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(54) Title **COMPOSITION FOR TREATING CANCER**

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Enclosed is a translation of the patent claims in Norwegian. Please note that as per the Norwegian Patents Acts, section 66i the patent will receive protection in Norway only as far as there is agreement between the translation and the language of the application/patent granted at the EPO. In matters concerning the validity of the patent, language of the application/patent granted at the EPO will be used as the basis for the decision. The patent documents published by the EPO are available through Espacenet (<http://worldwide.espacenet.com>) or via the search engine on our website here: <https://search.patentstyret.no/>

Patentkrav

1. Kombinasjonen av

- (a) de fire WT1-peptidene YMFPNAPYL (SEQ IDNO:124),
RSDELVRHHNMHQ RNMTKL (SEQ ID NO:1),
PGCNKRYFKLSHLQMHSRKHTG (SEQ ID NO: 2) og
SGQAYMFPNAPYLPSCLES (SEQ ID NO: 125), og
(b) minst én antistoff-PD1-inhibitor,

til anvendelse ved behandling, reduksjon av forekomsten av eller fremkallelse av en immunrespons mot en WT1-uttrykkende kreft.

2. Kombinasjonen til anvendelse ifølge krav 1, hvori WT1-peptidene administreres med en bærer, et hjelpestoff eller en fortynner, eller en adjuvans som QS21, Montanide, Freund's komplett eller ufullstendig adjuvans, aluminiumfosfat, aluminiumhydroksid, BCG, cytokin eller alun.
3. Kombinasjonen til anvendelse ifølge krav 1, hvori PD1-inhibitoren er nivolumab, pembrolizumab, pidilizumab eller MEDI0680 (AMP-514).
4. Kombinasjonen til anvendelse ifølge krav 1, hvori ett eller flere av WT1-peptidene administreres separat.
5. Kombinasjonen til anvendelse ifølge krav 1, hvori to eller flere av WT1-peptidene administreres sammen i den samme formuleringen.
6. Kombinasjonen til anvendelse ifølge krav 1, hvori WT1-peptidene og PD1-inhibitoren er tilstede i den samme sammensetningen.
7. Kombinasjonen til anvendelse ifølge krav 1, hvori WT1-peptidene og PD1-inhibitoren administreres samtidig eller i en overlappende plan, eller hvori den siste administrasjonen av WT1-peptidene er før den første administrasjonen av PD1-inhibitoren.

8. Kombinasjonen til anvendelse ifølge krav 1, hvori kreften er eggstokkreft, brysthinnekreft, leukemi, Wilms' tumor, akutt myelogenleukemi (AML), kronisk myeloid leukemi (CML), myelodysplastisk syndrom (MDS), melanom, magekreft, prostatakreft, gallegangskreft, kreft i urinveiene, glioblastom, bløtvevsarkom, osteosarkom eller ikke-småcellet lungekreft (NSCLC).

9. Kombinasjonen til anvendelse ifølge krav 1, hvori ett eller flere av peptidene NLMNLGATL (SEQ ID NO:21), WNLMNLGATLKGVA (SEQ ID NO:26), LVRHHNMHQ RNMTKL (SEQ ID NO:3), NKRYFKLSHLQMHSR (SEQ ID NO:4), SGQARMFPNAPYLPSCLES (SEQ ID NO:5), QARMFPNAPYLPSC (SEQ ID NO:6), RMFPNAPYL (SEQ ID NO:7), SLGEQQYSV (SEQ ID NO:8), ALLPAVPSL (SEQ ID NO:9), NLGATLKG (SEQ ID NO:10), DLNALLPAV (SEQ ID NO:11), GVFRGIQDV (SEQ ID NO:12), KRYFKLSHL (SEQ ID NO:13), ALLLRTPYS (SEQ ID NO:14), CMTWMQMNL (SEQ ID NO:15), NMHQ RNMTK (SEQ ID NO:16), QMNLGATL (SEQ ID NO:17), FMCAYPGCNK (SEQ ID NO:18), KLSHLQMHSR (SEQ ID NO:19), QAYMFPNAPYLPSC (SEQ ID NO:126), YLGEQQYSV (SEQ ID NO:127), YLLPAVPSL (SEQ ID NO:128), YLGATLKG (SEQ ID NO:129), YLNALLPAV (SEQ ID NO:130), GLRRGIQDV (SEQ ID NO:131), KLYFKLSHL (SEQ ID NO:132), ALLLRTPYV (SEQ ID NO:133), YMTWNQMNL (SEQ ID NO:134), NMYQRNMTK (SEQ ID NO:135), NMHQ RVMTK (SEQ ID NO:136), NMYQ RVMTK (SEQ ID NO: 137), QMYLGATL (SEQ ID NO:138), QMNLGVTLK (SEQ ID NO:139), QMYLGVTLK (SEQ ID NO: 140), FMYAYPGCNK (SEQ ID NO:141), FMCAYPFCNK (SEQ ID NO:142), FMYAYPFCNK (SEQ ID NO:143), KLYHLQMHSR (SEQ ID NO:144), KLSHLQMHSK (SEQ ID NO:145), KLYHLQMHSK (SEQ ID NO:146), NQMNLGATL (SEQ ID NO:20), NYMNLGATL (SEQ ID NO:22), CMTWNQMNLGATLKG (SEQ ID NO:23), CMTWNLMNLGATLKG (SEQ ID NO:24), WNQMNLGATLKGVA (SEQ ID NO:25), MTWNQMNLGATLKG (SEQ ID NO:27), TWNQMNLGATLKGVA (SEQ ID NO:28), MTWNLMNLGATLKG (SEQ ID NO:30), TWNLMNLGATLKGVA (SEQ ID NO:31), MTWNYMNLGATLKG (SEQ ID NO:33), TWNYMNLGATLKGVA (SEQ ID NO:34), CMTWNQMNLGATLKGVA (SEQ ID NO:35), WNQMNLGAT (SEQ ID NO:36),

TWNQMN LGA (SEQ ID NO:37), MTWNQMNLG (SEQ ID NO:38),
 CMTWNLMNLGATLKGVA (SEQ ID NO:39), WNLMLNLGAT (SEQ ID NO:40),
 MNLGATLKG (SEQ ID NO:41), CMTWNYMNLGATLKGVA (SEQ ID NO:43),
 GALRNPTAC (SEQ ID NO:46), GYLRNPTAC (SEQ ID NO:47), GALRNPTAL
 (SEQ ID NO:48), YALRNPTAC (SEQ ID NO:49), GLLRNPTAC (SEQ ID NO:50),
 RQRPHPGAL (SEQ ID NO:51), RYRPHPGAL (SEQ ID NO:52), YQRPHPGAL
 (SEQ ID NO:53), RLRPHPGAL (SEQ ID NO:54), RIRPHPGAL (SEQ ID NO:55),
 QFPNHSFKHEDPMGQ (SEQ ID NO:61), HSFKHEDPM (SEQ ID NO:63),
 HSFKHEDPY (SEQ ID NO:64), HSFKHEDPK (SEQ ID NO:65),
 KRPFMCAYPGCYKRY (SEQ ID NO:66), SEKRPFMCAYPGCNK (SEQ ID
 NO:67), KRPFMCAYPGCNK (SEQ ID NO:68), FMCAYPGCN (SEQ ID NO:69),
 FMCAYPGCY (SEQ ID NO:70), and FMCAYPGCK (SEQ ID NO:71) også
 administreres.

10. Kombinasjonen til anvendelse ifølge krav 11, hvori 200 mcg av hvert peptid emulgeres med Montanide ISA 51 VG og administreres subkutan i uke 0, 2, 4, 6, 8 og 10.
11. Kombinasjonen til anvendelse ifølge krav 10, hvori 3 mg/kg nivolumab administreres intravenøst i uke 0, 2, 4, 6, 8, 10 og 12.
12. Kombinasjonen til anvendelse ifølge krav 1, hvori behandlingen, reduksjonen i forekomsten av eller fremkallelse av en immunrespons mot en WT1-uttrykkende kreft er større enn det som oppnås ved å administrere WT1-peptidene for seg selv eller den minst ene PD1-inhibitoren for seg selv.
13. Sammensetning omfattende de fire WT1-peptidene YMFPNAPYL (SEQ ID NO:124), RSDLVRRHHNMHQRNMTKL (SEQ ID NO:1), PGCNKRYFKLSHLQMHSRKHTG (SEQ ID NO: 2) og SGQAYMFPNAPYLPSCLES (SEQ ID NO:125), og minst én antistoff-PD1-inhibitor.
14. Sammensetningen ifølge krav 13, hvori PD1-inhibitoren er nivolumab, pembrolizumab, pidilizumab eller MEDI0680 (AMP-514).