

The 5th International Electronic Conference on Applied Sciences



04-06 December 2024 | Online

Effect of sonication-assisted water extraction on the total antioxidant parameters of medicinal plants

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INTRODUCTION & AIM

Medicinal plants are one of the sources of biologically active compounds that determine their therapeutic effect. Water infusions and decoctions, as well as tinctures and extracts, are currently used in phytotherapy and part of complex treatment of various human diseases.

Sonication treatment is an effective approach to increasing the efficiency of active component extraction from plant material. Application of sonication reduces the time and consumption of extractant, as well as uses mild conditions. Sonication-assisted water extraction was applied to the various medicinal plants traditionally used in phytotherapy. Water extracts from herbs, leaves, bark, infructescences, flowers, roots, and rhizomes were studied using total antioxidant parameters obtained by constant-current coulometry.

RESULTS & DISCUSSION

Effect of sonication time on the antioxidant parameters of the extracts



METHOD

Plant material under study and extracts preparation

Extract type	Dlant material	Plant : solvent	
	Flant material	ratio	
	Quercus cortex	1:10	
	Bergeniae rhizomata	1:10	
Decoctions and	Potentillae rhizomata	1:10	
sonication-assisted	Chamomillae flores	1:33	
water extracts	Sanguisorbae rhizomata et radices	1:10	
	Frangulae cortex	1:10	
	Uvae Ursi folia	1:10	
	Alni fructus	1:10	
Infusions and	Leonuri herba	1:10	
sonication-assisted	Salviae folia	1:10	
water extracts	Urticae folia	1:33	
	Tiliae flores	1:20	



sample addition

t / min

30 -

20 -

10 -

0 -

0

Αμ / Ι



Maximum TAC and FRP achieved for 30 min of sonication treatment.

Comparison of TAC and FRP of the extracts pbtained by sonication and traditional method (*n* = 5; *P* = 0.95)

Plant material	Sonication for 30 min	RSD	Traditional method	RSD		
TAC / C 100 mL ⁻¹						
Alni fructus	304±6	0.02	810±10	0.01		
Quercus cortex	429±4	0.006	537±11	0.01		
Frangulae cortex	280±9	0.03	444±7	0.01		
Uvae Ursi folia	1178±31	0.02	1532±34	0.02		
Potentillae rhizomata	854±22	0.02	1494±26	0.01		
Bergeniae rhizomata	1390±60	0.04	1438±59	0.02		
Sanguisorbae rhizomata et radices	1432±52	0.02	2271±120	0.04		
Leonuri herba	105±5	0.03	132±3	0.01		
Tiliae flores	123±4	0.03	245±11	0.05		
Salviae folia	196±12	0.04	265±9	0.03		
Urticae folia	52±2	0.04	64±2	001		
Chamomillae flores	89±3	0.03	133±2	0.01		
FRP / C 100 mL ⁻¹						
Alni fructus	293±10	003	558±15	0.02		
Quercus cortex	188±7	0.03	214±5	0.024		
Frangulae cortex	34±1	0.03	59±3	0.05		
Uvae Ursi folia	643±17	0.02	922±44	0.03		
Potentillae rhizomata	578±12	0.02	912±32	0.03		
Bergeniae rhizomata	647±24	0.02	1044±25	0.01		
Sanguisorbae rhizomata et radices	691±38	0.02	1001±30	0.02		
Leonuri herba	54±1	0.02	67±2	0.03		
Tiliae flores	64±1	0.01	133±6	0.03		
Salviae folia	114±3	0.02	162±5	0.01		
Urticae folia	16.8±0.8	0.04	20.6±0.8	0.03		
Chamomillae flores	16.3±0.6	0.01	41±2	0.03		



Coulometric titration with electrogenerated bromine and ferricyanide ions

Total antioxidant capacity (TAC)

Anode: $2Br^- - 2e^- = Br_2$

Solution: Oxidation reactions Electrophilic substitution in aromatic systems Electrophilic addition to multiple bonds

Ferric reducing power (FRP)

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Anode: [Fe(CN)_6]^{4-} - e^- = [Fe(CN)_6]^{3-}
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Solution: Oxidation reactions

CONCLUSION

The effect of sonication on the antioxidant parameters of water extracts from medicinal plants was evaluated on the basis of coulometric data. The TAC and FRP of samples are are increased with the growth of sonication time. The best results are achieved within 30 min treatment. Further extension of the sonication time is inadvisable from practical point of view. The comparison of the antioxidant parameters for sonicated extracts with that ones for infusions and decoctions shows has been performed. Traditional technology of decoctions and infusions preparation provides higher antioxidant parameters due to the conditions used for preparation (boiling for 30 and 15 min respectively). TAC and FRP for different methods are more consistent in the case of finely ground plant materials, such as leaves and herb while bark and rhizomes show more diverse data.

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