## **International Nonproprietary Names (INN) for biological and biotechnological substances**

(a review)



## **International Nonproprietary Names (INN) for biological and biotechnological substances**

(a review)



International Nonproprietary Names (INN) Programme

Technologies Standards and Norms (TSN) Regulation of Medicines and other Health Technologies (RHT) Essential Medicines and Health Products (EMP)

# International Nonproprietary Names (INN) for biological and biotechnological substances

(a review)

FORMER DOCUMENT NUMBER: INN Working Document 05.179

#### © World Health Organization 2014

All rights reserved. Publications of the World Health Organization are available on the WHO website (<u>www.who.int</u>) or can be purchased from WHO Press, World Health Organization, 20 Avenue Appia, 1211 Geneva 27, Switzerland (tel.: +41 22 791 3264; fax: +41 22 791 4857; e-mail: <u>bookorders@who.int</u>).

Requests for permission to reproduce or translate WHO publications –whether for sale or for noncommercial distribution– should be addressed to WHO Press through the WHO website (www.who.int/about/licensing/copyright\_form/en/index.html).

The designations employed and the presentation of the material in this publication do not imply the expression of any opinion whatsoever on the part of the World Health Organization concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement.

The mention of specific companies or of certain manufacturers' products does not imply that they are endorsed or recommended by the World Health Organization in preference to others of a similar nature that are not mentioned. Errors and omissions excepted, the names of proprietary products are distinguished by initial capital letters.

All reasonable precautions have been taken by the World Health Organization to verify the information contained in this publication. However, the published material is being distributed without warranty of any kind, either expressed or implied. The responsibility for the interpretation and use of the material lies with the reader. In no event shall the World Health Organization be liable for damages arising from its use.

#### Printed by the WHO Document Production Services, Geneva, Switzerland

### CONTENTS

0.	INT	RODUCTIONiv
1.		ARMACOLOGICAL CLASSIFICATION OF BIOLOGICAL AND DTECHNOLOGICAL SUBSTANCES
2.		RRENT STATUS OF EXISTING STEMS OR SYSTEMS FOR BIOLOGICAL D BIOTECHNOLOGICAL SUBSTANCES
2	2.1.	Groups with respective stems
2	2.2.	Groups with respective pre-stems
2	2.3.	Groups with INN schemes
2	2.4.	Groups without respective stems / pre-stems and without INN schemes
3.		NERAL POLICIES FOR BIOLOGICAL AND BIOTECHNOLOGICAL BSTANCES
3	8.1.	General policies for blood products
3	8.2.	General policies for fusion proteins
3	3.3.	General policies for gene therapy products
3	8.4.	General policies for glycosylated compounds7
3	8.5.	General policies for immunoglobulins fractionated from plasma
3	8.6.	General policies for monoclonal antibodies
3	8.7.	General policies for non-glycosylated compounds 11
3	8.8.	General policies for skin substitutes 11
3	8.9.	General policies for transgenic products

	3.10.	General policies for vaccines	. 12
	3.11.	General policies for cell therapy products	. 12
4		MMARY OF INN ASSIGNED TO BIOLOGICAL AND DTECHNOLOGICAL SUBSTANCES	. 13
	4.1.	Antimicrobial, bactericidal permeability increasing polypeptides	. 13
	4.2.	Antisense oligonucleotides	. 13
	4.3.	Antithrombins	. 13
	4.4.	Blood coagulation cascade inhibitors	. 14
	4.5.	Blood coagulation factors	. 14
	4.6.	Colony stimulating factors	. 15
	4.7.	Enzymes	. 16
	4.8.	Erythropoietin type blood factors	. 20
	4.9.	Gene therapy products	. 21
	4.10.	Gonadotropin-releasing-hormone (GnRH) inhibitors, peptides	. 21
	4.11.	Growth factors	. 21
	4.12.	Growth hormone (GH) derivatives	. 23
	4.13.	Growth hormone antagonists	. 23
	4.14.	Heparin derivatives including low molecular mass heparins	. 23
	4.15.	Hirudin derivatives	. 24
	4.16.	Insulins	. 24
	4.17.	Interferons	. 25
	4.18.	Interleukin receptor antagonists	. 26
	4.19.	Interleukin type substances	. 26

4	.20.	Monoclonal antibodies	. 27		
4	.21.	Oxytocin derivatives	. 30		
4	.22.	Peptides and glycopeptides	. 30		
4	.23.	Peptide vaccines / recombinant vaccines	. 33		
4	.24.	Pituitary / placental glycoprotein hormones	. 34		
4	.25.	Pituitary hormone-release stimulating peptides	. 35		
4	.26.	Receptor molecules, native or modified	. 35		
4	.27.	Synthetic polypeptides with a corticotropin-like action	. 36		
4	.28.	Thrombomodulins	. 36		
4	.29.	Toxins	. 37		
4	.30.	Vasoconstrictors, vasopressin derivatives	. 37		
4	.31.	Various	. 37		
5.	CU	RRENT CHALLENGES	. 46		
RE	REFERENCES				
AN	NEX	1	49		
Т	The lis	at of INN for composite proteins published	. 49		
		2			
Т	Transl	iteration of Greek letters in English, French and Spanish	. 69		
AN	ANNEX 3				
Т	The previous naming scheme for monoclonal antibodies70				

### **0. INTRODUCTION**

More than 50 years ago, WHO established the International Nonproprietary Name (INN) Expert Group / WHO Expert Committee on Specifications for Pharmaceutical Preparations, to assign nonproprietary names to medicinal substances, so that each substance would be recognized globally by a unique name. These INNs do not give proprietary rights, unlike a trade mark, and can be used freely as they are public property.

INNs have been assigned to biological products since the early days of the INN Programme. As well as many names for individual substances, animal insulin preparations were given an INN in Recommended list 3 in 1959. In the period up to 1980, names were assigned to antibiotics, synthetic peptides, hormones and other proteins. In names of compounds related by structure and / or function, specific letter groups, called stems, are included to aid recognition by health professionals. The *-actide* for synthetic polypeptides with a corticotrophin-like action is an example.

In 1982, the name *insulin human* was proposed for the recombinant protein identical to natural human insulin, and since then names have been assigned to a growing number of recombinant products. Within the INN Programme, names have not been assigned to natural human blood products or vaccines. For those groups of biological products, the WHO Expert Committee on Biological Standardization (ECBS) has been adopting the scientific names of the biological products within the definitions of respective requirements.

Since the time when *insulin human* became the first recommended INN (rINN) for a recombinant product, the range of biological / biotechnological products has increased in size and complexity. For example, new stems have been introduced for tissue plasminogen activators (*-plase*) among other groups. Analogues of recombinant glycosylated proteins produced in different cell systems have been classified using Greek letters as indicators in the sequence of product introduction: erythropoietin (*epoetin alfa*, *beta* and so on) and glycoprotein hormones (*follitropin*) are examples. In the 1990s, a systematic scheme for naming monoclonal antibodies was implemented based on the

### 预览已结束, 完整报告链接和二维码如下:

https://www.yunbaogao.cn/report/index/report?reportId=5\_27976

