

Isolation and identification of psychrophilic *Chlamydomonas* **sp. ArF0032 from Arctic sea ice**

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ABSTRACT

Microalgae are able to convert solar energy and carbon dioxide to energy storage compounds like oil and starch, which can be exploited for food and biofuels. The Arctic microalgae (ArF0032) isolated from sea ice near Dasan Station in Ny-Ålesund was analyzed to evaluate the optimal growth condition and lipid contents. ArF0032 has an optimal growth temperature of 4 $^{\circ}$ C, reaching densities up to 1.7 × 10⁷ cells/mL. Lipid body formation was visualized by Nile red staining and fluorescence microscopy. Fatty acid methyl ester (FAME) was analyzed using gas chromatography. The ArF0032 showed fatty acid contents, which were dominated by palmitic acid (C16:0), 5,8,11-heptadecatrienoic acid (C17:3), oleic acid (C18:1), linoleic acid (C18:2) and α -linolenic acid (C18:3). The ArF0032 presented maximum percentage (20%) of oleic acid and the content of unsaturated fatty acids (72.1%) was higher than that of saturated fatty acid (9%). Chlamydomonas sp. ArF0032 grows and produces fatty acids at low temperature and may represent a good source for biodiesel production in cold environments.

INTRODUCTION

Arctic's climate is characterized by cold winter with the lowest temperature on earth. Despite the harsh conditions, polar region has a rich diversity of microalgal flora. With regards to potential abilities in Arctic microalgae, there have been trying to search polar algae for use in alternative energy and wastewater treatment. We performed screening for desirable candidates for lipid production and found *Chlamydomonas* species in Arctic ocean. In the present study, potential of *Chlamydomonas* species as a source of biodiesel was evaluated in terms of fatty acid content and growth temperature range.



Fig. 2. Growth curves at various temperatures. Microalgae grew in BBM medium and stationary cultured at 4 °C, 12 °C and 20 °C under white light intensity of 40 μ mol m⁻² s⁻¹



MATERIALS & METHODS

Samples were collected from sea ice near the Dasan station located in Ny-Alesund, Spitsbergen, Norway (78° 50' N, 11° 56' E). The algae were purified by serial dilution followed by plating on agar. Individual green colonies were isolated and inoculated into liquid Bold's basal medium (BBM). To detect lipid droplets, cells were stained with Nile red (1 μ g/ml, 9-diethylamino-5H-benzo[α]phenoxazine-5-one) in acetone. Lipids were extracted in hexane:methyl tert-butyl ether (1:1). Fatty acid methyl esters were analyzed by gas chromatography.

RESULTS

Fig. 3. Detection of lipid body in cell using Nile red staining. Cells cultured in BBM were stained with Nile red and viewed for yellow-gold fluorescence at 488 nm. A) ArF0032, B) *Chlamydomonas reinhardtii* CC-125. Yellow and red colors indicate lipid droplets and chlorophyll, respectively (scale bar = $20 \mu m$).

 Table 1. Fatty acid composition (% total fatty acid) of Arctic Chlamydomonas sp.

 ArF0032.

Fatty acid	ArF0032 (%)	FAME composition : C16:0, Palmitic acid; C16:1 Palmitoleic acid; C16:2, Hexadecadienoic acid; C17:1, cis 10-Heptadecenoic acid; C17:3, 5,8,11-Heptadecatrienoi acid; C18:1, Oleic acid; C18:2, Linoleic acid; C18:3, o Linolenic acid
C16:0	8.97	
C16:1	8.86	
C16:2	5.59	
C17:1	5.75	
C17:3	12.69	
C18:1	20.13	
C18:2	9.28	
C18:3	9.79	
Others	18.94	



Fig. 1. Phylogenetic tree based on nuclear SSU rDNA sequence data. The Arctic green microalgae analyzed for the SSU alignment are shown in bold. SSU sequences from the Arctic green microalga ArF0032 was newly completed for this study and have been deposited in GenBank.

Total 100

CONCLUSION

The Arctic microalga ArF0032 was shown to be psychrophilic microorganisms from its cardinal temperatures for growth. Unsaturated fatty acids (UFAs) were dominant over 70% from ArF0032. It was indicated that the survival of the cells was partly due to the predominance of UFAs, which increased membrane fluidity at low temperature. Arctic *Chlamydomonas* sp. ArF0032 may suggest economic feasibility for lipid production.

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