

**Table S2** Inhibitors of CDKs

<b>Name</b>	<b>Inhibitor</b>	<b>Clinical stage</b>	<b>Delivery</b>	<b>Mechanism and synergistic effect</b>	<b>Reference</b>
<b>CDK1</b>	BEY-1107, Roniciclib	Phase I/II	PO	Kinase activity, VEGFR	(132)
<b>CDK2</b>	Inditinib, FN-1501, Milciclib, PF-06873600, CYC-065, Seliciclib, Roniciclib	Phase I/II	IV/PO	Kinase activity, Src family	(132)
<b>CDK3</b>	Roniciclib	Phase I/II	PO	Kinase activity, VEGFR	(132)
<b>CDK4</b>	Palbociclib, Ribociclib, Abemaciclib, Trilaciclib, Lerociclib, SHR-6390, MM-D37K, PF-06873600, Birociclib, BPI-16350, FCN-437, BEBT-209, TQB-3616, Roniciclib, AMG-925	Phase I/II	IV/PO	Kinase activity, FLT3	(132)
<b>CDK6</b>	Palbociclib, Ribociclib, Abemaciclib, Trilaciclib, Lerociclib, SHR-6390, MM-D37K, PF-06873600, Birociclib, BPI-16350, FCN-437, BEBT-209, TQB-3616, AMG-925	Phase I/II	IV/PO	Kinase activity, FLT3	(132)
<b>CDK7</b>	C7001, SY-1365, Roniciclib	Phase I/II	PO/IV	Kinase activity	(132)

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<b>CDK8</b>	BCD-115	Phase II	PO	Kinase activity	(132)
<b>CDK9</b>	Flavopiridol, Zotiraciclib, CYC-065, Seliciclib, AT-7519, AZD-4573, TP-1287, Voruciclib, BAY-1251152, Roniciclib, Atuveciclib	Phase I/II	IV/PO	Kinase activity, FLT3, JAK2, MAPK7, GSK3 $\beta$ , VEGFR	(132)
<b>CDK12</b>	Dinaciclib, THZ1, THZ531, E9, CDK12-IN-3, SR 4835, Procatenol	Phase I	IV	Kinase activity, IPA, DDR, MYC, SEATF, p-RNA POL II, RNA POL II, MCL1, PARPc, DDC, PARPi	(28,132)
<b>CDK19</b>	BCD-115	Phase II	PO	Kinase activity	(132)

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Abbreviation: PO: per os, IV: Injection of vein. DDC: DNA damage chemotherapy, RNA POL II: RNA polymerase II, IPA: Intronic polyadenylation, PARPi: PARP inhibitors, PARPc: PARP cleavage, SEATF: SE-associated transcription factors.