CLAIMS

1. A liquid pharmaceutical composition comprising:
   0.5-10 mg/ml semaglutide,
   0.0% (w/w) to-0.1 % (w/w) phenol, and
   8.2 to 8.9 mg/ml sodium chloride.

2. The liquid pharmaceutical composition according to claim 1 comprising: 0.5 to 3.5 mg/ml semaglutide.

3. The liquid pharmaceutical composition according to claim 1 comprising: 3.2 mg/ml semaglutide.

4. The liquid pharmaceutical composition according to claim 1 further comprising: 0.01 to 50 mM of a buffer.

5. The liquid pharmaceutical composition according to claim 4, wherein the buffer is a phosphate buffer.

6. The liquid pharmaceutical composition according to claim 5, wherein the phosphate buffer is present in a concentration of 1 to 2 mg/ml.

7. The liquid pharmaceutical composition according to claim 1, wherein the pharmaceutical composition has a pH in the range of 6.0 to 10.0.

8. The liquid pharmaceutical composition according to claim 1, wherein the pharmaceutical composition is a parenteral composition.

9. The liquid pharmaceutical composition according to claim 1 further comprising:
   0.5 to 3.5 mg/ml semaglutide,
   0.01 to 50 mM buffer, and
   pH in the range of 7 to 8.

10. The liquid pharmaceutical composition according to claim 1 further comprising:
    0.5 to 3.5 mg/ml semaglutide,

1 to 2 mg/ml of phosphate buffer, and
pH in the range of 7 to 8.

11. The liquid pharmaceutical composition according to claim 1 further comprising:
0.5 to 3.5 mg/ml semaglutide,
8.25 mg/ml sodium chloride, and
1.42 mg/ml disodium hydrogen phosphate dihydrate.

12. The liquid pharmaceutical composition according to claim 1 further comprising:
histidine.

13. A kit comprising a liquid pharmaceutical composition according to claim 1, wherein the kit further comprises an injection device for administration of the composition to a subject in need of such pharmaceutical composition.

14. The kit according to claim 13, wherein the injection device comprises a needle and pre-filled syringe or syringe cartridge.

15. Use of an effective amount of the liquid pharmaceutical composition according to claim 1 in the manufacture of a medicament for the treatment of diabetes, obesity and nonalcoholic steatohepatitis (NASH).

16. A liquid pharmaceutical composition consisting of:
0.5 to 3.5 mg/mL semaglutide,
0% (w/w) to 0.1% (w/w) phenol,
buffer,
6.5 to 12 mg/mL sodium chloride,
at least 60% water,
optionally histidine, and
optionally one or more agents for adjusting pH.

17. The liquid pharmaceutical composition according to claim 16, wherein sodium chloride is in the amount of 8.1 to 12 mg/mL.

18. The liquid pharmaceutical composition according to claim 16, wherein sodium chloride is in the amount of 8.2 to 8.9 mg/mL.

19. The liquid pharmaceutical composition according to claim 16, wherein the buffer is a phosphate buffer.

20. The liquid pharmaceutical composition according to claim 19, wherein the buffer is in the amount of 0.01 to 50 mM.

21. The liquid pharmaceutical composition according to claim 19, wherein the buffer is in the amount of 1 to 2 mg/mL.

22. Use of an effective amount of the liquid pharmaceutical composition according to claim 16 in the manufacture of a medicament for the treatment of diabetes, obesity and nonalcoholic steatohepatitis (NASH).

23. A liquid pharmaceutical composition consisting of:
3.2 mg/mL semaglutide,
0% (w/w) to 0.1% (w/w) phenol,
1.42 mg/mL disodium hydrogen phosphate dihydrate buffer,
8.25 mg/mL sodium chloride,
pH of 7.4,
at least 60% water, and
optionally one or more agents for adjusting pH.

24. The liquid pharmaceutical composition according to claim 23, wherein the pharmaceutical composition is a parenteral composition.

25. The liquid pharmaceutical composition according to claim 23, wherein the pharmaceutical composition has a visual analogue scale (VAS) score below 15 on a 100 mm scale.

26. Use of an effective amount of the liquid pharmaceutical composition according to claim 23 in the manufacture of a medicament for the treatment of diabetes, obesity, Alzheimer's disease, nonalcoholic steatohepatitis (NASH).