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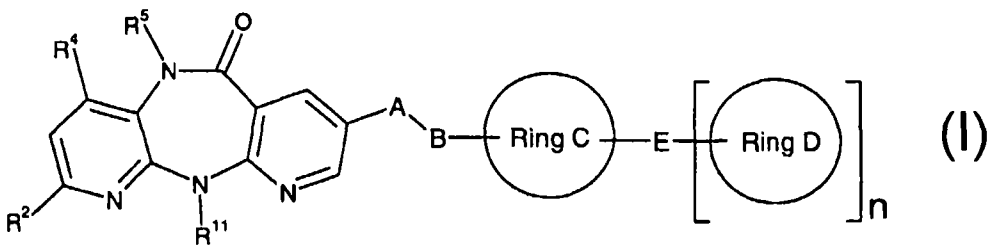
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(54) Title: NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITORS



(57) Abstract: Compounds represented by formula (I), wherein R^2 is selected from the group consisting of H, (C_{1-4})alkyl, halo, haloalkyl, OH, (C_{1-6})alkoxy, NH$(C_{1-4}$alkyl) or N$(C_{1-4}$alkyl)$_2$; R^4 is H or Me; R^5 is H or Me; R^{11} is H, (C_{1-4})alkyl, (C_{3-4})cycloalkyl and (C_{1-4})alkyl-(C_{3-4})cycloalkyl; A is a connecting chain of (C_{1-3})alkyl; B is O or S; n is 0 or 1; wherein when n is 0: Ring C is 6- or 10-membered aryl or 5- or 6-membered heterocycle having from 1 to 4 heteroatoms selected from the group consisting of O, N, and S, said aryl and said heterocycle being optionally substituted; and E is $CONR^{12}R^{13}$; $CONHNR^{14}R^{15}$; $NR^{16}COR^{17}$; $NR^{18}SO_2(C_{1-6})$alkyl; $SO_2NR^{19}R^{20}$; or SO_2R^{21}; or when n is 1: Ring C is as defined above and E is a single bond or a connecting group; and Ring D is 6- or 10-membered aryl or 5- or 6-membered heterocycle having from 1 to 4 heteroatoms selected from the group consisting of O, N, and S, said aryl and said heterocycle being optionally substituted with from 1 to 5 substituents; or a salt or a prodrug thereof are provided as inhibitors of HIV reverse transcriptase.

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