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1. Oral pharmaceutical compositions for use in dyslipidemias

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By Barranco Hernandez, Gustavo; Senosiain Pelaez, Juan Pablo; Garcia-Salgado Lopez, Enrique Raul; Luna Guiza, Maria del Coral
From Mex. Pat. Appl. (2014), MX 2013006332 A 20141219. | Language: Spanish, Database: CAPLUS

The invention relates to a solid oral pharmaceutical compn. contg. a statin and another antilipidemic agent, to a method for the prodn. of said compn., and to the use of said combination for producing a pharmaceutical formulation that can be used to treat metabolic syndrome, type II diabetes, or other diseases. The invention further relates to the use of a pharmaceutical combination formed by atorvastatin and fenofibrate, for producing a medicament that can be used to increase the levels of HDL2a and HDL2b, and to reduce the levels of HDL3a, HDL3b and HDL3c.

2. Simple and rapid method for preparation of atorvastatin hemi-calcium

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By Duan, Yuqiang
From Faming Zhu
Patent No. Kind Language
CN 104447487 A Chinese 150325. | Language: Chinese, Database: CAPLUS

The present invention provides a method for preparing atorvastatin hemi-calcium prepn., which consists of six steps to obtain atorvastatin hemi-calcium pure product. The special thing is that on the one hand the hydrolysis of atorvastatin ester and transformation into calcium salt are simultaneously carried out in one step in one-pot, on another aspect obtained crude atorvastatin hemi-calcium can be purified by heat refluxing in Et acetate, Pr acetate, or Bu acetate, cooling, and crystn. The present invention simplifies the process steps, shortens the reaction time, makes the reaction more thorough, c...

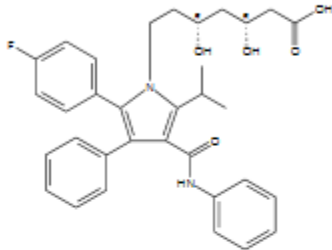
3. Preparation of atorvastatin derivatives



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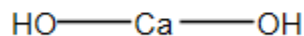
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CAS RN 134395-00-9


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(19) 中华人民共和国国家知识产权局



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(10) 申请

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(72) 发明人 段玉强 王利叶 贾玉香 罗明

(51) Int. Cl.

C07D 207/34(2006.01)

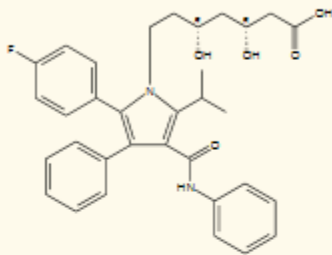


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HO—Ca—OH

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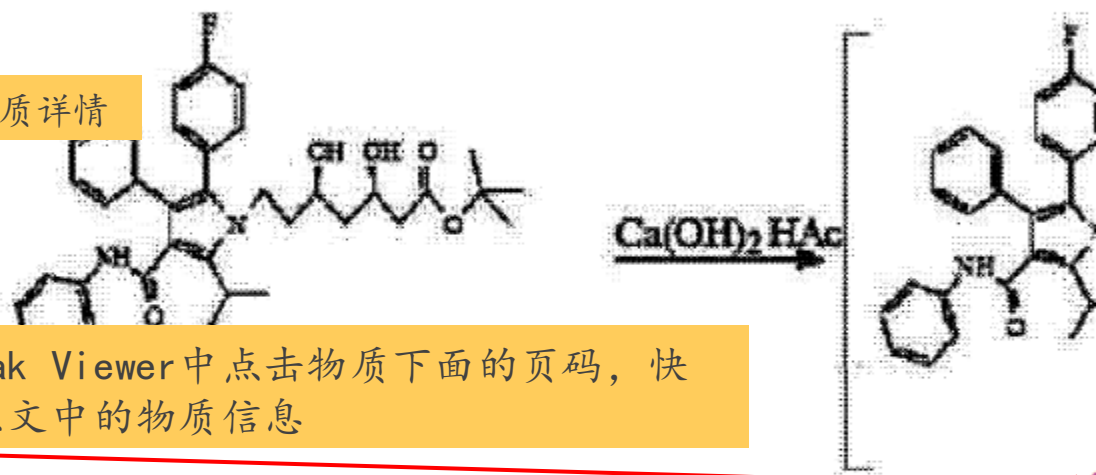
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实施例 1

向反应瓶中加入 100 ml 水、15ml 乙醇、20 g R-(R*, R*)-2-(4-基-5-(1-异丙基)-3-苯基-4-((苯胺)羰基)-1H-吡咯-1-庚及 4 g 氢氧化钙(0.10mol),将混合物搅拌并加热混合物至 45℃,然中滴加 5g (0.08mol)醋酸,用 HPLC 跟踪反应,7 小时左右反应即可达



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4 / 5

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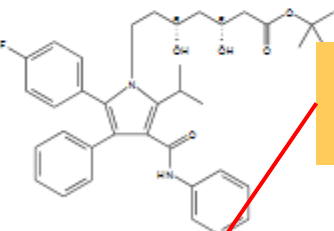
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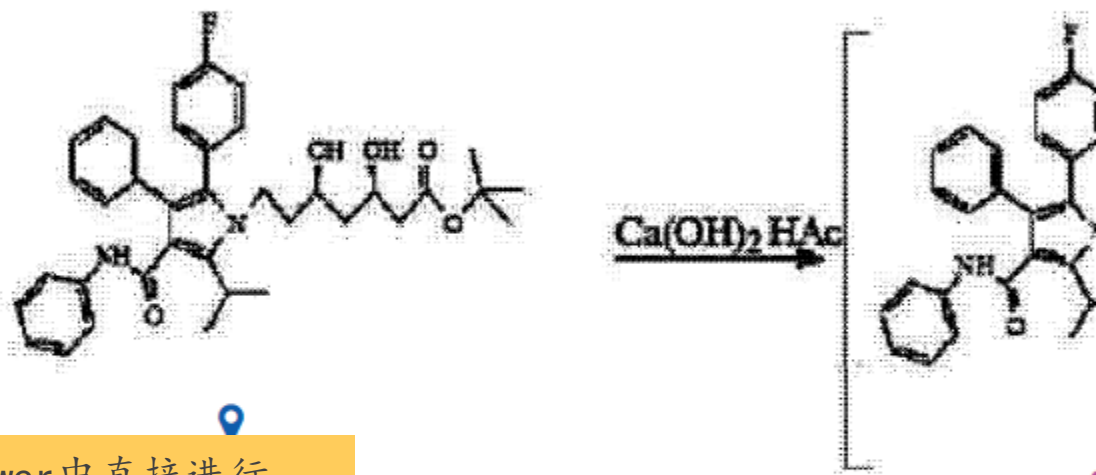
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 [0016] 本发明的合成路线如下:



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向反应瓶中加入 100 ml 水、15ml 乙醇、20 g R-(R*, R*)-2-(4-基-5-(1-异丙基)-3-苯基-4-((苯胺)羰基)-1H-吡咯-1-庚及 4 g 氢氧化钙(0.10mol),将混合物搅拌并加热混合物至 45℃,然中滴加 5g (0.08mol)醋酸,用 HPLC 跟踪反应,7 小时左右反应即可达

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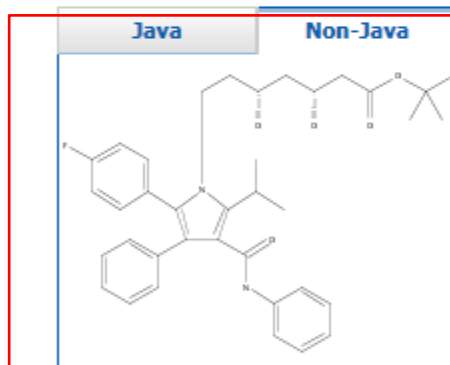
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