



**World Health
Organization**

INN Working Document 16.392 rev.
ENGLISH ONLY
11/07/2016

***Pre-stems*:
Suffixes used in the selection of INN
April 2016***

Programme on International Nonproprietary Names (INN)

Technologies Standards and Norms (TSN)

Regulation of Medicines and other health technologies (RHT)

***World Health Organization,
Geneva***

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*The prestems given have been flagged because they may be selected as official stems ("The use of stems in the selection of International Nonproprietary Names for Pharmaceutical Substances", 2013, WHO/EMP /RHT/TSN/2013.1). At present, they are made available for information and potential guidance to the applicants.

<i>stem</i>	<i>definition</i>
-suffix	
-infix-	
In bold:	new pre-stems selected during the last Consultation.
<u>In bold and underlined:</u>	newly promoted stems
<u>Underlined:</u>	modified pre-stem definition
<hr/>	
-algron	α_1 -adrenoreceptor agonists
-ampator	α -amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid (AMPA) receptor modulators
-axomab	see <i>mab</i>
-berel	beta estrogen receptor agonists
-brutinib	see <i>tinib</i>
-caftor	cystic fibrosis transmembrane regulator (CFTR) protein modulators
-calcet/-calcet-	calcium-sensing receptors (CaSR) agonists
-camra	intracellular adhesion molecule (ICAM-1) derivatives
-casan	caspase inhibitors
-caserin	serotonin receptor agonists (mostly 5-HT ₂)
-catib	cathepsin inhibitors
-cetrapib	cholesteryl transfer protein (CETP) inhibitors
-closporin	ciclosporin derivatives
-codar	see <i>dar</i>
-corat	glucocorticoid receptor agonists
-cridar	see <i>dar</i>
-dacin	antibiotics, DNA gyrase and topoisomerase IV inhibitors
-dar	<i>drugs used in multidrug resistance</i>
-cri-	acridinecarboxamide derivatives
-co-	pipecolate derivatives
-spo-	ciclosporin D derivatives

<u>-degib</u>	SMO receptor antagonists
-depsin	depsipeptide derivatives
-dil	vasodilators
-sudil	Rho protein kinase inhibitors
-domide	antineoplastics, thalidomide derivatives
-dustat	see <i>stat</i>
-ectedin	ecteinascidin derivatives
-espib	heat shock protein (HSP) 90 inhibitors (other than <i>-mycin</i>), antineoplastics
-estrant	estrogen antagonists
-fadine	monoamine transport inhibitors
-farnib	farnesyl transferase inhibitors
-fibatide	see <i>tide</i>
-fulven	antineoplastic, acylfulvene derivatives
-gacestat	see <i>-stat</i>
-ganan	antimicrobial, bactericidal permeability increasing polypeptides
-gepant	calcitonin gene-related peptide receptor antagonists
-gapil	neuronal apoptosis inhibitors, GAPDH
-imepodib	inosine monophosphate dehydrogenase inhibitors
-imod	immunomodulators, both stimulant/suppressive and stimulants
<u>-tolimod</u>	<u>toll-like receptors (TLR) agonists</u>
-inurad	urate transporter inhibitors
<u>-isant</u>	<u>histamine H₃ receptor antagonists</u>
-ixibat	ileal bile acid transporter (IBAT) inhibitors, bile acid reabsorption inhibitors

- <i>kalner</i>	openers of calcium-activated (maxi-K) K ⁺ -channels
- <i>laner</i>	antagonists of GABA (gamma-aminobutyric acid) regulated chloride channels, antiparasitic agents
- <i>leptin(e)</i>	leptin derivatives
- <i>mab</i> under species	<i>monoclonal antibodies</i>
- <i>axo</i> -	rat-murine hybrid antibodies
- <i>vet</i> -	veterinary use
under targets	
- <i>am(i)</i> -	serum amyloid protein (SAP)/amyloidosis
- <i>gr(o)</i> -	skeletal muscle mass related growth factors and receptors
- <i>melanotide</i>	<i>see tide</i>
- <i>metinib</i>	<i>see tinib</i>
- <i>moren</i>	non-peptidic growth hormone secretagogues
- <i>nesib</i>	kinesin inhibitors
- <i>neurin</i>	neurotrophins
- <i>nexor</i>	nuclear export inhibitors
- <i>nil</i> - <i>punil</i>	<i>benzodiazepine receptor antagonists/agonists</i> mitochondrial benzodiazepine receptor (MBR)- selective agonists also partial or inverse (purine derivatives)
- <i>opran</i>	μ-opioid receptors antagonists
- <i>orexant</i>	orexin receptor antagonists
- <i>osuran</i>	urotensin receptor antagonists
- <i>otilate</i>	hepatoprotectants, diisopropyl-1,3-dithiol-malonate derivatives
- <i>parantag</i>	antagonists of heparin and/or low-molecular weight heparins (LMWH)
- <i>paxar</i>	protease activated receptor type 1 (PAR1) antagonists
- <i>piridine</i>	serotonin receptor antagonists

<i>-plasimin</i>	inhibitors of plasminogen activator inhibitors-type 1 (PAI-1)
<u>-prazan</u>	proton pump inhibitors, not dependent on acid activation
<i>-prinin</i>	nootropic agents, purine derivatives
<i>-protafib</i>	protein tyrosine phosphatase (HPTP) inhibitors
<i>-punil</i>	see <i>nil</i>
<i>-reotide</i>	see <i>-tide</i>
<i>-siban</i>	oxytocin antagonists
<i>-sidenib</i>	isocitrate dehydrogenase inhibitors
<i>-spodar</i>	see <i>dar</i>
<i>-stat/-stat</i>	<i>enzymes inhibitors</i>
<i>-dustat</i>	hypoxia inducible factor (HIF) prolyl hydroxylase inhibitors
<i>-gacestat</i>	gamma-secretase inhibitors
<i>-stinel</i>	<u>NMDA receptor antagonists/partial agonists</u>
<i>-sudil</i>	see <i>dil</i>
<i>-sulind</i>	antineoplastics, sulindac metabolites <i>exisulind</i> (80)(42)
<u>-tansine</u>	maytansinoide derivatives, antineoplastics
<i>-tegravir</i>	see <i>vir</i>
<i>-terone</i>	<i>antiandrogens</i>
<i>-teronel</i>	non-steroid antiandrogens

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_26925

