

SUPPLEMENTARY INFORMATION

Preparation of amphiphilic poly(divinylbenzene-co-N-vinylpyrrolidone)-functionalized polydopamine magnetic nanoadsorbents for enrichment of synthetic cannabinoids in wastewater

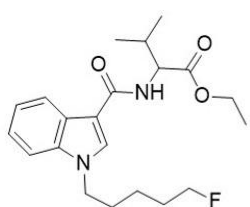
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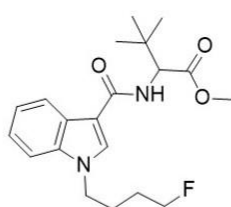
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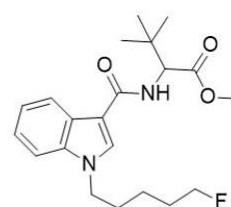
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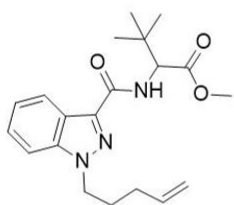
5F-EMB-PICA



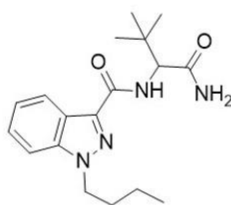
4F-MDMB-BUTICA



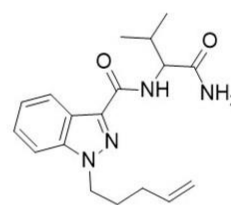
5F-MDMB-PICA



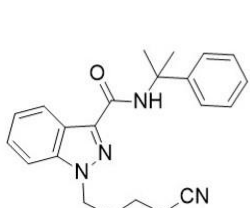
MDMB-4en-PINACA



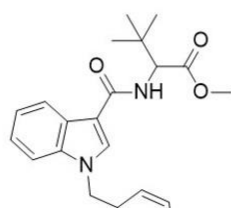
ADB-BUTINACA



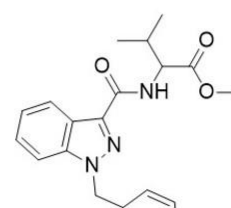
ADB-4en-PINACA



4CN-CUMYL-BUTINACA



MDMB-FUBICA



AMB-FUBINACA

Fig. S1. The structures of nine synthetic cannabinoids.

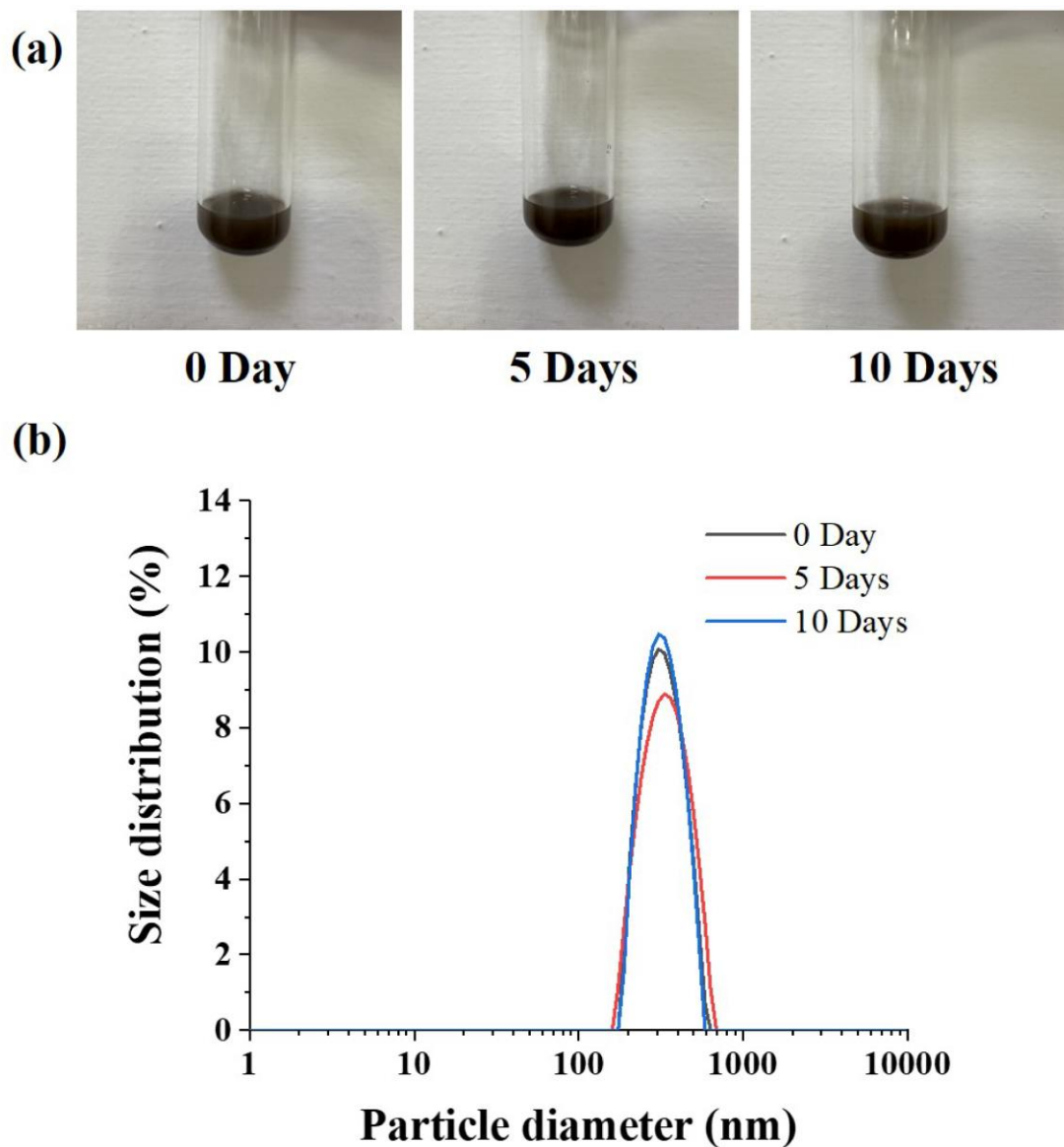


Fig. S2. (a) Digital photographs of $\text{Fe}_3\text{O}_4@\text{PDA}@\text{poly}(\text{DVB-co-NVP})$ MNPs dispersion in ethanol for ten days. (b) Dynamic light scattering (DLS) of $\text{Fe}_3\text{O}_4@\text{PDA}@\text{poly}(\text{DVB-co-NVP})$ MNPs in ethanol for ten days.

The mean hydrodynamic diameters for $\text{Fe}_3\text{O}_4@\text{PDA}@\text{poly}(\text{DVB-co-NVP})$ MNPs dispersed in ethanol were 339 nm on day 0, 354 nm on day 5 and 329 nm on day 10, respectively. As the particle diameter and visual appearance exhibited negligible difference within ten days, the synthesized nanoadsorbents are supposed to be stable during storage.

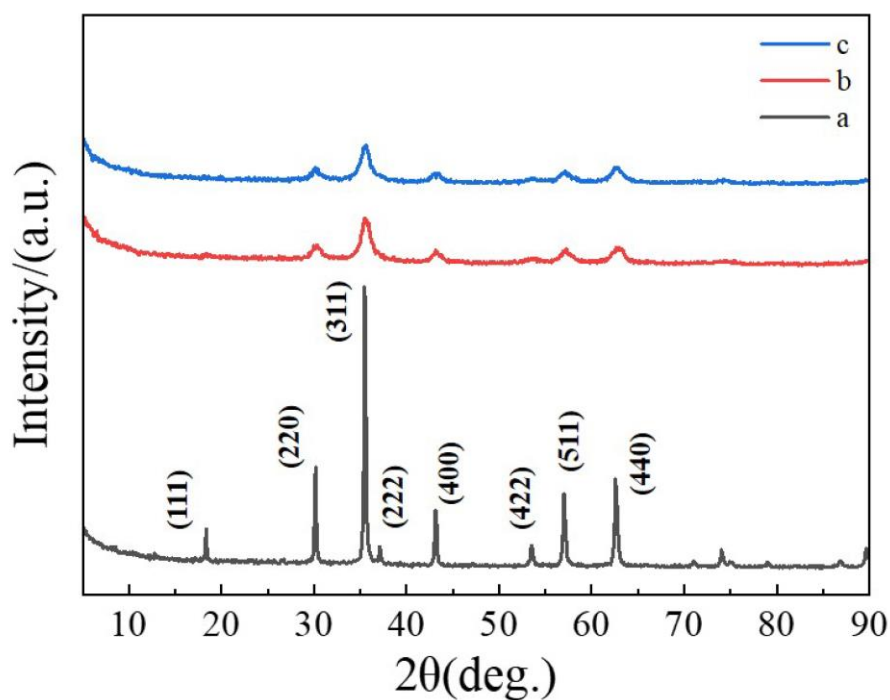


Fig. S3. XRD patterns of (a) Fe_3O_4 MNPs, (b) $\text{Fe}_3\text{O}_4@PDA$ MNPs and (c) $\text{Fe}_3\text{O}_4@PDA@poly(DVB-co-NVP)$ MNPs.

The XRD patterns of Fe_3O_4 , $\text{Fe}_3\text{O}_4@PDA$ and $\text{Fe}_3\text{O}_4@PDA@poly(DVB-co-NVP)$ MNPs are shown in Fig. S3. Common peaks were observed at 18.1° , 30.2° , 35.6° , 36.2° , 43.4° , 53.8° , 57.2° , and 62.8° , which were assigned to (111), (220), (311), (222), (400), (422), (511), and (440) planes, confirming the cubic crystallinity of the magnetite Fe_3O_4 nanoparticle core compared with standard XRD peaks (JCPDS No.19-0629)[1]. Surface modifications with PDA did not result in phase change of the Fe_3O_4 particles[2,3]. However, the XRD peaks of $\text{Fe}_3\text{O}_4@PDA@poly(DVB-co-NVP)$ were broader and lower compared with those of $\text{Fe}_3\text{O}_4@PDA$. This phenomenon is likely because the additional poly(DVB-co-NVP) coating on the surface reduce and cover up the intensities of the corresponding peaks[4]. It can be concluded that the magnetic particles were successfully fabricated.

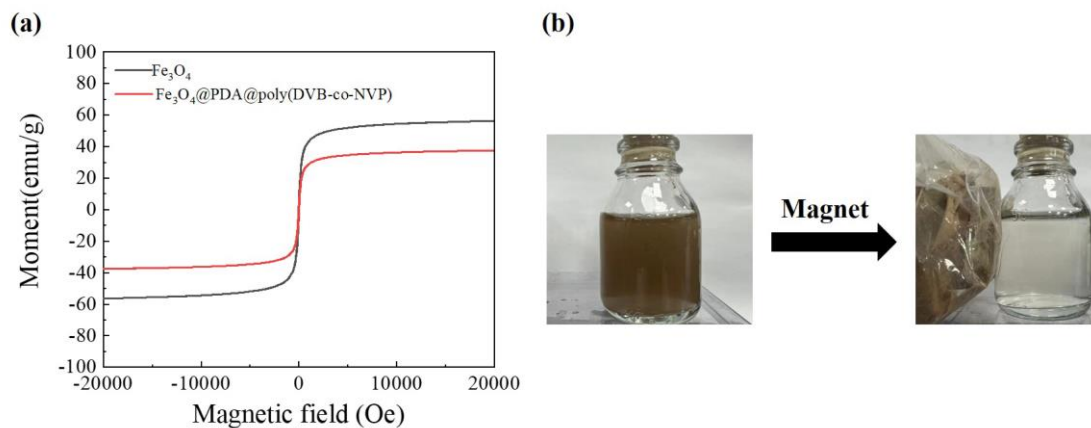


Fig. S4. (a) magnetization curves of Fe_3O_4 MNPs and $\text{Fe}_3\text{O}_4@\text{PDA}@\text{poly}(\text{DVB-co-NVP})$ MNPs. (b) Magnetic separation of $\text{Fe}_3\text{O}_4@\text{PDA}@\text{poly}(\text{DVB-co-NVP})$ MNPs.

Fig. S4(a) shows the magnetic properties of Fe_3O_4 MNPs and $\text{Fe}_3\text{O}_4@\text{PDA}@\text{poly}(\text{DVB-co-NVP})$ MNPs measured by VSM at a temperature of 300K. The practical saturation magnetization of Fe_3O_4 MNPs and $\text{Fe}_3\text{O}_4@\text{PDA}@\text{poly}(\text{DVB-co-NVP})$ is calculated at 56.33 and 37.38 emu/g. From the data, it can be seen that after coating and fabrication of $\text{Fe}_3\text{O}_4@\text{PDA}@\text{poly}(\text{DVB-co-NVP})$ MNPs, the actual saturation magnetization showed a tendency of decrease. Though $\text{Fe}_3\text{O}_4@\text{PDA}@\text{poly}(\text{DVB-co-NVP})$ MNPs exhibited a compromised magnetism compared to Fe_3O_4 , both of them could be readily separated from solution by applying a magnet as shown in Fig. S4(b), supporting their application as MSPE adsorbents.

Table S1. Real sample analysis

Samples	MDMB- FUBICA	4CN-CUMYL- BUTINACA	5F-MDMB- PICA	MDMB-4en- PINACA	ADB-4en- PINACA	5F-EMB- PICA	AMB- FUBINACA	ADB- BUTINACA	4F-MDMB- BUTICA
S1	-	-	-	-	-	-	-	-	-
S2	-	-	-	-	-	-	-	-	-
S3	-	-	-	-	-	-	-	-	-
S4	-	-	-	-	-	-	-	-	-
S5	-	0.15	-	-	-	-	-	-	-
S6	-	-	-	-	-	-	-	-	-
S7	-	-	-	-	-	-	-	-	-
S8	-	-	-	-	-	-	-	-	-
S9	-	0.11	-	-	-	-	-	-	-
S10	-	<LOQ	-	-	-	-	-	-	-
S11	-	-	<LOQ	<LOQ	-	-	-	-	-
S12	-	-	<LOQ	<LOQ	-	-	-	-	-

References

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