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STABLE FREEZE-DRIED PHARMACEUTICAL FORMULATION

Abstract of the technical content of the invention

The subject of the invention is a freeze-dried formulation consisting of an amorphous phase and a crystalline phase, which is pharmaceutically acceptable, comprising at least one nonprotein active ingredient, characterized in that it contains mannitol and alanine in a ratio R of between 0.1 and 1, R representing the mass of mannitol over the mass of alanine.

Figure none