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(54) Title      **COMPOUNDS AND COMPOSITIONS FOR USE IN A METHOD OF TREATMENT OF PARAINFLUENZA VIRUS**  
(56) References Cited:      US-A1- 2004 081 624, US-A1- 2011 171 132, US-A1- 2007 190 163  
                                CHIEN, MD, MS, JW ET AL.: 'Chapter 22: Influenza And Other Viral Respiratory Tract Infections.' EXPERT GUIDE TO INFECTIOUS DISEASES. 2008, page 424, XP055161994  
                                BROEDERS, ME ET AL.: 'Inhalation Profiles In Asthmatics And COPD Patients: Reproducibility And Effect Of Instruction.' J AEROSOL MED. SUMMER vol. 16, no. 2, 2003, pages 131 - 141, XP055080980  
                                G. B. TRIANA-BALTZER ET AL: "DAS181, a sialidase fusion protein, protects human airway epithelium against influenza virus infection: an in vitro pharmacodynamic analysis", JOURNAL OF ANTIMICROBIAL CHEMOTHERAPY, vol. 65, no. 2, 26 November 2009 (2009-11-26), pages 275-284, XP055206288, ISSN: 0305-7453, DOI: 10.1093/jac/dkp421  
                                MATTHEWS ABIGAIL A. ET AL: "Developing inhaled protein therapeutics for lung diseases", MOLECULAR BIOMEDICINE, vol. 1, no. 1, 1 December 2020 (2020-12-01), XP055893710, DOI: 10.1186/s43556-020-00014-z Retrieved from the Internet:  
                                URL:<https://link.springer.com/content/pdf/10.1186/s43556-020-00014-z.pdf>  
                                ANNE MOSCONA ET AL: "A Recombinant Sialidase Fusion Protein Effectively Inhibits Human Parainfluenza Viral Infection In Vitro and In Vivo", THE JOURNAL OF INFECTIOUS DISEASES, vol. 202, no. 2, 15 July 2010 (2010-07-15) , pages 234-241, XP055206285, ISSN: 0022-1899, DOI: 10.1086/653621

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 en terapeutisk effektiv mengde av et protein omfattende SEQ ID NO: 1 eller SEQ ID NO: 2 for anvendelse i en fremgangsmåte for behandling av parainfluensa- eller influensavirusinfeksjon, fremgangsmåten omfattende å administrere sammensetningen til luftveiene til en pasient, hvori sammensetningen er i flytende form og administreringen er med forstøver, hvori 0,1 – 10 mg av proteinet administreres til pasienten per dag.
2. Sammensetningen for anvendelse ifølge krav 1, hvori pasienten er en immunsvekket pasient.
3. Sammensetningen for anvendelse ifølge krav 2, hvori den immunsvekkede pasienten lider av en primær immunsvikt.
4. Sammensetningen for anvendelse ifølge krav 2, hvori den immunsvekkede pasienten lider av en sekundær immunsvikt.
5. Sammensetningen for anvendelse ifølge krav 2, hvori den immunsvekkede pasienten blir eller har blitt behandlet med en immunsuppressiv terapi.
6. Sammensetningen for anvendelse ifølge krav 2, hvori den immunsvekkede pasienten blir eller har blitt behandlet med et kjemoterapeutisk middel.