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Corrigendum

Corrigendum to “Synthesis of new fused and substituted benzo and pyrido carbazoles via C-2 (het)arylindoles” [Tetrahedron 64(49) (2008) 11012–11019]

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
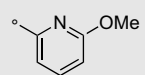
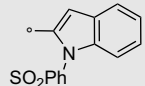


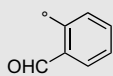
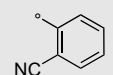
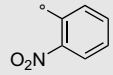
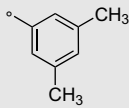
The authors regret that an error occurred in Table 1. The correct Table 1 is shown on the next page. The authors apologise for any inconvenience caused.

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Table 1
Experimental conditions for Stille reactions using **3**

Entry	(Het)ArX	Equiv	Conditions	Time	(Het)Ar	Yield ^a	
1	a	2-Bromopyridine	Pd(PPh ₃) ₄ (10%), CuI (20%), THF, reflux Pd(PPh ₃) ₄ (10%), CuI (20%), DMA, reflux Pd(PPh ₃) ₄ (10%), CuI (20%), DMA, 100 °C	3 days		4 (65%)	
	b			5 h		4 (65%)	
	c			6 h		4 (quant.)	
2	2-Bromo-6-methoxy-pyridine	3.0	Pd(PPh ₃) ₄ (10%), CuI (20%), DMA, 100 °C	14 h		5 (70%)	
3	2-Bromo-pyridine-N-oxide	3.0	Idem	12 h		6 (46%)	
4	2-Chloro-6-(O-MOM)pyridine ^b	3.0	Idem	24 h		6 (46%)	
5	a	Bromobenzene	10.0	Idem		7 (quant.)	
	b	Iodobenzene	3.0			4 h	7 (78%)
6	a	1-Bromo-2-fluorobenzene	3.0	Idem		8 (25%)	
	b	1-Bromo-2-fluorobenzene	10.0			16 h	8 (75%)
	c	1-Fluoro-2-iodobenzene	3.0			6 h	8 (82%)
7	a	2-Bromobenzaldehyde	3.0	Idem		9 (45%)	
	b					16 h	9 (89%)
8	2-Bromobenzonitrile	3.0	Idem	16 h		10 (84%)	
9	2-Bromonitrobenzene	3.0	Idem	16 h		11 (88%)	
10	1-Bromo-3,5-dimethylbenzene	3.0	Idem	12 h		12 (78%) ^c	

^a Yields are given for isolated products, which were fully characterized by IR, MS, ¹H and ¹³C NMR.

^b This compound was selectively obtained from 6-chloro-1*H*-pyridin-2-one and MOMCl (1.1 equiv) in THF at room temperature in presence of NaH (1.5 equiv) during 2 h in 72% yield.

^c A small amount of **12** was lost during the purification step.