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(54)	Title	EXON SKIPPING COMPOSITIONS FOR TREATING MUSCULAR DYSTROPHY
(56)	References Cited:	WO-A1-2010/048586, US-A1- 2010 168 212, WO-A1-2012/029986, JERRY R. MENDELL ET AL: "Gene therapy for muscular dystrophy: Lessons learned and path forward", NEUROSCIENCE LETTERS, vol. 527, no. 2, 17 May 2012 (2012-05-17), pages 90-99, XP055128775, ISSN: 0304-3940, DOI: 10.1016/j.neulet.2012.04.078 VAN VLIET LAURA ET AL: "Assessment of the feasibility of exon 45-55 multiexon skipping for duchenne muscular dystrophy", BMC MEDICAL GENETICS, BIOMED CENTRAL, LONDON, GB, vol. 9, no. 1, 1 December 2008 (2008-12-01), page 105, XP021042784, ISSN: 1471-2350, DOI: 10.1186/1471-2350-9-105 FOSTER H ET AL: "Genetic therapeutic approaches for Duchenne muscular dystrophy", HUMAN GENE THERAPY, MARY ANN LIEBERT, NEW YORK ,NY, US, vol. 23, no. 7, 30 May 2012 (2012-05-30), pages 676-687, XP002688790, ISSN: 1043-0342, DOI: 10.1089/HUM.2012.099 [retrieved on 2012-05-30] MOULTON H M ET AL: "Morpholinos and their peptide conjugates: Therapeutic promise and challenge for Duchenne muscular dystrophy", BIOCHIMICA ET BIOPHYSICA ACTA (BBA) - BIOMEMBRANES, ELSEVIER, AMSTERDAM, NL, vol. 1798, no. 12, 17 February 2010 (2010-

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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/>

1. Antisenseoligonukleotid som har følgende struktur:



$A =$ 
 $C =$ 
 $G =$ 
 $T = m^5U =$ 

hvor $\wedge$ = stereokjemien til fosforsenteret ikke er definert; eller et farmasøytisk

2. Farmasøytisk sammensetning som omfatter antisenseoligonukleotidet ifølge krav 1 og en farmasøytisk akseptabel bærer.

15 **3.** Sammensetningen ifølge krav 2 for anvendelse ved behandling av Duchenne-muskeldystrofi.