

Antioxidant Bioactivity of Seaweed Lipid Extracts

Julia Lach, Shane O'Reilly - Atlantic Technological University Sligo, May 2023

Introduction

- Seaweed is a macroalgae which is found in seawater and rocky coastal areas.
- There are multiple bioactive compounds present in seaweed;

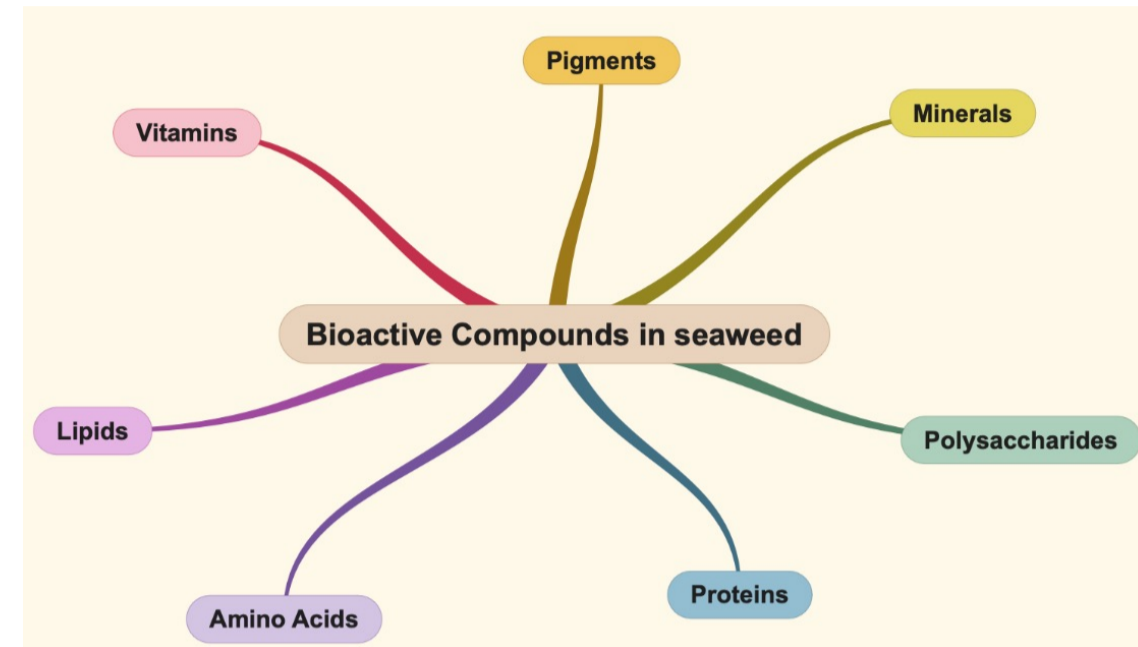
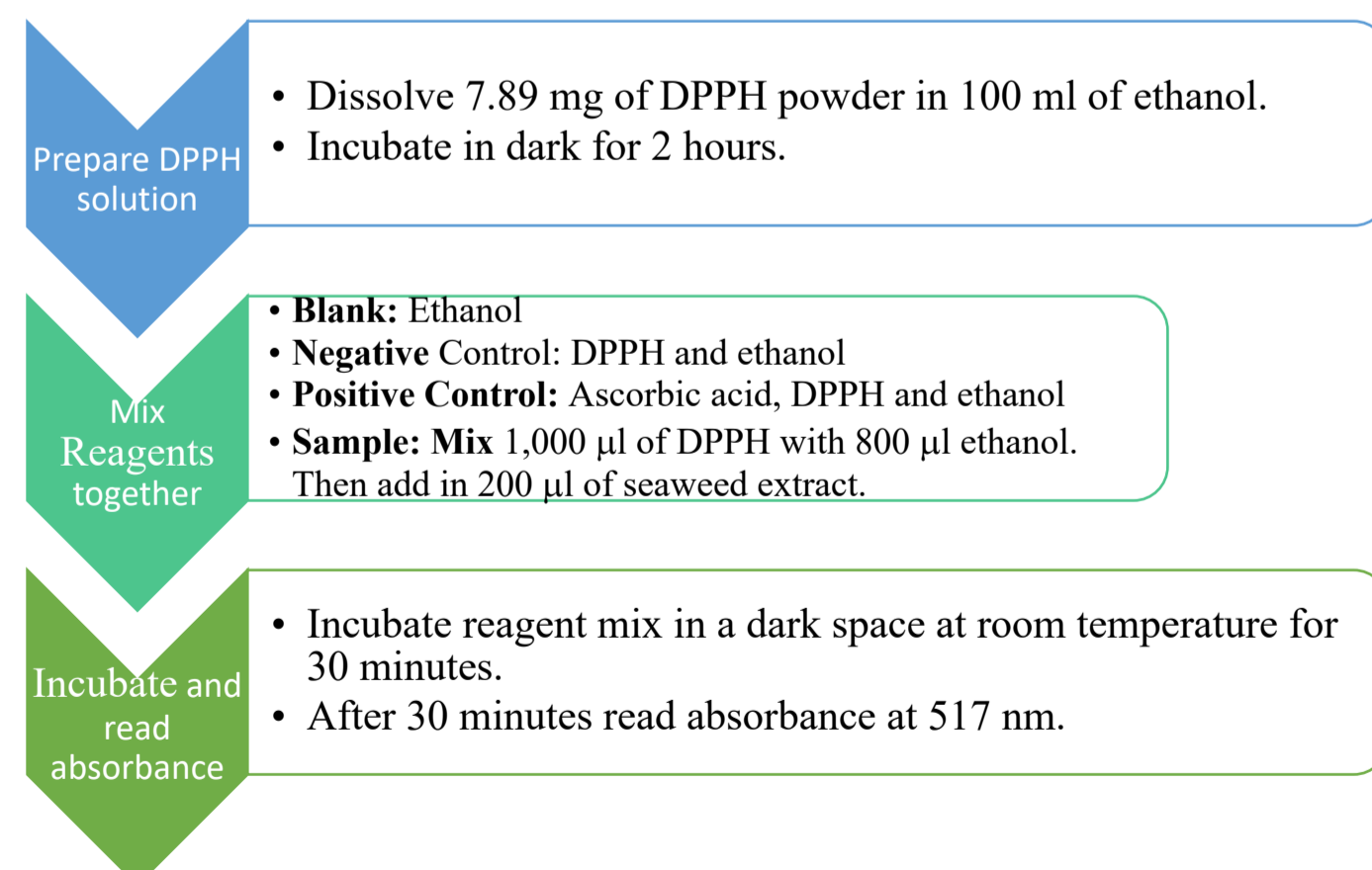


Figure 1. Bioactive compounds that seaweed produce, adapted from El-Beltagi et al (2022)¹.

- These bioactive compounds can generate different bioactive properties; anti-inflammatory, antioxidant, antitumoral, anticancer, antibacterial, e.t.c.

Methods

- 2-diphenyl-1-picrylhydrazil (DPPH) radical scavenging assay² was used to test antioxidant activity.
- Seaweed extracts were taken from original liquid form, dried down using Techne Dri-Block Digital Heater at 42°C .
- Then they were reconstituted with ethanol.



RSA is measured using Radical Scavenging Activity (%) Equation:

$$\text{Radical Scavenging Activity (\%)} = \frac{A_0 - A_1}{A_0} \times 100$$

Where A0 = Absorbance of negative control, A1 = Absorbance of sample.

Results

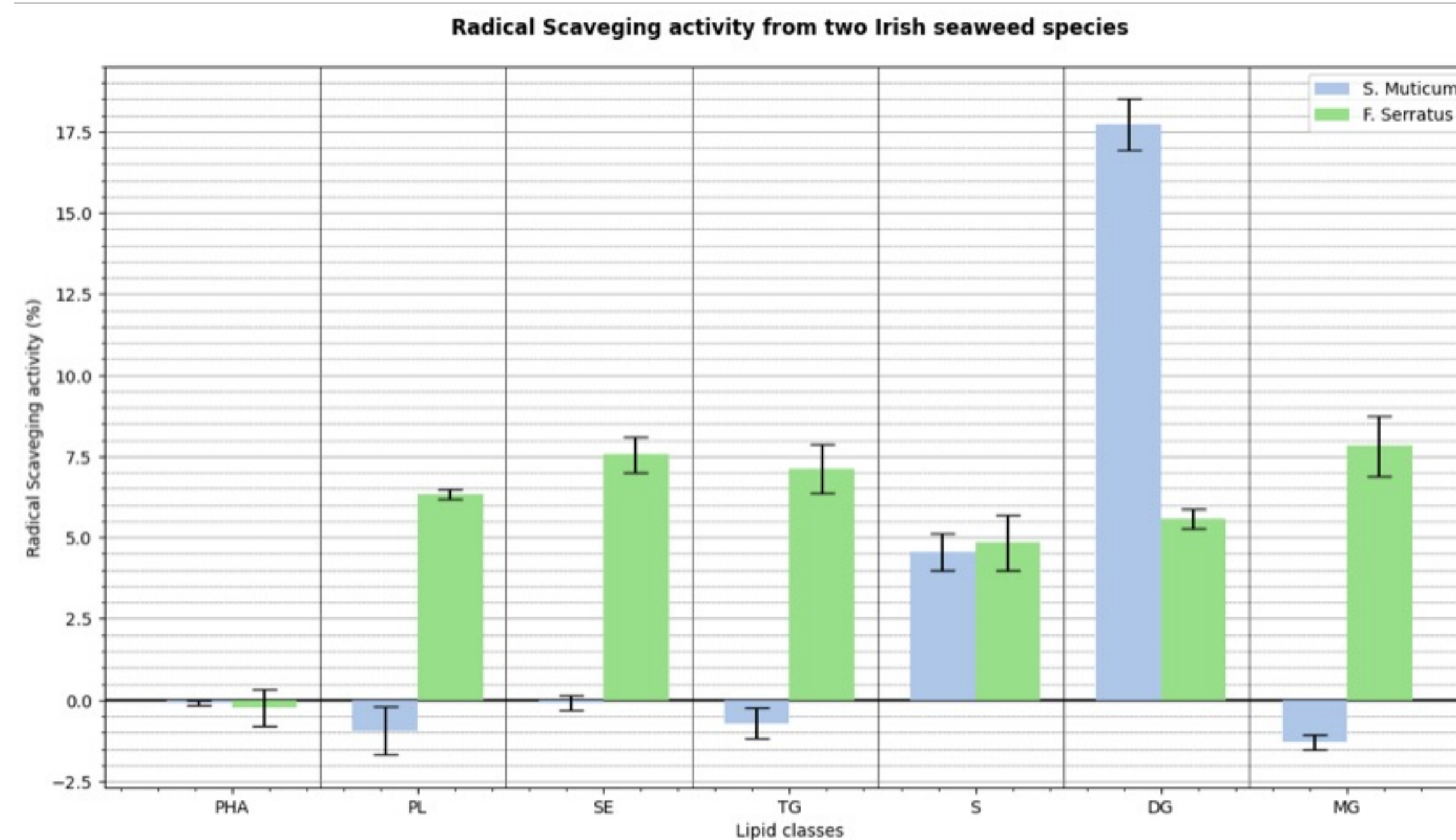


Figure 2. Comparison of Radical Scavenging Activity (%) of *S. Muticum* lipid fractions (blue) and *F. Serratus* lipid fractions (green); PHA= polyhydroxvalcanoate, PL= polar lipids, SE= steryl esters, TG= triglycerides, S= sterols, DG= diacylglycerols and MG= monoglycerides.

The most abundant lipid fractions for RSA was; Diacylglycerols (17.7%) for *S. Muticum* species, Monoglycerides (7.83%) for *F. Serratus* species, and PHA lipid fraction had no RSA for both species.

Table 1. Radical scavenging activity of three Irish Seaweed species.

Seaweed Species	Radical Scavenging Activity (RSA %) Mean ± Standard Deviation	
	0.03g of total Seaweed extract (2g)	0.1g of total Seaweed extract (2g)
<i>S. Muticum</i>	-2.06 ± 0.97	0.111 ± 0.27
<i>L. Digitata</i>	-1.77 ± 1.93	N/A
<i>F. Serratus</i>	3.84 ± 0.27	15.8 ± 3.23

The highest Radical Scavenging Activity was observed from *F. Serratus* at both 0.03g and 0.1g of total Seaweed extract. Low RSA was detected from 0.1g of *S. Muticum* while no RSA from the 0.03g extract. No RSA was seen from *L. Digitata* species.

Conclusion

Some limitations within the research project include;

- Light entering the test tubes and affecting the RSA results
- Results *in vivo* and *in vitro* can be different

Further testing can be performed;

- Increasing concentrations of total seaweed extract.
- Verify composition of each lipid fraction by gas chromatography
- Test other compounds in seaweed

Significance of results

- Total seaweed extract concentration influences antioxidant activity.
- Different lipid fractions display varying amounts of antioxidant activity.

References

- El-Beltagi, H.S., Mohamed, A.A., Mohamed, H.I., Ramadan, K.M.A., Barqawi, A.A. and Mansour, A.T. (2022) 'Phytochemical and Potential Properties of Seaweeds and Their Recent Applications: A Review', *Marine Drugs*, 20(6), 49, available: <http://dx.doi.org/10.3390/md20060342> [Accessed 15 Dec. 2022].
- Kedare, S.B. and Singh, R.P. (2011). Genesis and development of DPPH method of antioxidant assay. *Journal of Food Science and Technology*, [online] 48(4), pp.412–422. doi:<https://doi.org/10.1007/s13197-011-0251-1> [Accessed 15 Jan. 2023].