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(21) International Application Number: PCT/EP98/04192 ✓ (22) International Filing Date: 7 July 1998 (07.07.98) ✓ (30) Priority Data: ① 97202118.2 ✓ 10 July 1997 (10.07.97) ✓ EP ✓ (71) Applicant (for all designated States except US): ① JANSSEN PHARMACEUTICA N.V. [BE/BE]; Turnhoutseweg 30, B-2340 Beerse (BE). ✓ (72) Inventors; and (75) Inventors/Applicants (for US only): ① LACRAMPE, Jean, Fernand, Armand [FR/FR]; Janssen-Cilag S.A., 1, rue Camille Desmoulins, TSA 91003, F-92787 Issy-les-Moulineaux Cedex 9 (FR). FREYNE, Eddy, Jean ② Edgard [BE/BE]; Janssen Pharmaceutica N.V., Turnhoutseweg 30, B-2340 Beerse (BE). VENET, Marc, Gaston [FR/FR]; Janssen-Cilag S.A., 1, rue Camille Desmoulins, TSA 91003, F-92787 Issy-les-Moulineaux Cedex 9 (FR). ④ BOECKX, Gustaaf, Maria [BE/BE]; Janssen Pharmaceutica N.V., Turnhoutseweg 30, B-2340 Beerse (BE). ✓	(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, ③ SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published With international search report. ✓ Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments. 1 appl. 4 invs.	
(54) Title: IL-5 INHIBITING 6-AZAUACIL DERIVATIVES ✓		
(I)		
(57) Abstract The present invention is concerned with the compounds of formula (I) the N-oxides, the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof, p and q are 0, 1, 2, 3 or 4 and q is also 5; X is O, S, NR³ or a direct bond; R¹ is hydrogen, hydroxy, halo, optionally substituted amino, optionally substituted C₁₋₆alkyl, C₁₋₆alkyloxy, C₃₋₇cycloalkyl or aryl; R² is aryl, Het¹, C₃₋₇cycloalkyl, optionally substituted C₁₋₆alkyl; and if X is O, S or NR³, then R² may also be a carbonyl or thiocarbonyl linked substituent; R³ is hydrogen or C₁₋₄alkyl; R⁴ and R⁵ independently are optionally substituted C₁₋₆alkyl, halo, hydroxy, mercapto, C₁₋₆alkyloxy, C₁₋₆alkylthio, C₁₋₆alkylcarbonyloxy, aryl, cyano, nitro, Het³, R⁶ or NR⁷R⁸; R⁶ is substituted sulfonyl or sulfinyl; R⁷ and R⁸ are hydrogen, optionally substituted C₁₋₄alkyl, aryl, a carbonyl or thiocarbonyl linked substituent, C₃₋₇cycloalkyl, Het³ and R⁶; R⁹ and R¹⁰ are each independently selected from hydrogen, optionally substituted C₁₋₄alkyl, phenyl, a carbonyl or thiocarbonyl linked substituent, C₃₋₇cycloalkyl, Het³ and R⁶; R¹¹ is hydroxy, mercapto, cyano, nitro, halo, trihalomethyl, C₁₋₄alkyloxy, carboxyl, C₁₋₄alkyloxycarbonyl, trihaloC₁₋₄alkylsulfonyloxy, R⁶, NR⁷R⁸, C(=O)NR⁷R⁸, aryl, aryloxy, arylcarbonyl, C₃₋₇cycloalkyl, C₃₋₇cycloalkyloxy, phthalimide-2-yl, Het³ and C(=O)Het³; R¹² and R¹³ are each independently selected from hydrogen, optionally substituted C₁₋₄alkyl, phenyl, a carbonyl or thiocarbonyl linked substituent, C₃₋₇cycloalkyl and R⁶; aryl is optionally substituted phenyl; Het¹, Het² and Het³ are optionally substituted heterocycles; to processes for their preparation and compositions comprising them. It further relates to their use as a medicine.		