

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

Programme on International Nonproprietary Names (INN)

Medicines and Health Products

***World Health Organization,
Geneva***

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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|---------------------------|---|
| -algron | α_1 -adrenoreceptor agonists |
| -alkib | <i>ALK (anaplastic lymphoma kinase) inhibitors</i> |
| -ampator | α -amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid (AMPA) receptor modulators |
| -ase -liase | enzymes lyases (EC class 4) |
| -atovir | see <i>vir</i> |
| -bactam -borbactam | β -lactamase inhibitors β -lactamase inhibitors, boronic acid derivatives |
| -batinib | see <i>-tinib</i> |
| -berel | beta estrogen receptor agonists |
| -borbactam | see <i>-bactam</i> |
| -caltamide | T-type calcium channel blockers |
| -camra | intracellular adhesion molecule (ICAM-1) derivatives |
| -camtiv | cardiac myosin activators |
| -capavir | see <i>vir</i> |
| -casan | caspase inhibitors |
| -caserin | serotonin receptor agonists (mostly 5-HT ₂) |
| -cept -tacicept | receptor molecules or membrane ligands, native or modified TACI (TNFRSF13B)-derived TNF receptors |

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| <i>-cianine</i> | indocyanine fluorescence dye group |
| <i>-closporin</i> | ciclosporin derivatives |
| <i>-codar</i> | see <i>dar</i> |
| <i>-cridar</i> | see <i>dar</i> |
| <i>-corilant</i> | glucocorticoid receptor antagonists (non-steroidal) |
| <i>-corvir</i> | see <i>vir</i> |
| <i>-dacin</i> | antibiotics, DNA gyrase and topoisomerase IV inhibitors |
| <i>dar</i> <i>-codar</i> <i>-cridar</i> <i>-spodar</i> | <i>drugs used in multidrug resistance</i> pipercolinate derivatives acridinecarboxamide derivatives ciclosporin D derivatives |
| <i>-delpar</i> | PPAR delta agonists |
| <u>-demstat</u> | <u>see stat</u> |
| <i>-depsin</i> | depsipeptide derivatives |
| <i>-dirsen</i> | see <i>-rsen</i> |
| <i>-dutide</i> | see <i>-tide</i> |
| <i>-ectedin</i> | ecteinascidin derivatives |
| <u>-enatide</u> | <u>see -tide</u> |
| <i>-fadine</i> | monoamine transport inhibitors |
| <i>-farnib</i> | farnesyl transferase inhibitors |
| <i>-fenicol</i> | antibacterial antibiotics, chloramphenicol analogues |
| <i>-fibatide</i> | see <i>tide</i> |
| <i>-folastat</i> | see <i>-stat</i> |

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|--------------------------------|---|
| <i>-fulven</i> | antineoplastics, acylfulvene derivatives |
| <i>-ganan</i> | from antimicrobials, bactericidal permeability increasing polypeptides to antimicrobials, permeability increasing peptides |
| <i>-gapil</i> | neuronal apoptosis inhibitors, GAPDH |
| <i>-gli</i> <i>-gliatin</i> | antihyperglycaemics glucokinase activators |
| <i>-gratinib</i> | see <i>-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 ⁺ -channels |
| <i>-leptin(e)</i> | leptin derivatives |
| <i>-liase</i> | see <i>-ase</i> |
| <i>-lintide</i> | see <i>-tide</i> |
| <i>mab</i> <i>-ami-</i> | <i>monoclonal antibodies</i> serum amyloid protein (SAP)/amyloidosis |
| <i>-melanotide</i> | see <i>-tide</i> |
| <i>-metkib</i> | MET (mesenchymal epithelial transition factor) kinases inhibitors |
| <i>-moren</i> | non-peptidic growth hormone secretagogues |
| <i>-nersen</i> | see <i>-rsen</i> |
| <i>-nesib</i> | kinesin inhibitors |
| <i>-neurin</i> | neurotrophins |
| <i>-nexor</i> | nuclear export inhibitors |

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|------------------------------|---|
| <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>-(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>-perten</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>-prinim</i> | nootropic agents, purine derivatives |
| <i>-prodil</i> | <i>N</i> -methyl-D-aspartate (NMDA) receptor antagonists |
| <i>-protafib</i> | protein tyrosine phosphatase (HPTP) inhibitors |
| <i>-pultide</i> | see <i>-tide</i> |

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_23281

