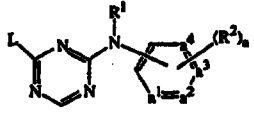




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(21) International Application Number: PCT/EP99/08688 (22) International Filing Date: 4 November 1999 (04.11.99) (30) Priority Data: 60/107,799 10 November 1998 (10.11.98) US 99203128.6 24 September 1999 (24.09.99) EP (71) Applicant (for all designated States except US): JANSSEN PHARMACEUTICA N.V. [BE/BE]; Patent Department, Turnhoutseweg 30, B-2340 Beerse (BE). (72) Inventors; and (75) Inventors/Applicants (for US only): DE CORTE, Bart [BE/US]; 1590 Winding Road, Southampton, PA 18966 (US). DE JONGE, Marc, René [NL/NL]; Hontenissestraat 3, NL-5045 CA Tilburg (NL). HEERES, Jan [NL/BE]; Leemskuiten 18, B-2350 Vosselaar (BE). JANSSEN, Paul, Adriaan, Jan [BE/BE]; Antwerpsesteenweg 20, B-2350 Vosselaar (BE). KAVASH, Robert, W. [US/US]; 148 N. Keswick Avenue, Glenside, PA 19038 (US). KOYMANS, Lucien, Maria, Henricus [NL/BE]; Stationstraat 68 bus 4, B-2300 Turnhout (BE). KUKLA, Michael, Joseph [US/US]; 1551 Oak Hollow Drive, Maple Glen, PA 19002 (US). LUDOVICI, Donald, William [US/US]; 1390 Masi Road, Quakertown, PA 18951 (US). VAN AKEN, Koen,	Jeanne, Alfons [BE/BE]; Tuinbouwstraat 22, bus 2, B-2300 Turnhout (BE). (74) Agent: BOULT WADE TENNANT; 27 Fumival Street, London EC4A 1PQ (GB). (81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, 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, GW, ML, MR, NE, SN, TD, TG). Published <i>Without international search report and to be republished upon receipt of that report.</i>	
(54) Title: 2,4-DISUBSTITUTED TRIAZINE DERIVATIVES		
 (1)		
(57) Abstract This invention concerns the use of the compounds of formula (I), the <i>N</i>-oxides, the pharmaceutically acceptable addition salts, quaternary amines and the stereochemically isomeric forms thereof, wherein $-a^1-a^2-a^3-a^4-$ forms a phenyl, pyridinyl, pyrimidinyl, pyridazinyl or pyrazinyl with the attached vinyl group; <i>n</i> is 0 to 4; and where possible 5; <i>R</i>¹ is hydrogen, aryl, formyl, C₁-alkylcarbonyl, C₁-alkyl, C₁-alkyloxycarbonyl or substituted C₁-alkyl; each <i>R</i>² independently is hydroxy, halo, optionally substituted C₁-alkyl, C₂-alkenyl or C₂-alkynyl, C₃-cycloalkyl, C₁-alkyloxy, C₁-alkyloxycarbonyl, carboxyl, cyano, nitro, amino, mono- or di(C₁-alkyl)amino, polyhalomethyl, polyhalomethyloxy, polyhalomethylthio, $-S(=O)_pR^4$, $-NH-S(=O)_pR^4$, $-C(=O)R^4$, $-NHC(=O)H$, $-C(=O)NHNH_2$, $-NHC(=O)R^4$, $-C(=NH)R^4$ or a 5-membered heterocyclic ring; <i>p</i> is 1 or 2; <i>L</i> is optionally substituted C₁-alkyl, C₂-alkenyl, C₂-alkynyl or C₃-cycloalkyl; or <i>L</i> is $-X-R^3$ wherein <i>R</i>³ is optionally substituted phenyl, pyridinyl, pyrimidinyl, pyrazinyl or pyridazinyl; <i>X</i> is $-NR^1-$, $-NH-NH-$, $-N=N-$, $-O-$, $-C(=O)-$, $-CHOH-$, $-S-$, $-S(=O)-$ or $-S(=O)_2-$; aryl is optionally substituted phenyl; for the manufacture of a medicine for the treatment of subjects suffering from HIV (Human Immunodeficiency Virus) infection.		