

Table of contents

Table of contents.....	1
1. General remarks.....	2
2. ^1H and ^{13}C NMR spectra.....	3
3. Crystal packing for compound 9	7

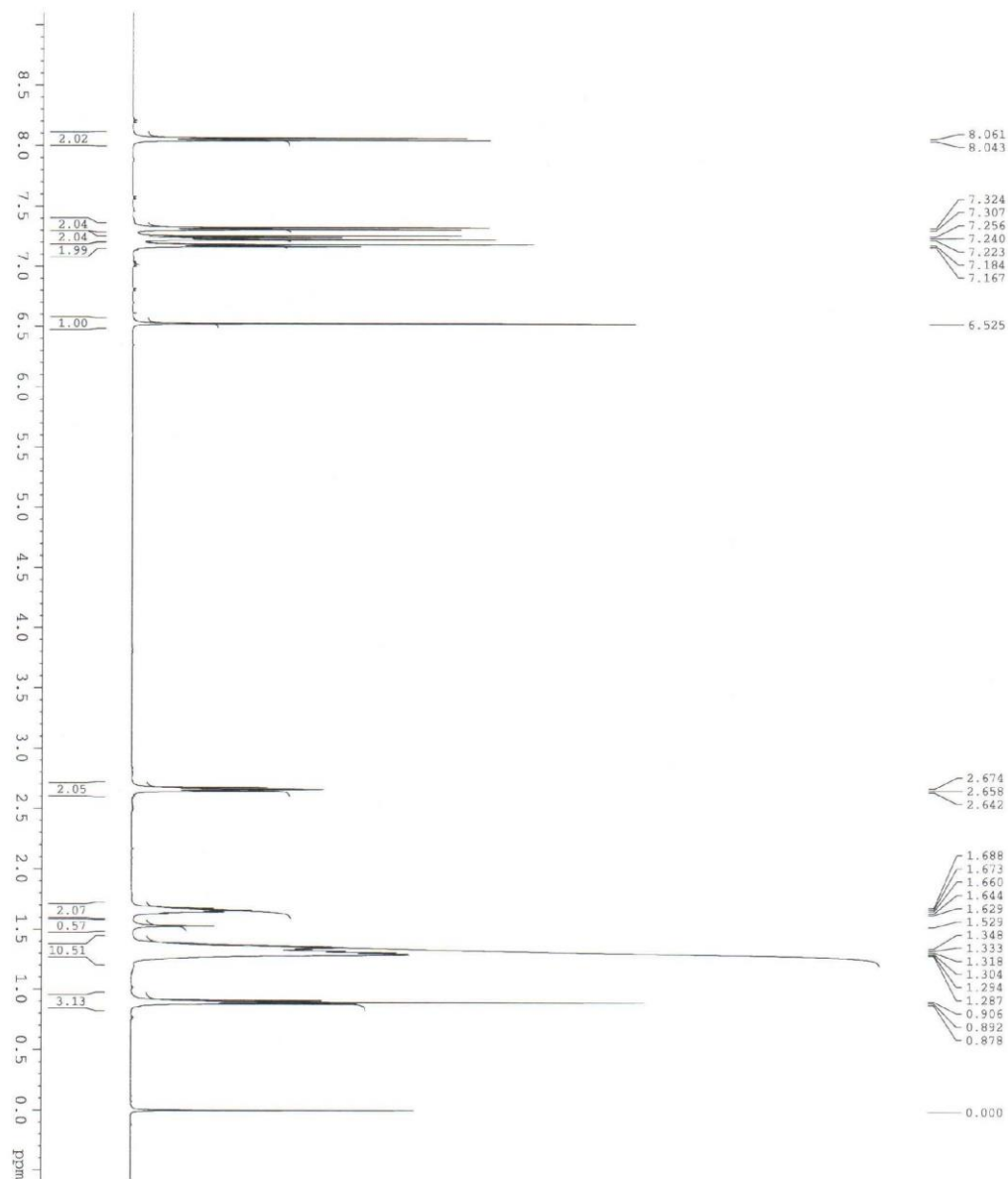
1. General remarks.

All reagents and solvents were purchased from commercial sources and were used as received unless otherwise noted. Reagent grade solvents (CH_2Cl_2 , hexanes) were distilled prior to use. DMF was dried over magnesium sulfate, then distilled and stored under argon. Transformations with moisture and oxygen sensitive compounds were performed under a stream of argon. The reaction progress was monitored by means of thin layer chromatography (TLC), which was performed on aluminium foil plates, covered with Silica gel 60 F₂₅₄ (Merck) or Aluminium oxide 60 F₂₅₄ (neutral, Merck). Products purification was done by means of column chromatography with Kieselgel 60 (Merck) or Aluminium oxide (Fluka). Occasionally, dry column vacuum chromatography (DCVC) for purification of products obtained was performed using Silica gel Type D 5F. The identity and purity of prepared compounds were proved by ¹H NMR and ¹³C NMR spectrometry as well as by MS-spectrometry (*via* EI-MS or ESI-MS). NMR spectra were measured on Bruker AM 500 MHz, Bruker AM 600 MHz, Varian 600 MHz, Varian 400 MHz or Varian 200 MHz instruments with TMS as internal standard. All chemical shifts are given in ppm. All melting points for crystalline products were measured with automated melting point apparatus EZ-MELT and were given without correction. The absorbance and fluorescence spectra were measured in dichloromethane on Perkin – Elmer Lambda 25 UV/VIS and Hitachi F-7000 respectively.

Linear optical measurements

Steady-state fluorescence measurements were performed with dilute solutions (10^{-6}M , optical density < 0.1) contained in standard 1 cm quartz cuvettes at room temperature. Compounds were dissolved in dichloromethane unless otherwise noted. Emission spectra were obtained, for each compound, under excitation at $\lambda=350$ nm. Fluorescence quantum yields were measured by using quinine hemisulphate monohydrate in 0.5M sulfuric acid as a standard.

2. ^1H and ^{13}C NMR spectra



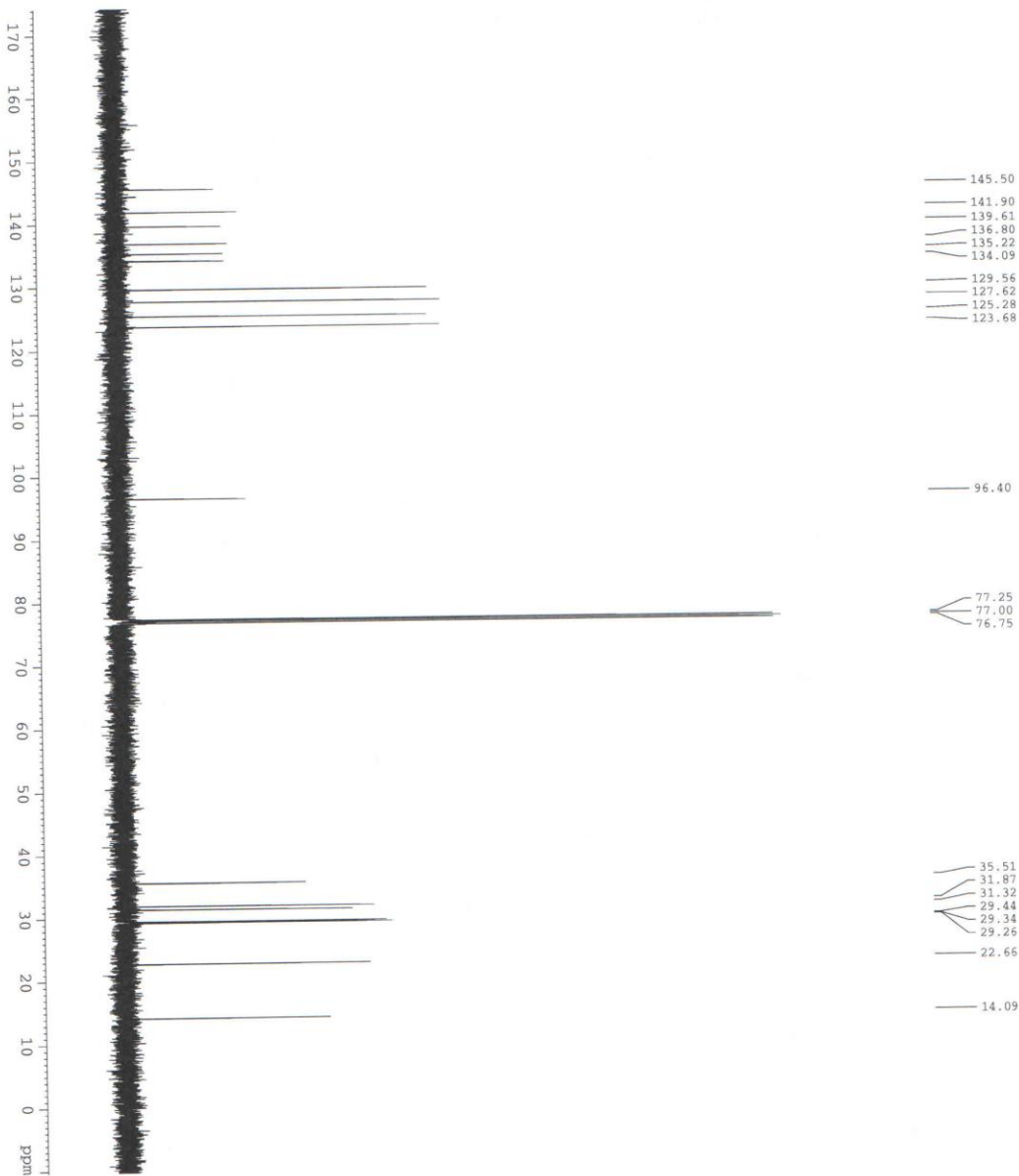
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 SOLVENT CDCl3
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 FIDRES 0.157632 Hz
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 RG 114
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 DE 6.78 usec
 TE 303.0 K
 D1 1.00000000 sec
 T10 1

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F2 - Processing parameters
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^1H NMR spectrum of compound 6



Current Data Parameters
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PROCNO 1

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NS 750
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FIDRES 0.498653 Hz
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TE 303.0 K
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d11 0.03000000 sec
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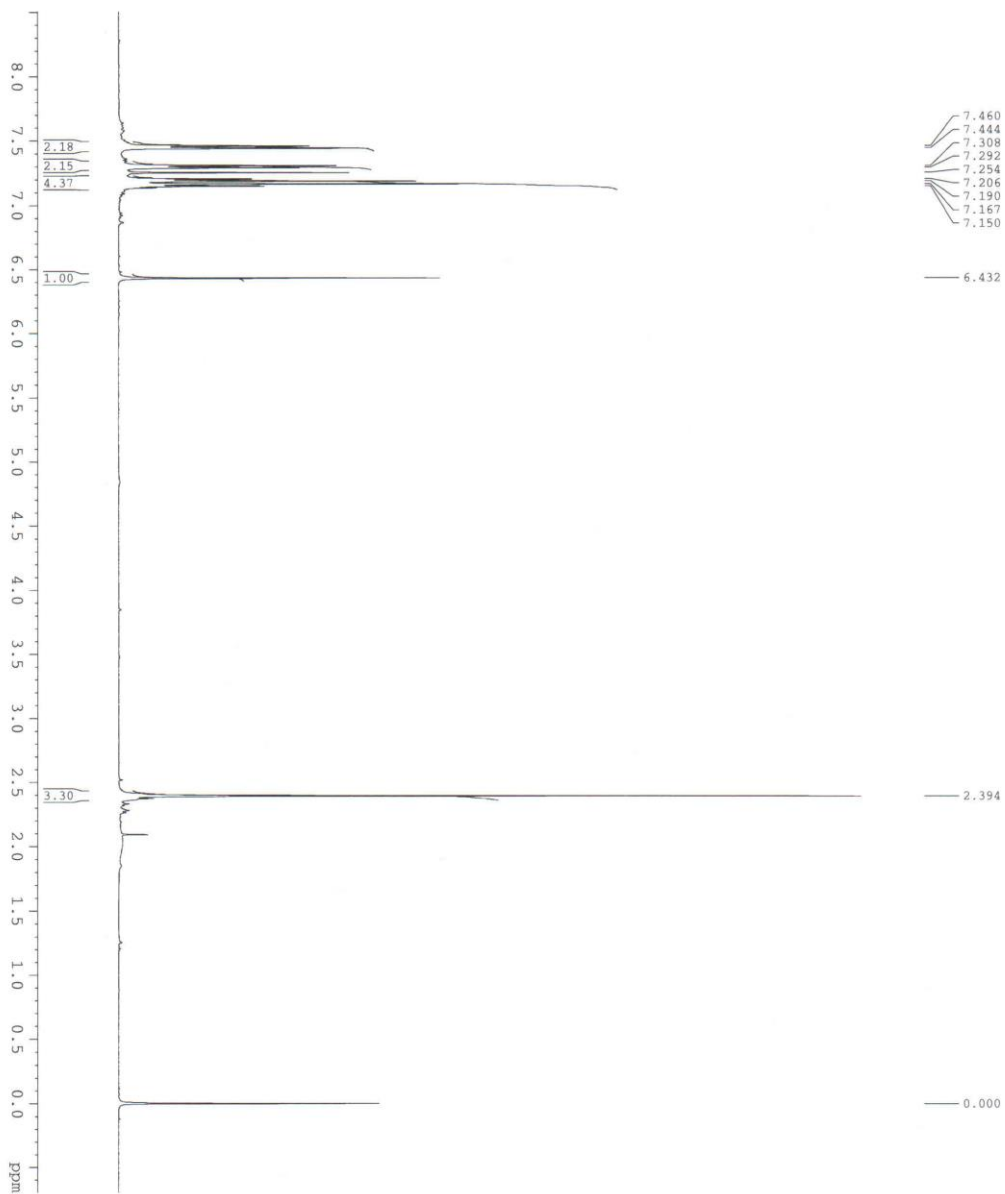
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F1 - Acquisition Parameters
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TD 128
SFO1 500.132 MHz
FIDRES 7.8121414 Hz
SFO 1.999 ppm
P1MODE OP

F2 - Processing Parameters
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LB 0.50 Hz
GB 0
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F1 - Processing Parameters
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LB 0.30 Hz
GB 0.1

¹³C NMR spectrum of compound 6



Current Data Parameters
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 PROCNO 1

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 FIDRES 0.157633 Hz
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 TE 303.0 K
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F2 - Processing parameters
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 LB 0.00 Hz
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 PC 1.00

¹H NMR spectrum of compound 7

137.06
136.97
136.13
135.01
132.68
130.01
127.91
127.80
127.65
125.21
125.13
125.10
125.07

95.36

77.25
77.00
76.75

21.04

-0.02



Current Data Parameters
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PROCNO 1

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Time 11.14

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PULPROG zgpg30

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NS 1550

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RG 15.300 usec

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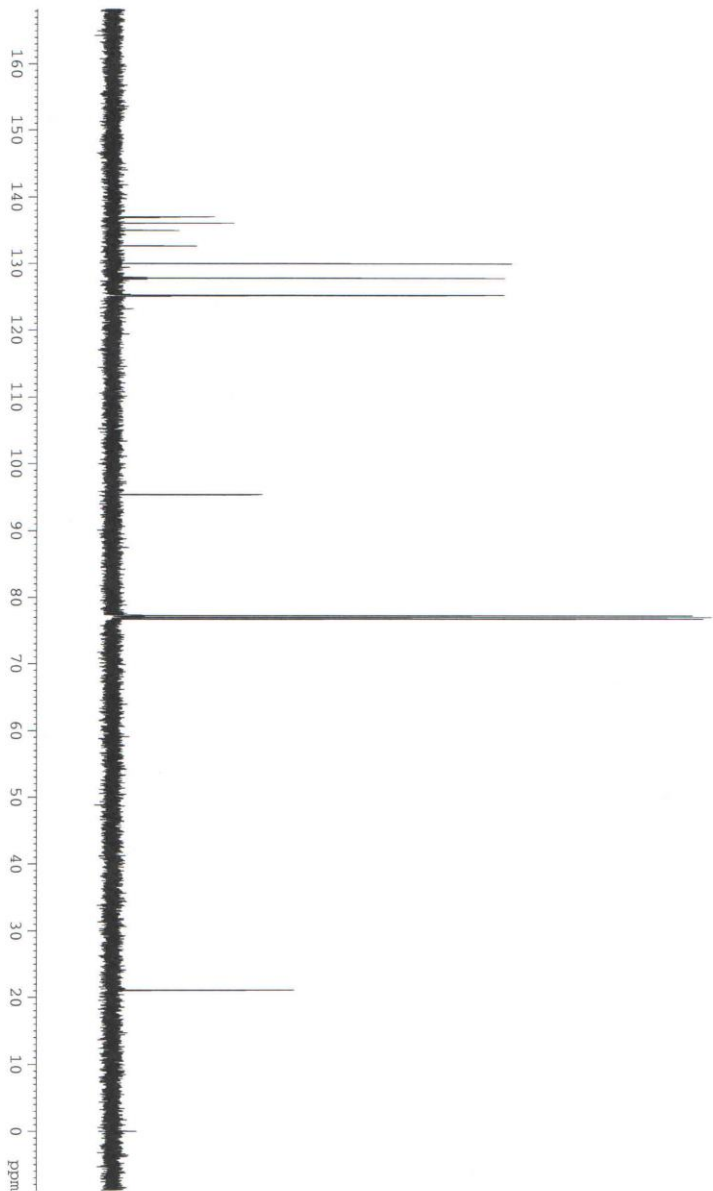
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===== CHANNEL F1 =====
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PL1 -3.00 dB
SFO1 125.7703643 MHz

===== CHANNEL F2 =====
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NUC2 1H
PCPD2 98.00 usec
PL2 2.00 dB
PL3 32.00 dB
SFO2 500.1320005 MHz

F1 - Acquisition parameters
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SFO1 500.132 MHz
FIDRES 7.812500 Hz
SW 1.999 ppm
PULPROG OP

F2 - Processing parameters
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SF 500.1300000 MHz
WDW SINE
SSB 0
LB 0.30 Hz
GB 0.1



¹³C NMR spectrum of compound 7

3. Crystal packing for compound 9.

