Supporting Information

Efficient synthesis of new indenopyridotriazine [4.3.3]propellanes and spiroindenopyridotriazine-4*H*-pyran derivatives

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Experimental Section

General remarks:

All commercially available reagents and other solvents were purchased from Aldrich and Merck Chemical Co. and used without further purification. The NMR spectra were recorded with a Bruker DRX-300 AVANCE instrument (300 MHz for ¹H and 75.4 MHz for ¹³C) with DMSO- d_6 as solvent. Chemical shifts are given in ppm (*d*) relative to the internal TMS, and the coupling constant (*J*) reported in hertz (Hz). Melting points were measured with an electrothermal 9100 apparatus. Mass spectra were recorded with an Agilent 5975C VL MSD with a Triple-Axis detector operating at an ionization potential of 70 eV. IR spectra were measured with a Bruker Tensor 27 spectrometer.



Figure 1. Structure of all products 6a-l



Figure 2. Structure of all products 8a-d

The structures all of the products **6a-l** and **8a-d** were deduced from their IR, ¹H NMR, ¹³C NMR and Mass spectra (see the Supporting Information).



¹H NMR of 3a



¹³C NMR of 3a



IR of 3a





¹H NMR of 6a (D₂O exchangeable)





File :C:\MSDCHEM\3\DATA\Snapshot\140110151.D Operator : Acquired : 2 Jan 2007 00:54 using AcqMethod test.M Instrument : MSD Sample Name: T8 Misc Info : Vial Number: 1



MS of 6a



¹H NMR of 6b



¹³C NMR of 6b



IR of 6b



MS of 6b





¹H NMR of 6c (D₂O exchangeable)







IR of 6c



MS of 6c







IR of 6d







IR of 6e



MS of 6e







IR of 6f



MS of 6f







IR of 6g



MS of 6g



¹H NMR of 6h













IR of 6i



¹H NMR of 6j



¹³C NMR of 6j



¹³C NMR of 6k

IR of 6k

¹³C NMR of 6l

¹H NMR of 8a

¹H NMR of 8a (D₂O exchangeable)

IR of 8a

MS of 8a

¹H NMR of 8b

IR of 8b

MS of 8b

¹³C NMR of 8c

IR of 8c

