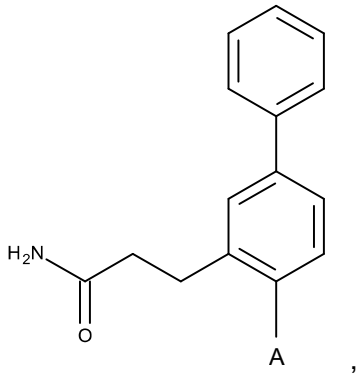
THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:-

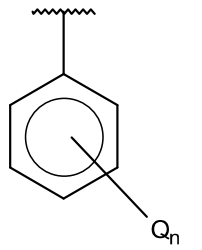
1. A compound of the formula



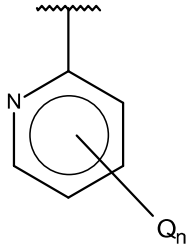
,

wherein:

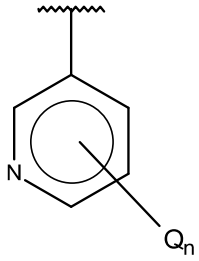
A is selected from the group consisting of:



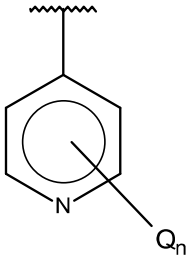
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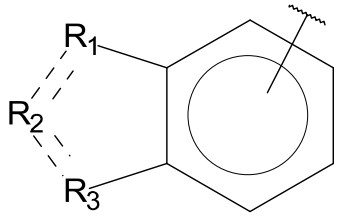
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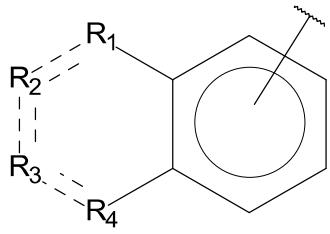


,



,





and

,

Q is independently selected from halo, alkyl, hydroxy, amino and substituted amino;

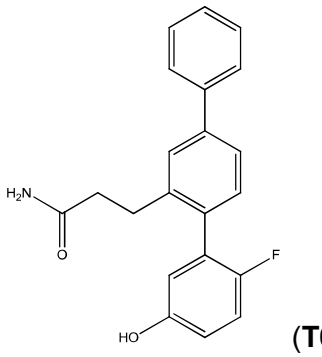
n is 0, 1, 2, 3, 4 or 5;

R_1, R_3 and R_4 are independently C, CH, CH_2, O, N, NH or S, and

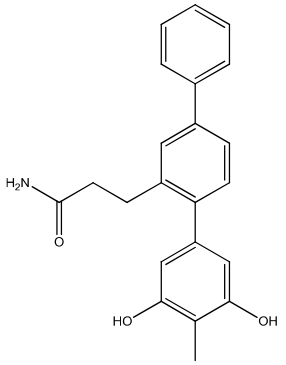
R_2 is C, CH, CH_2, N, NH, C-CF_3, CH-CF_3 or C=O,

or a stereoisomer or pharmaceutically acceptable salt thereof,

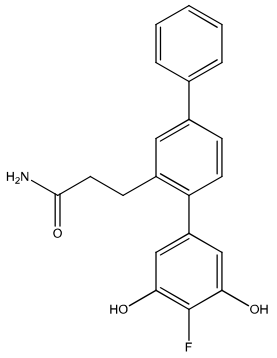
wherein when n is 1, Q cannot be hydroxy.



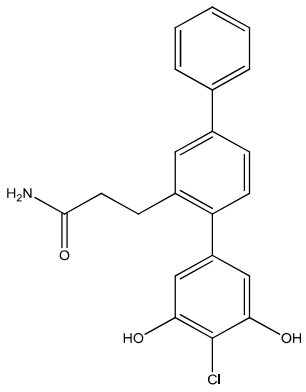
(**T66**),



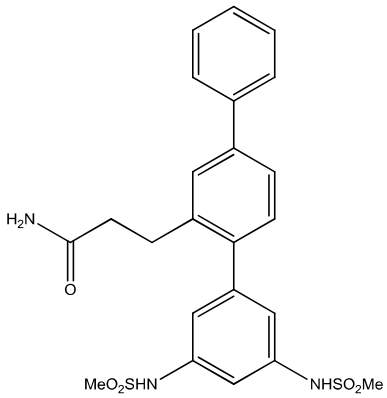
(**T67**),



(**T68**),



(**T69**) and



(**T70)**.

8. A pharmaceutical composition comprising a compound, or a stereoisomer or pharmaceutically acceptable salt thereof, according to any one of claims 1 to 7 and a pharmaceutically-acceptable excipient.

9. Use of a compound, or a stereoisomer or pharmaceutically acceptable salt thereof, according to any one of claims 1 to 7 for the manufacture of a medicament for the therapeutic treatment of hypertension or prehypertension.

10. Use of a compound, or a stereoisomer or pharmaceutically acceptable salt thereof, according to any one of claims 1 to 7 for the manufacture of a medicament for the prophylactic treatment of fibrosis.

11. Use of a compound, or a stereoisomer or pharmaceutically acceptable salt thereof, according to any one of claims 1 to 7 for the manufacture of a medicament for the therapeutic treatment of fibrosis.

12. Use of a compound, or a stereoisomer or pharmaceutically acceptable salt thereof, according to any one of claims 1 to 7 for the manufacture of a medicament for the therapeutic treatment of hypertension and fibrosis.

13. Use of a compound, or a stereoisomer or pharmaceutically acceptable salt thereof, according to any one of claims 1 to 7 for the manufacture of a medicament for the therapeutic treatment of prehypertension and fibrosis.