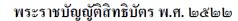


# วิธีการสืบคันข้อมูลสิทธิบัตรเบื้องตัน (Patent search)



ทันยธร เขตต์สุพรรณ
เจ้าหน้าที่ทรัพย์สินทางปัญญา
ฝ่ายจัดการทรัพย์สินทางปัญญา
สำนักบริหารงานวิจัยและนวัตกรรมพระจอมเกล้าลาดกระบัง

# ความใหม่ (Novelty)



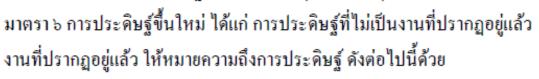
### แก้ไขเพิ่มเติมโดย พ.ร.บ. สิทธิบัตร (ฉบับที่ ๓)



ภูมิพลอคุลยเคช ป.ร.

ให้ไว้ ณ วันที่ ๑๑ มีนาคม พ.ศ. ๒๕๒๒





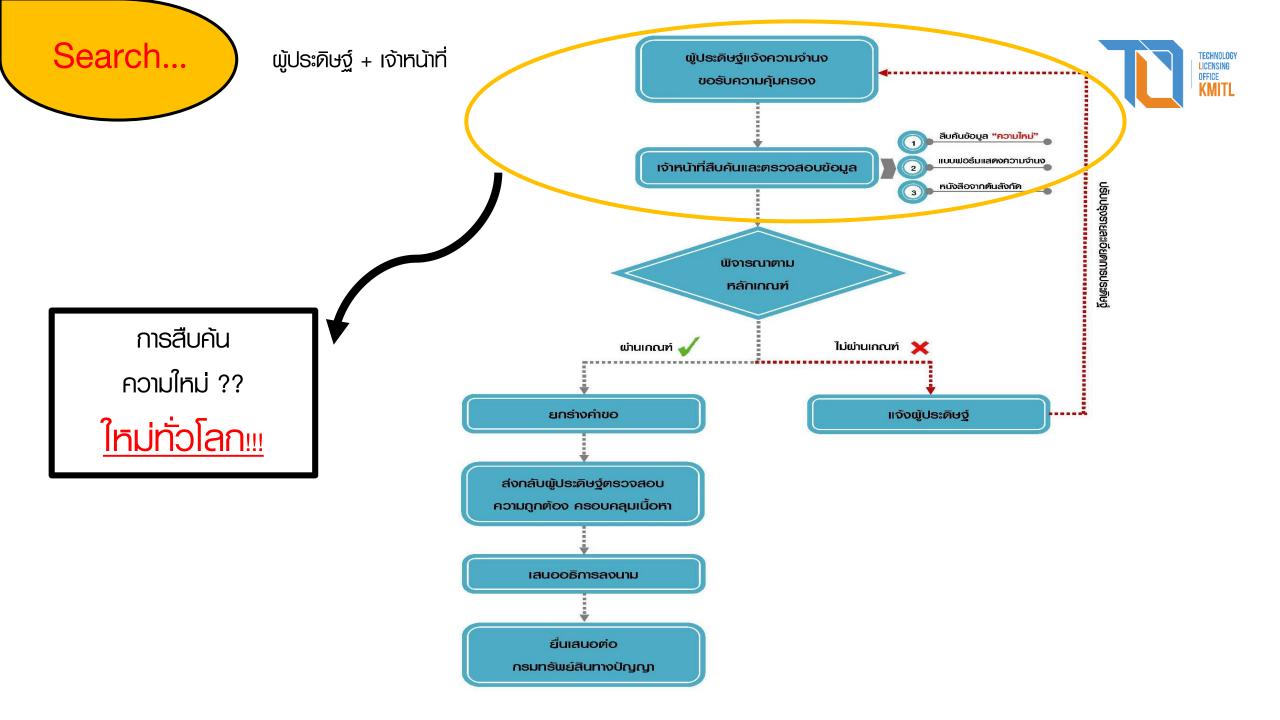
- (๑) การประคิษฐ์ที่มีหรือใช้แพร่หลายอยู่แล้วในราชอาณาจักรก่อนวันขอรับสิทธิบัตร
- (๒) การประดิษฐ์ที่ได้มีการเปิดเผยสาระสำคัญหรือรายละเอียดในเอกสาร หรือสิ่งพิมพ์ที่ได้เผยแพร่อยู่แล้วไม่ว่าในหรือนอกราชอาณาจักรก่อนวันขอรับสิทธิบัตร และ ไม่ว่าการเปิดเผยนั้นจะกระทำโดยเอกสาร สิ่งพิมพ์ การนำออกแสดง หรือการเปิดเผยต่อสาธารณชนด้วยประการใด ๆ
  - (ന)

การประดิษฐ์ที่ได้รับสิทธิบัตรหรืออนุสิทธิบัตรแล้วไม่ว่าในหรือนอกราชอาณาจักรก่อนวันขอรับสิทธิบัตร

(d)

- การประคิษฐ์ที่มีผู้ขอรับสิทธิบัตรหรืออนุสิทธิบัตรไว้แล้วนอกราชอาณาจักรเป็นเวลาเกินสิบแปคเคือน ก่อนวันขอรับสิทธิบัตรแต่ยังมิได้มีการออกสิทธิบัตรหรืออนุสิทธิบัตรให้
- (๕) การประดิษฐ์ที่มีผู้ขอรับสิทธิบัตรหรืออนุสิทธิบัตรไว้แล้วไม่ว่าในหรือนอกราชอาณาจักร และได้ประกาศโฆษณาแล้วก่อนวันขอรับสิทธิบัตรในราชอาณาจักร





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#### C12N15/1003?

Extracting or separating nucleic acids from biological samples, e.g. pure separation or isolation methods; Conditions, buffers or apparatuses therefor

#### Method and apparatus for DNA extraction



Grant US5989431A . Timothy Martin Evans . Progen Industries Ltd

Priority 1995-06-08 • Filing 1996-06-11 • Grant 1999-11-23 • Publication 1999-11-23

Methods and apparatus for the extraction of DNA from a suspension of cells are described. The methods utilise a hollow membrane filter to separate DNA from cellular debri after lysis of cells. The suspension of cells can be a suspension of ...

#### Method for obtaining human skin DNA samples with an adhesive sheet

Grant US6355439B1 · Yeon Bo Chung · I.D. Gene, Inc.



Priority 1998-09-23 • Filing 1999-09-22 • Grant 2002-03-12 • Publication 2002-03-12

Provided is a method for obtaining human DNA for genetic analysis, by taking the epidermis of testee by means of an adhesive sheet, and by extracting DNA from the epidermis stuck on the adhesive sheet. Provided are also combined sheets for ...

#### Method for purification and manipulation of nucleic acids using paramagnetic ...



Grant US5973138A • Matthew P. Collis • Becton Dickinson And Company

Priority 1998-10-30 • Filing 1998-10-30 • Grant 1999-10-26 • Publication 1999-10-26

2. The method of claim 1 wherein said at least one paramagnetic particle comprises iron. 3. The method of claim 1 wherein said at least one paramagnetic particle is selected from the group consisting of an iron oxide, iron sulfide and iron ...



Search within classification C12N15/1003 (1,138,114 results)

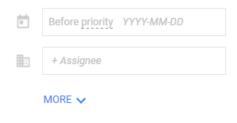
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# SEARCH TERMS 3 dna extraction x + Synonym method x + Synonym

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#### **BACK TO 1.6M RESULTS**

#### Method for purification and manipulation of nucleic acids using paramagnetic particles

#### Abstract

The present invention relates to a composition which is useful for the reversible binding of a nucleic acid molecule. The composition, which may be packaged in a kit, includes a paramagnetic particle in an acidic solution.

#### Images (7)



#### Classifications

C12N15/1013 Extracting or separating nucleic acids from biological samples, e.g. pure separation or isolation methods; Conditions, buffers or apparatuses therefor by means of a solid support carrier, e.g. particles, polymers by using magnetic beads

View 1 more classifications

#### US5973138A

**US Grant** 



Legal status: Active

Application number: US09183127

Inventor: Matthew P. Collis

Current Assignee: Becton Dickinson and Co

Original Assignee: Becton Dickinson and Co

Priority date: 1998-10-30

Filing date: 1998-10-30

Publication date: 1999-10-26

Grant date: 1999-10-26

Info: Patent citations (26), Non-patent citations (2), Cited by (125), Also published as (15), Legal events, Similar documents

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**BACK TO 1.6M RESULTS** 

+ Assignee

#### Description

#### BACKGROUND OF THE INVENTION

Access to cellular components such as nucleic acids is imperative to a variety of molecular biology methodologies. Such methodologies include nucleic acid sequencing, direct detection of particular nucleic acid sequences by nucleic acid hybridization and nucleic acid sequence amplification techniques.

The preparation and purification of high-purity double-stranded (ds) plasmid DNA, single-stranded (ss) phage DNA, chromosomal DNA, agarose gel-purified DNA fragments and RNA is of critical importance in molecular biology. Ideally, a method for purifying nucleic acids should be simple, rapid and require little, if any, additional sample manipulation. Nucleic acids rendered by such a method should be immediately amenable to transformation, restriction analysis, litigation or sequencing. A method with all of these features would be extremely attractive in the automation of nucleic acid sample preparation, a goal of research and diagnostic laboratories.

Typically, the preparation of plasmid DNA from crude alcohol precipitates is laborious, most often utilizing CsCl gradients, gel filtration, ion exchange chromatography, or RNase, proteinase K and repeated alcohol precipitation steps. These methods also require considerable downstream sample preparation to remove CsCl and other salts, ethidium bromide and alcohol. Similar arguments extend when using any of these methods for purifying DNA fragments. A further problem with these methods is that small, negatively-charged cellular components can co-purify with the DNA. Thus, the DNA can have an undesirable level of contamination.

Nucleic acids can also be purified using solid phases. Conventional solid phase extraction techniques have utilized surfaces which either (1) fail to attract and hold sufficient quantities of nucleic acid molecules because of surface design to permit easy recovery of the nucleic acid molecules during elution, or (2) excessively adhere nucleic acid molecules to the surface, thereby hindering recovery of the nucleic acid molecules during elution. Conventional metal surfaces which cause these problems when utilized in solid phase extraction include certain silica surfaces such as glass and Celite. Adequate binding of nucleic acids to these types of surfaces can be

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#### What is claimed is:

Claims (5)

- 1. A method for reversibly binding at least one nucleic acid molecule to at least one paramagnetic particle comprising:
  - (a) providing a suspension of at least one paramagnetic particle in an acidic solution; and
  - (b) combining said suspension with at least one nucleic acid molecule such that said at least one nucleic acid molecule is reversibly bound to said at least one paramagnetic particle.
    - The method of claim 1 wherein said at least one paramagnetic particle comprises iron.
    - The method of claim 1 wherein said at least one paramagnetic particle is selected from the group consisting of an iron oxide, iron sulfide and iron chloride.
    - 4. The **method** of claim 3 wherein the iron oxide is ferric hydroxide or ferrosoferric oxide.
    - 5. The method of claim 1 further comprising:
      - (c) eluting said at least one nucleic acid molecule from said at least one paramagnetic particle.

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DNA extraction

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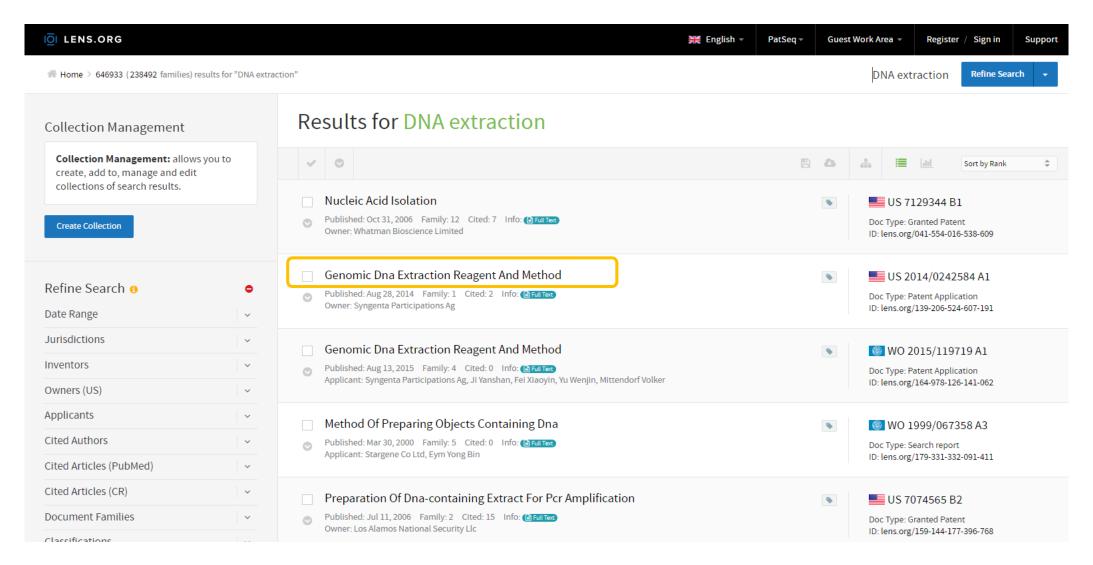




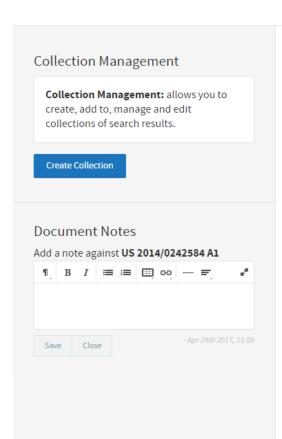
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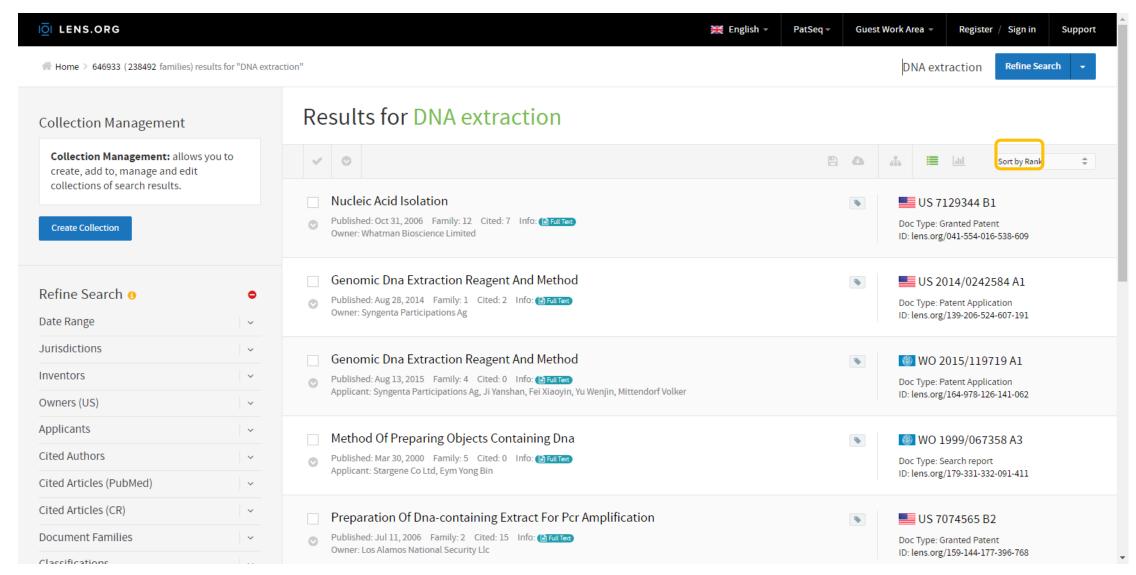
Full-text Citations Family Info Legal Info Notes 0 Summary Genomic Dna Extraction Reagent And Method Published: Aug 28, 2014 Family: 1 Cited: 2 Cites: 3 Non Patent Citations: 4 Info: Full Text Abstract ® The present invention is directed to a genomic DNA extraction reagent and method for improved extraction of DNA from biological tissue. The extraction reagent of the invention is mixed with disrupted biological tissue to form a DNA extraction solute which is incubated in a DNA extraction step. The extraction reagent includes an alkali component to maintain the DNA extraction solute at a pH of about 10 to 14 substantially throughout the extraction step. The extraction solute is centrifuged to clarify the supernatant. The supernatant containing the extracted DNA is diluted with a neutralizing buffer resulting in a high throughout method of generating high quantities of high quality DNA. Major PCR inhibitors are managed with the unique chemical combinations of the DNA extraction reagent designed and optimized for extraction of DNA from plant tissue and cells. Claims 📎 1. A DNA extraction reagent comprising an alkali, a surfactant, and a salt, wherein the pH of the reagent is at least 10. 2. The DNA extraction reagent of claim 1, wherein the alkali is NaOH. 3. The DNA extraction reagent of claim 1, wherein the surfactant is SDS. 4. The DNA extraction reagent of claim 1, wherein the salt is NH₄Ac. 5. The DNA extraction reagent of claim 1, further comprising a polyphenol absorbing compound. 6. The DNA extraction reagent of claim 5, wherein the polyphenol absorbing compound is PVP-40 7. The DNA extraction reagent according to claim 1, wherein the alkali is NaOH at a concentration of at least 0.1M, the surfactant is SDS at a concentration of at least 0.1%, and the salt is NH₄Ac at a concentration of at least 0.3M. 8. The DNA extraction reagent according to claim 1, wherein the pH of the reagent is at least 11. 9. The DNA extraction reagent according to claim 1, wherein the pH of the reagent is at least 12 10. ...Read More Owners (US) 📎 Applicants 📎 

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**US 2014/0242584 A1** 

Doc Type: Patent Application







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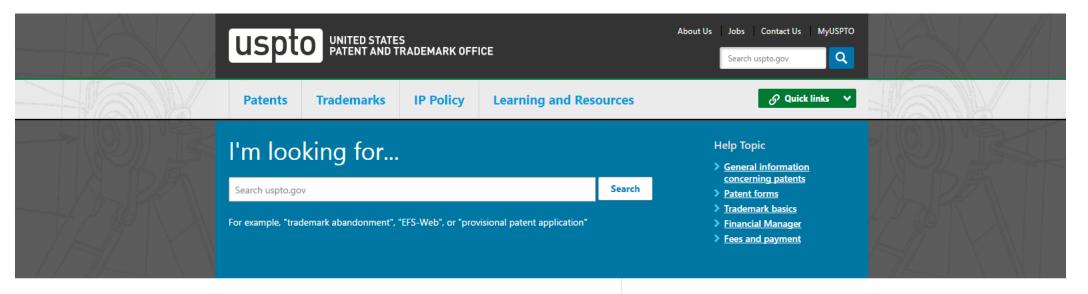
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		Soldering station with automatic soldering connection validation
		■ Datacenter in-row cooling units
		■ Electronic liquid cooling system including a bypass
		Refillable apparatus for aligning and depositing solder columns in a column grid array
		Reflow treating unit and substrate treating apparatus
		■ Intelligent soldering cartridge for automatic soldering connection validation
		Printed circuit board, ball grid array package and wiring method of printed circuit board
13	3 <u>9,629,239</u>	Resin composition, and prepreg as well as laminate using the same
		Touch panel having a sensing electrode and a printing electrode
1:	5 <u>9,629,220</u>	Sensor-based controllable LED lighting system with repositionable components and method
10	6 <u>9,629,216</u>	■ LED tube lamp
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48 7,582,031 Tennis ball holder

49 D594,702 Tennis ball shaped beverage container 50 D594,700 Tennis ball shaped beverage container

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#### Title 1 9.623.288 Table tennis ball and table tennis ball-use thermoplastic resin composition 2 9.592.429 Tennis court ball retainer 3 9,469,945 Ball collection and court drying system for a tennis court or the like 4 9.440,120 Ball holder for table tennis tables 5 D759,969 Tennis ball holder for use with dog collar 6 9,339,698 Tennis bag with ball retrieving and dispensing unit 7 D752,548 Tennis ball speaker 8 9.259.636 Tennis racquet airfoil training device 9 9.248.362 Table tennis paddle and ball holder 10 RE45,866 Tennis racquet with replaceable playing surface 11 D740,017 Tennis ball cane tip 12 9,114,285 Tennis ball pick-up cart 13 D729,891 String bed design for a tennis racquet 14 8,973,534 Pop up tennis ball pet tov 15 8.920.101 Tennis ball collection device 16 D719,227 Tennis racquet 17 D715,383 Tennis ball cat 18 D714,885 Tennis ball dog 19 8.771.375 Antimicrobial tennis ball 20 D706.027 Backpack having tennis ball material and texture 21 D704.567 Tennis ball bottle 22 D702,022 Tennis ball novelty headwear 23 D698,879 TString bed design for a tennis racquet 24 8.616.270 Tennis ball conditioner 25 8.602.711 Combination tennis ball cart and mower 26 8,556,565 Tennis ball retrieval device 27 8.534.726 Table tennis ball collector and dispenser 28 8.435.141 Tennis ball management system 29 8,414,431 Table tennis ball storage apron 30 8.313.396 Tennis ball vacuum collector 31 8.257,204 Automatic ball collection system for table tennis 32 8.192,308 Tennis racquet with replaceable playing surface 33 8.105.183 Celluloid-free table-tennis ball 34 8.028.345 Tennis garment with ball sleeves 35 8,002,651 Tennis racquet frame, its method of manufacture, and racquet comprising such a frame 36 7.922,608 Tennis ball retrieval, storage and dispensing device 37 7.896,760 Tennis ball delivery device 38 7.871.343 Tennis ball retriever 39 D627,020 Table tennis ball 40 D627.019 Table tennis ball 41 D627,018 Table tennis ball 42 D621,457 Automatic table tennis ball server 43 7,722,485 Tennis serve ball machine cum training device 44 7.690.543 Tennis ball holder 45 D611.707 Tennis ball bag 46 7.658.211 Tennis ball recharging apparatus method

47 D604,546 T Floating baseball, tennis ball, golf ball, or hockey puck display case



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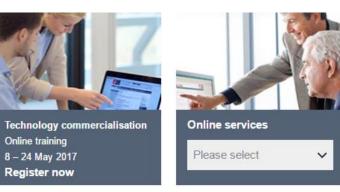


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Canadian Patent Office as well as PCT applications. The bibliographic and full text-coverage tables have been improved to indicate



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Quick help –	<b>★</b> → <b>= =</b>	(! i CPC		A»			
→ What is the Cooperative Patent Classification system?	Symbol	Classification and description					
→ How do I enter classification	A	HUMAN NECESSITIES	នាំ				
symbols?  → What do the different buttons	В	PERFORMING OPERATIONS; TRANSPORTING	s	i			
mean? → Can I retrieve a classification	C	CHEMISTRY; METALLURGY	s	i			
using keywords?  → Can I start a new search using	D	TEXTILES; PAPER	ន				
the classifications listed?	E	FIXED CONSTRUCTIONS	ន				
<ul> <li>→ Where can I view the description of a particular CPC class?</li> <li>→ What is the meaning of the stars</li> </ul>	F	MECHANICAL ENGINEERING; LIGHTING; HEATING; WEAPONS; BLASTING ENGINES OR PUMPS	s	i			
in front of the classifications found?	☐ <b>G</b>	PHYSICS	នាំ	i			
→ What does the text in brackets	H	ELECTRICITY	នាំ	i			
Mean? Selected classifications	Y	GENERAL TAGGING OF NEW TECHNOLOGICAL DEVELOPMENTS; GENERAL TAGGING OF CROSS-SECTIONAL TECHNOLOGIES SPANNING OVER SEVERAL SECTIONS OF THE IPC; TECHNICAL SUBJECTS COVERED BY FORMER USPC CROSS-REFERENCE ART COLLECTIONS [XRACs] AND DIGESTS	S	i			
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#### **Cooperative Patent Classification**

Search for DNA extraction

Œ	→ :::	□ I CPC	[] 2×0 2000
	Symbol	Classification	and description
•	****	C12Q 1/00	Measuring or testing processes involving enzymes, {nucleic acids} 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
•	statatata <del>k</del>	C12N 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, per se C12N 1/00, C12N 5/00, C12N 7/00; new plants per se A01H; plant reproduction by tissue culture techniques A01H 4/00; new animals per se 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)
-		B01L 3/00	Containers or dishes for laboratory use, e.g. laboratory glassware (bottles B65D; apparatus for enzymology or microbiology {specially adapted for culturing} C12M 1/00; Droppers (receptacles for volumetric purposes G01F)
•		B01L 2300/00	Additional constructional details
•		B01L 2400/00	Moving or stopping fluids
-		C12Q 2600/00	Oligonucleotides characterized by their use (not used, see subgroups)
-		☐ B01L 7/00	Heating or cooling apparatus (evaporators B01D 1/00; drying gases or vapours, e.g. desiccators, B01D 53/26; autoclaves B01J 3/04; drying ovens F26B; furnaces, ovens F27); Heat insulating devices
~		G06F 19/00	Digital computing or data processing equipment or methods, specially adapted for specific applications (G06F 17/00 takes precedence; data processing systems or methods specially adapted for administrative, commercial, financial, managerial, supervisory or forecasting purposes G06Q)
~		☐ B01J 20/00	Solid sorbent compositions or filter aid compositions; Sorbents for chromatography; Processes for preparing, regenerating or reactivating thereof (use of sorbent compositions in liquid separation B01D 15/00, use of filter aid compositions B01D 37/02; use of sorbent compositions in gas separation B01D 53/02, B01D 53/14)
•		C12N 1/00	Micro-organisms, e.g. protozoa; Compositions thereof (medicinal preparations containing material from micro-organisms A61K 35/66; 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

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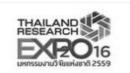
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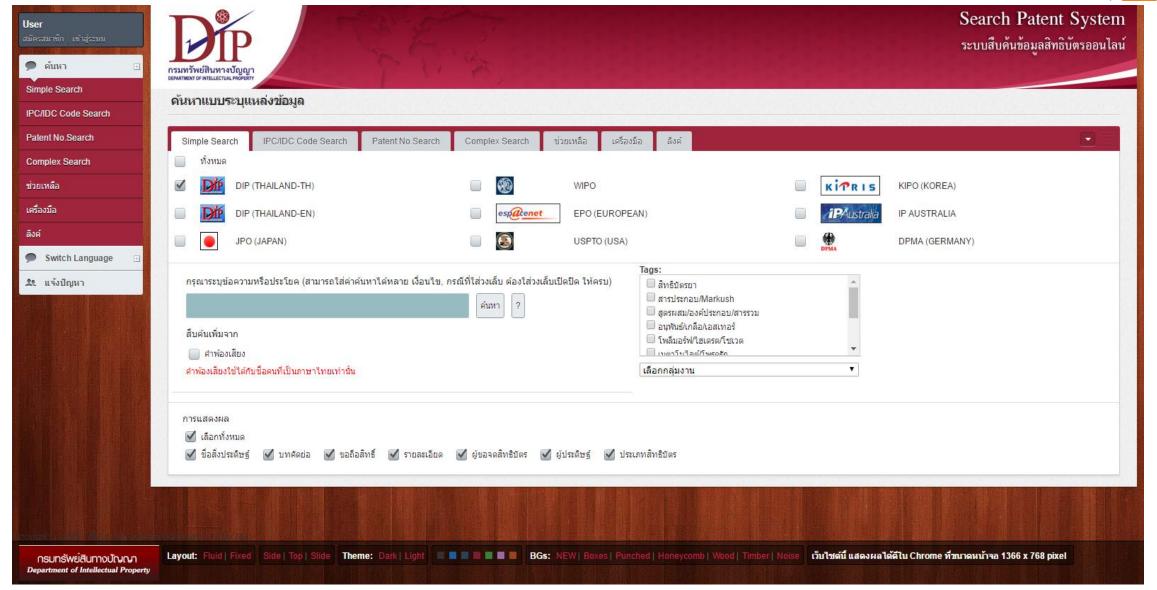


ศูนย์ข้อมูลข่าวสาร กรมทรัพย์สินทางปัญญา











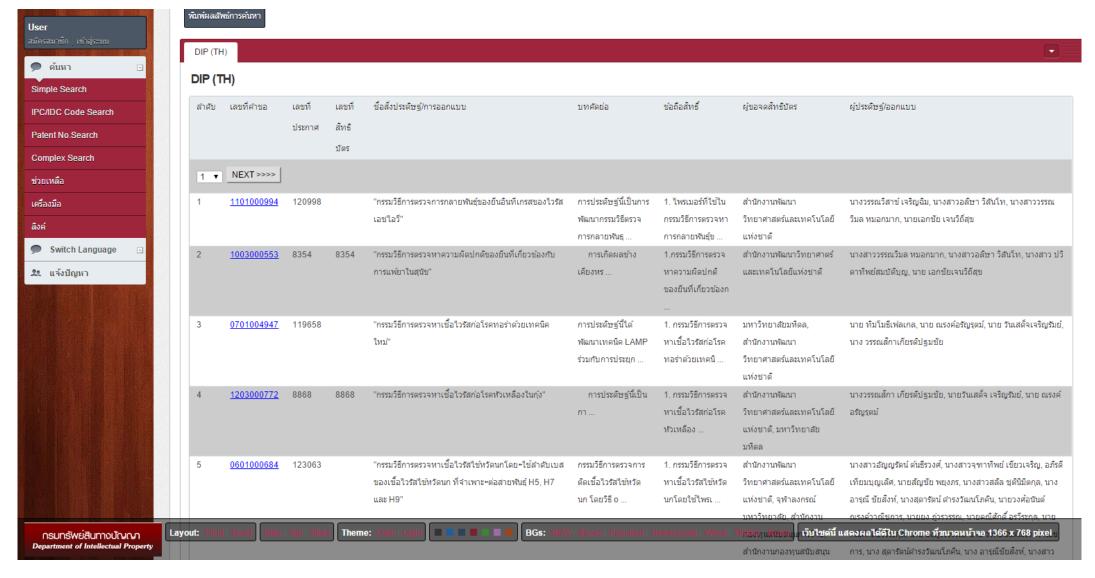
ศาพ้องเสียงใช้ได้กับชื่อคนที่เป็นภาษาไทยเท่านั้น	เลือกกลุ่มงาน	•
คำพ้องเสียง	<ul> <li>การใช้ครั้งที่ 2 หรือมากกว่าด้านเภสัชภัณฑ์</li> </ul>	~
สืบคันเพิ่มจาก	🔲 การใช้ครั้งแรกด้านเภสัชภัณฑ์	
สารพันธุกรรม คันหา ?	<ul> <li>ลักษณะทางกายภาพของเภสัชภัณฑ์</li> <li>กระบวนการผลิตของเภสัชภัณฑ์</li> </ul>	
THE TOTAL DESIGNATION OF THE THE THE THE THE TOTAL PROPERTY OF THE	🔲 เมตาโบ'ไลต์/โพรดรัก	
กรุณาระบุข้อความหรือประโยค (สามารถใส่ค่าค้นหาได้หลาย เงื่อนไข, กรณีที่ใส่วงเล็บ ต้องใส่วงเล็บเปิดปิด ให้ครบ)	Tags:  — เพลมอรพ/เฮเดรต/เซเวต	

# TOP 10 (Export Excel)

หมายเหตุ เฉพาะรายการที่คันได้จากฐานไทย DIP (TH) เท่านั้น

PC		Application Year	r	Publication Ye	ar	Registration Y	ear
IPC	Total	Year	Total	Year	Total	Year	Total
C12Q	32	2556	9	2556	13		58
C12N	19	2554	8	2559	13	2556	5
A61K	8	2550	6		10	2558	3
G01N	5	2553	5	2557	6	2557	2
C07K	3	2559	5	2558	6	2559	2
A01H	2	2557	5	2555	5	2553	1
C07H	2	2558	5	2554	3	2541	1
C12P	1	2549	4	2535	3	2555	1
C120	1	2552	4	2549	3	2560	1
A01K	1	2533	3	2548	2	2554	1
		Graph	Gra	aph	Gra	ph	Graph







ลำดับ	เลขที่คำขอ	เลขที	เลขที่	ชื่อสิ่งประดิษฐ์/การออกแบบ	บทศัดย่อ	ข้อถือสึทธิ์	ผู้ขอจดสิทธิบัตร	ผู้ประดิษฐ์/ออกแบบ
		ประกาศ	ส์ทธิ					
			บัตร					
1 🔻	NEXT >>>>							
-	TTEXT -							
1	<u>1101000994</u>	120998		"กรรมวิธีการตรวจการกลายพันธุ์ของยีนอินท์เกรสของไวรัส	การประดิษฐ์นี้เป็นการ	1. ไพรเมอร์ที่ใช้ใน	สำนักงานพัฒนา	นางวรรณวิสาข์ เจริญฉิม, นางสาวอลิษา วิสันโท, นางสาววรรณ
				เอชใอวี"	พัฒนากรรมวิธีตรวจ	กรรมวิธีการตรวจหา	วิทยาศาสตร์และเทคโนโลยี	วิมล หมอกมาก, นายเอกชัย เจนวิถีสุข
					การกลายพันธุ	การกลายพันธุ์ข	แห่งชาติ	
2	<u>1003000553</u>	8354	8354	"กรรมวิธีการตรวจหาความผิดปกติของยืนที่เกี่ยวข้องกับ	การเกิดผลข้าง	1.กรรมวิธีการตรวจ	สำนักงานพัฒนาวิทยาศาตร์	นางสาววรรณวิมล หมอกมาก, นางสาวอลิษา วิสันโท, นางสาว ปวิ
				การแพ้ยาในสุนัข"	เคียงหร	หาความผิดปกติ	และเทคโนโลยีแห่งชาติ	ตาทิพย์สมบัติบุญ, นาย เอกชัยเจนวิถีสุข
						ของยีนที่เกี่ยวข้องก		
3	0701004947	119658		"กรรมวิธีการตรวจหาเชื้อไวรัสก่อโรคทอร่าด้วยเทคนิค	การประดิษฐ์นี้ใด้	1. กรรมวิธีการตรวจ	มหาวิทยาลัยมหิดล,	นาย ทิมโมธีเฟลเกล, นาย ณรงค์อรัญรุตม์, นาย วันเสด็จเจริญรัมย์,
				ใหม่"	พัฒนาเทคนิค LAMP	หาเชื้อไวรัสก่อโรค	สำนักงานพัฒนา	นาง วรรณสึกาเกียรติปฐมชัย
					ร่วมกับการประยุก	ทอร่าด้วยเทคนิ	วิทยาศาสตร์และเทคโนโลยี	
							แห่งชาติ	
4	1203000772	8868	8868	"กรรมวิธีการตรวจหาเชื้อไวรัสก่อโรคหัวเหลืองในกุ้ง"	การประดิษฐ์นี้เป็น	1. กรรมวิธีการตรวจ	สำนักงานพัฒนา	นางวรรณสึกา เกียรติปฐมชัย, นายวันเสด็จ เจริญรัมย์, นาย ณรงค์
					กา	หาเชื้อไวรัสก่อโรค	วิทยาศาสตร์และเทคโนโลยี	อรัญรุตม์
						หัวเหลือง	แห่งชาติ, มหาวิทยาลัย	
							มหิดล	
5	<u>0601000684</u>	123063		"กรรมวิธีการตรวจหาเชื้อไวรัสไข้หวัดนกโดย÷ใช้ลำดับเบส	กรรมวิธีการตรวจการ	1. กรรมวิธีการตรวจ	สำนักงานพัฒนา	นางสาวอัญญรัตน์ ต้นธีรวงศ์, นางสาวจุฑาทิพย์ เขียวเจริญ, อภิรดี
			1	ของเชื้อไวรัสไข้หวัดนก ที่จำเพาะ÷ต่อสายพันธุ์ H5, H7	ติดเชื้อไวรัสไข่หวัด	หาเชื้อไวรัสไข้หวัด	วิทยาศาสตร์และเทคโนโลยี	เทียมบุญเลิศ, นายสัญชัย พยุงภร, นางสาวสลิล ชุตินิมิตกุล, นาง
				และ H9"	นก โดยวิธี o	นกโดยใช้ไพรเ	แห่งชาติ, จุฬาลงกรณ์	อารุณี ชัยสังห์, นางสุดารัตน์ ตำรงวัฒนโภคิน, นายวงศ์อนันต์
							มหาวิทยาลัย, สำนักงาน	ณรงค์วาณิชการ, นายยง ภู่วรวรรณ, นายคณิศักดิ์ อรวีระกุล, นาย
							กองทุนสนับสนุนการวิจัย,	คณิศักดิ์อรวีระกุล, นาย ยงภู่วรวรรณ, นาย วงศ์อนันต์ณรงค์วาณิช
							สำนักงานกองทุนสนับสนุน	การ, นาง สุดารัตน์ดำรงวัฒนโภคิน, นาง อารุณีขัยสิงห์, นางสาว
							การวิจัย, จุฬาลงกรณ์	สลิลชุตินิมิตกุล, นาย สัญชัยพยุงภร, นางสาว อภิรดีเทียมบุญเลิศ,

# http://www.wipo.int/portal/en/index.html





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Home > Reference > International Classifications > International Patent Classification

## International Patent Classification (IPC)

The International Patent Classification (IPC), established by the Strasbourg Agreement 1971, provides for a hierarchical system of language independent symbols for the classification of patents and utility models according to the different areas of technology to which they pertain. A new version of the IPC enters into force each year on January 1. Find out more about the IPC.

#### Resources

- General information on the IPC
- Guide to the IPC PDF
- IPC statistics
- Frequently Asked Questions

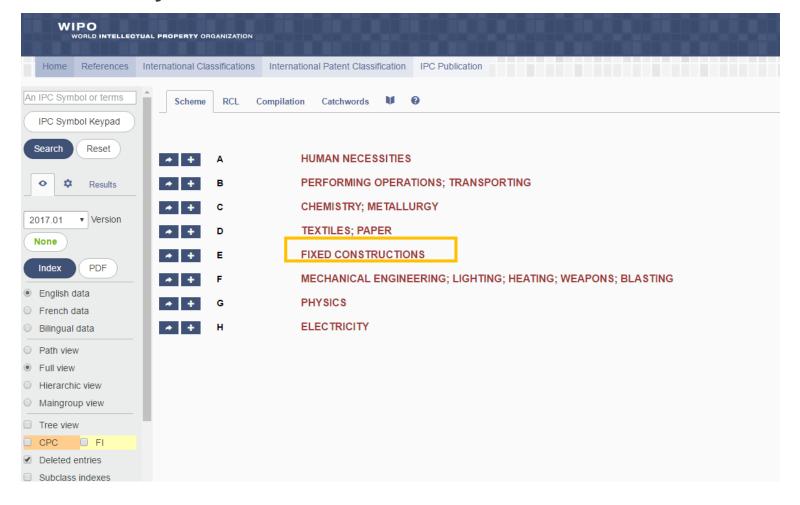
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# **International Patent Classification (IPC)**



The IPC divides technology into eight sections with approximately **70,000** subdivisions.





		CHEMISTRY
* +	C01	INORGANIC CHEMISTRY
* + I	C02	TREATMENT OF WATER, WASTE WATER, SEWAGE, OR SLUDGE
* + I	C03	GLASS; MINERAL OR SLAG WOOL
* + I	C04	CEMENTS; CONCRETE; ARTIFICIAL STONE; CERAMICS; REFRACTORIES [4]
* + I	C05	FERTILISERS; MANUFACTURE THEREOF [4]
* +	C06	EXPLOSIVES; MATCHES
* +	C07	ORGANIC CHEMISTRY [2]
* +	C08	ORGANIC MACROMOLECULAR COMPOUNDS; THEIR PREPARATION OR CHEMICAL WORKING-UP; COMPOSITIONS BASED THEREON
<b>*</b> +	C09	DYES; PAINTS; POLISHES; NATURAL RESINS; ADHESIVES; COMPOSITIONS NOT OTHERWISE PROVIDED FOR; APPLICATIONS OF MATERIALS NOT OTHERWISE PROVIDED FOR
4 +	C10	PETROLEUM. GAS OR COKE INDUSTRIES: TECHNICAL GASES CONTAINING CARBON MONOXIDE: FUELS: LUBRICANTS: PEAT
* +	C11	ANIMAL OR VEGETABLE OILS, FATS, FATTY SUBSTANCES OR WAXES; FATTY ACIDS THEREFROM; DETERGENTS; CANDLES
* +	C12	BIOCHEMISTRY; BEER; SPIRITS; WINE; VINEGAR; MICROBIOLOGY; ENZYMOLOGY; MUTATION OR GENETIC ENGINEERING
* +	C13	SUGAR INDUSTRY [4]
* +	C14	SKINS; HIDES; PELTS; LEATHER
		METALLURGY
* +	C21	METALLURGY OF IRON
* +	C22	METALLURGY; FERROUS OR NON-FERROUS ALLOYS; TREATMENT OF ALLOYS OR NON-FERROUS METALS
* +	C23	COATING METALLIC MATERIAL; COATING MATERIAL WITH METALLIC MATERIAL; CHEMICAL SURFACE TREATMENT; DIFFUSION TREATMENT OF METALLIC MATERIAL; COATING BY VACUUM EVAPORATION, BY SPUTTERING, BY ION IMPLANTATION OR BY CHEMICAL VAPOUR DEPOSITION, IN GENERAL; INHIBITING CORROSION OF METALLIC MATERIAL OR INCRUSTATION IN GENERAL [2]
* +	C25	ELECTROLYTIC OR ELECTROPHORETIC PROCESSES; APPARATUS THEREFOR [4]
* + I	C30	CRYSTAL GROWTH [3]

* -	<u>C12</u>	BIOCHEMISTRY; BEER; SPIRITS; WINE; VINEGAR; MICROBIOLOGY; ENZYMOLOGY; MUTATION OR GENETIC ENGINEERING
		Note(s) [5]  1. Between subclasses C12M-C12Q, and within each of these subclasses, the last place priority rule is applied, i.e. at each hierarchical level, 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 condition-responsive control is classified in subclass C12Q.
		2. In this class, viruses, undifferentiated human, animal or plant cells, protozoa, tissues and unicellular algae are considered as microorganisms.
		<ol><li>In this class, unless specifically provided for, undifferentiated human, animal or plant cells, protozoa, tissues and unicellular algae are classified together with microorganisms. Sub-cellular parts, unless specifically provided for, are classified with the whole cell.</li></ol>
		<ol> <li>The codes of subclass C12R are <u>only</u> for use as indexing codes associated with subclasses C12C-C12Q, so as to provide information concerning the microorganisms used in the processes classified in these subclasses.</li> </ol>
* <b>+</b>	C12C	BREWING OF BEER (cleaning of raw materials A23N; pitching or depitching machines, cellar tools C12L; propagating yeasts C12N 1/14)
* +	C12F	RECOVERY OF BY-PRODUCTS OF FERMENTED SOLUTIONS; DENATURING OF, OR DENATURED, ALCOHOL [6]
* +	C12G	WINE; OTHER ALCOHOLIC BEVERAGES; PREPARATION THEREOF (beer C12C)
* +	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]
* +	C12J	VINEGAR; ITS PREPARATION
* +	C12L	PITCHING OR DEPITCHING MACHINES; 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; brewing apparatus C12C; fermentation apparatus for wine C12G; apparatus for preparing vinegar C12J 1/10) [3]
* +	C12N	MICROORGANISMS OR ENZYMES; COMPOSITIONS THEREOF (biocides, pest repellants or attractants, or plant growth regulators containing microorganisms, viruses, microbial fungi, enzymes, fermentates, or substances produced by, or extracted from, microorganisms or animal material A01N 63/00; medicinal preparations A61K; fertilisers C05F); PROPAGATING, PRESERVING, OR MAINTAINING MICROORGANISMS; MUTATION OR GENETIC ENGINEERING; CULTURE MEDIA (microbiological testing media C12Q 1/00) [3]



+ +	C12P	FERMENTATION OR ENZYME-USING PROCESSES TO SYNTHESISE A DESIRED CHEMICAL COMPOUND OR COMPOSITION OR TO SEPARATE OPTICAL ISOMERS FROM A RACEMIC MIXTURE [3]
* ±	C12Q	MEASURING OR TESTING PROCESSES INVOLVING ENZYMES OR MICROORGANISMS (immunoassay G01N 33/53); COMPOSITIONS OR TEST PAPERS THEREFOR; PROCESSES OF PREPARING SUCH COMPOSITIONS; CONDITION-RESPONSIVE CONTROL IN MICROBIOLOGICAL OR ENZYMOLOGICAL PROCESSES [3]
* + I	C12R	INDEXING SCHEME ASSOCIATED WITH SUBCLASSES C12C-C12Q, RELATING TO MICROORGANISMS [3]

+ C12P FERMENTATION OR ENZYME-USING PROCESSES TO SYNTHESISE A DESIRED CHEMICAL COMPOUND OR COMPOSITION OR TO SEPARATE OPTICAL ISOMERS FROM A RACEMIC MIXTURE [3]



* -	C12P	FERMENTATION OR ENZYME-USING PROCESSES TO SYNTHESISE A DESIRED CHEMICAL COMPOUND OR COMPOSITION OR TO SEPARATE OPTICAL ISOMERS FROM A RACEMIC MIXTURE [3]
		Note(s) [6]  1. This subclass <u>covers</u> both major and minor chemical modifications.
		<ol> <li>Group C12P 1/00 covers processes for producing organic compounds not sufficiently identified to be classified in groups C12P 3/00-C12P 37/00. Compounds identified only by their empirical formulae are not considered to be sufficiently identified.</li> </ol>
		3. Attention is drawn to Notes (1) to (3) following the title of class C12.
		4. If a particular reaction is considered of interest, it is also classified in the relevant chemical compound class, e.g. C07, C08.
		5. In this subclass:
		metal or ammonium salts of a compound are classified as that compound.
		compositions are classified in the relevant compound groups.
		6. In this subclass, it is desirable to add the indexing codes of subclass C12R.
<b>*</b> -	C12P 1/00	Preparation of compounds or compositions, not provided for in groups C12P 3/00-C12P 39/00, by using microorganisms or enzymes; General processes for the preparation of compounds or compositions by using microorganisms or enzymes [2006.01]
*	C12P 1/02	• by using fungi [2006.01]
*	C12P 1/04	• by using bacteria [2006.01]
*	C12P 1/06	• by using actinomycetales [2006.01]
*	C12P 3/00	Preparation of elements or inorganic compounds except carbon dioxide [2006.01]
* -	C12P 5/00	Preparation of hydrocarbons [2006.01]
*	C12P 5/02	• acyclic [2006.01]

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The invention r	elates	to a novel	I metho	d for th	e synth	nesis o	nfan e	rythro	ocvte	protei	in in wi	nich said ni	roteir	n is synth	nesised	lin an a	cellula	r system	for the pro	duction of
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2. WO/2017/06	4373	DEVICE E	OR DRO	DHCI	NGAI	MIXTI	IRE O	S S DI	RIIII	IN A IN	I SEAW	ATER IN I	IVE	CHITHR	F			WO	20.04.201	7
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The invention of	oncer	ns a meth	od for n	roducir	na a mi	ixture	of spir	ulina	in se	awate	er in live	culture co	oneie	ting of a	rrangir	a spiruli	ina in s	eawatei	in live cult	ure in a
receiving mear	ıs (2),	said spirul	lina in s	eawate	er in liv	e culti	ure ha	ving	a pre	define	ed deve	lopment sta	ate; k	keeping t	the spi	rulina in	seawa	ater in liv	e culture in	said
predefined dev with a second of					ig mea	ns (2)	; addir	ig se	awat	er in n	notion t	o a contain	ier (4	); mixing	a first	quantity	of spir	ulina in	seawater in	live culture
with a second (	quandi	y or seawa	ater III II	iouon.																
3. WO/2017/06	5203	ANTI-CAN	IINE TA	RC AN	ITIBOD	OY US	ED FO	R TE	REAT	MENT	TAND	DIAGNOSI	s of	CANINE	E ATO	PIC		wo	20.04.201	7
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Provided is an	anti-ca	anine TAR	C antib	ody use	ed for t	reatm	ent an	d dia	gnos	is of c	anine a	topic derm	atitis	and a m	nethod	for treat	ment o	r diagno	sis of canin	e atopic

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# **Exercises**

- Prior Art Search
  - Find international patent application WO/2006/076067 using esp@cenet, uspto, patentscope and patent lens
  - Find number of application filed by Toyota using
    - Patentscope
    - esp@cenet
    - Uspto
    - JPO/IPDL
    - and patent lens
  - · Find who is the leading company in the field of electric car



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