

Supporting document S1

Effect of ultrasound on physical properties and processing of biopolymers-A review

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Table S1

Effect of ultrasound on the extraction and properties of three major biopolymers

Biopolymer	Extraction					Physical properties	Ref.
	Method of extraction	Step(s) involved US	US intensity	Time (min)	Temp (°C)		
Chitin and Chitosan	Chemical+ US in 40% (w/w) NaOH	Deacetylation	52.6 Wcm ⁻²	50	60	<ul style="list-style-type: none"> Chitosan molecular weight decreased to (912,000 g/mol) Degree of polymerization decreased from 6865 to 4889 Degree of acetylation falls from 80.7% to 4.3% Dispersity of 1.3 is found 	2016 ¹
	Chemical+ US in 40% (w/w) NaOH	Deacetylation	37 kHz	360	80	<ul style="list-style-type: none"> Molecular weight decreased from 378 to 361 kDa at 60% NaOH Molecular weight decreased from 439 to 421 kDa at 50% NaOH 	2017 ²
	US under oil-water emulsion	Emulsification	300-600 W	3 to 5	10	<ul style="list-style-type: none"> One step process requires less time and energy High intensity decreases droplet size 	2018 ³
	Chemical+ dual US in 10% NaOH	Deacetylation	15 kHz and 20 kHz	15	25	<ul style="list-style-type: none"> Molecular weight falls from 20000 g/mol to 5000 g/mol for dual frequency transducer at parallel position Degree of deacetylation (DD) increases from approximately 	2019 ⁴

						30% to 80%	
	US in 1.0 M NaOH under ice bath	Deproteinization	750W and 20 kHz	41.46	30-40	<ul style="list-style-type: none"> Viscosity reduced to 12 cP from 85 cP 	
	US in deionized water	Deproteinization	-	15	-	<ul style="list-style-type: none"> Less amount of protein is left after treatment Reduce extraction time with 34 % chitin after deproteinization Not suitable for fat absorption purpose 	2019 ⁵
				25		<ul style="list-style-type: none"> Molecular weight 73.61 kDa DD = 80.60 % Particle size = 35.70 μm Molecular weight 86.82 kDa DD = 92.86 % Particle size = 25.51 μm Molecular weight 55.66 kDa DD = 55.66 % Particle size = 20.10 μm 	2020 ⁶
				40			
Starch	US in starch solution + Enzyme		7.20 W/mL	10		<ul style="list-style-type: none"> Molecular weight decreased by 80.19 % Solubility raised by 136.50 % 	2017 ⁷
	US under water		450 W (25 kHz) with 70 % amplitude	15		<ul style="list-style-type: none"> Yield increase from 29.85 % to 32.09 % Amorphous region damaged but crystalline region remain fixed. 	2018 ⁸
	US under water		(20 and 45 kHz) (15.29,			<ul style="list-style-type: none"> High amylose content starch is obtained 	2019 ⁹

			20.38, 22.93, 24.46 and 25.38 kW/m ²)				
	US under water		150, 300, 450 and 600 W	20		<ul style="list-style-type: none"> • Granule size did not change but their homogeneity has increases • Peak and breakdown viscosity increases 	2019 ¹⁰
	US in starch suspension	Nano and Microparticle formation	20 kHz	30		<ul style="list-style-type: none"> • Starch nanoparticle and microparticle fabrication without chemicals and additional purification steps • High amylose content starch source provides smaller nano and microparticle 	2019 ¹¹
Cellulose	Acid hydrolysis and US in cellulose suspension to get NCC	Acid hydrolysis	60 Hz	10 (max)	-	<ul style="list-style-type: none"> • Molecular weight decreases • Disintegrate NCC aggregates and degrade nanocrystals • Increase optical clarity of fabricated material 	2016 ¹²

		37 Hz	120	45	<ul style="list-style-type: none"> • High crystallinity, thermally stable spherical shaped NCC is obtained 	2017 ¹³
US with Water and Fenton reagent	MCC enzymatic hydrolysis pretreatment	800 W, 21-23 kHz	150	25	<ul style="list-style-type: none"> • Decrease DP and aspect ratio of MCC 	2016 ¹⁴
US in NCC suspension	Modifying properties	20 kHz and 60% amplitude	Up to 10	RT	<ul style="list-style-type: none"> • Increase transparency and tensile strength of US treated NCC based film 	2017 ¹²
US in 2M NaOH	Cellulose and lignin extraction	500 W, 20 kHz	Up to 40	90	<ul style="list-style-type: none"> • Higher thermal stability of cellulose 	2017 ¹⁵
US with enzymatic hydrolysis		500 W, 20 kHz, 16.2-43.4 Wcm ⁻²	90	50	<ul style="list-style-type: none"> • Less cellulose extraction time • Enzymatic hydrolysis is more effective with US compared to conventional method • Size and form of raw material, position of the US radiation source are equally important to improve process efficiency 	2017 ¹⁶
US under water with cellulose	Cellulose nanofibers fabrication	25 kHz	-	-	<ul style="list-style-type: none"> • High thermal stability 	2016 ¹⁷
		400-1200 W			<ul style="list-style-type: none"> • Ionic repulsion between the fabricated fiber makes easy separation of fiber 	2017 ¹⁸

			400 W			<ul style="list-style-type: none"> • High yield of fiber 	2018 ¹⁹
US in MCC suspension	To produce fibrillated MCC			120		<ul style="list-style-type: none"> • Hydrogels from high aspect fiber is obtained 	2018 ²⁰
US applied to cellulose pulp	Improving swelling and dissolution behaviour		320 W, 37 kHz	20	30	<ul style="list-style-type: none"> • Lowers the crystallinity under a low temperature treatment • Decreases the particle size and dissolution time of cellulose 	2018 ²¹
US applied to homogenized cellulose pulp	Cellulose micro and nanofiber fabrication		500 W	40		<ul style="list-style-type: none"> • High aspect ratio small fiber is obtained 	2019 ²²
US of cellulose in water	Depolymerizing cellulose to glucose		525 kHz	180	60	<ul style="list-style-type: none"> • Selective depolymerization of cellulose is recorded without catalyst and extensive heating 	2020 ²³
US with NaOH solution	Cellulose nanocrystal fabrication		120 W, 20 kHz	105		<ul style="list-style-type: none"> • Increased crystallinity, lower particle size and high thermal stability are obtained 	2021 ²⁴

US= Ultrasound, NCC= Nanocrystalline cellulose, MCC= Microcrystalline cellulose, RT= Room temperature

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