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Botulism, caused by exposure to one or more of the eight serotypes of botulinum neurotoxins (BoNTs) (BoNT/A through H), is often fatal without rapid treatment. Botulism occurs sporadically all over the world, due to environmental exposure, contaminated food, wound infection, or overdose from genuine or counterfeit BoNT-based drug products.

In the past few decades, cosmetic and medical uses for an ever-expanding list of disease indications, from depression to spasticity [1], for BOTOX® [2] and other [3] approved BoNT-based drugs, such as Myobloc®, Xeomin®, and others, has resulted in an explosive growth in both the availability of BoNT and number of persons treated. Furthermore, intentional misuse of BoNTs by bioterrorists [4] or rogue nations [5] remains a risk. With increasing BoNT production, and use and mis-use, comes the need for treatment and prevention of BoNT intoxication.

In this special issue of “Botulinum Neurotoxins (BoNTs)—Antibody and Vaccine”, two review articles and ten original research papers on medical countermeasures for botulism, and therapeutic use of a new BoNT/A-based drug, are reported. Briefly, Grace Sundeen and Joseph T. Barbieri [6] reviewed the current status of DNA-based, viral vector-based, and recombinant protein-based vaccines against botulism. A vaccine is needed since the pentavalent toxoid vaccine is no longer available. Christine Rasett-Escargueil, Arnaud Avril, and their colleagues [7] reviewed the status of the EU Framework program for developing humanized AntibotABE antibodies. In this issue, we, along with our collaborators at UCSF, report engineering of a tri-epitopic antibody that recapitulates the neutralizing activity of a combination of three antibodies to BoNT/A, potentially offering a simpler route to an antibody-based therapeutic [8]. Along with Consuelo Garcia-Rodriguez, we also report development of a three-antibody combination that potently neutralizes BoNT/E [9], analogous to that reported for BoNT/A and/F. Yagmur Derman, Katja Selby, and their collaborators across the EU report on a humanized scFv-FC fusion, derived from immunized macaques and its potential in the neutralization of BoNT/E [10]. Osnat Rosen, Amram Torgeman, Ran Zichel, and their team from Israel, reported their development of an in vitro potency assay for selected anti-BoNT antibodies [11] and their research on the role of homologous Fc fragment in the potency and efficacy of anti-BoNT antibody [12]. Fetweh H. Al-Saleem, Rashmi Sharma, Scott K. Dessain, and their co-workers from the United States of America, presented the development of a fusion protein for red blood cell adherence of immune complexes containing BoNT, to improve neutralization and macrophage uptake [13]. Nicola Bak, Dorothea Sesardic, and their team from the United Kingdom, reported on the application of SiMa cells for cell-based neutralization test for BoNT/A and BoNT/E [14]. On the vaccine development side, Robert P. Webb, Theresa J. Smith, and their team from USAMRIID, reported on the production of non-toxic holoproteins and recombinant BoNT toxin domain subunits as vaccine candidates against multiple serotypes [15], Denis Y. Otaka, Felipe M. Salvatani, and their team from Brazil, reported on the humoral response of buffalos to a recombinant vaccine against BoNT/C and D [16].
Hyun Jung Chang, Bo Young Hong, Jeong-Yi Kwon, and their teams from South Korea, report results from a clinical trial on the efficacy and safety of Botulax for the treatment of dynamic equinus foot deformity in children with cerebral palsy [17].

We hope the papers presented here will provide new insight into this intriguing protein, and prevention or treatment of botulism. We appreciate the contributions of BoNT researchers from around the world to this special issue.

Conflicts of Interest: The authors declare no conflicts of interest.

References
1. Sifferlin, A. Botox: The Drug That’s Treating Everything. Time, 4 January 2017. Available online: http://time.com/4623409/botox-drug-treating-everything/ (accessed on 24 November 2018).
2. Brin, M.F. Basic and clinical aspects of BOTOX®. Toxicon 2009, 54, 676–682. [CrossRef] [PubMed]
3. Jankovic, J. Disease-oriented approach to botulinum toxin use. Toxicon 2009, 54, 614–623. [CrossRef] [PubMed]
4. Broad, W.J. SOWING DEATH: A Special Report; How Japan Germ Terror Alerted World. The New York Times, 26 May 1998. Available online: http://www.nytimes.com/1998/05/26/world/sowing-death-a-special-report-how-japan-germ-terror-alerted-world.html (accessed on 24 November 2018).
5. Arnon, S.A.; Schecter, R.; Inglesby, T.V.; Henderson, D.A.; Bartlett, J.G.; Ascher, M.S.; Etizen, E.; Fine, A.D.; Hauer, J.; Layton, M.; et al. Botulinum toxin as a biological weapon. JAMA 2001, 285, 1059–1070. [CrossRef] [PubMed]
6. Sundeen, G.; Barbieri, J.T. Vaccines against Botulism. Toxins 2017, 9, 268. [CrossRef] [PubMed]
7. Rasetti-Escargueil, C.; Avril, A.; Miethe, S.; Mazuet, C.; Derman, Y.; Selby, K.; Thuiller, P.; Pelat, T.; Urbain, R.; Fontayne, A.; et al. The European AntibotABE Framework Program and Its Update: Development of Innovative Botulinum Antibodies. Toxins 2017, 9, 309. [CrossRef] [PubMed]
8. Lou, J.; Wen, W.; Conrad, F.; Meng, Q.; Dong, J.; Sun, Z.; Garcia-Rodriguez, C.; Farr-Jones, S.; Cheng, L.W.; Henderson, T.D.; et al. A Single Tri-Epitopic Antibody Virtually Recapitulates the Potency of a Combination of Three Monoclonal Antibodies in Neutralization of Botulinum Neurotoxin Serotype A. Toxins 2018, 10, 84. [CrossRef] [PubMed]
9. Garcia-Rodriguez, C.; Razai, A.; Geren, I.N.; Lou, J.; Conrad, F.; Wen, W.H.; Farr-Jones, S.; Smith, T.J.; Brown, J.L.; Skerry, J.C.; et al. A Three Monoclonal Antibody Combination Potently Neutralizes Multiple Botulinum Neurotoxin Serotype E Subtypes. Toxins 2018, 10, 105. [CrossRef] [PubMed]
10. Derman, Y.; Selby, K.; Miethe, S.; Frenzel, A.; Liu, Y.; Rasetti-Escargueil, C.; Avril, A.; Pelat, T.; Urbain, R.; Fontayne, A.; et al. Neutralization of Botulinum Neurotoxin Type E by a Humanized Antibody. Toxins 2016, 8, 257. [CrossRef] [PubMed]
11. Rosen, O.; Ozeri, E.; Barnea, A.; David, A.B.; Zichel, R. Development of an Innovative in Vitro Potency Assay for Anti-Botulinum Antitoxins. Toxins 2016, 8, 276. [CrossRef] [PubMed]
12. Torgeman, A.; Ozeri, E.; Ben David, A.; Diamant, E.; Rosen, O.; Schwartz, A.; Barnea, A.; Makovitzki, A.; Minman, A.; Zichel, R. Role of Homologous Fc Fragment in the Potency and Efficacy of Anti-Botulinum Antibody Preparations. Toxins 2017, 9, 180. [CrossRef] [PubMed]
13. Al-Saleem, F.H.; Sharma, R.; Puligedda, R.D.; Elias, M.; Kattala, C.D.; Simon, P.M.; Simpson, L.L.; Dessain, S.K. RBC Adherence of Immune Complexes Containing Botulinum Toxin Improves Neutralization and Macrophage Uptake. Toxins 2017, 9, 173. [CrossRef] [PubMed]
14. Bak, N.; Rajagopal, S.; Stickings, P.; Sesardic, D. SiMa Cells for a Serotype Specific and Sensitive Cell-Based Neutralization Test for Botulinum Toxin A and E. Toxins 2017, 9, 230. [CrossRef]
15. Webb, R.P.; Smith, T.J.; Smith, L.A.; Wright, P.M.; Guernieri, R.L.; Brown, J.L.; Skerry, J.C. Recombinant Botulinum Neurotoxin Hc Subunit (BoNT Hc) and Catalytically Inactive Clostridium botulinum Holoproteins (ciBoNT HPs) as Vaccine Candidates for the Prevention of Botulism. Toxins 2017, 9, 269. [CrossRef] [PubMed]
16. Otaka, D.Y.; Barbosa, J.D.; Moreira, C.; Ferreira, M.R.A.; Cunha, C.E.P.; Brito, A.R.S.; Donassolo, R.A.; Moreira, Â.N.; Conceição, F.R.; Salvarani, E.M. Humoral Response of Buffaloes to a Recombinant Vaccine against Botulism Serotypes C and D. *Toxins* 2017, 9, 297. [CrossRef] [PubMed]

17. Chang, H.J.; Hong, B.Y.; Lee, S.J.; Lee, S.; Park, J.H.; Kwon, J.Y. Efficacy and Safety of Letibotulinum Toxin A for the Treatment of Dynamic Equinus Foot Deformity in Children with Cerebral Palsy: A Randomized Controlled Trial. *Toxins* 2017, 9, 252. [CrossRef] [PubMed]

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