Conditional amendments to EP (UK) No. 2 944 306 ("EP 306"):

1. A pre-filled syringe comprising an ophthalmic formulation of a vascular endothelial growth factor (VEGF) antagonist, comprising:

(a) 40-50~~1-100~~ mg/ml of the VEGF antagonist ~~comprising~~being a VEGF-specific fusion protein, wherein the VEGF-specific fusion protein comprises amino acids 27-457 of SEQ ID NO:4, and is glycosylated at Asn residues 62, 94, 149, 222 and 308;

(b) 0.01-5% of one or more organic co-solvent(s) which is one or more of polysorbate, polyethylene glycol (PEG), and propylene glycol;

(c) 30-150 mM of a tonicity agent selected from sodium chloride and potassium chloride;

(d) 5-40 mM of sodium phosphate buffer; and

(e) 1.0-7.5% of a stabilizing agent selected from the group consisting of sucrose, sorbitol, glycerol, trehalose, and mannitol;

having a pH of 5.8-7.0, wherein the formulation is suitable for intravitreal administration.

~~2. The pre-filled syringe according to claim 1, wherein the VEGF-specific fusion protein comprises a receptor component comprising an immunoglobulin-like (Ig) domain 2 of a first VEGF receptor and Ig domain 3 of a second VEGF receptor, and a multimerizing component.~~

~~3. The pre-filled syringe according to claim 2, wherein the first VEGF receptor is Flt1 and the second VEGF receptor is Flkl or Flt4.~~

~~4. The pre-filled syringe according to any one of claims 1-3, wherein the VEGF-specific fusion protein comprises amino acids 27-457 of SEQ ID NO:4, and is glycosylated at Asn residues 62, 94, 149, 222 and 308.~~

2~~5~~. The pre-filled syringe according to ~~any one of~~ claim~~s~~ 1-4, wherein the liquid stable ophthalmic formulation comprises 40 mg/mL of the VEGF specific fusion protein, 10 mM sodium phosphate, 40 mM NaCl, 0.03% polysorbate 20, and 5% sucrose, pH 6.2-6.3.