## 1-(Phenanthrolo[5,6-d]imidazol-2-yl)-3-(acenaphthro[4,5-d]imidazol-2-yl)benzene

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2-(3-formylphenyl)imidazo[4,5-f][1,10]phenanthroline was prepared by a previously published method [1]. A mixture of 2-(3-formylphenyl)imidazo[4,5-f][1,10]phenanthroline ( $0.2 \mathrm{~g}, 0.62 \mathrm{mmol}$ ), acenaphthenequinone ( $0.124 \mathrm{~g}, 0.68 \mathrm{mmol}$ ), ammonium acetate ( $0.713 \mathrm{~g}, 9.3 \mathrm{mmol}$ ) and glacial acetic acid ( $10 \mathrm{~cm}^{3}$ ) was refluxed for about 3 h . The cooled solution was filtered and then poured into $100 \mathrm{~cm}^{3}$ ether. The crude product was collected and purified on alumina with ethanol-toluene ( $1: 4 \mathrm{v} / \mathrm{v}$ ) as eluent to give the title compound as a yellow powder. ( 0.159 g , yield: 52.7\%).
${ }^{1}{ }^{H}$ NMR ( $500 \mathrm{MHz}, d_{6}$-DMSO): 13.84 (br, 2H), 9.24 (s, 1H), 8.93 (d, 2H, $J=8$ ), 8.91 (d, 2H, $J=7.5$ ), 8.69 (d, 1H, $J=8$ ), $8.39(\mathrm{~d}, 1 \mathrm{H}, J=8), 8.21(\mathrm{~d}, 1 \mathrm{H}, J=8), 7.85(\mathrm{~m}, 2 \mathrm{H}), 7.78(\mathrm{dd}, 2 \mathrm{H}, J=5), 7.73(\mathrm{dd}, 2 \mathrm{H}, J=4)$, 7.68 (t, 1H, $J=8$ ), $7.60(\mathrm{t}, 1 \mathrm{H}, \mathrm{J}=7$ ).
${ }^{13}$ C NMR (125 MHz, $d_{6}$-DMSO): 150.2, 147.9, 143.7, 136.8, 135.7, 131.7, 131.2, 130.9, 129.9, 129.5, 129.1, 128.8, 128.1, 127.8, 126.6, 125.9, 125.4, 123.6, 123.0, 122.8, 119.5.

IR (KBr. $\mathrm{cm}^{-1}$ ): 3392s, 1603s, 1593m, 1542m, 1474m, 1438m, 1402m, 1278s, 1202m, 1073m, 1015m, 806 m , 777s, 739s, 678w, 624w,

UV-Vis (l, nm, in ethanol): 227, 290, 319.
FAB-MS ([M+1] $\left.{ }^{+}\right): 487$.
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## References and Notes

1. Chao, H.; Ye, B.-H.; Li, H.; Li, R.-H.; Zhou, J.-Y.; Ji, L.-N. Polyhedron 2000, 19, 1975.



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