

Corrigendum

Corrigendum to “2,6,9-Trisubstituted Purines: Optimization Towards Highly Potent and Selective CDK1 Inhibitors”[†] (*Bioorg. Med. Chem. Lett.* 1999, 9, 91–96)

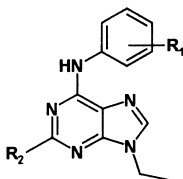
Patricia Imbach,^{a,*} Hans-Georg Capraro,^a Pascal Furet,^{a,b} Helmut Mett,^a
Thomas Meyer^a and Jürg Zimmermann^a

^aOncology Research, Novartis Pharmaceuticals, Therapeutic Area Oncology, Novartis Limited, CH-4002 Basel, Switzerland

^bMolecular Modelling, Novartis Pharmaceuticals, Therapeutic Area Oncology, Novartis Limited, CH-4002 Basel, Switzerland

On page 93, Table 1, R₂ is not correct for some of the examples (5–11). The correct text should read:

Table 1. Optimization of 2,6,9-trisubstituted purine derivatives



Compd no.	R ₁	R ₂	CDK1 ^h IC ₅₀ (μM)	T24 ⁱ IC ₅₀ (μM)	FAB-MS (M + H) ⁺	mp (°C)
5	3-Cl	-NHCH ₂ CH ₂ OH	0.33	17.5	333	99–102
6	3-Cl	-NHCH ₂ CH ₂ NH ₂	0.08	3.16	332	103
7 ^a	4-F	-NHCH ₂ CH ₂ NH ₂	0.45	8.3	316	>250
8 ^a	3-CN	-NHCH ₂ CH ₂ NH ₂	0.56	5.24	323	>175 dec.
9	3-OCH ₃	-NHCH ₂ CH ₂ NH ₂	0.55	9.9	328	Oil
10 ^a	3-CF ₃	-NHCH ₂ CH ₂ NH ₂	0.7	6.08	366	>250 dec.
11 ^a	3-F	-NHCH ₂ CH ₂ NH ₂	0.60	6.5	316	>250 dec.

*Corresponding author. Tel.: +4161-696-6256; fax: +4161-696-6246; e-mail: patricia.imbach@pharma.novartis.com

[†]PII of original article: S0960-894X(98)00691-X.