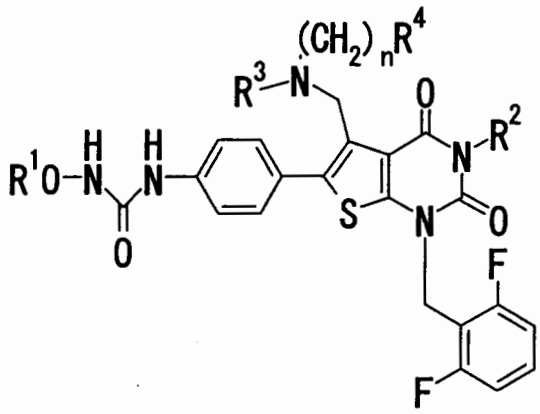
Abstract

The present invention provides a compound represented by the formula:



wherein R^1 is a C_{1-4} alkyl; R^2 is (1) a 5- to 7-membered nitrogen-containing heterocyclic group which may have a substituent selected from the group consisting of (1') a halogen, (2') a hydroxy group, (3') a C_{1-4} alkyl and (4') a C_{1-4} alkoxy, (2) a phenyl which may have a substituent selected from the group consisting of (1') a halogen, (2') a C_{1-4} alkoxy-C_{1-4} alkyl, (3') a mono-C_{1-4} alkyl-carbamoyl-C_{1-4} alkyl, (4') a C_{1-4} alkoxy and (5') a mono-C_{1-4} alkylcarbamoyl-C_{1-4} alkoxy, or the like; R^3 is a C_{1-4} alkyl; R^4 is a C_{1-4} alkoxy, or the like; n is an integer of 1 to 4; or a salt thereof, as a thienopyrimidine compound having gonadotropin-releasing hormone antagonistic activity.