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Influence of a facile pretreatment process on lipid extraction from *Nannochloropsis* sp. through an enzymatic hydrolysis reaction†

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Microalgal lipid is being considered as an alternative source of ω -3 polyunsaturated fatty acids (ω -3 PUFAs); however, the extraction of ω -3 PUFA-rich algal lipid still needs further research. In this study, single enzyme and their combinations were screened based on their cell wall disrupting capability and lipid recovery yield, and economical and feasible extraction parameters were determined. Lipid recovery could reach 22.18 ± 0.26 wt% when algae was treated with an 8 : 1 (w/w) algae/cocktail enzyme ratio and a 2 : 1 cellulase/lysozyme mass ratio at 50 °C under stirring for 5 h at a pH of 4.0, and the content of PUFAs was as high as 23.30%. In addition, alkaline pretreatment efficiently facilitates the degradation of algal cell wall in the enzymatic hydrolysis, and it has been verified that cell wall ruptures *via* two steps: swelling by a weak alkali pretreatment and decomposition by enzymatic hydrolysis. Furthermore, all the extraction operations can take place under facile conditions. These results imply that algal lipid extracted by a weak alkali pretreatment aiding enzymatic hydrolysis, is the most appropriate raw material for ω -3 PUFAs.

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1. Introduction

Polyunsaturated fatty acids (ω -3 PUFAs) are health-beneficial nutrients that are widely used in the food and pharmaceuticals fields.¹ Besides lowering cholesterol, they can also promote the immune system, prevent cancer and cardiovascular diseases, and contribute to the development of brain and cognition health.^{2,3} Although representative compounds, including γ -linolenic acid (ALA), eicosapentaenoic acid (EPA), and docosahexaenoic acid (DHA), of ω -3 PUFAs mainly exist in marine fish and microalgae, their commercial source is mainly deep-sea fish.^{2,4} However, ω -3 PUFAs derived from fish oils have many limitations such as fishy smell, heavy metal contamination, and inadequate supplies.⁵ Therefore, the lipid extracted from

microalgae has become a promising alternative source for ω -3 PUFAs owing to the high content of ω -3 PUFAs in it; thus, these microalgae have attracted significant attention and interest in academia and industry.⁶ However, ω -3 PUFAs extracted from algal lipid *via* practical processes are greatly restricted because of their poor stability and easy oxidation in the traditional extraction methods.^{7,8} Furthermore, the extraction of algal lipid is particularly difficult since the lipid is stored in lipid vesicles,³ which are protected by robust cell walls. Cell walls of microalgae are composed of microfibril networks of polysaccharides, including cellulose and pectin, and algaenans, with complicated interactions between them, and the entangled network is implanted in a matrix of pectin along with cross-linked algaenans;^{9,10} thus, they severely hamper conventional solvent access to the lipid vesicles. Therefore, it is essential to develop more efficient and effective approaches to break the algal cell wall for lipid extraction.

Compared to the conventional disruption technologies,¹¹ enzymatic hydrolysis is being considered as a promising wall-breaking method for algal lipid extraction because of its inherent advantages¹² such as mild conditions, low-energy, and ability to preserve the most labile compounds. However, it is difficult to obtain a desirable extraction efficiency for an individual enzyme. For example, the yield of the lipid extracted with free and immobilized cellulase from *Chlorella vulgaris* was 73% and 56%, respectively.^{13,14} To degrade the algae cell wall more completely and obtain more ω -3 PUFAs, cocktail enzymes have been employed in the hydrolysis reaction. Moreover, enzymatic hydrolysis in combination with a pretreatment, which includes

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† Electronic supplementary information (ESI) available: The compositional analysis of fatty acids in algal lipid extracted separately by EH and WAEH, morphology of algal cell, and the content of chlorophyll in algal lipid extracted separately by WAEH and EH can be found in the ESI, and detailed data can be found in Fig. S1–S3. See DOI: 10.1039/c7ra11483d

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hydrothermal reaction, ultra-sonication, and acidic or alkaline reaction, could cause an obvious increase in the lipid recovery.¹⁵ It was concluded that acidic environment could obviously enhance the disruption efficiency of algal cells in the enzymatic hydrolysis and improve the recovery of sugar,¹⁶ and the alkaline hydrothermal reaction could greatly increase the dissolution of cellulose present in cell wall in the enzymatic hydrolysis because of hydrogen bond rupture.^{17,18} Wu *et al.* developed a process *via* an alkaline pretreatment (at 110 °C and pH 10.5), which assisted enzymatic hydrolysis to extract algal lipid, and the yield exceeded 90%.¹⁹ However, oxidation easily occurs above 100 °C of the pretreatment temperature for algal lipids, which are released in the early stage owing to the presence of a great amount of unsaturated double bonds.²⁰ Moreover, significant research has been focused on the extraction of algal lipid, which is used as a feedstock for biodiesel. Furthermore, to the best of our knowledge, the extraction technology of algal lipid aimed at separating ω -3 PUFAs has not been investigated to date. In this study, the marine microalgae *Nannochloropsis* sp. was selected as the raw material for algal lipid extraction; the main reason for using this microalgae was its ability to accumulate large amounts of ω -3 PUFAs and EPA.²¹ The disruption process of algal cell wall *via* weak alkali pretreatment-aided enzymatic hydrolysis (WAEH) for the extraction of the lipid aimed at separating ω -3 PUFAs was developed, and the high efficiency cell wall-breaking of enzyme based on the yield of lipid recovery was investigated. The suitable reaction conditions, including reaction temperature and time, pH value, the ratio (w/w) of cocktail enzymes, and mass ratio between enzyme and algae, were determined, and the highest extraction yield of the algal lipid was obtained. Microalgal cells treated with different methods and lipids extracted by several methods were characterized using Fourier transform infrared spectroscopy (FT-IR), ultraviolet-visible spectroscopy (UV-Vis), scanning electron microscopy (SEM), and fluorescence microscopy. The composition and content of fatty acids obtained from algal lipid were analyzed using gas chromatography (GC). The effect of the weak alkali pretreatment (WAP) on the algal cell wall rupture and lipid extraction was investigated.

2. Materials and methods

2.1 Chemicals and reagents

Nannochloropsis sp. was purchased from Yantai Hairong Biology Technology Co. Ltd. (Yantai, China). Cellulase, pectinase, lysozyme, and neutral protease were obtained from Nanjing Odd-foni Biology Technology Co. Ltd. (Nanjing, China). Hexane, ethanol, methanol, chloroform, acetic acid, and sodium acetate were purchased from Sinopharm Chemical Reagent Co. Ltd. (Shanghai, China), and the purity of these solvents was analytical grade.

2.2 The extraction of algal lipid

2.2.1 The content of the lipid in *Nannochloropsis* sp.. In this study, algal lipid, by definition, is *n*-hexane soluble fractions that are obtained from the microalgae extract. The content

of the lipid in *Nannochloropsis* sp. was determined using two methods: Soxhlet extraction and Bligh–Dyer extraction, and the detailed procedures were employed according to the literature^{22,23} with slight modifications.

Soxhlet extraction was conducted using 1.0 g of microalgae and a mixed solvent of chloroform and methanol (2 : 1, v/v), and the temperature was set at 90 °C. The reaction continues until the solvent stored in the extractor is almost colorless and transparent. After the solvent is removed, the remnant is dissolved and filtered with *n*-hexane, and algal lipid, which is defined as the *n*-hexane soluble fraction, is acquired after the filtrate is obtained and evaporated.

On the other hand, in the Bligh and Dyer extraction, 100 mg of microalgae, which was ground using a high-pressure homogenizer, was added to a mixture of chloroform, methanol, and water (10 : 5 : 0.9, v/v/v) and stirred for 12 h. Then, the mixture was centrifuged, and the supernatant was transferred to a flask. The process was repeated two more times using a stirring time of 2 h. After removing the solvent from the above collected supernatant, the residual material was dissolved using *n*-hexane, and algal lipid was obtained from the filtrate by vacuum distillation. The weight of the algal lipid was measured gravimetrically, and the data were determined through the mean value obtained from the abovementioned two extraction methods.

2.2.2 Algal lipid extraction with the WAEH method.

Herein, two steps of weak alkali pretreatment and enzymatic hydrolysis (EH) are necessary for the lipid extraction using the WAEH method. Detailed procedures are as follows:

(1) *Weak alkali pretreatment.* Herein, 3.0 g of microalgae was added to a flask that contained 10 mL of deionized water. The pH value of the mixture was adjusted to 9.0 with sodium hydroxide. After being sealed with parafilm, the flask was incubated in an oil bath (DF-101S, Gongyi Yingyu Instrument Factory) with a magnetic stirrer for 5 h at 80 °C. After the reaction, the flask was removed from the oil bath and cooled to ambient temperature.

(2) *Enzymatic hydrolysis.* After WAP, the pH value of the reaction solution was adjusted to the acid range with acetic acid. Different dosages of single or cocktail enzymes were added to the mixture. Then, the algae suspension was stirred for various lengths of time at different temperatures. Then, the mixtures were heated in an oil bath at 95 °C for 10 min to deactivate the enzyme. After this, 15 mL of ethanol was added to the mixture prior to lipid extraction with *n*-hexane (30 mL) by centrifuging at 5000 rpm for 25 min. The obtained organic phases were evaporated by a rotary evaporator (Shanghai Yarong Biochemical Instrument Factory) under vacuum, and the residues were dried for 6 h under vacuum at 40 °C. Finally, the lipid was stored in a vacuum dryer.

2.2.3 Experimental designs. In the study of screening the enzymes, the wall-breaking ability of the tested enzymes, which include the single enzyme, binary mixed enzymes, ternary mixed enzymes, and quaternary mixed enzymes, was assessed based on the yield of lipid recovery in enzymatic hydrolysis. For the reaction of cocktail enzymes, 1 : 1 (w/w) ratio of the binary mixed enzymes was adopted, and the proportion of 1 : 1 : 1 was



adopted when three enzymes were involved in the reaction; moreover, the mass ratio of four enzymes participating in the reaction was 1 : 1 : 1 : 1.

The extraction conditions for algal lipid were optimized in enzymatic hydrolysis, and cocktail enzymes, including cellulase and lysozyme, were used as candidates. The influence of the reaction time on lipid recovery was investigated, and 1 h, 2 h, 3 h, 4 h, 5 h, 6 h, 7 h, 8 h... were set as the reaction times. Moreover, the temperature, pH value, cellulase/lysozyme mass ratio, and algae/enzyme mass ratio of the reaction was 50 °C, 4.0, 2 : 1, and 8 : 1, respectively. In addition, 30 °C, 35 °C, 40 °C, 45 °C, 50 °C, 55 °C, 60 °C, 65 °C, and 70 °C reaction temperatures were used for this research, and the other reaction conditions *i.e.* reaction time, pH, cellulase/lysozyme ratio, and algae/enzyme mass ratio were 7 h, 4.0, 2 : 1 (w/w), and 8 : 1, respectively. The effect of pH on lipid recovery was investigated when the reaction pH value was set as 3.0, 3.5, 4.0, 4.5, 5.0, 5.5, 6.0, 6.5, and 7.0, and other conditions of the reaction were 50, 7 h, 2 : 1 (w/w), and 8 : 1 (w/w). The relationship of mixed ratio with the binary enzymes and lipid recovery was investigated, and the mass ratio of cellulase and lysozyme was designed as 6 : 1, 5 : 1, 4 : 1, 3 : 1, 2 : 1, 1 : 1, 1 : 3, and 1 : 6, and other reaction conditions including reaction temperature, pH value, reaction time, and algae/enzyme mass ratio were 50 °C, 4.0, 7 h, and 8 : 1, respectively. The influence of enzyme dosage on the yield of lipid recovery was determined, and the quantity of mixed enzymes (cellulase and lysozyme) increases in the following order: 160 : 1, 80 : 1, 40 : 1, 20 : 1, 8 : 1, 2.6 : 1, and 2 : 1. Temperature, pH, time, and the cellulase/lysozyme mass ratio of the reaction were 50 °C, 4.0, 7 h, and 2 : 1, respectively. Moreover, the control test was performed without the pretreatment of WAP under the same conditions.

2.3 The yield of algal lipid recovery

The weight of algal lipid was then determined gravimetrically, and the yield of lipid recovery was calculated using the equation as follows:

$$\text{lipid yield (wt\%)} = \frac{m_1}{m_2} \times 100\%$$

where m_1 is the dry weight of lipid recovery, mg g^{-1} DCW (dry cell weight), and m_2 is the dry weight of microalgae, mg g^{-1} DCW.

2.4 Analytical methods

2.4.1 Fourier transform infrared spectroscopy characterization of algal lipid. The spectra of algal lipid extracted separately by EH and WAEH were obtained using a Nicolet Nexus FTIR spectrometer with KBr pellets. Test samples were scanned and investigated using FTIR spectroscopy by 32 scans at a resolution of 2 cm^{-1} .

2.4.2 Imaging of algal cell by fluorescence microscopy and scanning electron microscopy. Morphology features and particle sizes of the microalgal cell and microalgae cell treated separately with EH, WAEH, and WAP methods were measured using JEOL JSM-6390LV SEM with an acceleration voltage of 20

kV and Leica DMI 3000 B fluorescence microscopy in the bright field mode, respectively. To ensure uniform well dispersion of the individual algal cell on the slide, liquid samples were diluted 5 times with deionized water. In addition, solid samples were washed for 3 times with deionized water by centrifuge, and the acquisition was freeze-dried at $-78 \text{ }^\circ\text{C}$ under vacuum before being characterized.

2.4.3 Compositional analysis of algal lipid. To determine the compositions and content of ω -3 PUFAs in algal lipid, algal lipid treated *via* the methyl esterification reaction is used prior to sample injection in the gas phase chromatograph. Detailed procedure was performed according to a reference with a slight adjustment.²⁴

An acidic catalyst of the methyl esterification reaction was prepared by mixing acetyl chloride and methanol at a ratio of 1 : 10 (v/v) at ambient temperature, and the reaction mixture was stirred for a period of time until the reactant was cooled to atmospheric temperature. Then, 50 mg of lipid sample, 5 mL of *n*-hexane, and 4 mL of the abovementioned solution were added to a flask in which the mixture was refluxed for 2 h in an oil bath at $75 \text{ }^\circ\text{C}$. The reaction was quenched by adding 4 mL of water. The hexane layer was transferred into a pre-weighed glass tube and evaporated to dryness under a nitrogen stream at $37 \text{ }^\circ\text{C}$. Finally, FAMES were stored under a nitrogen atmosphere.

The composition of FAMES was determined using GC (6890A, HP-Agilent Technologies, USA) equipped with a DB-23 capillary column ($30 \text{ m} \times 0.25 \text{ mm} \times 0.25 \text{ }\mu\text{m}$). All the ingredients of FAMES were identified by comparing their retention times with a reference standard comprising 37 components. The injection and flame ionization detector (FID) temperature were maintained at $240 \text{ }^\circ\text{C}$, and the time-temperature program was as follows: $100\text{--}180 \text{ }^\circ\text{C}$ at $10 \text{ }^\circ\text{C min}^{-1}$ and $180\text{--}240 \text{ }^\circ\text{C}$ at $2 \text{ }^\circ\text{C min}^{-1}$. Helium was used as the carrier gas at a flow rate of 1 mL min^{-1} . The fatty acids content was determined by adding an internal standard of undecanoic acid (known quantity) to the oil sample before extraction, and the peak areas were quantified using Chromcard for Windows software. The areas of the peaks were summed and compared with the area of the internal standard (C11:0).

2.5 Statistical analysis

All experiments were conducted in triplicate. The data were evaluated using one-way analysis of variance (ANONA) followed by SPSS. P value ≤ 0.05 was regarded as significant, and data were presented as mean \pm standard deviation (SD).

3. Results and discussion

3.1 Screening enzymes

Considering two factors that cell walls rigidity of *Nannochloropsis* sp. is three times stronger than that of plant and the activity of enzyme is specificity, enzymes with stronger wall-breaking capability, which include cellulase, lysozyme, pectinase, and neutral protease, were selected as candidates for the screening enzyme.^{25,26} Moreover, a series of reactions of individual or cocktail enzymes was designed to screen the suitable



Table 1 Influence of individual and cocktail enzymes on lipid recovery yield^a

Enzymes				Lipid recovery yield (wt%)
Cellulase	Lysozyme	Pectinase	Neutral protease	
+	-	-	-	15.55 ± 0.23
-	+	-	-	18.16 ± 0.03
-	-	+	-	13.82 ± 0.15
-	+	-	+	10.18 ± 0.07
+	+	-	-	22.18 ± 0.26
+	-	+	-	20.49 ± 0.07
+	-	-	+	15.14 ± 0.19
-	+	+	-	18.55 ± 0.10
-	+	-	+	20.97 ± 0.27
-	-	+	+	13.40 ± 0.09
-	+	+	+	17.05 ± 0.13
+	+	+	-	22.46 ± 1.34
+	-	+	+	18.04 ± 0.22
+	+	-	+	21.17 ± 0.08
+	+	+	+	20.89 ± 0.06

^a Note: +, the enzyme was added; -, the enzyme was not added.

enzymes that have a better cell wall breaching efficiency. The most suitable enzymes were determined based on the lipid recovery after enzymatic hydrolysis, and the relevant results are shown in Table 1.

As seen in Table 1, for an individual enzyme, the lipid recovery followed the order lysozyme > cellulase > pectinase > neutral protease. The yields of lipid extracted separately with the binary, ternary, and quaternary mixed enzymes were higher than those of lipid extracted with a single enzyme, and higher lipid yields were obtained when lysozyme was among the cocktail enzymes, which was consistent with the reported results.¹⁹ Furthermore, 22.46 ± 1.34 wt% of the lipid recovery for cocktail enzymes, containing lysozyme, cellulase, and pectinase, was the highest among that of all the enzyme combinations. There are two possible reasons: one is that the cocktail enzymes containing lysozyme, cellulase, and pectinase can lead to a stronger degradation capability of algal cell wall in the hydrolysis reaction; the other reason is that the cocktail enzymes enable the fast liberation of the lipid that exists in the subcellular structure to the slurry when it is extracted by complex solvents.²⁷ However, it is noteworthy that the lipid recovery declined slightly in the presence of neutral protease. The reason for this is that the activity of neutral protease is inhibited to some extent, and the concentration of effective cocktail enzymes is reduced equivalently since the hydrolysis medium pH is 4.0, whereas the optimized pH value of neutral-protease hydrolysis is 7.0.²⁸ Although the lipid recovery yield obtained after utilizing the cocktail enzymes cellulase and lysozyme was 0.28 wt% lower than that obtained after using cellulase, lysozyme, and pectinase, the most suitable cocktail enzymes were chosen to be lysozyme and cellulase for lipids extracted from *Nannochloropsis* sp. taking into consideration the lipid recovery and economic feasibility.

3.2 The effect of enzymatic hydrolysis conditions on lipid recovery

Enzymatic hydrolysis conditions including enzyme dosage, temperature, time, pH, and mass ratio of cocktail enzymes were investigated (Fig. 1). As shown in Fig. 1, the yield of algal lipid recovery increased with an increase in enzymatic dosage and remained almost at the same level when the mass ratio between the microalgae and cocktail enzymes was larger than 2.6 : 1. The highest yield of lipid could reach 22.18 ± 0.26 wt% when the hydrolysis reaction took place at 50 °C for 5 h at a pH value of 4.0 with a 2 : 1 mass ratio of cellulase and lysozyme, and this result was more than the yield of the algal lipid extracted by the Blich-Dyer method and Soxhlet extraction; the data was 22.09 ± 0.18 wt%. The reason is that algal lipid extracted by WAEH contains more free fatty acids as compared to that extracted by the Blich and Dye and Soxhlet extraction, and the relevant evidence has been provided in Section 3.3. The abovementioned results indicate that sufficient quantity of cocktail enzymes is necessary for the complete degradation of algal cell walls. However, the curve of the lipid yield was almost flat when the mass ratio between the algae and cocktail enzymes was increased to 8 : 1. Therefore, 8 : 1 mass ratio between microalgae and cocktail enzymes was appropriate for the abovementioned enzymatic hydrolysis considering the extraction costs. Furthermore, lipid recovery increased with an increase in the mass ratio between cellulase and lysozyme when the ratio was higher than 2 : 1, whereas the yield declined with an increase in the ratio when this ratio was below 2 : 1. Based on the abovementioned discussion, the highest yield of lipid

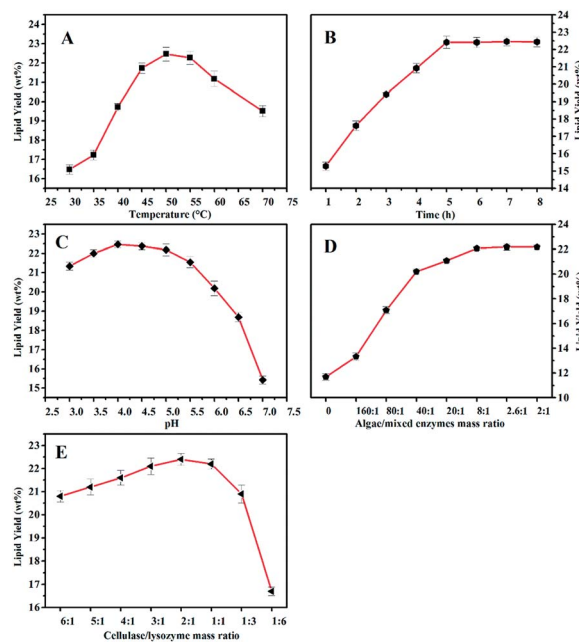


Fig. 1 The influence of enzymatic hydrolysis conditions on lipid recovery yield (mean ± SD; $n = 3$). (A) Effect of time; (B) effect of temperature; (C) effect of pH value; (D) effect of mass ratio between cellulase and lysozyme; and (E) effect of algae/mixed enzymes mass ratio.



recovery could be obtained under the following conditions: temperature: 50 °C, the reaction time: 5 h, pH: 4.0, the ratio of microalgae to enzyme: 8 : 1 (w/w), and mass ratio between cellulase and lysozyme: 2 : 1.

3.3 Analysis of the lipid

The compositions and content of fatty acids in algal lipid were measured by a GC with FID detector, and test samples were extracted separately by EH and WAEH. A detailed analysis of fatty acids is displayed in Table 2 and ESI Fig. S1.† It was observed that the content of total fatty acids obtained from algal lipid extracted by WAEH was much more than that extracted by EH, whereas the amount of unsaturated fatty acids was more than that of saturated fatty acids in the abovementioned lipids.²⁹ In addition, it is worth noting that EPA and PUFA account for 13.21 ± 0.06% and 23.3 ± 0.31% total fatty acids in algal lipid, respectively, and these results are higher than the data (10.30% and 17.10%) reported in previous literature.¹⁹ Furthermore, EPA, which accounts for 56.69% of PUFAs, is the most abundant ingredient in ω-3 PUFAs.³⁰ This suggests that the lipid extracted by WAEH is recognized as a promising feedstock for separating PUFAs. Moreover, the extraction yields of palmitic acid and palmitoleic acid are 33.23 ± 0.02% and 23.90 ± 0.04%, respectively, similar to the result reported in the

literature.³¹ The abovementioned results demonstrate that the weak alkaline hydrothermal treatment-aided enzymatic hydrolysis reaction is more suitable for the extraction of algal lipid, and the extracted lipid can be used as the feedstock for PUFA separation.

3.4 Influence of weak alkali pretreatment on cell wall disruption

It was reported that the cell wall disruption of microalgae that was processed by freeze-drying was difficult because the cell surface of the algae powder was covered with wrinkles and the algal cell was more unsusceptible to enzyme and water.³² However, an algal cell is more easily degraded by enzymatic hydrolysis after being treated by an acidic or alkaline hydrothermal reaction.³³ To investigate the influence of the weak alkali pretreatment on algal cell wall disruption and lipid extraction, morphologies of the algal cell treated with different methods were investigated (Fig. 2). Upon comparing Fig. 2A with Fig. 2B, it was deduced that the particle size of an algal cell shown in Fig. 2B was larger than that shown in Fig. 2A; this might be caused by the swelling of algal cells during the pretreatment process.³⁴ In addition, small amounts of cracks on the cell surface were found, as shown in Fig. 2B, since the swollen cell might be destroyed during the agitation process.

Table 2 Compositional analysis and content of algal lipid extracted separately by WAEH and EH^a

Classifications of fatty acids	Content of fatty acids in algal lipid (mg g ⁻¹ DCW)			Fraction (%)				
	WAEH	EH	Item	WAEH	EH			
Saturate	220.1 ± 0.97	71.1 ± 0.99	C6:0 caproic acid	0.05 ± 0.00	— ^b			
			C8:0 octanoic acid	0.25 ± 0.00	—			
			C10:0 decanoic acid	0.17 ± 0.00	0.12 ± 0.00			
			C12:0 dodecylic acid	0.88 ± 0.09	0.51 ± 0.00			
			C13:0 tridecanoic acid	0.31 ± 0.01	—			
			C14:0 tetradecanoic acid	7.22 ± 0.02	6.54 ± 0.00			
			C15:0 pentadecanoic acid	0.55 ± 0.02	0.56 ± 0.00			
			C16:0 palmitic acid	33.23 ± 0.02	30.21 ± 0.03			
			C17:0 heptadecanoic acid	0.38 ± 0.00	0.35 ± 0.00			
			C18:0 stearic acid	0.88 ± 0.01	0.75 ± 0.00			
			C20:0 arachidic acid	0.14 ± 0.00	—			
			C22:0 docosanoic acid	0.31 ± 0.03	—			
			Mono-unsaturated fatty acids (MUFAs)	160.3 ± 0.85	56.1 ± 0.70	C15:1 pentadecylenic acid	0.11 ± 0.00	—
						C16:1 palmitoleic acid	23.91 ± 0.04	22.61 ± 0.00
C17:1 heptadecenoic acid	0.58 ± 0.01	0.65 ± 0.00						
C18:1 <i>trans</i> -octadecenoic acid	6.78 ± 0.01	0.75 ± 0.00						
C18:1 <i>cis</i> -octadecenoic acid	0.91 ± 0.00	6.45 ± 0.03						
C20:1 eicosenoic acid	0.14 ± 0.00	—						
C22:1 erucic acid	—	0.21 ± 0.00						
Poly-unsaturated fatty acids (PUFAs)	115.9 ± 1.27	55.2 ± 1.45				C18:2 <i>trans</i> -octadecadienoic acid	3.65 ± 0.02	3.63 ± 0.03
			C18:2 <i>cis</i> -octadecadienoic acid	0.63 ± 0.08	—			
			C18:3 octadecatrienoic acid	1.73 ± 0.00	1.87 ± 0.01			
			C20:3 epoxyeicosatrienoic acid	0.22 ± 0.00	—			
			C20:4 arachidonic acid (AA)	3.76 ± 0.15	4.47 ± 0.00			
			C20:5 eicosapentaenoic acid (EPA)	13.21 ± 0.06	20.32 ± 0.03			
Total fatty acids (TFAs)	496.3 ± 3.09	182.4 ± 3.13						

^a Note: each sample was analyzed in duplicate, in addition to duplicate injections into the gas chromatograph, and mean values are shown. ^b Not detected or relative content less than 0.4%.



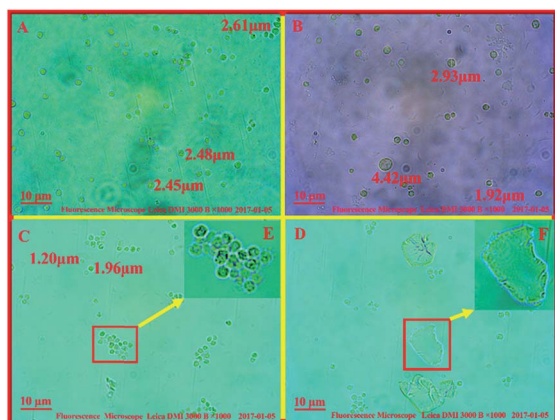


Fig. 2 Microalgae cell morphology was imaged by bright-field fluorescence microscopy. (A) Untreated algae cell; (B) algae cell treated by WAP; (C) algae cell treated by EH; (D) algae cell treated by WAEH; (E) the magnification of the image enclosed by a red box in plane (C); and (F) the magnification of the image enclosed by a red box in plane (D).

Compared to that of raw algal cell, there was no significant change in the morphology of the algal cell treated by EH, whereas a handful of cell debris was found, as shown in Fig. 2C and E, and the surface of freeze-dried algal powders after EH presented small amounts of collapse (ESI Fig. S2[†]). Herein, one explanation of this could be that algae cells were damaged by enzymes in the process of hydrolysis; then, the algal lipid was released from the broken cell with the permeation of organic solvent, and the rupture of algal cells with the internal cavity was caused by external forces during the agitation process.³⁵ Cell fragments increased significantly when algal cells were treated with weak alkali before the enzymatic hydrolysis as compared to those of the original algal cell and those after treatment by EH; the detailed information can be seen in Fig. 2C and F. Furthermore, the damage to tiny amounts of algal cells is severe such that the intracellular components are completely released, and only a layer of membrane remained in the slurry. In addition, it was observed that the particle size of the algal cell treated with WAEH decreased significantly as compared to that of the original algae. The reason for this is that the swelling of algal cells is more easily degraded by enzyme in the hydrolysis reaction, and the degree of algal cell rupture is enhanced remarkably after the liberation of the lipid. The abovementioned results suggested that the more complete degradation of the algal cell after WAEH was caused by the high degradation capabilities of the cocktail enzymes toward the algal cell, which were treated by weak alkali in the hydrolysis reaction.

Interestingly, as depicted in the ESI Fig. S3,[†] although greater than 10 wt% of the lipid could be extracted by EH, a larger amount of pigments, such as chlorophyll, were obtained as compared to the case of WAEH.³⁶ In addition, the content of total fatty acids obtained from algal lipid extracted by EH was much smaller than that obtained from algal lipid extracted by WAEH (Table 2); this further confirmed that EH could extract more pigments than WAEH. The possible reason is that a great quantity of algal cells are not disrupted after EH;

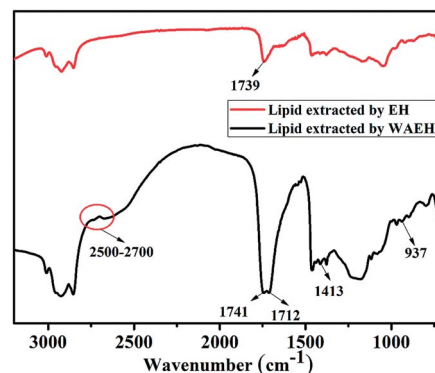


Fig. 3 The FTIR spectra of algal lipids extracted separately by EH and WAEH.

this hinders the permeation of the extracting agent into the cell and hampers the extraction of the lipid. Furthermore, the components of fatty acids in algal lipids extracted by WAEH were more than those extracted by EH, and the result was verified by FTIR spectra of the algal lipid extracted by EH and WAEH (see Fig. 3). It is observed that both the absorption peaks at about 1739 cm^{-1} and 1741 cm^{-1} can be assigned to the stretching vibration of the carbonyl group from the glyceryl ester, whereas the absorption at 1712 cm^{-1} may be associated with the vibration peak of the carbonyl group of free fatty acids.³⁷ This may be ascribed to the movement of the absorption peak of C=O towards lower wavenumbers because of the formation of fatty acid dimers in which hydrogen bonds are generated.³⁸ The absorption peaks at 1413 cm^{-1} and 937 cm^{-1} are separately attributed to the in-plane and out-of-plane bending vibration of O-H from the carboxylic acid dimer, and several weak peaks at 2500 cm^{-1} – 2700 cm^{-1} are assigned to the coupled vibration of frequency doubling and combination of frequency resulting from stretching and bending of the hydroxyl group in carboxylic acid.^{37,39} These results indicate that algal lipids extracted by WAEH contain a certain amount of free fatty acids,⁴⁰ whereas no or lower content is found in that extracted by EH. However, the compositions of algal lipid extracted separately by EH and WAEH are on the whole similar, and the relevant evidence is shown in the Table 2. Therefore, it is confirmed that algal cell swelling occurs in the process of WAP, following which the cell wall is decomposed *via* enzymatic hydrolysis. The constitutions of fatty acids obtained from algal lipid extracted separately by EH and WAEH are roughly same, even if there is a greater difference between their composition and content.

3.5 Investigation of the extraction technology for algal lipid

To further investigate the extraction process of algal lipid, more comprehensive understanding of the wall-breaking process by weak alkaline treatment-aided enzymatic hydrolysis is quite essential. Based on the abovementioned results, an extraction protocol of algal lipid by WAEH is schematically illustrated in Fig. 4. It can be seen that the disruption of cell wall is an imperative procedure to extract lipid from



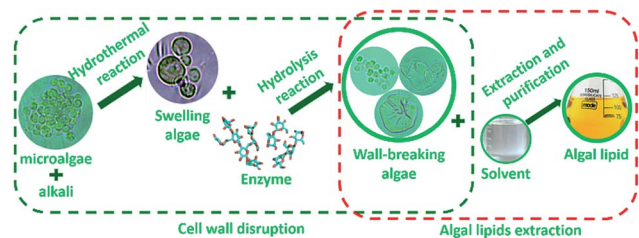


Fig. 4 Schematic of the process of algal lipid extracted by WAEH.

microalgae, which needs two steps of cell swelling and cell wall rupture. As far as the former is concerned, the surface of the original algal cell is not very smooth because of the loss of moisture during the drying process. However, the wrinkled cell walls gradually become smooth since microalgae absorb some amount of moisture during the reaction of alkaline hydrothermal, and subsequent algae cell begins to swell because hydrogen bond networks of native polysaccharide existing in the cell wall are destroyed by alkali.⁴¹ Furthermore, as is well known, swollen algal cells are more easily subjected to the targeted attack derived from enzyme.⁴² For this reason, as the fundamental structural unit building, the cell wall of microalgae, micro-fibrous network within a gel-like protein matrix, can be destroyed with cellulose and lysozyme breaking β (1–4) glucosidic linkages in the polysaccharide skeleton during the hydrolysis, which reduces polysaccharides to soluble oligosaccharides and monosaccharides.^{43,44} Furthermore, cell membrane consisting of phospholipid and glycolipid can be degraded in the hydrolysis reaction since cellulase is a composite enzyme containing multiple hydrolase.⁴⁵ After this, by adding the solvent to the reaction, damaged algal cells are disrupted or deconstructed under the interaction of the intracellular osmotic pressure,⁴⁶ and algal lipids are released into the slurry from the lipid vesicle when the complex solvent permeates into the damaged algal cell. Then, the released lipids are achieved *via* centrifuging the slurry and distilling under vacuum. These results suggest that weak alkali treatment can only make cell wall swell, whereas enzymatic hydrolysis is the crucial procedure for the disruption of algal cell wall. Furthermore, all experimental manipulations of algal lipid extracted by WAEH were carried out at lower temperatures, using a weaker base, and at atmospheric pressure. Therefore, the technology of WAEH makes algal lipid extraction become more complete and economically viable, and the extracted lipid is more suitable as the raw material of the separation of ω -PUFAs.

4. Conclusions

A conclusion that the disruption process of cell wall was composed of swelling by weak alkali and decomposition by enzyme was verified through exploring the effect of WAP on cell wall-breaking. Single and mixed enzymes were screened for superior disruption capacity, and the highest yield (22.18 ± 0.26 wt%) of algal lipid recovery was obtained when the

temperature, time, pH value, algae/enzyme ratio, and mass ratio of cellulase and lysozyme during the enzymatic hydrolysis were 50°C , 5 h, 4.0, 8 : 1 (w/w), and 2 : 1, respectively. Compositional analysis of fatty acids demonstrates that algal lipid extracted by WAEH is applicable as a raw material for the separation of ω -PUFAs.

Conflicts of interest

There are no conflicts to declare.

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