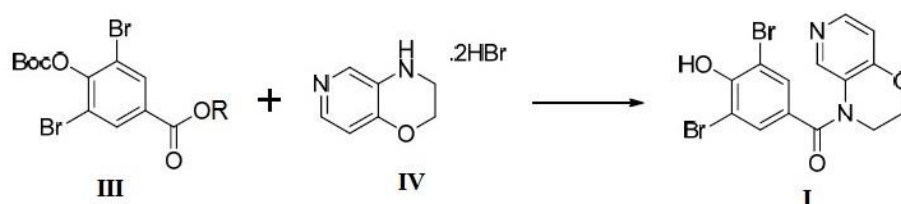


[Claims]

[Claim 1]

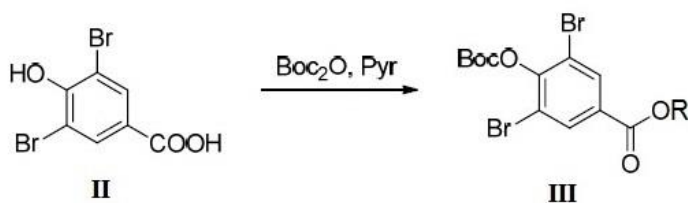
A process for preparing a compound of the following Formula I, or a pharmaceutically acceptable salt thereof or a hydrate thereof, comprising coupling-reacting a compound of the following Formula III with a compound of the following Formula IV:



wherein R is hydrogen or tert-butyloxycarbonyl (Boc).

[Claim 2]

The process according to Claim 1, wherein the compound of Formula III is obtained by reacting a compound of the following Formula II with di-tert-butyl dicarbonate and pyridine:



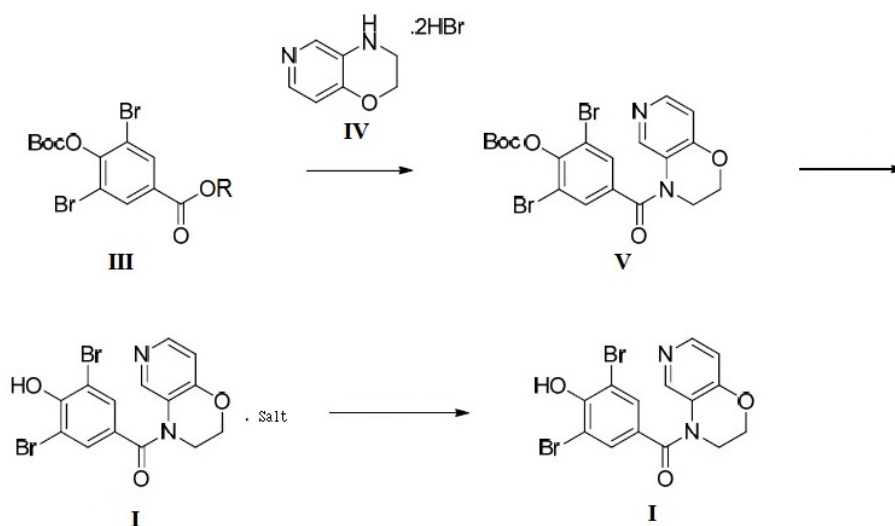
[Claim 3]

The process according to Claim 1, wherein said process comprises the following steps:

(1) reacting the compound of Formula III with the compound of Formula IV to obtain a compound of Formula V;

(2) reacting the compound of Formula V with an alcohol in the presence of an acid to obtain a salt of the compound of Formula I; and

(3) reacting the salt of the compound of Formula I with a base first and then with an acid secondarily:

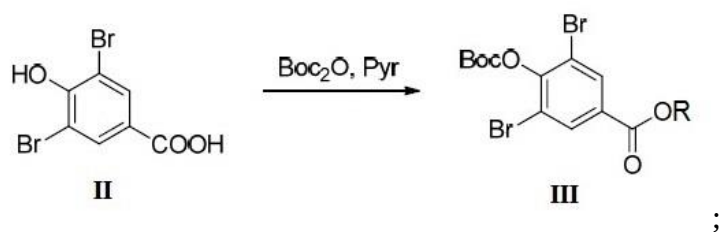


[Claim 4]

The process according to Claim 3, wherein Steps (1) and (2) are carried out as an *in situ* reaction.

[Claim 5]

The process according to Claim 3, wherein the compound of Formula III is obtained by reacting the compound of Formula II with di-tert-butyl dicarbonate and pyridine:



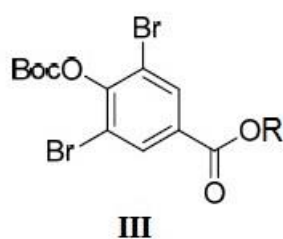
and the above step, and Steps (1) and (2) are carried out as an *in situ* reaction.

[Claim 6]

The process according to Claim 1, wherein the compound of Formula IV is obtained by reacting 3,4-dihydro-2*H*-pyrido[4,3-*b*][1,4]oxazine with bromic acid in acetic acid.

[Claim 7]

A compound of the following Formula III:



wherein R is hydrogen or Boc.

[Claim 8]

A compound of the following Formula IV:

