Claims

1. A liquid composition comprising a clostridial neurotoxin, a surfactant, an amino acid selected from tryptophan and tyrosine, a buffer comprising sodium, chloride and phosphate ions,
   wherein said liquid composition has a pH between 5.5 and 8,
   wherein said liquid composition is free of animal derived proteins, and
   wherein said liquid composition is stable for 2 months at 2-8 ^{o}C.
2. The liquid composition according to claim 1, wherein said surfactant is a non-ionic surfactant.
3. The liquid composition according to claim 2, wherein said non-ionic surfactant is a polysorbate.
4. The liquid composition according to claim 3, wherein said polysorbate is Polysorbate 20, Polysorbate 60 or Polysorbate 80.
5. The liquid composition according to any one of the preceding claims, wherein said amino acid is tryptophan.
6. The liquid composition according to claim 5, wherein said tryptophan is L-tryptophan.
7. The liquid composition according to any one of the preceding claims, wherein said buffer further comprises potassium ions.
8. The liquid composition according to any one of the preceding claims, wherein said liquid composition has a pH between 6.0 and 7.5.
9. The liquid composition according to any one of the preceding claims, wherein no more than 30% loss in extracellular proteolytic activity occurs over 2, 3, 6, 12, 18, 24 or 36 months at 5°C.
10. The liquid composition according to any one of the preceding claims, wherein said clostridial neurotoxin is a botulinum neurotoxin, selected from a natural botulinum neurotoxin in complex form, a high purity natural botulinum neurotoxin and a recombinant botulinum neurotoxin, wherein said high purity natural botulinum neurotoxin is a botulinum neurotoxin essentially free of neurotoxin-associated proteins (NAPs).
11. The liquid composition according to claim 10, wherein said botulinum neurotoxin is a recombinant botulinum neurotoxin selected from a botulinum neurotoxin A, B, C, D, E, F or G, a modified botulinum neurotoxin and a chimeric botulinum neurotoxin, wherein said modified botulinum neurotoxin is a botulinum neurotoxin having an

amino acid sequence that has at least 50% sequence identity with SEQ ID NO 1, 2, 3, 4, 5, 6 or 7.

12. The liquid composition according to claim 1, wherein said liquid composition comprises:
    - 4 to 10000 LD50 units of botulinum neurotoxin per mL,
    - 0.001 to 15% v/v polysorbate,
    - 0.1 to 5 mg/mL tryptophan,
    - 10 to 500 mM NaCl,
    - 1 to 50 mM KCl,
    - 1 to 100 mM Sodium phosphate,

    has a pH between 5.5 and 8, and is stable for 6 months at 5°C.
13. The liquid composition according to claim 12, wherein said liquid composition comprises:
    - 10 to 2000 LD50 units of botulinum neurotoxin per mL,
    - 0.05 to 0.2% v/v polysorbate 80,
    - 0.1 to 5 mg/mL tryptophan,
    - 25 to 300 mM NaCl,
    - 1 to 10 mM KCl,
    - 2 to 50 mM Sodium phosphate,

    has a pH between 6.0 and 7.5, and is stable for 12 months at 5°C.
14. The liquid composition according to any one of the preceding claims for use in therapy.
15. Use of the liquid composition according to any one of claims 1-13 in the manufacture of a medicament for treating or preventing muscular disorders, neuromuscular disorders, neurological disorders, ophthalmological disorders, pain disorders, psychological disorders, articular disorders, inflammatory disorders, endocrine disorders or urological disorders.
16. Use of the liquid composition according to any one of claims 1 to 13 in aesthetic medicine.
17. The use according to claim 16 for treating or preventing skin wrinkles.
18. The use according to claim 17, wherein said skin wrinkles include facial wrinkles such as facial frown lines, wrinkles of the contour of the eye, glabellar frown lines, downturned mouth, wrinkles of the neck (platysmal bands), wrinkles of the chin

(mentalis, peau d'orange, dimpled chin), forehead lines, "scratched skin" wrinkles, nasal lift treatment or sleep lines.

19. Use of an amino acid selected from tryptophan and tyrosine to protect a clostridial neurotoxin from degradation in a liquid composition which is free of animal derived proteins.
20. The use according to claim 19, wherein said amino acid is tryptophan.
21. The use according to claim 19 or 20, wherein said clostridial neurotoxin is a botulinum neurotoxin.
22. The use according to any one of claims 19-21, wherein said amino acid is used in combination with a surfactant and a buffer comprising sodium, chloride and phosphate ions, and said liquid composition has a pH between 5.5 and 8.