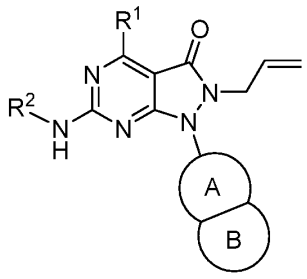
WHAT IS CLAIMED IS:

1. Use of a combination of compounds for treating a disease or condition, wherein the combination includes an effective amount of Compound (A) and an effective amount of one or more of Compound (B), or a pharmaceutically acceptable salt thereof, wherein:

the Compound (A) has the structure:



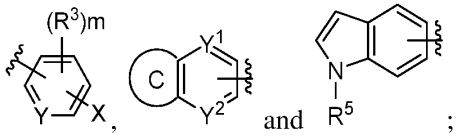
(A)

wherein:

R^1 is selected from the group consisting of hydrogen, halogen and a substituted or unsubstituted C_1-C_6 alkyl;

Ring A is selected from the group consisting of a substituted or unsubstituted phenyl and a substituted or unsubstituted 5-6 membered monocyclic heteroaryl;

Ring B is selected from the group consisting of a substituted or unsubstituted monocyclic 5-7 membered carbocyclyl and a substituted or unsubstituted 5-7 membered monocyclic heterocyclyl;

R^2 is selected from the group consisting of ;

m is 0, 1, 2 or 3;

R^3 is selected from the group consisting of halogen and a substituted or unsubstituted C_1-C_6 alkyl;

X is selected from the group consisting of hydrogen, halogen, hydroxy, cyano, a substituted or unsubstituted 4-6 membered monocyclic heterocyclyl, a substituted or unsubstituted amine(C_1-C_6 alkyl), a substituted or unsubstituted –NH-$(CH_2)_{1-6}$-amine, a mono-substituted amine, a di-substituted amine, an amino, a substituted or unsubstituted C_1-C_6 alkyl, a substituted or unsubstituted C_1-C_6 alkoxy, a substituted or unsubstituted C_3-C_6 cycloalkoxy, a substituted or unsubstituted (C_1-C_6 alkyl)acyl, a substituted or unsubstituted C-amido, a

substituted or unsubstituted N-amido, a substituted or unsubstituted C-carboxy, a substituted or unsubstituted O-carboxy, a substituted or unsubstituted O-carbamyl and a substituted or unsubstituted N-carbamyl;

Y is CH or N;

Y^1 is CR^{4A} or N;

Y^2 is CR^{4B} or N;

Ring C is selected from the group consisting of a substituted or unsubstituted C_6-C_{10} aryl, a substituted or unsubstituted monocyclic 5-10 membered heteroaryl, a substituted or unsubstituted monocyclic 5-7 membered carbocyclyl, a substituted or unsubstituted 5-7 membered monocyclic heterocyclyl and a substituted or unsubstituted 7-10 membered bicyclic heterocyclyl;

R^{4A} and R^{4B} are independently selected from the group consisting of hydrogen, halogen and an unsubstituted C_{1-4} alkyl; and

R^5 is a substituted or unsubstituted 5-7 membered monocyclic heterocyclyl; and

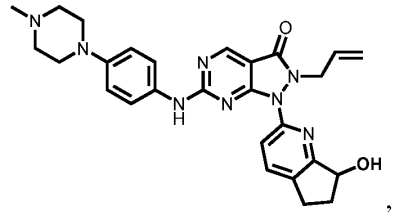
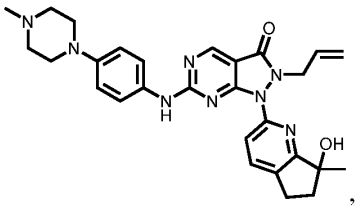
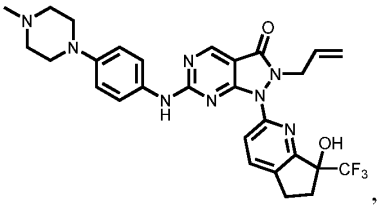
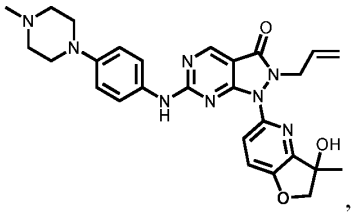
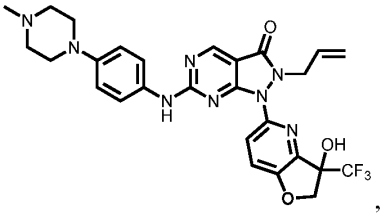
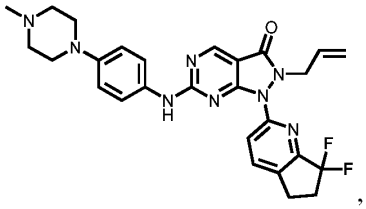
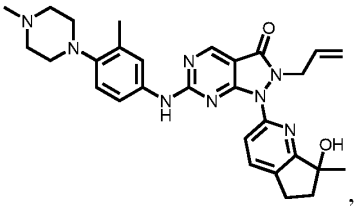
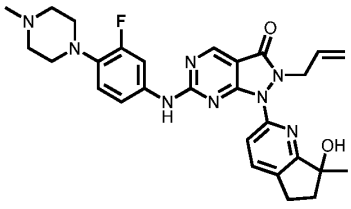
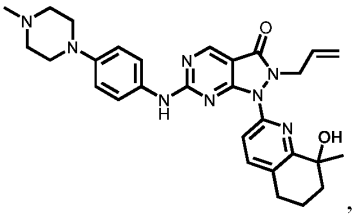
the one or more of Compound (B) is selected from the group consisting of a SERD compound and a SERM compound, or a pharmaceutically acceptable salt of any of the foregoing;

wherein the SERD compound is selected from the group consisting of fulvestrant, (E)-3-[3,5-Difluoro-4-[(1R,3R)-2-(2-fluoro-2-methylpropyl)-3-methyl-1,3,4,9-tetrahydropyrido[3,4-b]indol-1-yl]phenyl]prop-2-enoic acid (AZD9496), (R)-6-(2-(ethyl(4-(2-(ethylamino)ethyl)benzyl)amino)-4-methoxyphenyl)-5,6,7,8-tetrahydronaphthalen-2-ol (elacestrant, RAD1901), (E)-3-(4-((E)-2-(2-chloro-4-fluorophenyl)-1-(1H-indazol-5-yl)but-1-en-1-yl)phenyl)acrylic acid (Brilanestrant, ARN-810, GDC-0810), (E)-3-(4-((2-(2-(1,1-difluoroethyl)-4-fluorophenyl)-6-hydroxybenzo[b]thiophen-3-yl)oxy)phenyl)acrylic acid (LSZ102), (E)-N,N-dimethyl-4-((2-((5-((Z)-4,4,4-trifluoro-1-(3-fluoro-1H-indazol-5-yl)-2-phenylbut-1-en-1-yl)pyridin-2-yl)oxy)ethyl)amino)but-2-enamide (H3B-6545), (E)-3-(4-((2-(4-fluoro-2,6-dimethylbenzoyl)-6-hydroxybenzo[b]thiophen-3-yl)oxy)phenyl)acrylic acid (rintodestrant, G1T48), D-0502, SHR9549, ARV-471, 3-((1R,3R)-1-(2,6-difluoro-4-((1-(3-fluoropropyl)azetidin-3-yl)amino)phenyl)-3-methyl-1,3,4,9-tetrahydro-2H-pyrido[3,4-b]indol-2-yl)-2,2-difluoropropan-1-ol (giredestrant, GDC-9545), (S)-8-(2,4-dichlorophenyl)-9-(4-((1-(3-fluoropropyl)pyrrolidin-3-yl)oxy)phenyl)-6,7-dihydro-5H-benzo[7]annulene-3-carboxylic acid

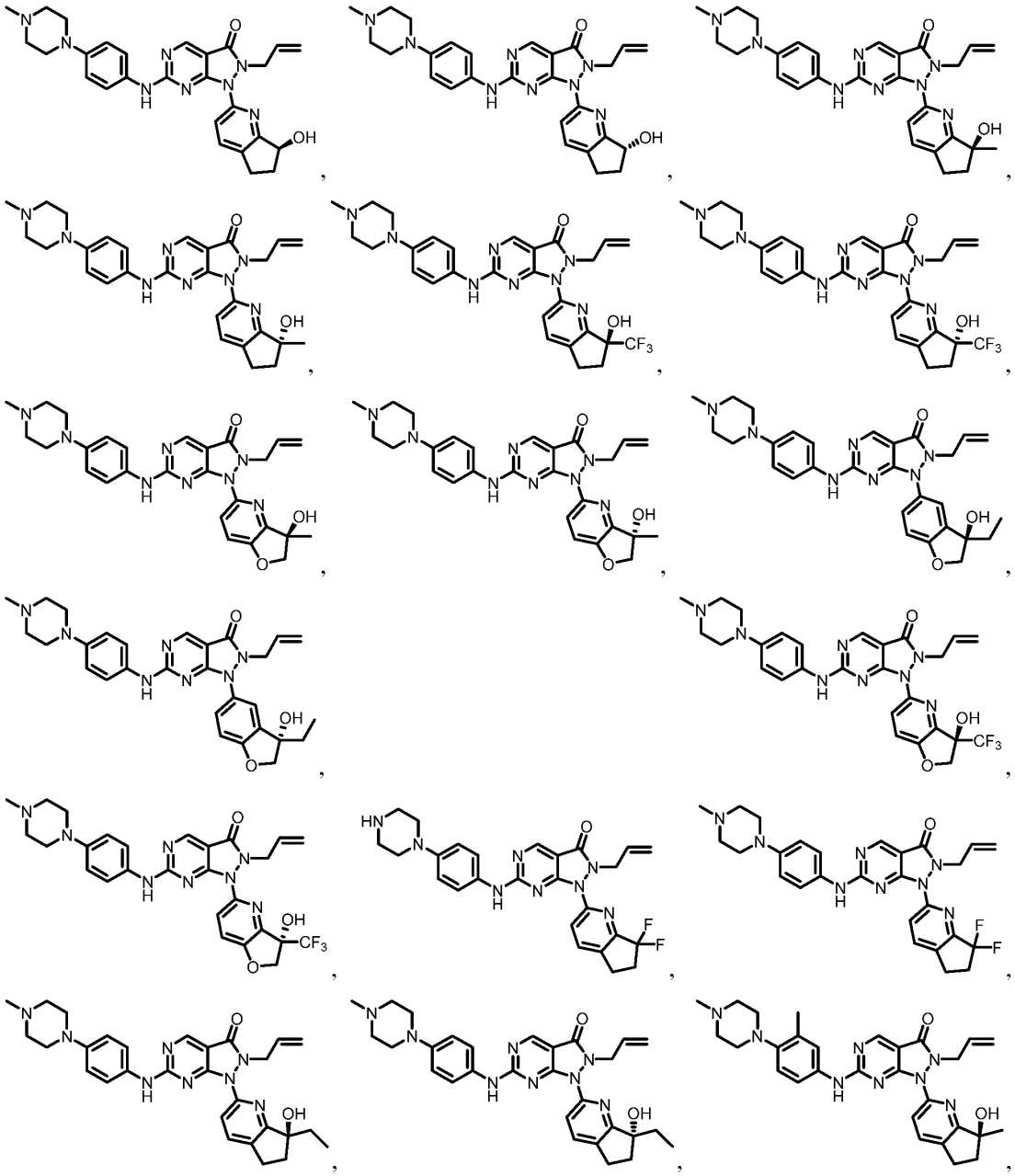
(SAR439859), N-[1-(3-fluoropropyl)azetidin-3-yl]-6-[(6S,8R)-8-methyl-7-(2,2,2-trifluoroethyl)-6,7,8,9-tetrahydro-3H-pyrazolo[4,3-f]isoquinolin-6-yl]pyridin-3-amine (AZD9833), OP-1250 and LY3484356, or a pharmaceutically acceptable salt of any of the foregoing; and

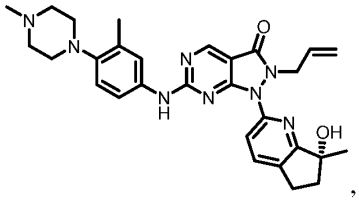
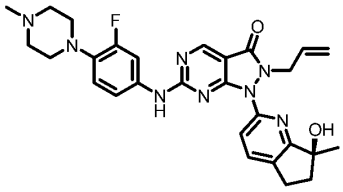
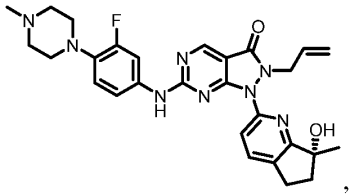
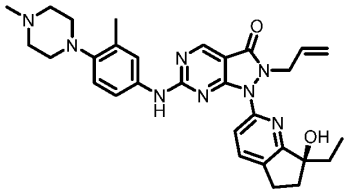
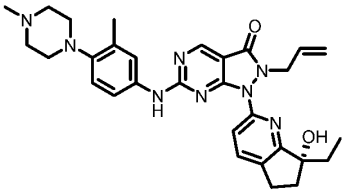
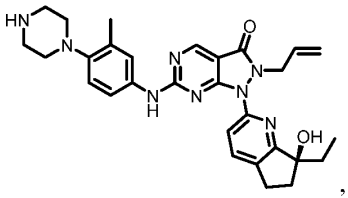
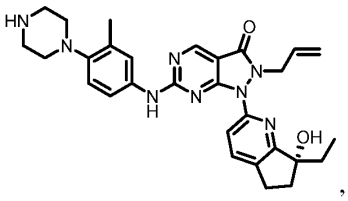
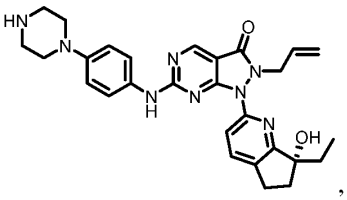
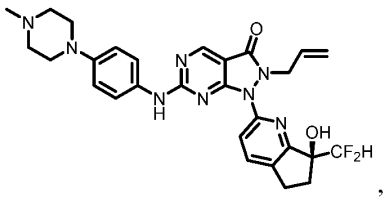
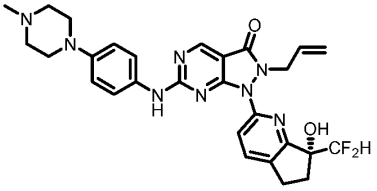
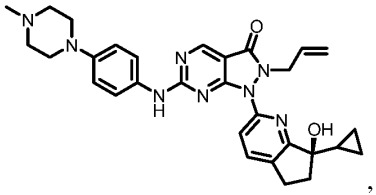
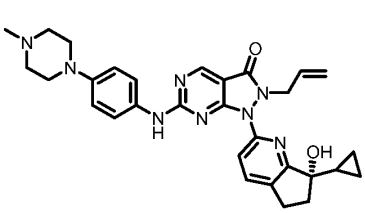
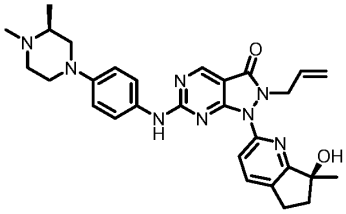
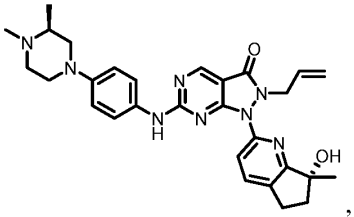
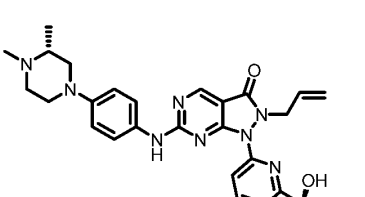
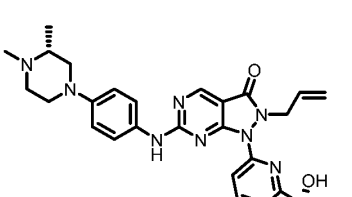
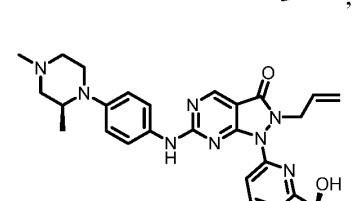
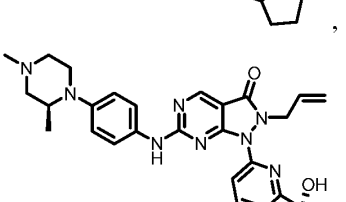
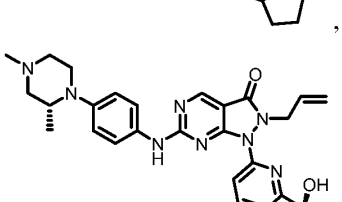
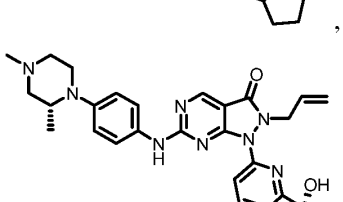
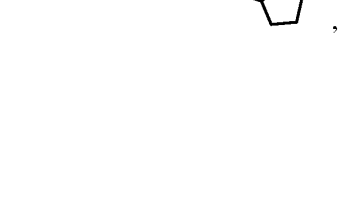
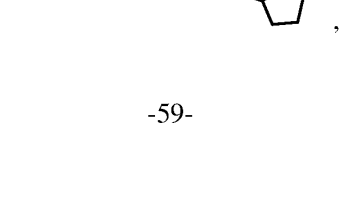
wherein the SERM compound is selected from the group consisting of tamoxifen, raloxifene, ospemifene, bazedoxifene, toremifene and lasofoxifene, or a pharmaceutically acceptable salt of any of the foregoing.

2. The use of Claim 1, wherein the Compound (A) is selected from the group consisting of:

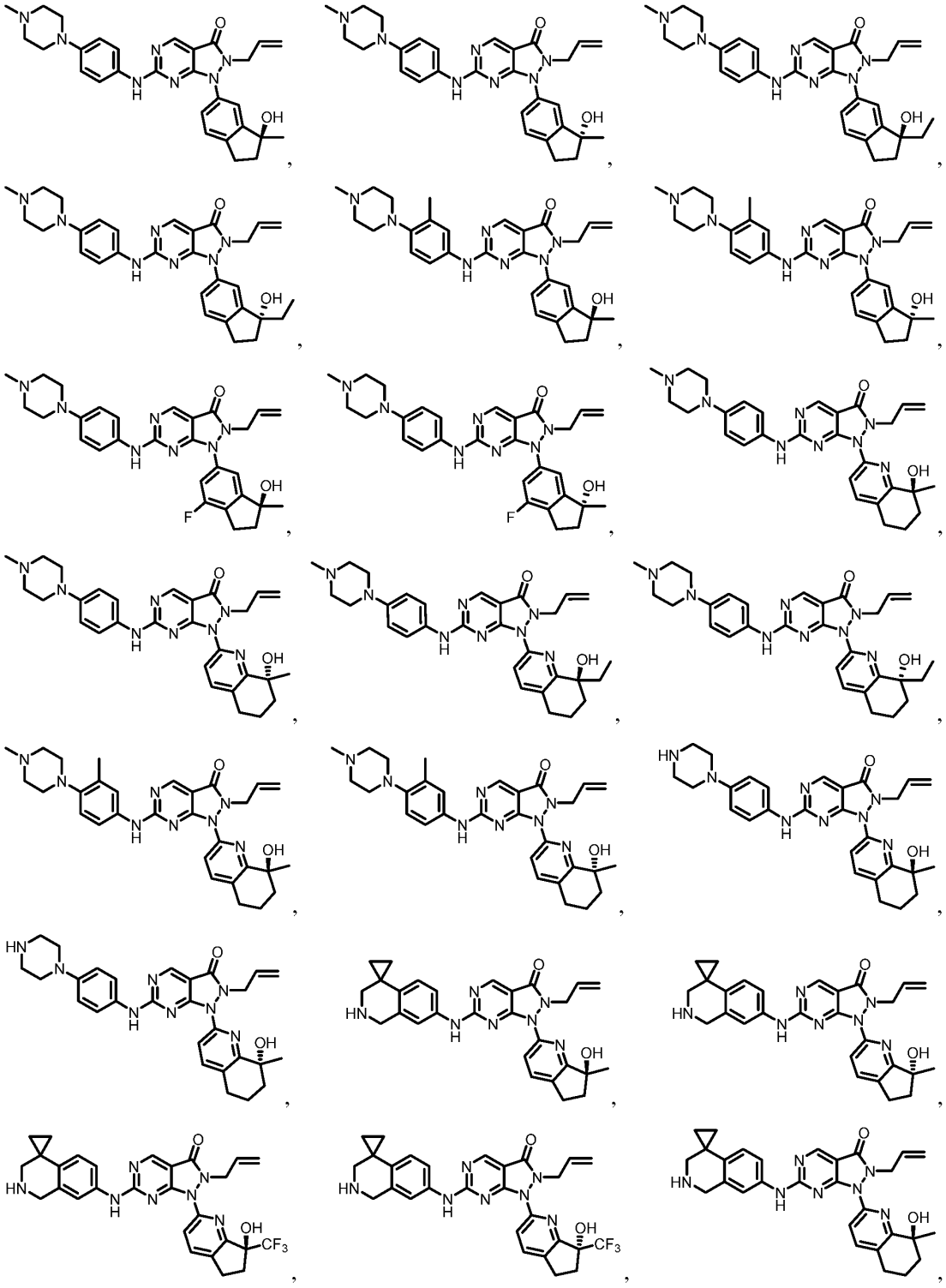

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, or a pharmaceutically acceptable salt of any of the foregoing.

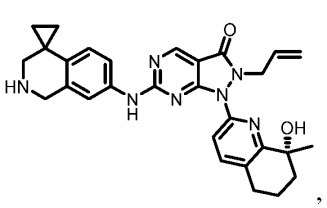
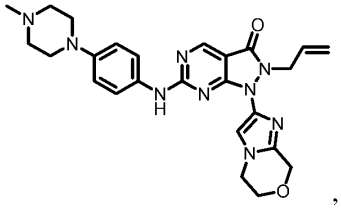
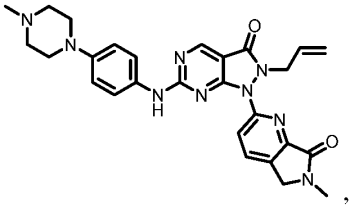
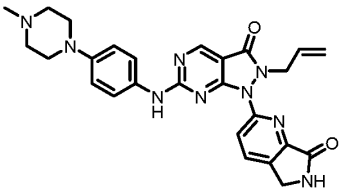
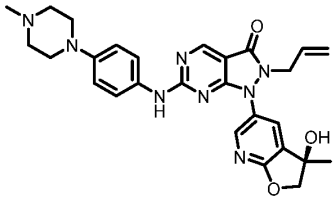
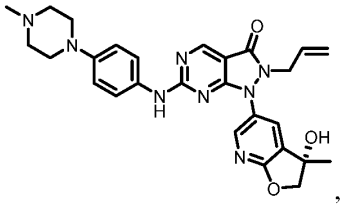
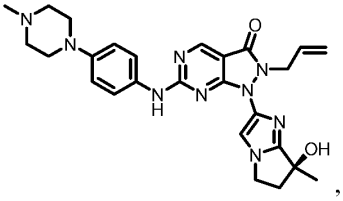
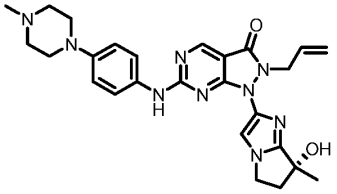
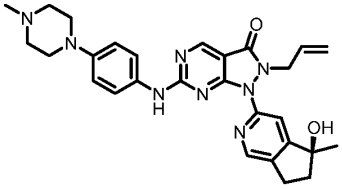
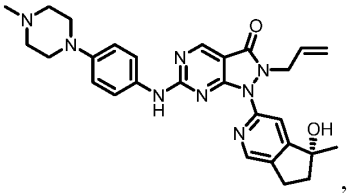
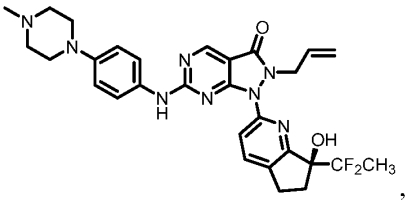
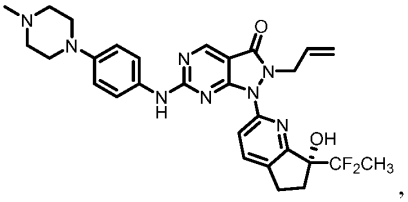
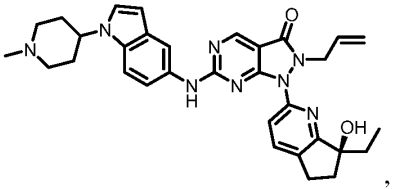
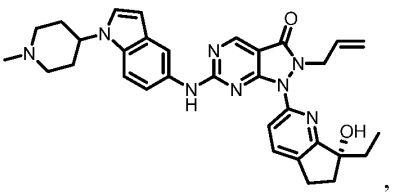
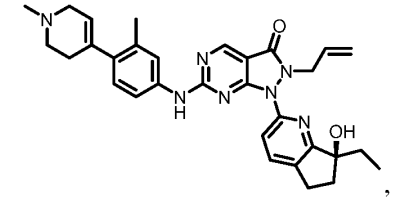
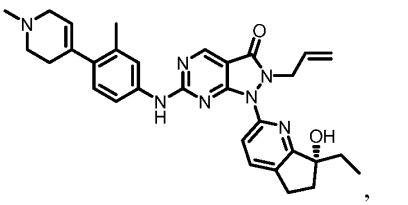
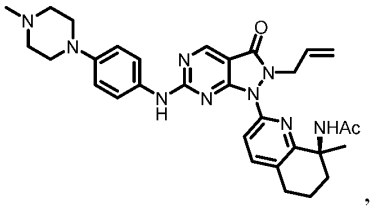
3. The use of Claim 1 or 2, wherein the Compound (A) is selected from the group consisting of:




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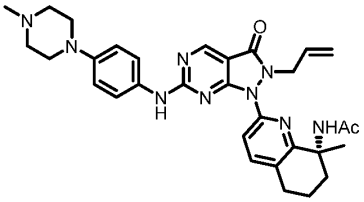
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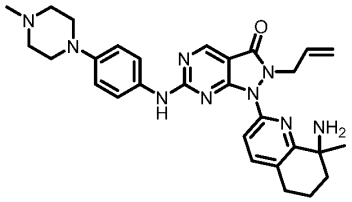

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and



, or a pharmaceutically acceptable salt of any of any of the foregoing.

4. The use of any one of Claims 1-3, wherein the Compound (B) is selected from the group consisting of fulvestrant, (E)-3-[3,5-Difluoro-4-[(1R,3R)-2-(2-fluoro-2-methylpropyl)-3-methyl-1,3,4,9-tetrahydropyrido[3,4-b]indol-1-yl]phenyl]prop-2-enoic acid (AZD9496), (R)-6-(2-(ethyl(4-(2-(ethylamino)ethyl)benzyl)amino)-4-methoxyphenyl)-5,6,7,8-tetrahydronaphthalen-2-ol (elacestrant, RAD1901), (E)-3-(4-((E)-2-(2-chloro-4-fluorophenyl)-1-(1H-indazol-5-yl)but-1-en-1-yl)phenyl)acrylic acid (Brilanestrant, ARN-810, GDC-0810), (E)-3-(4-((2-(2-(1,1-difluoroethyl)-4-fluorophenyl)-6-hydroxybenzo[b]thiophen-3-yl)oxy)phenyl)acrylic acid (LSZ102), (E)-N,N-dimethyl-4-((2-((5-((Z)-4,4,4-trifluoro-1-(3-fluoro-1H-indazol-5-yl)-2-phenylbut-1-en-1-yl)pyridin-2-yl)oxy)ethyl)amino)but-2-enamide (H3B-6545), (E)-3-(4-((2-(4-fluoro-2,6-dimethylbenzoyl)-6-hydroxybenzo[b]thiophen-3-yl)oxy)phenyl)acrylic acid (rintodestrant, G1T48), D-0502, SHR9549, ARV-471, 3-((1R,3R)-1-(2,6-difluoro-4-((1-(3-fluoropropyl)azetidin-3-yl)amino)phenyl)-3-methyl-1,3,4,9-tetrahydro-2H-pyrido[3,4-b]indol-2-yl)-2,2-difluoropropan-1-ol (giredestrant, GDC-9545), (S)-8-(2,4-dichlorophenyl)-9-(4-((1-(3-fluoropropyl)pyrrolidin-3-yl)oxy)phenyl)-6,7-dihydro-5H-benzo[7]annulene-3-carboxylic acid (SAR439859), N-[1-(3-fluoropropyl)azetidin-3-yl]-6-[(6S,8R)-8-methyl-7-(2,2,2-trifluoroethyl)-6,7,8,9-tetrahydro-3H-pyrazolo[4,3-f]isoquinolin-6-yl]pyridin-3-amine (AZD9833), OP-1250 and LY3484356, or a pharmaceutically acceptable salt of any of the foregoing.

5. The use of any one of Claims 1-3, wherein the Compound (B) is selected from the group consisting of tamoxifen, raloxifene, ospemifene, bazedoxifene, toremifene and lasofoxifene, or a pharmaceutically acceptable salt of any of the foregoing.

6. The use of any one of Claims 1-5, wherein the disease or condition is selected from the group consisting of a breast cancer, a cervical cancer, an ovarian cancer, an uterine cancer, a vaginal cancer, a vulvar cancer, a brain cancer, a cervicocerebral cancer, an esophageal cancer, a thyroid cancer, a small cell cancer, a non-small cell cancer, a lung cancer , a stomach cancer, a gallbladder/bile duct cancer, a liver cancer, a pancreatic cancer, a colon cancer, a rectal

cancer, a choriocarcinoma, an uterus body cancer, an uterocervical cancer, a renal pelvis/ureter cancer, a bladder cancer, a prostate cancer, a penis cancer, a testicular cancer, a fetal cancer, a Wilms' cancer, a skin cancer, a malignant melanoma, a neuroblastoma, an osteosarcoma, an Ewing's tumor, a soft part sarcoma, an acute leukemia, a chronic lymphatic leukemia, a chronic myelocytic leukemia, polycythemia vera, a malignant lymphoma, multiple myeloma, a Hodgkin's lymphoma, and a non-Hodgkin's lymphoma.

7. The use of any one of Claims 1-5, wherein the disease or condition is a breast cancer, a cervical cancer, an ovarian cancer, an uterine cancer, a vaginal cancer, and a vulvar cancer.

8. The use of Claim 7, wherein the disease or condition is a breast cancer.

9. The use of any one of Claims 6-8, wherein the breast cancer that does not include any point mutations ER mutations.

10. The use of any one of Claims 6-8, wherein the disease or condition is breast cancer that has at least one point mutation within the Estrogen Receptor 1 (ESR1) that encodes Estrogen receptor alpha (ERα), wherein the mutation is selected from the group consisting of: K303R, D538G, Y537S, E380Q, Y537C, Y537N, A283V, A546D, A546T, A58T, A593D, A65V, C530L, D411H, E279V, E471D, E471V, E523Q, E542G, F461V, F97L, G145D, G160D, G274R, G344D, G420D, G442R, G557R, H524L, K252N, K481N, K531E, L370F, L453F, L466Q, L497R, L536H, L536P, L536Q, L536R, L540Q, L549P, M388L, M396V, M421V, M437I, M522I, N156T, N532K, N69K, P147Q, P222S, P535H, R233G, R477Q, R503W, R555H, S282C, S329Y, S338G, S432L, S463P, S47T, S576L, V392I, V418E, V478L, V533M, V534E, Y537D and Y537H.

11. The use of any one of Claims 6-10, wherein the breast cancer is ER positive breast cancer.

12. The use of any one of Claims 6-10, wherein the breast cancer is ER positive/HER2-negative breast cancer.

13. The use of any one of Claims 6-12, wherein the breast cancer is local breast cancer.

14. The use of any one of Claims 6-12, wherein the breast cancer is metastatic breast cancer.

15. The use of any one of Claims 6-14, wherein the breast cancer is recurrent breast cancer.

16. The use of any one of Claims 6-15, wherein the breast cancer has been previously treated with an endocrine therapy.

17. The use of Claim 16, wherein the treatment was with a selective ER modulator (SERM).

18. The use of Claim 17, wherein the selective ER modulator is selected from the group consisting of tamoxifen, raloxifene, ospemifene, bazedoxifene, toremifene and lasofoxifene, or a pharmaceutically acceptable salt of any of the foregoing.

19. The use of Claim 16, wherein the treatment was with a selective ER degrader (SERD).

20. The use of Claim 19, wherein the selective ER degrader is selected from the group consisting of fulvestrant, (E)-3-[3,5-Difluoro-4-[(1R,3R)-2-(2-fluoro-2-methylpropyl)-3-methyl-1,3,4,9-tetrahydropyrido[3,4-b]indol-1-yl]phenyl]prop-2-enoic acid (AZD9496), (R)-6-(2-(ethyl(4-(2-(ethylamino)ethyl)benzyl)amino)-4-methoxyphenyl)-5,6,7,8-tetrahydronaphthalen-2-ol (elacestrant, RAD1901), (E)-3-(4-((E)-2-(2-chloro-4-fluorophenyl)-1-(1H-indazol-5-yl)but-1-en-1-yl)phenyl)acrylic acid (Brilanestrant, ARN-810, GDC-0810), (E)-3-(4-((2-(2-(1,1-difluoroethyl)-4-fluorophenyl)-6-hydroxybenzo[b]thiophen-3-yl)oxy)phenyl)acrylic acid (LSZ102), (E)-N,N-dimethyl-4-((2-((5-((Z)-4,4,4-trifluoro-1-(3-fluoro-1H-indazol-5-yl)-2-phenylbut-1-en-1-yl)pyridin-2-yl)oxy)ethyl)amino)but-2-enamide (H3B-6545), (E)-3-(4-((2-(4-fluoro-2,6-dimethylbenzoyl)-6-hydroxybenzo[b]thiophen-3-yl)oxy)phenyl)acrylic acid (rintodestrant, G1T48), D-0502, SHR9549, ARV-471, 3-((1R,3R)-1-(2,6-difluoro-4-((1-(3-fluoropropyl)azetidin-3-yl)amino)phenyl)-3-methyl-1,3,4,9-tetrahydro-2H-pyrido[3,4-b]indol-2-yl)-2,2-difluoropropan-1-ol (giredestrant, GDC-9545), (S)-8-(2,4-dichlorophenyl)-9-(4-((1-(3-fluoropropyl)pyrrolidin-3-yl)oxy)phenyl)-6,7-dihydro-5H-benzo[7]annulene-3-carboxylic acid (SAR439859), N-[1-(3-fluoropropyl)azetidin-3-yl]-6-[(6S,8R)-8-methyl-7-(2,2,2-trifluoroethyl)-6,7,8,9-tetrahydro-3H-pyrazolo[4,3-f]isoquinolin-6-yl]pyridin-3-amine (AZD9833), OP-1250 and LY3484356, or a pharmaceutically acceptable salt of any of the foregoing.

21. The use of Claim 16, wherein the treatment was with an aromatase inhibitor.

22. The use of Claim 21, wherein the aromatase inhibitor is a steroidal aromatase inhibitor.

23. The use of Claim 22, wherein the steroidal aromatase inhibitor is selected from the group consisting of exemestane and testolactone, or a pharmaceutically acceptable salt of any of the foregoing.

24. The use of Claim 21, wherein the aromatase inhibitor is a non-steroidal aromatase inhibitor.

25. The use of Claim 24, wherein the non-steroidal aromatase inhibitor is selected from the group consisting of anastazole and letrazole, or a pharmaceutically acceptable salt of any of the foregoing.

26. The use of any one of Claims 6-14, wherein the breast cancer has not been previously treated.