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CAplus includes complete International Patent Classification (IPC) reclassification data for the first quarter of 2015.

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AΒ Title derivs. I [R = H, C1-6 alkyl, C2-6 alkenyl, (C6-12 aryl)-C1-4 alkyl; R1 = H, (un)substituted C1-6 alkyl, C2-6 alkenyl, C5-9 cycloalkyl, C5-9 cycloalkenyl, etc.; R2 = H, C1-6 alkyl, C2-6 alkenyl, (C6-12 aryl)-C1-4 alkyl; R3 = H, OH, R4 = H; R3R4 = O; R5 = C1-6 alkyl, C2-6 alkenyl, C2-6 alkenyl, C5-9 cycloalkyl, (un)substituted C6-12 aryl; n = 0, 1] were prepd. as antihypertensives (no data) due to their ability to inhibit angiotensin-converting enzyme. Thus, cis-bicyclo[3.1.0]hexan-2-one was treated with H2NOSO3H and then subjected to the Beckman rearrangement to give cis-azabicyclo[4.1.0]heptane cis-II (R6 = R7 = H), which was chlorinated with PC15 to give cis-II (R6 = R7 = C1), which was dechlorinated by hydrogenation over Raney Ni to give cis-II (R6 = Cl, R7 = H). The latter was hydrolyzed in the presence of Ba(OH)2 to give cis-azabicyclo[3.1.0]hexane-3-carboxylate cis-III, which was sepd. into its exo and endo isomers. The latter were esterified with PhCH2OH via SOC12 to give the corresponding benzyl esters, which were condensed with (S)-PhCH2CH2CH(CO2Et)-L-Ala-OH by DCC/1-hydroxybenzotriazole to give the exo and endo isomers of title compd. cis-IV (R8 = Et, R9 = CH2Ph), which were sepd. into the 3S-endo, 3R-endo, 3S-exo, and 3R-exo isomers. The latter were debenzylated by hydrogenolysis over Pd/C and then treated with HCl/EtOH to give the corresponding cis-IV.HCl (R8 = Et, R9 = H). 3S-endo-cis-IV.HCl (R8 = Et, R9 = H) was sapond. to give 3S-endo-cis-IV (R8 = R9 = H); 3S-exo-cis-IV (R8 = R9 = H) was also prepd. OSC.G THERE ARE 8 CAPLUS RECORDS THAT CITE THIS RECORD (8 CITINGS)