

1 Abstract

2 **DABCO-functionalized nanoemulsions with antimicrobial**  
3 **properties for potential treatment of ocular myasthenia gravis**4 **Carlotta Coccolini<sup>1</sup>, Elisa Berselli<sup>1</sup>, Faezeh Fathi<sup>2</sup>, M. Beatriz P. P. Oliveira<sup>2</sup>, Patrick Masson<sup>3</sup>, Andrey V. Bogdanov<sup>4</sup>,**  
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13 Sciences, Kazan, 420088, Russian Federation; abogdanov@inbox.ru; tatyana\_pashirova@mail.ru;14 <sup>5</sup> Polythecnic Institut of Guarda, Rua da Cadeia, 6300-035 Guarda, Portugal.15 <sup>6</sup> UCIBIO, Department of Drug Sciences, Faculty of Pharmacy, University of Porto, Rua de Jorge Viterbo Fer-  
16 reira, 228, 4050-313 Porto, Portugal17 \* Correspondence: [karolline@ipg.pt](mailto:karolline@ipg.pt); [esouto@ff.up.pt](mailto:esouto@ff.up.pt)18 **Abstract:** Ocular myasthenia gravis (OMG) is an autoimmune disease in which Ab is produced  
19 against proteins at the neuromuscular junction in the ocular district, causing inability to contract  
20 extraocular and eyelid muscles and thus leading to muscle weakness, diplopia, ptosis, and therefore  
21 difficulty in vision. In cases where treatment with Acetylcholinesterase inhibitors fails, oral cortico-  
22 steroids are used. One way to avoid the side effects of systemic administration of these drugs is their  
23 local administration. However, by topical administration, the percentage of drug absorbed in the  
24 eye is less than 5%. The use of oil-in-water nanoemulsions (NEs) to deliver corticosteroids increases  
25 their bioavailability and improves their absorption. The use of DABCO as a cationic surfactant for  
26 the formulation of the NEs allows a controlled drug release over time, through electrostatic interac-  
27 tion with the negatively charged mucins in the tears. DABCO's antibacterial properties also allow it  
28 to act as a preservative, making it possible to avoid the use of preservatives in the formulation,  
29 which are often responsible for allergic reactions. In this work, DABCO S2-NEs were produced and  
30 characterised, leading to the definition of a delivery system akin to ocular delivery, supporting the  
31 hypothesis of their use in the treatment of OMG. It is also possible to consider functionalizing NEs  
32 with monoclonal antibodies (one of the latest treatments in the cure of the disease) to achieve a  
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