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(54) Title: MICROENABLED CHEMICAL REACTION IN MICROFLUIDIC CHIPS

(57) Abstract: A chemical process to obtain products of higher purity or in shorter time is provided by performing chemical reactions in a reactions volume less than 1 microliter. The advantages of this process are due to better mixing or heat exchanche (each 10 fold decrease in reaction volume leads to a 100-fold improvement mixing or heat exchange) or better surface to volume ratio. Better surface to volume ratio is a major factor in reactions critically dependent on surface mediated events such as heterogeneous catalysis. This increase in yield and speed of reaction can be conducted in parallel sets of reaction chambers provided by multilayer glass and silicon chips. Reagents can be added via a microfluidic chip incorporating channels and orthogonal row/column reagent delivery options. This discovery is amendable to solid supported synthesis as well as solution synthesis. Because of the better surface to volume ratio, this process is particularly useful with solid catalyst on the edges of the reaction chamber and can be used for hydrogenation, and hydroformulation. This process is demonstrated in reactions involving nucleophilic aromatic substitution on a solid support.

INTERNATIONAL SEARCH REPORT

International application No. PCT/US00/16056

A. CLASSIFICATION OF SUBJECT MATTER IPC(7) :C12Q 1/00, 1/68; A61K/38/00; C07H 21/00, 21/02, 21/04 US CL :Please See Extra Sheet. According to International Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) U.S. : 435/4, 6, DIGEST 46, DIGEST 49; 422/129, 187, 232, 530/333, 334, 338, 341; 536/25.3 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) WEST, DIALOG: terms: microreactor, chemical synthesis, catalyst, peptide, nucleotide, DNA, micorfludics, ultramicrotiter, array, library, combinatorial					
C. DOCUMENTS CONSIDERED TO BE RELEVANT					
Category*	Citation of document, with indication, where ap	propriate, of the relevant passages	Relevant to claim No.		
X Y	US 5,472,672 A (BRENNAN) 05 December 1995, see entire document, especially the abstract and claims.		1, 2, 5, 6, 9 3, 4, 7, 8, 10-14, 16-20		
Y	US 5,474,796 A (BRENNAN) 12 Ded document, especially column 7 lines 45	1-2, 5-12, 14-20			
Y Y	US 5,585,275 A (HUDSON et al) 17 E document, especially the abstract and f	igures.	1, 2, 6, 9, 11-13, 16-20 14		
X Further documents are listed in the continuation of Box C. See patent family annex.					
"A" do to to "E" ear cit sp. "O" do me	ecial categories of cited documents: cument defining the general state of the art which is not considered be of particular relevance rlier document published on or after the international filing date cument which may throw doubts on priority claim(s) or which is ed to establish the publication date of another citation or other ecial reason (as specified) cument referring to an oral disclosure, use, exhibition or other enas	"Y" later document published after the int date and not in conflict with the app the principle or theory underlying the document of particular relevance; the considered novel or cannot be considered to involve an inventive combined with one or more other such being obvious to a person skilled in document member of the same pater.	dication but cited to understand e invention ne claimed invention cannot be ered to involve an inventive step ne claimed invention cannot be e step when the document is sh documents, such combination the art		
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International application No.
PCT/US00/16056

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C (Continua	ation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages		Relevant to claim No
<i>(</i>	US 5,008,040 A (TSUKAMOTO et al) 16 April 1991, see entire document especially column 4 lines 44-50.		3, 4
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	US 5605,662 A (HELLER et al) 25 February 1997, see document especially the abstract and figures.	entire	1, 2, 6, 9, 11, 14, 15, 16, 17
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