedence) [3] . Enzyme inactivation by chemical treatment [3] Carrier-bound or immobilised enzymes; Carrier-bound or immobilised microbial cells; Preparation thereof [3] . Enzymes or microbial cells being immobilised on or in an organic carrier [3] . entrapped within the carrier, e.g. gel, hollow fibre [3] . attached to the carrier via a bridging agent [3] . the carrier being a synthetic polymer [3] . the carrier being a carbohydrate [3] . Cellulose or derivatives thereof [3] . Enzymes or microbial cells being immobilised on or in an inorganic carrier [3] . Enzymes or microbial cells being immobilised on or in a biological cell [3] . Multi-enzyme systems [3] Treatment of micro-organisms or enzymes with electrical or wave energy, e.g. magnetism, sonic

15/00	Mutation or genetic engineering; DNA or RNA	15/29	Genes encoding plant proteins,
	concerning genetic engineering, vectors, e.g. plasmids, or their isolation, preparation or	15/00	e.g. thaumatin [5]
	purification; Use of hosts therefor (mutants or	15/30	Genes encoding protozoal proteins, e.g. from
	genetically engineered micro-organisms C12N 1/00,	15/31	Plasmodium, Trypanosoma, Eimeria [5] Genes encoding microbial proteins,
	C12N 5/00, C12N 7/00; new plants A01H; plant	13/31	e.g. enterotoxins [5]
	reproduction by tissue culture techniques A01H 4/00;	15/32	Bacillus crystal proteins [5]
	new animals A01K 67/00; use of medicinal preparations	15/33	Genes encoding viral proteins [5]
	containing genetic material which is inserted into cells	15/34	Proteins from DNA viruses [5]
	of the living body to treat genetic diseases, gene therapy A61K 48/00; peptides in general C07K) [3,5,6]	15/35	Parvoviridae, e.g. feline panleukopenia
	710114 40/00, peptides in general Covity [3,5,6]		virus, human parvovirus [5]
<u>Note</u>		15/36	Hepadnaviridae [5]
	This can be a second of the se	15/37	Papovaviridae, e.g. papillomaviruses,
	This group <u>covers</u> processes wherein there is a modification of the genetic material which would not		polyomavirus, SV40 [5]
	normally occur in nature without intervention of man	15/38	Herpetoviridae, e.g. herpes simplex
	which produce a change in the gene structure which is		virus, varicella-zoster virus, Epstein-
	passed on to succeeding generations. [3]		Barr virus, cytomegalovirus, pseudorabies virus [5]
		15/39	Poxviridae, e.g. vaccinia virus, variola
15/01	. Preparation of mutants without inserting foreign	20,00	virus [5]
	genetic material therein; Screening processes	15/40	Proteins from RNA viruses,
15/00	therefor [5]		e.g. flaviviruses [5]
15/02	 Preparation of hybrid cells by fusion of two or more cells, e.g. protoplast fusion [5] 	15/41	Picornaviridae, e.g. rhinovirus,
15/03	. Bacteria [5]		coxsackie viruses, echoviruses,
15/04	Fungi [5]	15/42	enteroviruses [5] Foot-and-mouth disease virus [5]
15/05	. Plant cells [5]	15/42	Foot-and-mouth disease virus [5]
15/06	. Animal cells [5]	15/43	Orthomyxoviridae, e.g. influenza
15/07	Human cells [5]	13/44	virus [5]
15/08	Cells resulting from interspecies fusion [5]	15/45	Paramyxoviridae, e.g. measles virus,
15/09	. Recombinant DNA-technology [5]	20, 10	mumps virus, Newcastle disease virus,
15/10	Processes for the isolation, preparation or		canine distemper virus, rinderpest
	purification of DNA or RNA (chemical		virus, respiratory syncytial viruses [5]
	preparation of DNA or RNA C07H 21/00;	15/46	Reoviridae, e.g. rotavirus, bluetongue
	preparation of non-structural polynucleotides from	15/47	virus, Colorado tick fever virus [5]
15/11	micro-organisms or with enzymes C12P 19/34) [5] . DNA or RNA fragments; Modified forms thereof	15/47	Rhabdoviridae, e.g. rabies viruses, vesicular stomatitis virus [5]
13/11	(DNA or RNA not used in recombinant technology	15/48	Retroviridae, e.g. bovine leukaemia
	C07H 21/00) [5,2010.01]	13/10	virus, feline leukaemia virus [5]
15/113	Non-coding nucleic acids modulating the	15/49	Lentiviridae, e.g. immunodeficiency
	expression of genes, e.g. antisense		viruses such as HIV, visna-maedi
15/115	oligonucleotides [2010.01]		virus, equine infectious anaemia
15/115	Aptamers, i.e. nucleic acids binding a target molecule specifically and with high affinity	15/50	virus [5]
	without hybridising therewith [2010.01]	15/50	bronchitis virus, transmissible
15/117	Nucleic acids having immunomodulatory		gastroenteritis virus [5]
	properties, e.g. containing CpG-	15/51	Hepatitis viruses [5]
	motifs [2010.01]	15/52	Genes encoding for enzymes or proenzymes [5]
15/12	Genes encoding animal proteins [5]		
15/13	Immunoglobulins [5]	<u>Note</u>	
15/14	Human serum albumins [5]		In this group:
15/15	Protease inhibitors, e.g. antithrombin,		 genes encoding for proenzymes are classified with
15/16	antitrypsin, hirudin [5]		the corresponding genes encoding enzymes;
15/16 15/17	Hormones [5] Insulins [5]		- enzymes are generally categorised according to the
15/17	Growth hormones [5]		"Nomenclature and Classification of Enzymes" of the International Commission on Enzymes. Where
15/19	Interferons; Lymphokines; Cytokines [5]		appropriate, this designation appears in the groups
15/20	Interferons [5]		below in parenthesis. [5]
15/21	Alpha-interferons [5]		
15/22	Beta-interferons [5]	15/53	Oxidoreductases (1) [5]
15/23	Gamma-interferons [5]	15/54	Transferases (2) [5]
15/24	Interleukins [5]	15/55	Hydrolases (3) [5]
15/25	Interleukin-1 [5]	15/56	acting on glycosyl compounds (3.2),
15/26	Interleukin-2 [5]		e.g. amylase, galactosidase, lysozyme [5]
15/27	Colony stimulating factors [5]	15/57	acting on peptide bonds (3.4) [5]
15/28	Tumor necrosis factors [5]		

15/58	Plasminogen activators, e.g. urokinase,	15/72 Expression systems using regulatory
15/59	TPA [5] Chymosin [5]	sequences derived from the lac-operon [5] 15/73 Expression systems using phage lambda
15/60	Lyases (4) [5]	regulatory sequences [5]
15/61	Isomerases (5) [5]	15/74 Vectors or expression systems specially adapted
15/62	DNA sequences coding for fusion proteins [5]	for prokaryotic hosts other than E. coli,
	8	e.g. Lactobacillus, Micromonospora [5]
<u>Note</u>		Note
	In this group, the following term is used with the	
	meaning indicated:	This group <u>covers</u> the use of prokaryotes as hosts. [5]
	 "fusion" means the fusion of two different proteins. 	
	[5]	15/75 for Bacillus [5]
		15/76 for Actinomyces; for Streptomyces [5]
15/63	Introduction of foreign genetic material using	15/77 for Corynebacterium; for Brevibacterium [5]
	vectors; Vectors; Use of hosts therefor; Regulation	15/78 for Pseudomonas [5]
15/64	of expression [5] General methods for preparing the vector, for	15/79 Vectors or expression systems specially adapted
13/04	introducing it into the cell or for selecting the	for eukaryotic hosts [5]
	vector-containing host [5]	Note
15/65	using markers (enzymes used as markers	
	C12N 15/52) [5]	This group <u>covers</u> the use of eukaryotes as hosts. [5]
15/66	General methods for inserting a gene into a	
	vector to form a recombinant vector using	15/80 for fungi [5]
	cleavage and ligation; Use of non-functional linkers or adaptors, e.g. linkers containing the	15/81 for yeasts [5]
	sequence for a restriction endonuclease [5]	15/82 for plant cells [5]
<u>Note</u>	sequence for a resultation endonacteuse [5]	15/83 Viral vectors, e.g. cauliflower mosaic virus [5]
11000		15/84 Ti-plasmids [5]
	In this group, the following expression is used with the	15/85 for animal cells [5]
	meaning indicated: - "non-functional linkers" means DNA sequences	15/86 Viral vectors [5]
	which are used to link DNA sequences and which	15/861 Adenoviral vectors [7]
	have no known function of structural gene or	15/863 Poxviral vectors, e.g. vaccinia virus [7
	regulating function. [5]	15/864 Parvoviral vectors [7]
		15/866 Baculoviral vectors [7]
15/67	General methods for enhancing the	15/867 Retroviral vectors [7] 15/869 Herpesviral vectors [7]
	expression [5]	15/869 Herpesviral vectors [7] 15/87 Introduction of foreign genetic material using
15/68	Stabilisation of the vector [5]	processes not otherwise provided for, e.g. co-
15/69	Increasing the copy number of the vector [5]	transformation [5]
15/70	Vectors or expression systems specially adapted	15/873 Techniques for producing new embryos,
	for E. coli [5]	e.g. nuclear transfer, manipulation of totipoten
		cells or production of chimeric
		embryos [2010.01]
(1)	This group <u>covers</u> the use of E. coli as host. [5]	15/877 Techniques for producing new mammalian
(2)	Shuttle vectors also replicating in E. coli are classified	cloned embryos [2010.01] 15/88 using micro-encapsulation, e.g. using liposome
	according to the other host. [5]	vesicle [5]
15/71	Expression systems using regulatory	15/89 using micro-injection [5]
	sequences derived from the trp-operon [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]

⁽¹⁾ This subclass <u>covers</u> 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]

<u>Note</u>

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

Subclass index

BIOSYN SUBSTA	THESIS OF CHEMICAL NCES	Steroids 33/00
SUBSIA	Inorganic compounds	Heterocyclic organic compounds
	•	containing saccharide radicals
	Acyclic or carbocyclic organic compounds5/00 to 15/00	Riboflavin25/00
	peptides or proteins 21/00	Giberellin27/00
		Cephalosporin; penicillin35/00; 37/00
	Carotenes	SEPARATION OF OPTICAL ISOMERS41/00
	Tetracyclines	OTHER PROCESSES FOR BIOSYNTHESIS
	Prostaglandins	PREPARATIONS
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	7/38 Cyclopentanone- or cyclopentadione- containing products [3] 7/40 . containing a carboxyl group [3]
	processes for the preparation of compounds or	7/42 Hydroxy carboxylic acids [3]
	compositions by using micro-organisms or	
	enzymes [3]	j j i i
1/02	. by using fungi [3]	7/46 Dicarboxylic acids having four or less carbon atoms, e.g. fumaric acid, maleic acid [3]
1/04	by using bacteria [3]	7/48 Tricarboxylic acids, e.g. citric acid [3]
1/06	 by using actinomycetales [3] 	7/50 having keto groups, e.g. 2-ketoglutaric acid [3]
2 100		7/50
3/00	Preparation of elements or inorganic compounds	
	except carbon dioxide [3]	7/54 . Acetic acid (vinegar C12J) [3]
5/00	Preparation of hydrocarbons [3]	7/56 Lactic acid [3]
5/02	acyclic (producing methane by anaerobic treatment)	7/58 . Aldonic, ketoaldonic or saccharic acids (uronic acids C12P 19/00) [3]
	of sludge C02F 11/04) [3]	
		7/60 2-Ketogulonic acid [3]
7/00	Preparation of oxygen-containing organic	7/62 Carboxylic acid esters [3]
7.700	compounds [3]	7/64 • Fats; Fatty oils; Ester-type waxes; Higher fatty acids, i.e. having at least seven carbon atoms in an
7/02	. containing a hydroxy group [3]	unbroken chain bound to a carboxyl group; Oxidised
7/04	acyclic [3]	oils or fats [3]
7/06	Ethanol, i.e. non-beverage [3]	7/66 . containing the quinoid structure [3]
7/08	produced as by-product or from waste or cellulosic material substrate [3]	v community are quinous structure [e]
7/10		9/00 Preparation of organic compounds containing a
7/10	substrate containing cellulosic material [3]	metal or atom other than H, N, C, O, S, or halogen [3]
7/12 7/14	 substrate containing sulfite waste liquor or citrus waste [3] Multiple stages of fermentation; Multiple 	11/00 Preparation of sulfur-containing organic compounds [3]
// 14	types of micro-organisms or reuse for micro- organisms [3]	13/00 Preparation of nitrogen-containing organic
7/16	Butanols [3]	compounds [3] 13/02 . Amides, e.g. chloramphenicol [3]
7/18	polyhydric [3]	13/04 . Alpha- or beta-amino acids [3]
7/20	Glycerol [3]	•
7/22	aromatic [3]	13/06 Alanine; Leucine; Isoleucine; Serine; Homoserine [3]
7/24	containing a carbonyl group [3]	13/08 . Lysine; Diaminopimelic acid; Threonine;
7/26	Ketones [3]	Valine [3]
7/28	Acetone-containing products [3]	13/10 . Citrulline; Arginine; Ornithine [3]
7/30	produced from substrate containing	13/12 . Methionine; Cysteine; Cystine [3]
	inorganic compounds other than water [3]	13/14 Glutamic acid; Glutamine [3]
7/32	produced from substrate containing	13/16 using surfactants, fatty acids or fatty acid esters
	inorganic nitrogen source [3]	i.e. having at least seven carbon atoms in an
7/34	produced from substrate containing protein as nitrogen source [3]	unbroken chain bound to a carboxyl group or a carboxyl ester group [3]
7/36	produced from substrate containing grain or	13/18 using biotin or its derivatives [3]
	cereal material [3]	

13/20	Aspartic acid; Asparagine [3]	19/36	Dinucleotides, e.g. nicotineamide-adenine dinucleotide phosphate [3]
13/22	. Tryptophan; Tyrosine; Phenylalanine; 3,4-Dihydroxyphenylalanine [3]	19/38	Nucleosides [3]
13/24	Proline; Hydroxyproline; Histidine [3]	19/40	having a condensed ring system containing a six-membered ring having two nitrogen
15/00	Preparation of compounds containing at least three condensed carbocyclic rings [3]		atoms in the same ring, e.g. purine nucleosides [3]
17/00	Preparation of heterocyclic carbon compounds with	19/42	Cobalamins, i.e. vitamin B_{12} , LLD factor [3]
	only O, N, S, Se, or Te as ring hetero atoms (C12P 13/04 to C12P 13/24 take precedence) [3]	19/44	Preparation of O-glycosides, e.g. glucosides [3]
17/02	Oxygen as only ring hetero atoms [3]	19/46	 having an oxygen atom of the saccharide radical bound to a cyclohexyl radical,
17/04	 containing a five-membered hetero ring, e.g. griseofulvin [3] 	19/48	e.g. kasugamycin [3] the cyclohexyl radical being substituted by two
17/06	containing a six-membered hetero ring, e.g. fluorescein [3]	277.10	or more nitrogen atoms, e.g. destomycin, neamin [3]
17/08	containing a hetero ring of at least seven ring members, e.g. zearalenone, macrolide aglycons [3]	19/50	having two saccharide radicals bound through only oxygen to adjacent ring carbon
17/10	. Nitrogen as only ring hetero atom [3]		atoms of the cyclohexyl radical,
17/12	containing a six-membered hetero ring [3]	19/52	e.g. ambutyrosin, ribostamycin [3]
17/14	 Nitrogen or oxygen as hetero atom and at least one other diverse hetero ring atom in the same ring [3] 		containing three or more saccharide radicals, e.g. neomycin, lividomycin [3]
17/16	containing two or more hetero rings [3]	19/54	the cyclohexyl radical being bound directly to a nitrogen atom of two or more
17/18	 containing at least two hetero rings condensed among themselves or condensed with a common carbocyclic ring system, e.g. rifamycin [3] 		N-C-N ← radicals, e.g. streptomycin [3]
19/00	Preparation of compounds containing saccharide	19/56	having an oxygen atom of the saccharide radical
<u>Note</u>	radicals (ketoaldonic acids C12P 7/58) [3]	19730	directly bound to a condensed ring system having three or more carbocyclic rings, e.g. daunomycin,
11000	A	19/58	adriamycin [3] having an oxygen atom of the saccharide radical
	Attention is drawn to Note (3) following the title of subclass C07H, which defines the expression "saccharide radical". [3]		directly bound through only acyclic carbon atoms to a non-saccharide heterocyclic ring, e.g. bleomycin, phleomycin [3]
19/02	. Monosaccharides (2-ketogulonic acid C12P 7/60) [3]	19/60	having an oxygen of the saccharide radical directly
19/04	Polysaccharides, i.e. compounds containing more		bound to a non-saccharide heterocyclic ring or a condensed ring system containing a non-
	than five saccharide radicals attached to each other by glycosidic bonds [3]		saccharide heterocyclic ring, e.g. coumermycin, novobiocin [3]
19/06	Xanthan, i.e. Xanthomonas-type	19/62	the hetero ring having eight or more ring
19/08	heteropolysaccharides [3] . Dextran [3]		members and only oxygen as ring hetero atoms, e.g. erythromycin, spiramycin, nystatin [3]
19/10	Pullulan [3]	19/64	Preparation of S-glycosides, e.g. lincomycin [3]
19/12	. Disaccharides [3]	21/00	Preparation of peptides or proteins (single-cell
19/14	 produced by the action of a carbohydrase, e.g. by alpha-amylase [3] 	21/00	protein C12N 1/00) [3]
19/16	produced by the action of an alpha-1, 6-glucosidase, e.g. amylose, debranched amylopectin (non-	21/02	 having a known sequence of two or more amino acids, e.g. glutathione [3]
	biological hydrolysis of starch C08B 30/00) [3]	21/04	Cyclic or bridged peptides or polypeptides,
19/18	 produced by the action of a glycosyl transferase, e.g. alpha-, beta- or gamma-cyclodextrins [3] 		e.g. bacitracin (cyclised by $-S-S$ -bonds only C12P 21/02) [3]
19/20	produced by the action of an exo-1, 4 alpha-	21/06	 produced by the hydrolysis of a peptide bond, e.g. hydrolysate products (preparing foodstuffs by
19/22	glucosidase, e.g. dextrose [3] produced by the action of a beta-amylase,		protein hydrolysis A23J 3/00) [3]
	e.g. maltose [3]	21/08	. Monoclonal antibodies [5]
19/24	 produced by the action of an isomerase, e.g. fructose [3] 	23/00	Preparation of compounds containing a cyclohexene
19/26	Preparation of nitrogen-containing carbohydrates [3]		ring having an unsaturated side chain containing at least ten carbon atoms bound by conjugated double
19/28	N-glycosides [3]		bonds, e.g. carotenes (containing hetero-rings
19/30	Nucleotides [3]		C12P 17/00) [3]
19/32	having a condensed ring system containing a six-membered ring having two nitrogen	25/00	Preparation of compounds containing alloxazine or isoalloxazine nucleus, e.g. riboflavin [3]
	atoms in the same-ring, e.g. purine nucleotides, nicotineamide-adenine	27/00	Preparation of compounds containing a gibbane ring
10/04	dinucleotide [3]	_,, 00	system, e.g. gibberellin [3]
19/34	• • • Polynucleotides, e.g. nucleic acids, oligoribonucleotides [3]		

29/00	Preparation of compounds containing a naphthacene ring system, e.g. tetracycline (C12P 19/00 takes precedence) [3]	33/04 33/06 33/08	 Forming an aryl ring from A ring [3] Hydroxylating [3] at 11 position [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]	33/10 33/12 33/14 33/16 33/18 33/20 35/00	 at 11alpha-position [3] . Acting on D ring [3] Hydroxylating at 16 position [3] Acting at 17 position [3] Hydroxylating at 17 position [3] . containing heterocyclic rings [3] Preparation of compounds having a 5-thia-1-azabicyclo [4.2.0] octane ring system, e.g. cephalosporin [3]
33/00	Preparation of steroids [3]	35/02 35/04	 by desacylation of the substituent in the 7 position [3] by acylation of the substituent in the 7 position [3]
<u>Note</u>	Attention is drawn to Note (1) following the title of subclass C07J, which explains what is covered by the term "steroids". [3]	35/06 35/08 37/00	 Cephalosporin C; Derivatives thereof [3] disubstituted in the 7 position [3] Preparation of compounds having a 4-thia-1-azabicyclo [3.2.0] heptane ring system, e.g. penicillin [3]
<u>Note</u>	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]	37/02 37/04 37/06 39/00 41/00	 in presence of phenylacetic acid or phenylacetamide or their derivatives [3] by acylation of the substituent in the 6 position [3] by desacylation of the substituent in the 6 position [3] Processes involving micro-organisms of different genera in the same process, simultaneously [3] Processes using enzymes or micro-organisms to generate entirel improve from a reception mixture [4]
33/02	. Dehydrogenating; Dehydroxylating [3]		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]

- (1) This subclass <u>does not cover</u> 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]

C12Q - C12R

1/30 1/32	involving catalase [3]involving dehydrogenase [3]	1/56	• involving blood clotting factors, e.g. involving thrombin, thromboplastin, fibrinogen [3]
1/34	involving hydrolase [3]	1/58	. involving urea or urease [3]
1/37	involving peptidase or proteinase [5]	1/60	. involving cholesterol [3]
1/40	involving amylase [3]	1/61	. involving triglycerides [5]
1/42	involving phosphatase [3]	1/62	. involving uric acid [3]
1/44	involving esterase [3]	1/64	. Geomicrobiological testing, e.g. for petroleum [3]
1/46	involving cholinesterase [3]	1/66	. involving luciferase [3]
1/48	. involving transferase [3]	1/68	. involving nucleic acids [3]
1/50	involving creatine phosphokinase [3]	1/70	 involving virus or bacteriophage [3]
1/52	involving transaminase [3]	3/00	Condition-responsive control processes (apparatus
1/527	. involving lyase [5]	3/00	therefor C12M 1/36; controlling or regulating in general
1/533	. involving isomerase [5]		G05) [3]
1/54	. involving glucose or galactose [3]		/

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

- (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]

1/01	1/00 Micro-organisms [3]	1/26 Methylomonas [3]
1/02		,
1/025		
1/03		
1/04		
1/045	1/04 Actinomyces [3]	•
1/06	1/045 Actinoplanes [3]	1/30 Micromonospora chalcea [3]
1/065 Azotobacter [3]	1/05 Alcaligenes [3]	1/31 Micromonospora purpurea [3]
1/07	1/06 Arthrobacter [3]	1/32 Mycobacterium [3]
1/08 . Bacillus brevis [3] 1/34 . Mycobacterium smegmatis [3] 1/085 . Bacillus cercus [3] 1/35 . Mycoplasma [3] 1/09 . Bacillus circulans [3] 1/36 . Neisseria [3] 1/10 . Bacillus licheniformis [3] 1/365 . Nocardia [3] 1/11 . Bacillus megaterium [3] 1/37 . Proteus [3] 1/12 . Bacillus subtilis [3] 1/38 . Pseudomonas [3] 1/125 . Bacillus subtilis [3] 1/38 . Pseudomonas fluorescens [3] 1/13 . Brevibacterium [3] 1/39 . Pseudomonas fluorescens [3] 1/14 . Chainia [3] 1/40 . Pseudomonas putida [3] 1/145 . Clostridium [3] 1/41 . Rhizobium [3] 1/15 . Corynebacterium [3] 1/42 . Salmonella [3] 1/16 . Corynebacterium diphtheriae [3] 1/425 . Serratia [3] 1/165 . Corynebacterium pyogenes [3] 1/43 . Serratia marcescens [3] 1/17 . Corynebacterium pyogenes [3] 1/44 . Staphylococcus [3] 1/185 . Escherichia [3] 1/445 . Staphylococcus epidermidis [3] <tr< td=""><td>1/065 Azotobacter [3]</td><td>1/325 Mycobacterium avium [3]</td></tr<>	1/065 Azotobacter [3]	1/325 Mycobacterium avium [3]
1/085	1/07 Bacillus [3]	1/33 Mycobacterium fortuitum [3]
1/09	1/08 Bacillus brevis [3]	1/34 Mycobacterium smegmatis [3]
1/10 . Bacillus licheniformis [3] 1/365 . Nocardia [3] 1/11 . Bacillus megaterium [3] 1/37 . Proteus [3] 1/12 . Bacillus polymyxa [3] 1/38 . Pseudomonas [3] 1/125 . Bacillus subtilis [3] 1/385 . Pseudomonas aeruginosa [3] 1/13 . Brevibacterium [3] 1/39 . Pseudomonas fluorescens [3] 1/14 . Chainia [3] 1/40 . Pseudomonas putida [3] 1/145 . Clostridium [3] 1/41 . Rhizobium [3] 1/15 . Corynebacterium [3] 1/42 . Salmonella [3] 1/165 . Corynebacterium diphtheriae [3] 1/425 . Serratia [3] 1/165 . Corynebacterium poinsettiae [3] 1/43 . Serratia marcescens [3] 1/17 . Corynebacterium poinsettiae [3] 1/43 . Serratia marcescens [3] 1/17 . Corynebacterium poinsettiae [3] 1/44 . Staphylococcus [3] 1/18 . Erwinia [3] 1/44 . Staphylococcus [3] 1/18 . Escherichia [3] 1/45 . Staphylococcus epidermidis [3] 1/19 . Escherichia coli [3] 1/46 . Streptomyces [3] <td>1/085 Bacillus cereus [3]</td> <td>1/35 Mycoplasma [3]</td>	1/085 Bacillus cereus [3]	1/35 Mycoplasma [3]
1/11		1/36 Neisseria [3]
1/12	1/10 Bacillus licheniformis [3]	1/365 Nocardia [3]
1/125 Bacillus subtilis [3] 1/385 Pseudomonas aeruginosa [3] 1/13 . Brevibacterium [3] 1/39 Pseudomonas fluorescens [3] 1/14 Chainia [3] 1/40 Pseudomonas putida [3] 1/145 Clostridium [3] 1/41 Rhizobium [3] 1/15	1/11 Bacillus megaterium [3]	1/37 Proteus [3]
1/13 . Brevibacterium [3] 1/39 . Pseudomonas fluorescens [3] 1/14 . Chainia [3] 1/40 . Pseudomonas putida [3] 1/145 . Clostridium [3] 1/41 . Rhizobium [3] 1/15 . Corynebacterium [3] 1/42 . Salmonella [3] 1/16 . Corynebacterium diphtheriae [3] 1/425 . Serratia [3] 1/165 . Corynebacterium poinsettiae [3] 1/43 . Serratia marcescens [3] 1/17 . Corynebacterium pyogenes [3] 1/44 . Staphylococcus [3] 1/18 . Erwinia [3] 1/45 . Staphylococcus aureus [3] 1/185 . Escherichia [3] 1/45 . Staphylococcus epidermidis [3] 1/19 . Escherichia coli [3] 1/46 . Streptococcus [3] 1/20 . Flavobacterium [3] 1/46 . Streptomyces [3] 1/21 . Haemophilus [3] 1/47 . Streptomyces albus [3] 1/22 . Klebsiella [3] 1/48 . Streptomyces aureofaciens [3] 1/23 . Lactobacillus acidophilus [3] 1/49 . Streptomyces aureus [3] 1/24 . Lactobacillus brevis [3] 1/50 . Streptomyces candidus [3]	1/12 Bacillus polymyxa [3]	1/38 Pseudomonas [3]
1/14 . Chainia [3] 1/40 . Pseudomonas putida [3] 1/145 . Clostridium [3] 1/41 . Rhizobium [3] 1/15 . Corynebacterium [3] 1/42 . Salmonella [3] 1/16 . Corynebacterium diphtheriae [3] 1/425 . Serratia [3] 1/165 . Corynebacterium poinsettiae [3] 1/43 . Serratia marcescens [3] 1/17 . Corynebacterium pyogenes [3] 1/44 . Staphylococcus [3] 1/18 . Erwinia [3] 1/45 . Staphylococcus aureus [3] 1/185 . Escherichia [3] 1/45 . Staphylococcus epidermidis [3] 1/19 . Escherichia coli [3] 1/45 . Streptococcus [3] 1/20 . Flavobacterium [3] 1/46 . Streptomyces [3] 1/21 . Haemophilus [3] 1/47 . Streptomyces albus [3] 1/22 . Klebsiella [3] 1/48 . Streptomyces aureofaciens [3] 1/23 . Lactobacillus acidophilus [3] 1/49 . Streptomyces aureus [3] 1/24 . Lactobacillus brevis [3] 1/50 . Streptomyces candidus [3] 1/245 . Lactobacillus casei [3] 1/51 . Streptomyces candidus [3] </td <td>1/125 Bacillus subtilis [3]</td> <td>1/385 Pseudomonas aeruginosa [3]</td>	1/125 Bacillus subtilis [3]	1/385 Pseudomonas aeruginosa [3]
1/145 . Clostridium [3] 1/41 . Rhizobium [3] 1/15 . Corynebacterium [3] 1/42 . Salmonella [3] 1/16 . Corynebacterium diphtheriae [3] 1/425 . Serratia [3] 1/165 . Corynebacterium poinsettiae [3] 1/43 . Serratia marcescens [3] 1/17 . Corynebacterium pyogenes [3] 1/44 . Staphylococcus [3] 1/18 . Erwinia [3] 1/445 . Staphylococcus aureus [3] 1/185 . Escherichia [3] 1/45 . Staphylococcus epidermidis [3] 1/19 . Escherichia coli [3] 1/46 . Streptococcus [3] 1/20 . Flavobacterium [3] 1/465 . Streptomyces [3] 1/21 . Haemophilus [3] 1/47 . Streptomyces albus [3] 1/22 . Klebsiella [3] 1/48 . Streptomyces aureofaciens [3] 1/225 . Lactobacillus acidophilus [3] 1/49 . Streptomyces aureus [3] 1/24 . Lactobacillus brevis [3] 1/50 . Streptomyces candidus [3] 1/245 . Lactobacillus casei [3] 1/51 . Streptomyces candidus [3]	1/13 Brevibacterium [3]	
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1/165	1/15 Corynebacterium [3]	1/42 Salmonella [3]
1/17	1/16 Corynebacterium diphtheriae [3]	1/425 Serratia [3]
1/18 . Erwinia [3] 1/445 . Staphylococcus aureus [3] 1/185 . Escherichia [3] 1/45 . Staphylococcus epidermidis [3] 1/19 . Escherichia coli [3] 1/46 . Streptococcus [3] 1/20 . Flavobacterium [3] 1/465 . Streptomyces [3] 1/21 . Haemophilus [3] 1/47 . Streptomyces albus [3] 1/22 . Klebsiella [3] 1/48 . Streptomyces antibioticus [3] 1/225 . Lactobacillus [3] 1/485 . Streptomyces aureofaciens [3] 1/23 . Lactobacillus acidophilus [3] 1/49 . Streptomyces aureus [3] 1/24 . Lactobacillus brevis [3] 1/50 . Streptomyces bikiniensis [3] 1/245 . Lactobacillus casei [3] 1/51 . Streptomyces candidus [3]	1/165 Corynebacterium poinsettiae [3]	1/43 Serratia marcescens [3]
1/185 . Escherichia [3] 1/45 . Staphylococcus epidermidis [3] 1/19 . Escherichia coli [3] 1/46 . Streptococcus [3] 1/20 . Flavobacterium [3] 1/465 . Streptomyces [3] 1/21 . Haemophilus [3] 1/47 . Streptomyces albus [3] 1/22 . Klebsiella [3] 1/48 . Streptomyces antibioticus [3] 1/225 . Lactobacillus [3] 1/485 . Streptomyces aureofaciens [3] 1/23 . Lactobacillus acidophilus [3] 1/49 . Streptomyces aureus [3] 1/24 . Lactobacillus brevis [3] 1/50 . Streptomyces bikiniensis [3] 1/245 . Lactobacillus casei [3] 1/51 . Streptomyces candidus [3]	1/17 Corynebacterium pyogenes [3]	1/44 Staphylococcus [3]
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1/245 Lactobacillus casei [3] 1/51 Streptomyces candidus [3]		
1/25 Lactobacillus plantarum [3] 1/52 Streptomyces chartreusis [3]		
	1/25 Lactobacillus plantarum [3]	1/52 Streptomyces chartreusis [3]

1/525 Streptomyces diastatochromogenes [3]	1/73 Candida lipolytica [3]
1/53 Streptomyces filipinensis [3]	1/74 Candida tropicalis [3]
1/54 Streptomyces fradiae [3]	1/745 Cephalosporium [3]
1/545 Streptomyces griseus [3]	1/75 Cephalosporium acremonium [3]
1/55 Streptomyces hygroscopicus [3]	1/76 Cephalosporium coerulescens [3]
1/56 Streptomyces lavendulae [3]	1/765 Cephalosporium crotocinigenum [3]
1/565 Streptomyces lincolnensis [3]	1/77 Fusarium [3]
1/57 Streptomyces noursei [3]	1/78 Hansenula [3]
1/58 Streptomyces olivaceus [3]	1/785 Mucor [3]
1/585 Streptomyces platensis [3]	1/79 Paecilomyces [3]
1/59 Streptomyces rimosus [3]	1/80 Penicillium [3]
1/60 Streptomyces sparsogenes [3]	1/81 Penicillium brevi [3]
1/61 Streptomyces venezuelae [3]	1/82 Penicillium chrysogenum [3]
1/62 Streptosporangium [3]	1/825 Penicillium notatum [3]
1/625 Streptoverticillium [3]	1/83 Penicillium patulum [3]
1/63 Vibrio [3]	1/84 Pichia [3]
1/64 Xanthomonas [3]	1/845 Rhizopus [3]
1/645 . Fungi [3]	1/85 Saccharomyces [3]
1/65 Absidia [3]	1/86 Saccharomyces carlsbergensis [3]
1/66 Aspergillus [3]	1/865 Saccharomyces cerevisiae [3]
1/665 Aspergillus awamori [3]	1/87 Saccharomyces lactis [3]
1/67 Aspergillus flavus [3]	1/88 Torulopsis [3]
1/68 Aspergillus fumigatus [3]	1/885 Trichoderma [3]
1/685 Aspergillus niger [3]	1/89 . Algae [3]
1/69 Aspergillus oryzae [3]	1/90 . Protozoa [3]
1/70 Aspergillus ustus [3]	1/91 . Cell lines [3,7]
1/71 Aspergillus wentii [3]	1/92 . Viruses [5,7]
1/72 Candida [3]	1/93 Animal viruses [7]
1/725 Candida albicans [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]

- (1) This subclass <u>covers</u> 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.

- (2) Attention is drawn to Notes (2) and (3) following the title of class C12. [5]
- (3) In this subclass, in the absence of an indication to the contrary, classification is made in the last appropriate place. [2009.01]
- (4) The classification symbols of this subclass are not listed first when printed on 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]	3/04 3/06	Cellulose, e.g. plant fibres [5]Treatment of hemp or flax [5]
1/02	. Desulfurising [5]	3/08	in the production of paper pulp [5]
3/00	Treatment of animal or plant materials or micro- organisms [5]	3/10 3/12	Treatment of sugar or molasses [5]Treatment of pectin or starch [5]
3/02	. Recovery or purification of carbohydrate material [5]		

3/14	•	or purification of proteinaceous	5/00	Treatment of emulsions, gases or foams [5]	
3/16	material [:	en or gelatin [5]	7/00	Treatment of hides, e.g. depilating, bating [5]	
3/18	•	or purification of glyceridic oils, fats, esters or fatty acids [5]	9/00	Cleaning solid surfaces of materials [5]	
3/20	J 1	of nucleic acids from intact or disrupted	11/00	Treatment of textiles, e.g. cleaning [5]	
	cells [5]		99/00	Subject matter not provided for in other groups of this	
3/22	. Treatment	of blood fractions [5]	<i>3370</i> 0	subclass [2010.01]	
3/24	. Treatment	of animal secretions or organs [5]		50000055 [2010101]	