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(54)	Title	PARTICULATE COMPOSITIONS FOR DELIVERY OF POORLY SOLUBLE DRUGS
(56)	References Cited:	EP-A- 0 012 523 EP-A- 0 605 024 EP-A- 0 631 781 EP-A- 1 752 142 EP-A2- 0 205 282 WO-A-01/15699 WO-A-95/27482 WO-A-2007/008480 WO-A1-2005/074890 WO-A1-2007/003278 CA-A1- 987 588 GB-A- 1 456 618 RU-C2- 2 228 180 RU-C2- 2 241 458 RU-C2- 2 253 452 US-A- 5 145 684 US-A- 5 948 787 US-A1- 2001 007 665 US-A1- 2003 129 244 US-A1- 2004 110 781 US-A1- 2005 048 115 US-A1- 2007 020 339

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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. Sammensetning omfattende partikler som har en gjennomsnittlig diameter på mindre enn 2 mm, hvor
5 partiklene omfatter buprenorfin eller buprenorfin HCl tilstede i en mengde som varierer fra omtrent 0,2% til omtrent 4 masse% av partiklene, og polyetylglykol (PEG),
hvor buprenorfin eller buprenorfin HCl er innkapslet av PEG.
- 10 2. Sammensetning ifølge krav 1, hvor PEG har et gjennomsnittlig molekylvektområde fra omtrent 500 Dalton til omtrent 10.000 Dalton eller hvor PEG er PEG 3350.
- 15 3. Sammensetning ifølge krav 1, som videre omfatter et overflateaktivt middel.
4. Farmasøytisk sammensetning omfattende sammensetningen ifølge krav 1 og nalokson.
- 20 5. Farmasøytisk sammensetning ifølge krav 4, hvor sammensetningen er formulert for oral administrering.