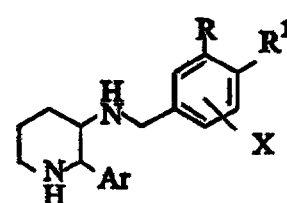


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(21) International Application Number: PCT/IB96/00572 (22) International Filing Date: 10 June 1996 (10.06.96) (30) Priority Data: PCT/IB95/00683 24 August 1995 (24.08.95) WO (34) Countries for which the regional or international application was filed: JP et al. (71) Applicant (for JP only): PFIZER PHARMACEUTICALS INC. [JP/JP]; Mitsui Building, 1-1, Nishi-shinjuku 2-chome, Shinjuku-ku, Tokyo 163 (JP). (71) Applicant (for all designated States except JP US): PFIZER INC. [US/US]; 235 East 42nd Street, New York, NY 10017 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): SATAKE, Kunio [JP/JP]; 4-58, Hakusan-cho, Handa-shi, Aichi-ken 475 (JP). SHISHIDO, Yuji [JP/JP]; 362, Aza-Giro, Oaza-Okuda, Mihama-cho, Chita-gun, Aichi-ken 470-32 (JP). WAKABAYASHI, Hiroaki [JP/JP]; 3-1, Sumiyoshi-cho, Kariya-shi, Aichi-ken 448 (JP).	(74) Agents: SPIEGEL, Allen, J. et al.; Pfizer Inc., 235 East 42nd Street, New York, NY 10017 (US). (81) Designated States: AU, BG, BR, BY, CA, CN, CZ, HU, IS, JP, KR, KZ, LK, LV, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TR, UA, US, UZ, VN, European patent (AT, BE, CH, 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.	
(54) Title: SUBSTITUTED BENZYLAMINOPIPERIDINE COMPOUNDS  (I) (57) Abstract This invention provides a compound of formula (I) and its pharmaceutically acceptable salts, wherein R is halo C₁-C₈ alkyl, halo C₂-C₈ alkenyl, halo C₂-C₈ alkynyl or halo C₁-C₈ alkyl substituted by hydroxy or C₁-C₈ alkoxy; R¹ is hydrogen, halo or C₁-C₆ alkoxy; or R and R¹, together with the two carbon atoms shared between the benzene ring and the R and R¹, complete a fused C₄-C₆ cycloalkyl wherein one carbon atom is optionally replaced by oxygen and wherein one or two of the carbon atoms are optionally substituted by up to five substituents selected from halo, C₁-C₆ alkyl and halo C₁-C₆ alkyl; X is C₁-C₆ alkoxy, halo C₁-C₆ alkoxy, phenoxy or halo; and Ar is phenyl optionally substituted by halo. These compounds are of use in treating a gastrointestinal disorder, a central nervous system (CNS) disorder, an inflammatory disease, emesis, urinary incontinence, pain, migraine, sunburn, diseases, disorders and adverse conditions caused by <i>Helicobacter pylori</i>, or angiogenesis especially CNS disorders in a mammalian subject, especially humans.		

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