



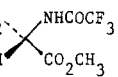

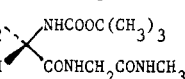


ERRATUM

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In the article "N<sup>6</sup>-Functionalized Congeners of Adenosine with High Potency at A<sub>2</sub>-Adenosine Receptors: Potential Ligands for Affinity Chromatography" by Kenneth A. Jacobson, Noboyiuki Yamada, Kenneth L. Kirk, John W. Daly, and R. A. Olsson, pages 1097—1102:

Page 1100, Table 1, the values for compounds 13-19 in the last three columns are incorrect. For the readers' convenience, the corrected Table 1 is reproduced on the following page.

Table 1. Potency of adenosine analogues at a coronary A<sub>2</sub>-adenosine receptor and at a brain A<sub>1</sub>-receptor

Analogue	A <sub>2</sub> Receptor		A <sub>1</sub> Receptor	EC <sub>50</sub> (A <sub>2</sub> )
	MPR <sup>a</sup>	Estimated Potency <sup>b</sup>	K <sub>i</sub> (nM) <sup>c</sup>	K <sub>i</sub> (A <sub>1</sub> )
	to Ado	EC <sub>50</sub> (nM)		
<u>1</u> NECA	150	8	5.1	1.6
<u>2</u> 2-chloroadenosine	27	44	6.7	6.6
<u>3</u> N <sup>6</sup> -methyladenosine	0.05	24,000	60	400
<u>4</u> R-PIA	4.3	280	1.2	230
<u>5</u> CHA	1.6	750	0.85	880
<u>6</u> N <sup>6</sup> -phenyladenosine	1.4	860	3.3	260
<u>7</u> N <sup>6</sup> -p-tolyladenosine	1.35	890	2.5	360
<u>8</u> OH		[27% @ 21μM] <sup>d</sup>	210	-
<u>9</u> NHCH <sub>3</sub>	4.9±1.3	240	16	15
<u>10</u> NH- 	2.1±1.3	570	1.7	340
<u>11</u> NH-  -CH <sub>2</sub> - 	3.0±0.7	400	18	22
<u>12</u> NH-  -CH <sub>2</sub> - 	4.4±0.4	270	13	21
R = NH-  -CH <sub>2</sub> -COR'				
<u>13</u> OCH <sub>3</sub>	3.2±1.6	380	2.5	150
<u>14</u> NHCH <sub>3</sub>	7.2±1.3	170	6.7	25
<u>15</u> NHNH <sub>2</sub>	3.5±0.8	340	4.5	80
<u>16</u> NH(CH <sub>2</sub> ) <sub>2</sub> NH <sub>2</sub>	7.8±2.5	150	0.85	180
<u>17</u> NH(CH <sub>2</sub> ) <sub>2</sub> NHCO(CH <sub>2</sub> ) <sub>2</sub> - 	2.8±0.1	430	4.5	96
<u>18</u> NH(CH <sub>2</sub> ) <sub>2</sub> NH-CO-biotin	3.0±0.2	400	11.4	35
<u>19</u> NH(CH <sub>2</sub> ) <sub>2</sub> NHCO(CH <sub>2</sub> ) <sub>5</sub> NH-CO-biotin	10.2±7.3	120	18	6.7

<sup>a</sup>Molar potency ratio relative to adenosine, which is set equal to 1.0.

<sup>b</sup>Estimated IC<sub>50</sub> values based on potency of adenosine (MPR = 1.0) of 1,200±150 nM (6).

<sup>c</sup>K<sub>i</sub> values for antagonism of binding of 1 nM [<sup>3</sup>H]N<sup>6</sup>-cyclohexyladenosine to rat cerebral cortical membranes (data from 3, 4, 12).

<sup>d</sup>Highest concentration of analogue did not raise coronary blood flow to a level x 50% of maximum possible increase. In such a case we report % increase in flow over control at the plasma nucleoside concentration achieved during infusion.