



**World Health  
Organization**

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***Pre-stems\*:  
Suffixes used in the selection of INN  
August 2021***

***Programme on International Nonproprietary Names (INN)***

***Health Products Policy and Standards (HPS)***

***Access to Medicines and Health Products (MHP)***

***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", 2018, WHO/EMP /RHT/TSN/2018.1). At present, they are made available for information and potential guidance to the applicants.

*stem*

*definition*

-*suffix*

-*infix*-

**In bold:** new pre-stems selected during the last Consultation.

**In bold and underlined:** pre-stems newly promoted as stems.

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- <i>adenant</i>	antagonists of adenosine receptors
- <i>algron</i>	$\alpha_1$ -adrenoreceptor agonists
- <i>alkib</i>	ALK (anaplastic lymphoma kinase) inhibitors
- <i>ampator</i>	$\alpha$ -amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid (AMPA) receptor modulators
- <i>atovir</i>	see <i>vir</i>
- <i>batinib</i>	see <i>tinib</i>
- <i>bactam</i>	$\beta$ -lactamase inhibitors
- <i>borbactam</i>	$\beta$ -lactamase inhibitors, boronic acid derivatives
<b><u>-bep</u></b>	<u>from engineered or synthetic protein scaffolds, non-immunoglobulin variable domain derived <b>to engineered or synthetic scaffolds proteins with non-immunoglobulin variable derived binding domains</b></u>
- <i>berel</i>	beta estrogen receptor agonists
- <i>borbactam</i>	see <i>-bactam</i>
<b><u>-bresib</u></b>	<u><b>inhibitors of the bromodomain and extra-terminal motif (BET) family of bromodomain (BRD) proteins, antineoplastics</b></u>
- <i>camra</i>	intracellular adhesion molecule (ICAM-1) derivatives
- <i>camtiv</i>	cardiac myosin activators
- <i>capavir</i>	see <i>vir</i>
- <i>casan</i>	caspase inhibitors
- <i>caserin</i>	serotonin receptor agonists (mostly 5-HT <sub>2</sub> )
- <i>cianine</i>	indocyanine fluorescence dye group

-closporin	ciclosporin derivatives
-codar	see <i>dar</i>
-corat	glucocorticoid receptor agonists
<b>-corilant</b>	<b>glucocorticoid receptor antagonists (<i>non-steroidal</i>)</b>
<b>-corvir</b>	see <i>vir</i>
-cridar	see <i>dar</i>
-dacin	antibiotics, DNA gyrase and topoisomerase IV inhibitors
-dar	<i>drugs used in multidrug resistance</i>
-cridar	acridinecarboxamide derivatives
-codar	pipecolinate derivatives
-spodar	ciclosporin D derivatives
<b>-delpar</b>	<b>PPAR delta agonists</b>
-demstat	see <i>stat</i>
-depsin	depsipeptide derivatives
-dil	<i>vasodilators</i>
<u><b>-sudil</b></u>	<u><b>Rho protein kinase inhibitors</b></u>
<b>-dutide</b>	see <b>-tide</b>
-ectedin	ecteinascidin derivatives
<b>-enatide</b>	see <b>-tide</b>
<u><b>-estrant</b></u>	<u><b>estrogen antagonists, including estrogen receptor down-regulators</b></u>
-fadine	monoamine transport inhibitors
-farnib	farnesyl transferase inhibitors
-fenicol	antibacterial antibiotics, chloramphenicol analogues
-fibatide	see <i>tide</i>
-folastat	see <i>-stat</i>

<i>-fulven</i>	antineoplastic, acylfulvene derivatives
<i>-ganan</i>	antimicrobial, bactericidal permeability increasing polypeptides
<i>-gapil</i>	neuronal apoptosis inhibitors, GAPDH
<i>-gratinib</i>	<i>see -tinib</i>
<i>-imepodib</i>	inosine monophosphate dehydrogenase inhibitors
<i>-inapant</i>	inhibitors of inhibition-of-apoptosis proteins (IAPs)
<i>-inurad</i>	urate transporter inhibitors
<i>-kalner</i>	openers of calcium-activated (maxi-K) K <sup>+</sup> -channels
<i>-leptin(e)</i>	leptin derivatives
<i>-lintide</i>	See <i>-tide</i>
<i>mab</i> under targets	<i>monoclonal antibodies</i>
<i>-ami-</i>	serum amyloid protein (SAP)/amyloidosis
<b><u>-madlin</u></b>	<b><u>E3 ubiquitin-protein ligase Mdm2 (Hdm2) inhibitors</u></b>
<i>-melanotide</i>	<i>see tide</i>
<b><u>-meran</u></b>	<b><u>messenger RNA (mRNA)</u></b>
<i>-metkib</i>	MET (mesenchymal epithelial transition factor) kinases inhibitors
<b><u>-metostat</u></b>	<i>see stat</i>
<i>-moren</i>	non-peptidic growth hormone secretagogues
<i>-nersen</i>	<i>see -rsen</i>
<i>-nesib</i>	kinesin inhibitors
<i>-neurin</i>	neurotrophins
<i>-nexor</i>	nuclear export inhibitors
<i>nil</i>	<i>benzodiazepine receptor antagonists/agonists</i>

<i>-punil</i>	mitochondrial benzodiazepine receptor (MBR)-selective agonists, also partial or inverse (purine derivatives)
<i>-opran</i>	$\mu$ -opioid receptors antagonists
<i>-(o)pterin</i>	pteridine derivatives
<i>-osuran</i>	urotensin receptor antagonists
<i>-otilate</i>	hepatoprotectants, di(propan-2-yl) 2-(2 <i>H</i> -1,3-dithiol-2-ylidene)propanedioate and analogues
<i>-parantag</i>	antagonists of heparin and/or low-molecular weight heparins (LMWH)
<i>-paxar</i>	protease activated receptor type 1 (PAR1) antagonists
<i>-pertin</i>	glycine transporter inhibitors
<i>-pirdine</i>	serotonin receptor antagonists
<i>-pixant</i>	purinoreceptor (P2X) antagonists
<i>-plasinin</i>	inhibitors of plasminogen activator inhibitors-type 1 (PAI-1)
<i>-plenib</i>	Spleen tyrosine kinase (Syk) inhibitors
<i>-prinin</i>	nootropic agents, purine derivatives
<b><i>-prodil</i></b>	<b><i>N</i>-methyl-D-aspartate (NMDA) receptor antagonists</b>
<i>-protafib</i>	protein tyrosine phosphatase (HPTP) inhibitors
<i>-pultide</i>	see -tide
<i>-punil</i>	see <i>nil</i>

预览已结束，完整报告链接和二维码如下：

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