32. The cell of claim 1, wherein the alpha2-3sialyltransferase gene is selected from the group consisting of St3gal1, St3gal2, St3gal3, St3gal4, St3gal5, St3gal6 gene and a combination thereof.

~~33. A glycoprotein produced by the cell of claim 1, wherein the glycoprotein comprises an increase in alpha2-6 sialic acid linkages and a decrease in alpha2-3 sialic acid linkages as compared with a glycoprotein produced in an isolated CHO cell; and wherein the glycoprotein is an organophosphorus-scavenger protein.~~

~~34~~33. The cell of claim 1, further comprising a nucleic acid sequence that encodes a heterologous protein.

~~35~~34. The cell of claim ~~34~~33, wherein the protein is a glycoprotein.

~~36~~35. The cell of claim 1, wherein the means for knock-out is a zinc finger nuclease or deletion, mutation or addition of nucleotides.

What is claimed is:

1. An isolated mammalian cell comprising a heterologous alpha2-6 sialyltransferase nucleic acid sequence and a nucleic acid sequence that decreases expression of or silences an alpha2-3sialyltransferase gene or means for knock-out of the alpha2-3sialyltransferase gene.

2. The cell of claim 1, wherein the cell is a CHO cell.

3. The cell of claim 1, further comprising a nucleic acid sequence that encodes for human butyrylcholinesterase (huBChE).

4. The cell of claim 1, wherein the nucleic acid sequence that silences the alpha2-3sialyltransferase gene is selected from the group consisting of antisense, siRNA and miRNA.

5. The cell of claim 1, wherein the cell further comprises the nucleic acid sequence encoding for the proline-rich attachment domain (PRAD) of the ColQ gene.

6. A glycoprotein produced by the cell of claim 1, wherein the glycoprotein comprises an increase in alpha2-6 sialic acid linkages and a decrease in alpha2-3 sialic acid linkages as compared with a glycoprotein produced in an isolated CHO cell; and wherein the glycoprotein is recombinant huBChE (rhuBChE).

7. The cell of claim 1, further comprising a nucleic acid sequence encoding for an enzyme that reduces or inhibits alpha2-6 sialic acid degradation.

8. A method for the biosynthesis of an alpha2-6-rich glycoprotein comprising culturing a cell of claim 1 under conditions to co-express a nucleic acid sequence that encodes a peptide or protein.

9. The method of claim 8, further comprising inhibiting expression of alpha2-3 sialyltransferase.

10. The method of claim 8, further comprising reducing or inhibiting degradation of alpha2-6 sialic acid.

11. The method of claim 10, wherein alpha2-6 sialic acid degradation is reduced or inhibited by increasing activity of an enzyme that prevents alpha2-6 sialic acid degradation.

12. The method of claim 11, wherein the enzyme is fucosyltransferase.

13. The method of claim 12, wherein the fucosyltransferase is an alpha3fucosyltransferase (alpha3FucT).

14. The method of claim 13, wherein alpha3FucT is encoded by a nucleic acid sequence selected from FUT4, FUT5, FUT6, FUT7, FUT8, and FUT9.

15. The method of claim 12, wherein the fucosyltransferase is alpha3,4 fucosyltransferase (FucTLe) or alpha2fucosyltransferase (FucTLe).

16. The method of claim 10, wherein alpha2-6 sialic acid degradation is reduced or inhibited by decreasing activity of an enzyme that promotes alpha2-6 sialic acid degradation.

17. The method of claim 16, wherein the enzyme is a sialidase or neuramidase.

18. The method of claim 8, further comprising increasing the number or the length of N-glycan branches.

19. The method of claim 18, wherein the number of N-glycan branches is increased by increasing activity of galactose transferases or GlcNAc-transferases.

20. The method of claim 18, wherein the length of branches is increased by increasing the number of polylactosamines.

21. The method of claim 20, wherein the number of polylactosamines is increased by increasing expression of beta3-GlcNAC transferase (iGnT) and/or Gal transferase.

22. The method of claim 8, further comprising increasing CMP-sialic acid content.

23. The method of claim 8, wherein the peptide is a biological protective agent.

24. The method of claim 8, wherein the peptide is an organophosphorus scavenger.

25. The method of claim 8, wherein the peptide is rhuBChE.

26. The method of claim 8, further comprising modifying the cell to co-express tetramer assembly chaperones, thereby generating glycoprotein tetramers.

27. The method of claim 26, wherein the chaperone is PRAD.

28. A glycoprotein comprising an increase in alpha2-6 sialic acid linkages and a decreased level of alpha2-3 sialic acid linkages for use in the prevention or treatment of neurotoxin poisoning; wherein the glycoprotein is a recombinant human butyrylcholinesterase (rhuBChE).

29. A method for producing rhuBChE comprising culturing a mammalian cell that co-expresses huBChE, alpha2-6 sialyltransferase and PRAD in cell culture medium, thereby producing rhuBChE.

30. The method of claim 29, wherein the rhuBChE is isolated from the culture medium.

31. The method of claim 29, wherein the medium is serum-free medium.

32. The cell of claim 1, wherein the alpha2-3sialyltransferase gene is selected from the group consisting of St3gal1, St3gal2, St3gal3, St3gal4, St3gal5, St3gal6 gene and a combination thereof.

33. The cell of claim 1, further comprising a nucleic acid sequence that encodes a heterologous protein.

34. The cell of claim 33, wherein the protein is a glycoprotein.

35. The cell of claim 1, wherein the means for knock-out is a zinc finger nuclease or deletion, mutation or addition of nucleotides.