

पेटेंट कार्यालय  
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पेटेंट कार्यालय का एक प्रकाशन  
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(54) Title of the invention : INSILICO STUDY, SYNTHESIS AND PHARMCOLOGICAL ACTIVITY OF AMINE BASIS ?ETA, ?ETA-DIPHENYL PROPIONIC ACID DERIVATIVE

<p>(51) International classification</p> <p>(31) Priority Document No</p> <p>(32) Priority Date</p> <p>(33) Name of priority country</p> <p>(86) International Application No Filing Date</p> <p>(87) International Publication No</p> <p>(61) Patent of Addition to Application Number Filing Date</p> <p>(62) Divisional to Application Number Filing Date</p>	<p>:A61P25/08, G16C20/60, A61K31/609, C07C231/02, C07C233/02, C07C51/347, A61K9/00, A61K47/00</p> <p>:NA</p> <p>:NA</p> <p>:NA</p> <p>:</p> <p>:01/01/1900</p> <p>: NA</p> <p>:NA</p> <p>:NA</p> <p>:NA</p> <p>:NA</p> <p>:NA</p>	<p>(71)<b>Name of Applicant :</b> <b>1)Shweta Verma</b> Address of Applicant :Pharmacy Academy, Faculty of Pharmacy, IFTM University, Moradabad, Uttar Pradesh, India – 244102 Uttar Pradesh India <b>2)Aalia Naaz</b></p> <p>(72)<b>Name of Inventor :</b> <b>1)Shweta Verma</b> <b>2)Aalia Naaz</b></p>
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## (57) Abstract :

The present invention relates to the in silico study, synthesis, and pharmacological evaluation of amine-based beta, beta-diphenyl propionic acid derivatives as anticonvulsant agents. Beta, beta-diphenyl propionic acid was synthesized via Friedel–Crafts reaction of cinnamic acid with dry benzene in the presence of anhydrous aluminum chloride at 40–45 degrees Celsius, yielding 89.18 percent with a melting point of 148–150 degrees Celsius. The compound was converted to N,3,3-triphenylpropanamide (C<sub>21</sub>H<sub>19</sub>ON, mp 125–127 degrees Celsius, yield 45.18 percent) using thionyl chloride and aniline. Structural characterization was confirmed by IR (C=O at 1690 cm<sup>-1</sup>, N–H at 3500–3350 cm<sup>-1</sup>) and <sup>1</sup>H NMR spectroscopy. Pharmacological studies using maximal electroshock and strychnine-induced seizure models in albino mice showed significant anticonvulsant activity at 30 mg/kg, comparable to phenytoin sodium, indicating a promising pharmacophore.

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