

Supporting Information for

Fe-Ni@ACC Nanocomposite for Magnetic Dispersive Micro Solid-phase Extraction of Cu (II) From Food and Hair Samples

Seda Duman^{a,b} and Mustafa Soylak^{a,b,c,*}

^a Erciyes University, Faculty of Sciences, Department of Chemistry, 38039 Kayseri-Turkiye

^b Technology Research and Application Center (ERU-TAUM), Erciyes University, 38039 Kayseri-Turkiye

^c Turkish Academy of Sciences (TUBA), Bayraktar Mahallesi, Vedat Dalokay Caddesi No: 112, 06670 Cankaya, Ankara, Turkiye

***Corresponding author**

E-mail: soylak@erciyes.edu.tr (M. Soylak);

Table S1. Application of the developed M-D- μ SPE method (pH: 9, Fe-Ni@ACC amount: 5 mg, eluent type: 2M HNO₃, eluent volume: 0.75 mL N=3).

Added ($\mu\text{g g}^{-1}$)	Dye-Free Hair		Colored Hair		Cigarette	
	Found ($\mu\text{g g}^{-1}$)	Recovery (%)	Found ($\mu\text{g g}^{-1}$)	Recovery (%)	Found ($\mu\text{g g}^{-1}$)	Recovery (%)
0	BDL	-	0.148 \pm 0.001	-	0.124 \pm 0.005	-
0.100	0.098 \pm 0.001	98	0.245 \pm 0.002	97	0.221 \pm 0.002	97
0.200	0.191 \pm 0.002	96	0.342 \pm 0.003	97	0.318 \pm 0.003	97
0.400	0.390 \pm 0.003	98	0.561 \pm 0.005	103	0.512 \pm 0.005	97

BDL: below the detection limit.

Table S2. Application of M-D- μ SPE method to some real samples (pH: 9, Fe-Ni@ACC amount: 5 mg, eluent type: 2M HNO₃, eluent volume: 0.75 mL N=3).

Added ($\mu\text{g g}^{-1}$)	Black Tea / Turkey		Black Tea / Indonesia		Tap water	
	Found ($\mu\text{g g}^{-1}$)	Recovery (%)	Found ($\mu\text{g g}^{-1}$)	Recovery (%)	Found ($\mu\text{g mL}^{-1}$)	Recovery (%)
0	0.248 \pm 0.004	-	0.442 \pm 0.006	-	BDL	-
0.200	0.442 \pm 0.010	97	0.635 \pm 0.010	97	0.192 \pm 0.005	96
0.400	0.636 \pm 0.020	97	0.822 \pm 0.030	95	0.377 \pm 0.010	94
0.800	1.025 \pm 0.020	97	1.268 \pm 0.030	103	0.781 \pm 0.020	98

BDL: below the detection limit.

Table S3. Comparison of the SPME method developed for the separation and preconcentration of Cu (II) with other methods in the literature.

Method	System / Reagents	LOD ($\mu\text{g L}^{-1}$)	LOQ ($\mu\text{g L}^{-1}$)	PF	RSD (%)	Instrument	Ref.
Solid Phase Microextraction	Pyrocatechol violet impregnated magnetic graphene oxide	4.0	13.3	20	4.93	FAAS	[2]
Ionic liquid-based ultrasound assisted microextraction	1-butyl-3-methylimidazolium hexafluorophosphate [C ₄ mim][PF ₆]	0.36	-	31	5	AAS	[6]
Solid Phase Microextraction	Bis(2-hydroxy acetophenone) ethylenediamine	0.27	-	-	0.90	AAS	[38]
Solidified floating organic drop microextraction	sodium dodecyl benzene sulfonate	0.18	0.58	541	2.7	FI-FAAS	[60]
Solid Phase Microextraction	bamboo fiber	0.5	1.66	33	2.70-2.79	FAAS	[61]
Dispersive liquid liquid microextraction	corrugated quartz tube	0.7	2.2	-	3.7-8.4	FAAS	[62]
Thin film microextraction	Poly m-phenylenediamine/CNT electrospun nanofiber	0.32	-	76	2.9-3.7	FAAS	[63]
Dispersive liquid liquid microextraction	No-Vanillidin-2-amino-p-cresol, 1-undecanol	0.93	-	20	-	FAAS	[64]
Ultrasound assisted liquid phase microextraction	Deep eutectic solvent	6.6	21.8	15	5	FAAS	[65]
Magnetic dispersive micro solid-phase extraction	Fe-Ni@ACC	0.69	2.29	40	1.18	FAAS	This work

LOD: Limit of detection, LOQ: Limit of quantification, PF: Preconcentration Factor, % RSD: Relative Standard Deviation,