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(54) Title: CYCLOCARBAMATE DERIVATIVES AS PROGESTERONE RECEPTOR MODULATORS			
(I)			
(57) Abstract			
This invention provides compounds of Formula (I) wherein R¹ and R² may be single substituents or fused to form spirocyclic or heterospirocyclic rings; R³ is H, OH, NH₂, C₁ to C₆ alkyl, substituted C₁ to C₆ alkyl, C₃ to C₆ alkenyl, substituted C₁ to C₆ alkenyl, alkynyl, or substituted alkynyl, COR^C; R^C is H, C₁ to C₃ alkyl, substituted C₁ to C₃ alkyl, aryl, substituted aryl, C₁ to C₃ alkoxy, substituted C₁ to C₃ alkoxy, C₁ to C₃ aminoalkyl, or substituted C₁ to C₃ aminoalkyl; R⁴ is H, halogen, CN, NO₂, C₁ to C₆ alkyl, substituted C₁ to C₆ alkyl, alkynyl, or substituted alkynyl, C₁ to C₆ alkoxy, substituted C₁ to C₆ alkoxy, substituted C₁ to C₆ alkoxy, amino, C₁ to C₆ aminoalkyl, or substituted C₁ to C₆ aminoalkyl; and R⁵ is selected from a trisubstituted benzene ring of a five or six membered ring with 1, 2, or 3 heteroatoms from the group including O, S, SO, SO₂ or NR⁶ and containing one or two independent substituents from the group including H, halogen, CN, NO₂, amino, and C₁ to C₃ alkyl, C₁ to C₃ alkoxy, C₁ to C₃ aminoalkyl, COR^F, or NR^GCOR^F; or pharmaceutically acceptable salt thereof, as well as pharmaceutical compositions and methods using the compounds as antagonists of the progesterone receptor.			