Supporting Information

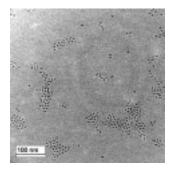


Figure 1. A TEM micrograph of nanocrystalline MnP synthesized using $Mn_2(CO)_{10}$ and TOP (trioctylphosphine) as the phosphide source. The synthesis was carried out at 280 °C with dodecylamine as the capping agent.

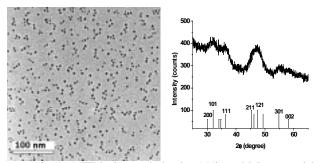


Figure 2. (A) A TEM micrograph showing 5.1(3) nm MnP nanoparticles synthesized at 220 ° C using $Mn_2(CO)_{10}$ and MA (Mn:MA = 1:0.4 mol) as capping agent at X40 K. (B) The corresponding powder X-ray diffraction pattern of MnP nanoparticles (the line diagram is for JCDPS # 07-0384).

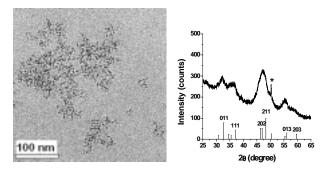


Figure 3. (A) A TEM micrograph showing 3.2(3) nm FeP nanoparticles synthesized using Fe(CO)₅ and DA as the capping agent at X40 K. (B) The corresponding powder X-ray diffraction pattern of FeP nanoparticles (the line diagram is for JCDPS # 39-0809). The (*) denotes a background peak from the sample holder.

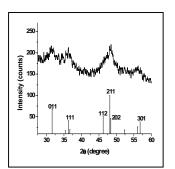


Figure 4. Powder X-ray diffraction pattern of CoP nanoparticles (the line diagram is for JCPDS # 29-0497).

Supporting Information

Table 1. Summary of magnetic data for MnP nanoparticle samples.

Size (nm)	T _b (K) 500 Oe	H _c (Oe) 5 K	M _s * (emu/mol) 5K	M _r (emu/mol) 5K
5.1(5)	60.8	6000	3420	890
6.7(3)	74.3	4860	7620	930

*Obtained from zero extrapolation of the plot of M vs. 1/H.