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(54)	Title	COMPOSITIONS COMPRISING CEFEPIME AND TAZOBACTAM
(56)	References Cited:	WO-A1-2007/086011, WO-A1-2007/086014, US-A1- 2010 197 650, A. ENDIMIANI ET AL: "Characterization of blaKPC-containing Klebsiella pneumoniae isolates detected in different institutions in the Eastern USA", JOURNAL OF ANTIMICROBIAL CHEMOTHERAPY, vol. 63, no. 3, 15 January 2009 (2009-01-15), pages 427-437, XP55017325, ISSN: 0305-7453, DOI: 10.1093/jac/dkn547, , 21 December 2017 (2017-12-21), Retrieved from the Internet: URL: https://www.merriam-webster.com/dictionary/composition [retrieved on 2017-12-21], MACGOWAN ALASDAIR; ROGERS CHRIS; BOWKER KAREN: "In vitro models, in vivo models, and pharmacokinetics: What can we learn from in vitro models?", CLINICAL INFECTIOUS DISEASES, vol. 33, no. Suppl. 3, 15 September 2001 (2001-09-15), pages S214-

S220,, JONES R N ET AL: "Antimicrobial spectrum of cefpirome combined with tazobactam against the *Bacteroides fragilis* group", DIAGNOSTIC MICROBIOLOGY AND INFECTIOUS DISEASES, ELSEVIER SCIENCE PUBLISHING CO., AMSTERDAM, NL, vol. 13, no. 5, 1 September 1990 (1990-09-01), pages 371-373, XP023790076, ISSN: 0732-8893, DOI: 10.1016/0732-8893(90)90004-F [retrieved on 1990-09-01], US-A1- 2010 029 604, VANSKOY BRIAN D ET AL.: "Pharmacokinetics-Pharmacodynamics of Tazobactam in Combination with Cefepime in an In Vitro Infection Model", ANTIMICROBIAL AGENTS AND CHEMOTHERAPY, vol. 61, no. 12, 1 December 2017 (2017-12-01), CN-A- 1 565 456, CN-A- 1 565 455

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. Farmasøytisk sammensetning omfattende (a) cefepim eller et farmasøytisk akseptabelt salt derav; og (b) tazobaktam eller et farmasøytisk akseptabelt salt

5 der sammensetningen er **karakterisert ved at** den omfatter 2 g cefepim og 2 g tazobaktam.

2. Den farmasøytiske sammensetningen ifølge krav 1, for anvendelse i behandling eller kontroll av en bakteriell infeksjon hos et individ.

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3. Den farmasøytiske sammensetningen ifølge krav 1, omfattende et farmasøytisk akseptabelt salt av tazobaktam, som er til stede som tazobaktamnatrium.

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4. Den farmasøytiske sammensetningen ifølge krav 1, hvor i sammensetningen er et pulver eller en oppløsning.

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5. Den farmasøytiske sammensetningen ifølge krav 1, der sammensetningen omfatter et farmasøytisk akseptabelt salt av cefepim, som er til stede som cefepimhydroklorid, og et farmasøytisk akseptabelt salt av tazobaktam, som er til stede som tazobaktamnatrium.