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(54) Title                   **IMMUNE CHECKPOINT INHIBITOR COMBINATIONS**

(56) References  
Cited: Berge et al: "Abstract 477: Long-term protection against B16F1 melanoma upon vaccination with tumor cell lysate combined with LTX-315 as a novel adjuvant.", Cancer Research, vol. 73 15 April 2013 (2013-04-15), page 477, XP002753069, Retrieved from the Internet:  
URL:[http://cancerres.aacrjournals.org/cont ent/73/8\\_Supplement/477](http://cancerres.aacrjournals.org/cont ent/73/8_Supplement/477) [retrieved on 2016-01-14]  
RIBAS: N ENGL J MED, 2012,

SCIENCE, vol. 348, 2015, pages 56-61,  
Phase I/ii Clinical Trial: C12-315-03 T-cell infiltration across different indications  
P. A. OTT ET AL: "CTLA-4 and PD-1/PD-L1 Blockade: New Immunotherapeutic Modalities with Durable Clinical Benefit in Melanoma Patients", CLINICAL CANCER RESEARCH, vol. 19, no. 19, 1 October 2013 (2013-10-01), pages 5300-5309, XP55200528, ISSN: 1078-0432, DOI: 10.1158/1078-0432.CCR-13-0143

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. Forbindelse med formelen Lys Lys Trp Trp Lys Lys Trp Dip Lys, eller et salt, ester  
eller amid derav;

for anvendelse i behandlingen av en tumor ved koadministrering med et

5 immunterapeutisk middel, hvor det immunoterapeutiske middelet er et anti-CTLA-4-  
antistoff.

2. Forbindelse for anvendelse ifølge krav 1, hvor anti-CTLA-4-antistoffet er valgt fra  
gruppen bestående av ipilimumab og tremelimumab.

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3. Forbindelse for anvendelse ifølge krav 1 eller krav 2, hvor det immunoterapeutiske  
middelet administreres før eller samtidig med forbindelsen.

4. Forbindelse for anvendelse ifølge krav 3, hvor det immunoterapeutiske middelet  
15 administreres før forbindelsen.

5. Produkt inneholdende:

- (i) en forbindelse som definert i krav 1; og
- (ii) et immunterapeutisk middel, hvor det immunterapeutiske middelet er et  
20 anti-CTLA-4-antistoff, som et kombinert preparat for separat, samtidig eller  
sekvensiell anvendelse i behandling av tumorer.

6. Produkt for anvendelse ifølge krav 5, hvor antistoffet er valgt fra gruppen  
bestående av ipilimumab og tremelimumab.

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