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Erratum

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ERRATUM

The publishers would like to apologise for an error that occurred in a recent issue of the *Journal of Enzyme Inhibition and Medicinal Chemistry*, 2009; 24(4): 972–985.

Synthesis of imidazole-containing analogues of farnesyl pyrophosphate and evaluation of their biological activity on protein farnesyltransferase

Laëtitia Coudray, Renata Marcia de Figueiredo, Stéphanie Duez, Sylvie Cortial, and Joëlle Dubois *Scheme 3 & 4 should have been displayed as they are below:*

Scheme 3. Second synthetic pathway. a) nBuLi, THF, $-78^{\circ}C$, 45 min then addition of 4a-c, $-78^{\circ}C$, 20 min to 2h30 to RT 1h (68-100%); b) MnO $_2$, THF, 0°C, 2h (80-100%); c) 9, THF, Zn, ultrasounds, $40^{\circ}C$, 5h (51-79%); d) POCl $_3$, Pyridine, $0^{\circ}C$ to RT 14h (75-85%); e) for a (R= SEM, n= 0) TFA, CH_2Cl_2 , RT, 5h, for b (R= CH_3 , n= 0) HCO $_2$ H RT, 19h and for c (R= CH_3 , n= 1) and d (R= CH_3 , n= 2) SiO_2 , toluene, reflux, 14h; f) Mg, MeOH, RT, 3h (62-80%); g) LDA, THF, $-78^{\circ}C$, 35 min then addition of 9, $-78^{\circ}C$, 4h (42-72%).

Scheme 4. Synthesis of farnesyl acids. a) NaH and 20, THF, 0° C, 10 min then 30 min RT, then addition of 4c, RT, 4h30 (74%); b) Mg, MeOH, RT, 4h (38%); c) CuI, LDA, THF, 2h, -110° C, then addition of 23, -110° C, 2h (69%); d) NaH and 25, THF, 0° C, 10 min then RT, 40 min, then addition of 4c, RT, 2h15 (66%); e) LDA, THF, -78 °C, 35 min then addition of 27, -78°C; f) NaOH 2M, EtOH, 70 °C, 15h (77-100%).