

Title (en)
PHARMACEUTICAL COMBINATIONS OF HISTONE DEACETYLASE INHIBITOR AND PROTEASOME INHIBITOR OR IMMUNOMODULATORY DRUG FOR THE TREATMENT OF HEMATOLOGICAL CANCER

Title (de)
PHARMAZEUTISCHE KOMBINATIONEN AUS EINEM HISTON-DEACETYLASE-HEMMER UND PROTEASOM-HEMMER ODER IMMUNOMODULATOR ZUR BEHANDLUNG VON BLUTKREBSEKANKUNGEN

Title (fr)
COMBINAISONS PHARMACEUTIQUES D'INHIBITEUR DE L'HISTONE DÉACÉTYLASE ET D'INHIBITEUR DU PROTÉASOME OU DE MÉDICAMENT IMMUNOMODULATEUR POUR LE TRAITEMENT D'UN CANCER HÉMATOLOGIQUE

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• KR 2017011015 W 20170929

Abstract (en)
[origin: WO2018066946A1] The present invention relates to a pharmaceutical combination for treating a hematological cancer comprising the histone deacetylase (HDAC) inhibitor of the chemical formula 1, a proteasome inhibitor or an immunomodulatory drug and a steroidal anti-cancer agent together. The pharmaceutical combination of the present invention can be useful for treating a hematological cancer such as multiple myeloma, by reducing toxicity which is the problem of the conventional HDAC inhibitor and exhibiting an equivalent level of pharmaceutical effects due to a complex inhibitory mechanism against the cancer of the compound of chemical formula 1 and its pharmaceutically acceptable salt, the proteasome inhibitor or the immunomodulatory drug and the steroidal anti-cancer agent.

IPC 8 full level
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Citation (search report)
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• [T] ANONYMOUS: "A Phase 1, Open-Label, Multi-Center Study of Alteminostat (CKD-581) in Combination with Lenalidomide and Dexamethasone in Patients with Previously Treated Multiple Myeloma (MM) | Blood | American Society of Hematology", 13 November 2019 (2019-11-13), XP055674271, Retrieved from the Internet <URL:https://ashpublications.org/blood/article/134/Supplement_1/1847/427755/A-Phase-1-OpenLabel-MultiCenter-Study-of> [retrieved on 20200306]
• See references of WO 2018066946A1

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