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(54)	Title	ANTIGEN-BINDING MOLECULE CAPABLE OF BINDING TO TWO OR MORE ANTIGEN MOLECULES REPEATEDLY
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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. Fremgangsmåte for å fremstille en farmasøytisk sammensetning omfattende antistoffet som isoleres ved screening-fremgangsmåten, omfattende trinnene:

- 5 (a) å bestemme den antigenbindende aktiviteten til et antistoff ved pH 6,7 til pH 10,0;
(b) å bestemme den antigenbindende aktiviteten til antistoffet ved pH 4,0 til pH 6,5; og
(c) å velge antistoffet hvis antigenbindende aktivitet ved pH 6,7 til pH 10,0 er
10 høyere enn den antigenbindende aktiviteten ved pH 4,0 til pH 6,5;
og å formulere antistoffet valgt i trinn (c) til en farmasøytisk sammensetning.

2. Fremgangsmåte for å fremstille en farmasøytisk sammensetning omfattende antistoffet som isoleres ved screening-fremgangsmåten, omfattende trinnene:

- 15 (a) å binde et antistoff til et antigen under en betingelse ved en fra pH 6,7 til pH 10,0;
(b) å plassere antistoffet bundet til antigenet i (a) under en betingelse ved en fra pH 4,0 til pH 6,5; og
(c) å oppnå antistoffet som dissosierer under betingelsen ved en fra pH 4,0 til
20 pH 6,5;
og å formulere antistoffet oppnådd i trinn (c) til en farmasøytisk sammensetning.

3. Fremgangsmåte for å fremstille en farmasøytisk sammensetning omfattende antistoffet som isoleres ved screening-fremgangsmåten, omfattende trinnene:

- 25 (a) å binde et antistoff til en antigenimmobilisert søyle under en betingelse ved en fra pH 6,7 til pH 10,0;
(b) å eluere antistoffet bundet til søylen under betingelse ved en fra pH 6,7 til pH 10,0 fra kolonnen under en betingelse ved en fra pH 4,0 til pH 6,5; og
(c) å samle inn det eluerte antistoffet;
30 og å formulere antistoffet samlet opp i trinn (c) til en farmasøytisk sammensetning.