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# ORGANISATION MONDIALE DE LA PROPRIÉTÉ INTELLECTUELLE GENÈVE

# UNION PARTICULIÈRE POUR LA CLASSIFICATION INTERNATIONALE DES BREVETS (UNION DE L'IPC)

# **COMITÉ D'EXPERTS**

Vingt-neuvième session Genève, 13 – 17 mars 2000

## DEMANDE DE RÉVISION DE LA CLASSIFICATION INTERNATIONALE DES BREVETS (CIB)

Document établi par le Bureau international

L'annexe du présent document contient une proposition concernant la création d'une nouvelle sous-classe relative à la chimie combinatoire élaborée par le Royaume-Uni suite à une demande de révision soumise au comité d'experts (voir l'annexe 1 du document IPC/CE/29/7).

[L'annexe suit]

#### ANNEX/ANNEXE

## C40B Combinatorial chemistry proposal

Please find attached detailed IPC revision proposal for C40B combinatorial chemistry. As discussed at the last meeting of the revision working group, the proposal only contains a very broad breakdown of main groups. I anticipate it will largely be necessary to classify elsewhere as well as in this subclass according to the actual chemical structures involved. I will attempt to define "line notes" (references to other areas of IPC) in due course.

Being a totally new subclass, I though it was a great opportunity to try our proposed new format for subclasses. I have therefore devised a very sparse subclass with allied classification definitions for the subclass and various main groups. It is my intention to hyperlink the various definitions to the relevant parts of the schedule to which they relate but I haven't actually made the links as I thought they would not translate to Word format.

I apologise that the details of this schedule have been submitted so long after the original notification of submission of a proposal. I'm sure you and all my colleagues in IPC will let me know what they think of a somewhat experimental project!

Best Regards

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### C40B Combinatorial chemistry

### **Classification subgroups**

1/00	synthesis characterised by chemical nature of the compound library produced
3/00	apparatus specially adapted for use in the synthesis of chemical libraries
5/00	screening, methods specially adapted for use in screening of chemical libraries
7/00	identification of active compounds present in chemical libraries
9/00	combinatorial chemistry subjects not covered by 1/00 to 7/00

## **C40B** Combinatorial chemistry

#### **Subclass definitions**

This subclass is primarily concerned with combinatorial chemistry, *ie* the creation of large populations of compounds *en masse* (known as a chemical or combinatorial library). Combinatorial synthesis may be followed by screening for desired activity and identification of individual compounds in the chemical library.

In this subclass the following terms are used with the following meanings:

Combinatorial chemistry: the systematic and repetitive connection of a set of different `building blocks` of varying structures to each other to yield a large array of diverse molecular entities

Screening: the process of determining whether compounds in a chemical library have a desired chemical or biological activity, *without* necessarily identifying the precise chemical nature of the compound(s) being screened

Iterative deconvolution: a method of identifying compounds in a chemical library involving the successive comparison (eg for activity) of an unknown compound with each of a set of sub-libraries, each sub-library having a fixed, known sub-unit in one of the positions, the set covering all possible sub-units for each other position.

#### Example:

Consider a library of 100 different esters of formula R₁-CO₂-R₂, formed by the reaction of 10 different carboxylic acids of formula R₁-CO₂H with 10 different alcohols of formula R₂-OH, and where one ester of highest activity is detected.

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Iterative deconvolution, is used to find the highest activity ester, as follows:

10 sub-libraries of formula  $R_1$ - $CO_2$ - $R_2$  are formed, each being formed from **one** of the 10 acids  $R_1$ - $CO_2$ H, and all 10 of the alcohols  $R_2$ -OH, thus each sub-library has 10 members and  $R_1$  is known ( $R_1$  is the fixed, known sub-unit). The most active sub-library tells us which  $R_1$  is likely to be present in the most active ester in the 100 member library.

Then 10 further sub-libraries are formed, each being formed from **one** of the 10 alcohols  $R_2$ -OH, and all 10 of the acids  $R_1$ -CO<sub>2</sub>H thus each sub-library has 10 members and  $R_2$  is known. The most active sub-library tells which  $R_2$  is likely to be present in the most active ester in the 100 member library.

Thus the identity of the highest activity ester is suggested by the result of the iterative deconvolution. (It should be noted that the most active ester need not be the ester corresponding to the  $R_1$  and  $R_2$  sub-libraries showing highest activity - iterative deconvolution is not necessarily definitive.)

#### C40B Combinatorial chemistry

Classification definitions for main group 1/00 - synthesis characterised by chemical nature of the compound library produced

Chemical natures including:-

inorganic libraries

peptide libraries (including polypeptide and protein libraries)

amide-linked oligomeric libraries

nucleic acid libraries

oligomeric or polymeric libraries

synthesis characterised by chemical nature of the compound library produced nonoligomeric organic libraries

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#### C40B Combinatorial chemistry

Classification definitions for subgroup 5/00 - screening, methods specially adapted for use in screening of chemical libraries

Screening is defined as the process of determining whether compounds in a chemical library have a desired chemical or biological activity, *without* necessarily identifying the precise chemical nature of the compound(s) being screened

### C40B Combinatorial chemistry

Classification definitions for subgroup 7/00 - identification of active compounds present in chemical libraries

7/00 includes identification through means dependent on the particular synthetic methodology used, e.g.:-

identification by position in space, eg by use of grids of phials

identification by tagging, ie association with an identifiable moiety (eg using nucleotides or radioactive markers)

identification by iterative deconvolution

7/00 also includes identification through means through means not dependent on particular synthetic methodology, e.g.:-

spectroscopy

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