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(54)	Title	LIQUID PHARMACEUTICAL COMPOSITION
(56)	References Cited:	WO-A1-2013/164837 WO-A2-2009/073569 WO-A2-2011/104381 US-A1- 2004 033 228 US-A1- 2010 137 213 US-A1- 2010 278 822 ZHENG J Y ET AL: "Influence of pH, buffer species, and storage temperature on physicochemical stability of a humanized monoclonal antibody LA298", INTERNATIONAL JOURNAL OF PHARMACEUTICS, ELSEVIER, NL, vol. 308, no. 1-2, 3 February 2006 (2006-02-03), pages 46-51, XP027972782, ISSN: 0378-5173 [retrieved on 2006-02-03]

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. En flytende farmasøytisk sammensetning bestående av:

- 5 - 50 mg/ml adalimumab;
- et sitratbuffersystem;
- en sukkerstabilisator;
- en tonisitetsmiddel;
- et overflateaktivt middel; og
- 10 - vann (for injeksjon);
- hvor nevnte adalimumab, sitratbuffersystem, sukkerstabilisator, tonisitetsmiddel og overflateaktivt middel er til stede i et molforhold på 1:14-40:288-865:28-576:0,1-3,2.