

35. The formulation of claim 34, wherein in the Avelumab glycosylation said A2 has a share of about 3.5% - about 4.5%, said A2G1 has a share of about 0.5% - about 3.5%, said A2G2 has a share of < 2.5% and said FA2G2 has a share of about 5.5% of all glycan species.

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36. The formulation of any one of claims 29-35, wherein said Avelumab has the heavy chain sequence of (SEQ ID NO:2) .

37. The formulation of any one of claims 1-36 which is for intravenous (IV) administration.

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38. A vial containing the formulation of claim 37.

39. The vial of claim 38 which contains 200 mg avelumab in 10 mL of solution for a concentration of 20 mg/mL.

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40. The vial of claims 38 or 39 which is a glass vial.

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