

## Additions and Corrections

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**Bruce E. Maryanoff,\* Michael J. Costanzo, Samuel O. Nortey, Michael N. Greco, Richard P. Shank, James J. Schupsky, Marta P. Ortegon, and Jeffrey L. Vaught:** Structure–Activity Studies on Anti-convulsant Sugar Sulfamates Related to Topiramate. Enhanced Potency with Cyclic Sulfate Derivatives.

Page 1329. The procedure for the synthesis of **6** should read "...concentrated in vacuo to a syrup, which was dissolved in toluene and concentrated to a solid. The solid was triturated with hexanes and dried in vacuo to give **6** (1.20 g, 11%) as a colorless powder: mp 50–52 °C; ...". Chromatography was not performed. The <sup>1</sup>H NMR resonance at  $\delta$  2.40 should read " $\delta$  3.40". The molecular formula should have "C<sub>12</sub>" instead of C<sub>11</sub>; the "0.1C<sub>7</sub>H<sub>8</sub>" represents residual toluene. The corresponding information on **6** in Table 1 (p 1317) should be corrected: % yield should read "11%" and "LC" should be deleted.

Page 1330. To the procedure for the synthesis of **8**, "and concentrated to a solid" should be added to the end of the phrase "... treated with acetic acid (3.32 g), stirred for 5 min,".

Page 1331. In the synthesis of **32**, the 10.0 g of intermediate **39** represents "0.029 mol".

Page 1332. The <sup>1</sup>H NMR resonance "3.75–4.05 (m, 2H, H<sub>6</sub>)" for **32** should be deleted, as it is redundant.

Page 1333. The procedure for the synthesis of **47** is missing some operations. It should read "... at 23 °C for 5 days and poured into ice–water (225 mL). The organic phase was separated, and the aqueous phase was washed with chloroform (3 × 75 mL). The combined organic solution was washed sequentially with NaHCO<sub>3</sub> (saturated aqueous) and brine, and then it was dried (MgSO<sub>4</sub>) and concentrated in vacuo. The residue was dissolved in methanol (180 mL), treated with NaHCO<sub>3</sub> (20.2 g, 0.24 mmol), and stirred for 16 h. After filtration, the solution was concentrated in vacuo to a yellow solid, which was partitioned between chloroform and water. The chloroform solution was washed with brine, dried (MgSO<sub>4</sub>), and concentrated in vacuo. The residue was purified..."

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