

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
3 September 2009 (03.09.2009)

PCT

(10) International Publication Number
WO 2009/108775 A3

(51) International Patent Classification:
A61K 9/20 (2006.01) *A61M 36/14* (2006.01)

(21) International Application Number:
PCT/US2009/035276

(22) International Filing Date:
26 February 2009 (26.02.2009)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
61/032,298 28 February 2008 (28.02.2008) US

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(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report (Art. 21(3))

(88) Date of publication of the international search report:
3 December 2009

(54) Title: PROCESS TO MINIMIZE POLYMORPHISM

(57) Abstract: The commercial formulation of fast dispersing dosage forms (FDDF) requires substantial holding times during which large quantities of pharmaceutically active substance are formed into individual dosage units. During this holding time, pharmaceutical agents with a propensity to polymorphism in an aqueous environment may crystallize into various, and sometimes unpredictable forms. These crystalline forms may affect the efficacy of the pharmaceutical agent. Previous attempts to control this process have included attempts to direct crystallization into a stable form. The instant invention acts to suppress crystallization, by utilizing a combination of standard molecular weight fish gelatin and a low processing temperature, to suppress crystallization to a degree that is not accomplished by either the use of standard molecular weight fish gelatin or low processing temperatures individually.



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