What is claimed is:

1. An isolated mammalian cell comprising a heterologous alpha2-6 sialyltransferase ~~(ST6GAL1)~~ nucleic acid sequence.

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 butyrlcholinesterase (huBChE).

4. The cell of claim 1, further comprising a nucleic acid sequence that decreases expression of or silences an alpha2-3sialyltransferase gene ~~(St3gal1)~~.

5. The cell of claim 4, wherein the nucleic acid sequence that silences ~~St3gal1~~ the alpha2-3sialyltransferase gene is selected from the group consisting of antisense, siRNA and miRNA.

6. The cell of claim 1, further comprising means for knock-out of ~~St3gal1~~ the alpha2-3sialyltransferase gene.

7. 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.

8. A ~~recombinant~~ glycoprotein comprising an increase in alpha2-6 sialic acid linkages and a decreased level of alpha2-3 sialic acid linkages.

9. The glycoprotein of claim 8, wherein the glycoprotein is in a tetrameric assembly state.

10. The glycoprotein of claim 8, wherein the glycoprotein is recombinant huBChE (rhuBChE).

11. The glycoprotein of claim 8, wherein the glycoprotein has an extended circulatory half-life or mean residence time (MRT).

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

13. 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 ~~for a peptide~~ a peptide or protein.

14. The method of claim 13, further comprising inhibiting expression of alpha2-3 sialyltransferase.

15. The method of claim 13, further comprising reducing or inhibiting degradation of alpha2-6 sialic acid.

16. The method of claim 15, wherein alpha2-6 sialic acid degradation is reduced or inhibited by increasing activity of an enzyme that prevents alpha2-6 sialic acid degradation.

17. The method of claim 16, wherein the enzyme is fucosyltransferase.

18. The method of claim 17, wherein the fucosyltransferase is an alpha3fucosyltransferase (alpha3FucT).

19. The method of claim 18, wherein alpha3FucT is encoded by a nucleic acid sequence selected from FUT4, FUT5, FUT6, FUT7, FUT8, and FUT9.

20. The method of claim 17, wherein the fucosyltransferase is alpha3,4 fucosyltransferase (FucTLe) or alpha2fucosyltransferase (FucTLe).

21. The method of claim 15, wherein alpha2-6 sialic acid degradation is reduced or inhibited by decreasing activity of an enzyme that promotes alpha2-6 sialic acid degradation.

22. The method of claim 21, wherein the enzyme is a sialidase or neuramidase.

23. The method of claim 13, further comprising increasing the number or the length of N-glycan branches.

24. The method of claim 23, wherein the number of N-glycan branches is increased by increasing activity of galactose transferases or GIcNAc-transferases.

25. The method of claim 23, wherein the length of branches is increased by increasing the number of polylactosamines.

26. The method of claim 25, wherein the number of polylactosamines is increased by increasing expression of beta3-GlcNAC transferase (iGnT) and/or Gal transferase.

27. The method of claim 13, further comprising increasing CMP-sialic acid content.

28. The method of claim 13, wherein the peptide is a biological protective agent.

29. The method of claim 13, wherein the peptide is an OP scavenger.

30. The method of claim 13, wherein the peptide is rhuBChE.

31. The method of claim 13, further comprising modifying the cell to co-express tetramer assembly chaperones, thereby generating glycoprotein tetramers.

32. The method of claim 31, wherein the chaperone is PRAD.

33. A glycoprotein comprising an increase in alpha2-6 sialic acid linkages and a decreased level of alpha2-3 sialic acid linkages ~~A method~~ for use in the prevention or treatment ~~in a subject~~ of neurotoxin poisoning ~~comprising administering to the subject a therapeutically effective amount of the glycoprotein of claim 8~~.

34. 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.

35. The method of claim 34, wherein the rhuBChE is isolated from the culture medium.

36. The method of claim 34, wherein the medium is serum-free medium.

37. The cell of claim 4, wherein the alpha2-3sialyltransferase gene is selected from the St3gal1 St3gal2, St3gal3, St3gal4, St3gal5, St3gal6 gene and a combination thereof.

What is claimed is:

1. An isolated mammalian cell comprising a heterologous alpha2-6 sialyltransferase nucleic acid sequence.

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 butyrlcholinesterase (huBChE).

4. The cell of claim 1, further comprising a nucleic acid sequence that decreases expression of or silences an alpha2-3sialyltransferase gene.

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

6. The cell of claim 1, further comprising means for knock-out of the alpha2-3sialyltransferase gene.

7. 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.

8. A glycoprotein comprising an increase in alpha2-6 sialic acid linkages and a decreased level of alpha2-3 sialic acid linkages.

9. The glycoprotein of claim 8, wherein the glycoprotein is in a tetrameric assembly state.

10. The glycoprotein of claim 8, wherein the glycoprotein is recombinant huBChE (rhuBChE).

11. The glycoprotein of claim 8, wherein the glycoprotein has an extended circulatory half-life or mean residence time (MRT).

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

13. 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.

14. The method of claim 13, further comprising inhibiting expression of alpha2-3 sialyltransferase.

15. The method of claim 13, further comprising reducing or inhibiting degradation of alpha2-6 sialic acid.

16. The method of claim 15, wherein alpha2-6 sialic acid degradation is reduced or inhibited by increasing activity of an enzyme that prevents alpha2-6 sialic acid degradation.

17. The method of claim 16, wherein the enzyme is fucosyltransferase.

18. The method of claim 17, wherein the fucosyltransferase is an alpha3fucosyltransferase (alpha3FucT).

19. The method of claim 18, wherein alpha3FucT is encoded by a nucleic acid sequence selected from FUT4, FUT5, FUT6, FUT7, FUT8, and FUT9.

20. The method of claim 17, wherein the fucosyltransferase is alpha3,4 fucosyltransferase (FucTLe) or alpha2fucosyltransferase (FucTLe).

21. The method of claim 15, wherein alpha2-6 sialic acid degradation is reduced or inhibited by decreasing activity of an enzyme that promotes alpha2-6 sialic acid degradation.

22. The method of claim 21, wherein the enzyme is a sialidase or neuramidase.

23. The method of claim 13, further comprising increasing the number or the length of N-glycan branches.

24. The method of claim 23, wherein the number of N-glycan branches is increased by increasing activity of galactose transferases or GlcNAc-transferases.

25. The method of claim 23, wherein the length of branches is increased by increasing the number of polylactosamines.

26. The method of claim 25, wherein the number of polylactosamines is increased by increasing expression of beta3-GlcNAC transferase (iGnT) and/or Gal transferase.

27. The method of claim 13, further comprising increasing CMP-sialic acid content.

28. The method of claim 13, wherein the peptide is a biological protective agent.

29. The method of claim 13, wherein the peptide is an OP scavenger.

30. The method of claim 13, wherein the peptide is rhuBChE.

31. The method of claim 13, further comprising modifying the cell to co-express tetramer assembly chaperones, thereby generating glycoprotein tetramers.

32. The method of claim 31, wherein the chaperone is PRAD.

33. 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.

34. 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.

35. The method of claim 34, wherein the rhuBChE is isolated from the culture medium.

36. The method of claim 34, wherein the medium is serum-free medium.

37. The cell of claim 4, wherein the alpha2-3sialyltransferase gene is selected from the St3gal1 St3gal2, St3gal3, St3gal4, St3gal5, St3gal6 gene and a combination thereof.