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(54)	Title	CRYSTALLINE FORM OF 4-[5-(PYRIDINE-4-YL)-1H-1,2,4-TRIAZOLE-3-YL]PYRIDINE-2-CARBONITRILE
(56)	References Cited:	EP-A1- 1 471 065, JP-A- 2003 519 698, WO-A1-2005/009991, EP-A1- 1 650 204, WO-A1-03/064410, JP-A- 2005 041 802 CAIRA M R: "CRYSTALLINE POLYMORPHISM OF ORGANIC COMPOUNDS", TOPICS IN CURRENT CHEMISTRY, SPRINGER, BERLIN, DE, vol. 198, 1 January 1998 (1998-01-01), pages 163-208, XP001156954, ISSN: 0340-1022, DOI: 10.1007/3-540-69178-2_5

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

- 5 **1.** Type I-krystaller til 4-[5-(pyridin-4-yl)-1H-1,2,4-triazol-3-yl]pyridin-2-karbonitril **karakterisert ved** et røtgendpulverdiffraksjoneringspektrum som vist i figur 1 og utvisende karakteristiske topper i røtgendpulverdiffraksjon ved diffraksjonsvinkler 2θ på omtrent $10,1^\circ$, $16,0^\circ$, $20,4^\circ$, $25,7^\circ$ og $26,7^\circ$, hvori røtgendpulverdiffraksjoneringspektret refererer til et spektrum målt ved hjelp av Mini Flex, produkt av Rigaku Corporation, ved å anvende Cu som røntgenkilde.
- 10 **2.** Fremgangsmåte for fremstilling av type I-krystaller til 4-[5-(pyridin-4-yl)-1H-1,2,4-triazol-3-yl]pyridin-2-karbonitril som angitt i krav 1, der fremgangsmåten omfatter behandling av et syresalt til 4-[5-(pyridin-4-yl)-1H-1,2,4-triazol-3-yl]pyridin-2-karbonitril med en base og etterfølgende nøytralisering av det behandlede produktet med en syre.
- 15 **3.** Farmasøytisk sammensetning omfattende type I-krystaller som angitt i krav 1, og en farmasøytisk akseptabel bærer.