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(54)	Title	<b>ANTI-PD-L1 ANTIBODIES AND THEIR USE TO ENHANCE T-CELL FUNCTION</b>
(56)	References Cited:	EP-A1- 1 537 878, WO-A2-2006/042237, WO-A2-2007/005874, WO-A2-2008/071447 WO-A2-2008/085562, US-A1- 2003 039 653, HIRANO FUMIYA ET AL: "Blockade of B7-H1 and PD-1 by monoclonal antibodies potentiates cancer therapeutic immunity", CANCER RESEARCH, AMERICAN ASSOCIATION FOR CANCER REREARCH, US, vol. 65, no. 3, 1 February 2005 (2005-02-01), pages 1089 - 1096, XP002419626, ISSN: 0008-5472, DONG HAIDONG ET AL: "Tumor-associated B7-H1 promotes T-cell apoptosis: a potential mechanism of immune evasion", NATURE MEDICINE, NATURE PUBLISHING GROUP, NEW YORK, NY, US, vol. 8, no. 8, 24 June 2002 (2002-06-24), pages 793 - 800, XP002397368, ISSN: 1078-8956, BARBER DANIEL L ET AL: "Restoring function in exhausted CD8 T cells during chronic viral infection", NATURE, NATURE PUBLISHING GROUP, LONDON, GB LNKD- DOI:10.1038/NATURE04444, vol. 439, no. 7077, 9 February 2006 (2006-02-09), pages 682 - 687, XP002419629, ISSN: 0028-0836 LATCHMAN YVETTE ET AL: "PD-L2 is a second ligand for PD-1 and inhibits T cell activation", NATURE IMMUNOLOGY, NATURE PUBLISHING GROUP, GB LNKD- DOI:10.1038/85330, vol. 2, no. 3, 1 March 2001 (2001-03-01), pages 261 - 268, XP001064842, ISSN: 1529-2908

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.** Et isolert, blokkerende anti-PD-L1-antistoff full-lengde som omfatter en tung kjede og en lett kjede variabel region sekvens, hvori:

(a) den tunge kjeden omfatter en HVR-H1, HVR-H2 og HVR-H3, hvori videre:

- (i) HVR-H1-sekvensen er GFTFSX<sub>1</sub>SWIH (SEKV ID NR:1);
- (ii) HVR-H2-sekvensen er AWIX<sub>2</sub>PYGGSX<sub>3</sub>YYADSVKG (SEKV ID NR:2);
- (iii) HVR-H3-sekvensen er RHWPGGF DY, og (SEKV ID NR:3);

(b) den lette kjeden omfatter en HVR-L1, HVR-L2 og HVR-L3, hvori videre:

- (iv) HVR-L1-sekvensen er RASQX<sub>4</sub>X<sub>5</sub>X<sub>6</sub>TX<sub>7</sub>X<sub>8</sub>A (SEKV ID NR:8);
- (v) HVR-L2-sekvensen er SASX<sub>9</sub>LX<sub>10</sub>S (SEKV ID NR:9);
- (vi) HVR-L3-sekvensen er QQX<sub>11</sub>X<sub>12</sub>X<sub>13</sub>X<sub>14</sub>PX<sub>15</sub>T (SEKV ID NR:10);

hvor X<sub>1</sub> = D, X<sub>2</sub> = S og X<sub>3</sub> = T, X<sub>4</sub> = D, X<sub>5</sub> = V, X<sub>6</sub> = S, X<sub>7</sub> = A og X<sub>8</sub> = V, X<sub>9</sub> = F og X<sub>10</sub> = Y, X<sub>11</sub> = Y, X<sub>12</sub> = L, X<sub>13</sub> = Y, X<sub>14</sub> = H og X<sub>15</sub> = A, og videre omfattende:

(c) variabel region tung kjede rammeverksekvenser sidestilt mellom HVR-ene i henhold til formelen: (HC-FR1)-(HVR-H1)-(HC-FR2)-(HVR-H2)-(HC-FR3)-(HVR-H3)-(HC-FR4), og

(d) variabel region lett kjede rammeverksekvenser sidestilt mellom HVR-ene i henhold til formelen: (LC-FR1)-(HVR-L1)-(LC-FR2)-(HVR-L2)-(LC-FR3)-(HVR-L3)-(LC-FR4),

hvor rammeverksekvensene til (c) og (d) er avledet fra humane konsensus-rammeverksekvenser, hvori konsensus-VH-rammeverksekvensen er en Kabat-undergruppe III konsensus-rammeverksekvens og konsensus-VL-rammeverksekvensen er en Kabat kappa I-konsensus-rammeverksekvens.

**2.** En sammensetning omfattende det isolerte, blokkerende anti-PD-L1-antistoffet ifølge krav 1 og minst én farmasøytisk akseptabel bærer.