# New acetylacetone-polymer modified nanoparticles as magnetically separable complexing agents. 

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## Supporting information

## 4-vinylbenzyliodide

Sodium iodide ( $534 \mathrm{mg}, 3.56 \mathrm{mmol}$ ) was placed in Schlenk flask under argon atmosphere. Subsequently 4 ml of butan-2-one and $0,1 \mathrm{ml}(0.71 \mathrm{mmol})$ of 4 -vinylbenzylchloride were added. The reaction mixture was stirred at inert atmosphere for 4 days in room temperature with absence of sunlight. After completion of the reaction solvent was evaporated and then diethyl ether was added ( 15 ml ). Afterwards mixture was washed 3 times with water. Organic layer were then dried under anhydrous Na 2 SO 4 and filtrated. Filtrate was evaporated and the product as yellow oil was obtained (yield 85\%).
${ }^{1} \mathrm{HNMR}\left(400 \mathrm{MHz}, \mathrm{CD}_{3} \mathrm{Cl}, \delta, \mathrm{ppm}\right): 7.35(\mathrm{~s}, 4 \mathrm{H}), 6.70\left(\mathrm{dd}, 1 \mathrm{H}, \mathrm{J}_{1}=17.57 \mathrm{~Hz}, \mathrm{~J}_{2}=10.85 \mathrm{~Hz}\right), 5.77(\mathrm{~d}, 1 \mathrm{H}$, $\mathrm{J}=17.60 \mathrm{~Hz}), 5.27(\mathrm{~d}, 1 \mathrm{H}, \mathrm{J}=10.92 \mathrm{~Hz}), 4.48(\mathrm{~s}, 2 \mathrm{H}) ;{ }^{13} \mathrm{CNMR}\left(100 \mathrm{MHz}, \mathrm{CD}_{3} \mathrm{Cl}, \delta, \mathrm{ppm}\right): 138.8,137.2$, 136.3, 129.0, 126.7, 114.5, 5.86; FT-IR (ATR, v) cm²: 3083, 3020, 2921, 2848, 1682, 1508, 1406, 1153, 1081, 987, 840, 720, 580.

## IR Spectra:






MNP10

## UV spectra (normalized to 1)





## Luminescence spectra:



## Normalised to 1



SEM/EDX:



${ }^{1} \mathrm{H}$ and ${ }^{13} \mathrm{C}$ NMR
Monomer 1



Compound 2a


4-vinylbenzyliodide



Monomer 2


Monomer 3




## TEM and SEM photographs

MNP 5


MNP 11


MNP5


TGA, DTG and DSC curves
Determination of Tg temperature for polymer:






