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dosage form in an amount of 0.3% to 0.6%, and SEQ ID NO.: 10 is present in the unit dosage form in an amount of 0.1%.

9. The method of claim 1, wherein the human's hypotension is associated with vasodilatory shock.

10. The method of claim 9, wherein the administration provides to the human from about 0.01 units of vasopressin or the pharmaceutically-acceptable salt thereof per minute to about 0.07 units of vasopressin or the pharmaceutically-acceptable salt thereof per minute.

11. The method of claim 1, wherein the impurities comprise a plurality of peptides, wherein the impurities are determined based on:

(a) injecting the unit dosage form into a high pressure liquid chromatography apparatus, wherein the apparatus comprises:

(i) a chromatography column containing adsorbent particles as a stationary phase;

(ii) a first mobile phase passing through the chromatography column, wherein the first mobile phase is phosphate buffer at pH 3; and

(iii) a second mobile phase passing through the chromatography column, wherein the second mobile phase is a 50:50 acetonitrile:water solution;

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(b) running the unit dosage form through the chromatography column for 55 minutes;

(c) eluting the vasopressin and the plurality of peptides from the chromatography column using a gradient of the first mobile phase, and a gradient of the second mobile phase, wherein each of the first and second mobile phase are run at a flow rate of 1 mL/min through the chromatography column;

(d) passing the eluted vasopressin and the plurality of peptides through a UV detector to generate a UV spectrum of the eluted vasopressin and the plurality of peptides;

(e) identifying a peptide of the plurality of peptides based on a retention time of the peptide of the plurality of peptides relative to a standard; and

(f) calculating an amount of the peptide of the plurality of peptides based on an integration of a peak obtained for the peptide of plurality of peptides from the UV spectrum.

12. The method of claim 1, wherein the unit dosage form further comprises sodium acetate.

13. The method of claim 1, the unit dosage form further comprising a pH adjusting agent.

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