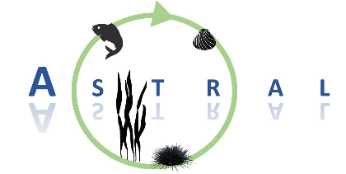
Assessing the commercial potential of the unexploited South African Cape sea urchin, *Parechinus angulosus*, for the seafood market.

****

Aimee Cloete

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Dr Marissa Brink-Hull (UCT Postdoctoral Research Fellow)

Minor Dissertation submitted in partial fulfilment of the requirements for the degree of Master of Science in Applied Ocean Science A purple and orange sea urchin

Description automatically generated

*A close-up of a sea urchin

Description automatically generated*University of Cape Town, Department of Biological Sciences

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# **Declaration**

I declare that this project is my own, unaided work and has not been previously submitted, in whole or in part, for the award of any degree. Where use has been made of the research of others, it has been duly acknowledged in the text. This project is carried out in the Department of Biological Sciences under the supervision of Dr Marissa Brink-Hull, Dr Brett Macey and Professor John Bolton.

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# **Acknowledgements**

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(acknowledgement of family, friends etc who have assisted..)

# **Abstract**

Wild *Parechinus angulosus* were collected in the intertidal and shallow subtidal on a rocky shore in front of the DFFE Marine Research Aquarium in Sea Point, Cape Town (33° 55' 6.492'' S, 18° 22' 52.572'' E) to assess their feasibility as a new market product for South Africa.

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# **List of acronyms**

ASTRAL All Atlantic Ocean Sustainable, Profitable and Resilient Aquaculture

AAEC Aquaculture Animal Ethics Committee

CIE International Commission on Illumination

DEFF Department of Environment, Forestry and Fisheries

DOM Dissolved Organic Matter

GSI Gonadal Somatic Index

IMTA Integrated Multi-Trophic Aquaculture

ISA

MRA Marine Research Aquarium

POM Particulate Organic Matter

SA South Africa

SGR Specific Growth Rate

# **Introduction**

## *Parechinus angulosus*

The Cape sea urchin, *Parechinus angulosus* (Leske, 1778), is endemic to southern Africa and belongs to the Parechinidae family. It is widely distributed from Lüderitz, Namibia to Umhlali, Kwa-Zulu Natal, South Africa (SA) with dense concentrations around the Cape Peninsula (Fricke, 1980). The Cape Peninsula is a region with a strong seawater temperature gradient (~12oC - 17 oC) (Muller et al., 2012; Smit et al., 2013), due to the overlap of the cool Benguela (SA west coast) and warm Agulhas (SA south coast) provinces, and the co-occurrence of species from both marine provinces results in a high species diversity (Leliaert et al., 2000). The region of overlap, referred to as the ‘Western Overlap’ in previous literature, is characterized by kelp beds created by *Ecklonia maxima* (to depths of 8m) and *Laminaria pallida* (predominantly 8-14m and up to 20m depths) wherever there are rocky substrata in the shallow subtidal (Leliaert et al., 2000). The distribution of the Cape urchin is closely associated with the southern African kelp which characterize its habitat. They are prey, although not preferentially, for rock lobsters, *Jasus lalandii* (Van Zyl et al., 2003). Due to low predatory activity their population density is high (\*idea of population density value). The Cape urchin is an important grazer in the southern African kelp bed ecosystem (Greenwood, 1980) where they can feed opportunistically on marine invertebrates and biofilm but primarily feed on macroalgae (Freeman, 2006; Loiderios & Gracía, 2006; Russell et al., 2018; Saucede et al., 2006; Ma et al., 2021), primarily *Ecklonia maxima* (Fricke, 1979).

In contrast to urchins from kelp bed ecosystems in other regions, e.g. *Stronglyocentrotus spp,* the Cape urchin does not sever kelp stipes or form feeding fronts which decimate kelp populations to the point of a regime shift from an algal-dominated ecosystem to an urchin-dominated alternative stable state known as an “urchin barren”(Anderson et al., 1997). Urchin barrens have much lower primary productivity and habitat structural complexity, threatening the health of kelp bed ecosystems in many areas of the world (Filbee-Dexter & Scheibling, 2014). The Cape urchin regulates kelp density by grazing on the young sporophytes and kelp debris (Fricke, 1979). Although Cape urchins can climb up and actively graze on the kelp stipes under calm conditions (Fricke, 1979; Morris & Blamey, 2018) this behavior is uncommon due to the generally turbulent sea conditions which produce kelp detritus and provide an ample supply of organic matter for the urchins, reducing the likelihood of active grazing (Fricke, 1979). Although sea urchins have a reputation for destruction in some regions, they are keystone species in the structuring of kelp forest communities (Estes & Palmisano, 1974; Scheibling et al., 1999; Ma et al., 2021).

Sea urchins are a key trophic intermediate in kelp bed ecosystems, through shredding, sloppy feeding and egestion of kelp litter they make kelp detritus more accessible to benthic consumers which are important resources for fish and other predators (Yorke et al., 2019). By grazing on young sporophytes, they also control the rate of kelp settlement (Branch, 2013), increasing the available surface area for other species within the kelp bed ecosystem. Thus, *P. angulosus*, influences the habitat structure and transfer of energy for a guild of consumers with the kelp bed ecosystem. (reference). The disappearance of sea urchins would rapidly increase macroalgae recruitment, decreasing the available surface area for benthic fauna. In addition, without the sea urchin trophic linkage the macroalgae debris would be inaccessible to detritovores, increasing competition for resources and impacting the diversity and trophic structure of the ecosystem. *Parechinus angulosus* maintains the balance and productivity of kelp bed ecosystems in South Africa, their ecological significance is clear. However, their economic and cultural significance requires further study.

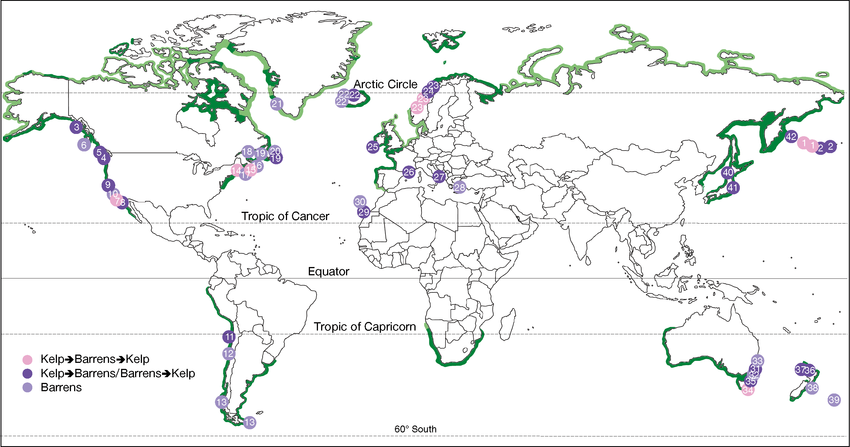


Figure 1 Global occurrence of sea urchin barrens documented throughout the range of kelp. Numbered locations (colored circles) indicate areas where urchin barrens have been documented: (pink) through the course of multiple phase shifts between kelp beds and barrens, (dark purple) following a single phase shift from a kelp to a barrens state, or vice versa, and (light purple) in areas that might otherwise support kelp, although a phase shift has not been observed. Observed range (dark green) and potential range (light green) of kelp, based on light and temperature requirements, is also shown (adapted from (Filbee-Dexter & Scheibling, 2014).

## The sea urchin market

Sea urchins are harvested for their gonads (roe) which is a prized delicacy in Asia, Europe (France, Italy & Spain), South America (Chile), North America and the Mediterranean (Stefánsson et al., 2017). The market is mostly traditional, dominated by Japan which accounts for ~90% of the world demand (Stefánsson et al., 2017). In Japan only a small portion of sea urchin demand is for household consumption, it is mostly served in sushi bars and at special celebrations such as wedding banquets and parties (Sun & Chiang, 2015). The demand for sea urchin roe in Japan is therefore seasonal, with the largest amount being consumed over the Japanese holidays in December (Sonu, 2017). Domestic markets exist in many sea urchin harvesting countries (e.g., Chile, New Zealand and the Philippines) (Stefánsson et al., 2017). Sea urchin gonads are used as a basic ingredient in several Mediterranean dishes, where it is blended into sauces, with pasta and breads and demand is constantly increasing (Furesi et al., 2016; Stefánsson et al., 2017).

Japan was the largest global harvester until 1987, since then Japanese harvests declined significantly due to overfishing and to meet the increasing demand Japan became dependent on imports (Sonu, 2017). Key suppliers to Japan are Chile, Russia, North Korea, the United States and Canada (Stefánsson et al., 2017). Global landings peaked at ~120 000 tonnes in 1995 (FAO, 2016; Stefánsson et al., 2017) since then demand has increased, resulting in overfishing and stock decline. (Johnson et al., 2012; Medellín-Ortiz et al., 2020; Nane & Paramata, 2020; Stefánsson et al., 2017). Where appropriate harvest management strategies have not been put in place, sea urchin fisheries, e.g., in France (Mediterranean and Atlantic), Iceland, Ireland, South Korean (inshore) and the Philippines, have collapsed (Micael et al., 2009). Increased demand and decreased supply have influenced the value of sea urchin roe. From 1975 to 2016, Japanese imports of sea urchin products increased more than six-fold in volume and nine-times in value (Sonu, 2017; Stefánsson et al., 2017).

Global harvest volumes of sea urchin are difficult to estimate as reporting by FAO often lumps all echinoderms together. Global harvests were estimated at ~76 000 tonnes in 2014 (FAO, 2016) and it is assumed not all of it could be sold due to damage or unfavourable size (Stefánsson et al., 2017). The sea urchin market is dominated by only a handful of species (Figure 2) with green (*Strongylocentrotus pulcheriius*), red (*Strongylocentrotus franciscanus*), and purple sea urchins (*Strongylocentrotus intermedius*) in highest demand (Stefánsson et al., 2017). Thus, there is potentially an unmet demand on the Japanese market for good quality sea urchin products at the appropriate price, particularly with less current supply to the market, indicating an opportunity for new entrants into the market (Stefánsson et al., 2017).

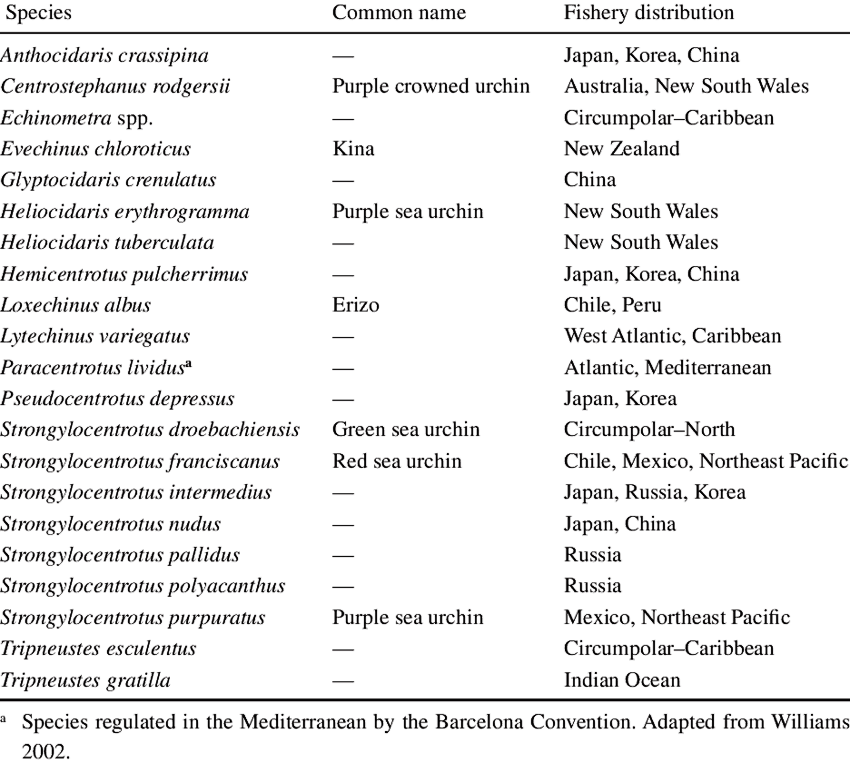


Figure . Sea urchin species with a major contribution to the modern fishery (adapted from Micael et al., 2009)

## Research rationale

The collector sea urchin, *Tripneutes gratilla* (Linnaeus, 1758), a sought-after species on the sea urchin market which is distributed on the eastern coast of Southern Africa, has been identified as a species with potential for aquaculture production in South Africa, due to it’s fast growth rate (\*) and marketable roe production (Brink et al., 2018; Brink-Hull et al., 2022; Cyrus et al., 2013; Scholtz et al., 2013). The feasibility of producing this warm – water species is limited however, to regions of the South African coastline that have temperatures above \* or to farms which can afford the high heating costs (\*) (how much of the coast is warm vs cold?). *Parechinus angulosus* has a much wider distribution and copes with the strong temperature gradient along the South African coast, making it a more accessible and resilient species for South African sea urchin aquaculture investors and existing farms in areas along the cooler regions of the coast.

The powerhouse of South African aquaculture is abalone, *Haliotis midae,* which is produced primarily for export to Asia and outshines all other South African aquaculture products in terms of product value, employment and production volume with a 76% share of the overall value generated by the aquaculture sector (Britz & Venter, 2016). The high value abalone species and the Cape sea urchin, *Parechinus angulosus,* have a similar preferred temperature range (12 – 20 °C) (Fricke 1980; Britz et al. 1997; Day and Branch 2002a) and commonly occur together in nature, particularly during the juvenile stages of the abalone life cycle (Day and Branch 2000, 2002a).

Laboratory experiments by Day & Branch (2002a) showed that juvenile abalone prefer to shelter beneath urchins rather than under rocks and crevices. There is insufficient microalgae growth under rocks and crevices to meet the dietary requirements of juvenile abalone (Day & Branch, 2002a) and therefore, the juvenile abalone need to leave their shelter and expose themselves to predators, such as octopus, rock lobster and predatory fish while grazing (Nepgen, 1982; Smith, 1999; Mayfield et al., 2000), when grazing. However, juvenile abalone that shelter beneath urchins can graze on the uneaten algae made available by the sea urchins through their sloppy feeding, shredding and egestion which reduces or potentially eliminates their exposure to predators while grazing (\*). Previous work done for my honours research project (2022), by the same research group, studied the impacts of urchin waste products on abalone growth and found that supplementing hatchery-reared juvenile abalone diets with Cape sea urchin faecal matter enhanced the growth rates of juvenile abalone (Cloete, 2022). Considering the co-habitation of sea urchins and abalone in natural environments, as well as the potential symbiotic relationships that exist between them, they could be co-cultured on existing abalone farms as a method of improving juvenile abalone health through the trophic transfer of microbial communities and as a method to improve the sustainability of the South African abalone industry by introducing a commercial product. However, the feasibility of the Cape sea urchin as an additional value-added product has not been investigated as yet.



Figure 3. Image of juvenile abalone (*Haliotis midae*) sheltering beneath Cape sea urchins (*Parechinus angulosus*) in Simon’s Town, Cape Town, South Africa (Peter Southwood, 2005).

## Research aim and objectives

One of the major factors influencing the marketability of sea urchins is their gonad colour and texture (Shpigel et al., 2005). The effects of different temperatures and feeding regimes on the growth performance, optimal gonad colour and gonadal somatic index (GSI) of this species has not been assessed, this project aims to address these knowledge gaps. The spinal colour variation of the Cape sea urchin may potentially impact their gonad colour and thus, may add commercial interest to the species.

* + 1. **Research Aim**

The aim of the study is to assess the potential of the Cape sea urchin, *Parechinus angulosus*, as a new entrant for the sea urchin market.

* + 1. **Research Objectives:**

Assess the effects of different dietary and temperature treatments on:

1. Survival rate (%)
2. Somatic growth rate in terms of weight and size
3. Gonad quantity (GSI)
4. Gonad quality (gonad colour)
5. Gonad development
6. FCR
7. Effects of test colour on gonad colour

Dietary treatments

* *Ulva lacinulata* (U),
* *Ecklonia maxima* kelp (K),
* 16% Ulva inclusion formulated feed (F) (Cyrus et al., 2013)
* combination of the forementioned diets (U, K, F) rotated on a weekly basis to form a mixed diet (M)

Temperature treatments

* Ambient sea water temperature at Sea Point DFFE mariculture laboratory on the Cape southwest coast
* Sea water warmed to 18°C.

# **Literature review**

## Aquaculture’s contribution to seafood supply

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Description automatically generated with medium confidenceAquatic foods play a crucial role in ensuring food and nutrition security, particularly for vulnerable coastal populations, by providing accessible and affordable sources of proteins and micronutrients (FAO, 2022). Human population growth has contributed to an increase in marine exploitation and subsequent declines in biologically sustainable fishing stocks (Pham et al., 2023). (add more evidence or detail about reduced fisheries capacity) The increasing demand for seafood products, coupled with the diminishing productivity of wild-caught marine fish stocks, mostly due to the overexploitation and illegal exploitation of fish stock, positions the aquaculture industry as a significant contributor to the global seafood supply (Granada et al. 2016) (Figure 3). Aquaculture production significantly supplements capture fisheries production (Longo et al., 2019) contributing 49.2 % to the global production of aquatic animals in 2020, but despite the great diversity in farmed aquatic species, only a small number of “staple” species dominate aquaculture production (FAO, 2022). Further development of the aquaculture industry is necessary to meet growing demand for seafood products.

Figure . World capture fisheries and aquaculture production excluding aquatic mammals, crocodiles, alligators and caimans. Data expressed in live weight equivalent. (FAO,2022).

## Aquaculture sustainability

Food security is an on-going concern within the seafood industry in the Global South, while the Global North tends to prioritize the sustainability of the industry (Belton et al., 2020). A balance between these two important goals is critical if the industry wishes to meet future demands for seafood. The aquaculture industry is known to be an important consumer of wild-caught marine fish stocks by using fishmeal in aquafeeds, raising concerns about the sustainability of the industry. Aquaculture effluent is another growing concern for the rapidly expanding industry. Generally, (with perhaps the exception of some extractive species?) aquaculture practices require large amounts of water of good quality. Uneaten feed and waste products, result in the accumulation of suspended solids, otherwise known as particulate organic matter (POM) and dissolved organic substances (DOM), which contain compounds such as nitrogen and phosphorus, and possibly also other chemicals that are used to prevent diseases of aquatic species (Jegatheesan et al., 2011). This effluent water is discharged into the environment, which if untreated can be detrimental to the surrounding environmental and human health. Thus, intensive development of aquaculture has raised a range of environmental concerns such as effluent discharge, excessive use of resources and dependence on commercial feed with high fishmeal content (Granada et al., 2016) placing a spotlight on the need for research on sustainable aquaculture systems and practices.

Integrated multi-trophic aquaculture (IMTA) is an advanced form of aquaculture considered a suitable approach to limit aquaculture nutrients and organic matter outputs through biomitigation (Granada et al., 2016). In IMTA systems, nutrients from uneaten feed and excreted waste from fed species become food for extractive species (FAO, 2022). Converting the waste products from one species into a valuable resource for another reduces the amount of nutrients released into the environment while enhancing overall productivity. The extractive species in IMTA systems are both traded as a commodity and used as a biofiltration system, which increases their value to the farm, this is especially important for extractive species with low commercial value or species which are new entrants to the market. The implementation of IMTA systems can increase the efficiency of aquaculture systems and contribute to the development of a sustainable aquaculture industry, particularly when species that are ecologically compatible are co-cultured (Kang et al. 2003; Kim et al. 2015). IMTA has the potential to reduce environmental impacts, increase profitability and diversify commercial production in a sustainable way.

(Why is diversification important? how can IMTA contribute towards this? importance of low trophic species)

(Describe SA implementation of IMTA: abalone – ulva system) Considering the co-habitation of *Parechinus angulosus* and *Haliotis midae* in natural environments, they could be co-cultured within an IMTA system on existing abalone farms. Further improving the sustainability of the most valuable aquaculture sector in South Africa. However, for IMTA systems to succeed, both species being co-cultured should have commercial potential.

## Echinoculture research and practices

The primary driver of sea urchin market value is gonad quality (Teck et al., 2018). Gonad quality is determined by a number of factors including colour, texture, taste and gonad somatic index (GSI) (Cyrus et al., 2015). Quality of the sea urchin fishery is highly variable and dependent on the reproductive state of the organism (Teck et al., 2018). The price differential paid for sea urchin roe across varying reproductive stages can be substantial, with early stages (prior to spawning) being preferred what stage is preferred and why? (Teck et al., 2018). However, good health is required for good gonad production. The health and well-being of sea urchins is also determined by a number of factors incl. …

* (What are factors that have affected gonad quality/growth rate/feeding rate for other species?)

Live sea urchin products are in the highest demand. Japan imported ~ 11 000 tonnes of live sea urchin in 2016 valued at 183 million US dollars, a six-fold increase in volume and nine-fold increase in value since 1975 (Sonu, 2017). (processed) Sea urchin product (live and processed) prices are dependent on a number of factors including appearance (color, quality), origin (species, region of harvest), palatability (flavour, texture), demand, distribution, form and processing(Stefánsson et al., 2017) .

Although quality of roe is the most important factor in determining prices, total supply (domestic and imported roe) is also significant (Sonu, 2017).

* Has anyone done anything similar? What did they find?
* Do sea urchins have other uses besides consumption?



Figure 4: Image of Cape urchin, *P. angulosus*, harvested by Veld and Sea for consumption

## Project expectations

This project

* Motivation for the factors I am considering & methodology followed:
  + Feeds chosen: is it what they naturally eat? (e.g., *Ecklonia maxima* characterizes the environments usually populated by *P. angulosus*)
  + Temperatures chosen
  + How does my methodology compare with other studies?
* Project hypotheses

# **Materials and methods**

## Ethics statement

Wild *Parechinus angulosus* were collected in the intertidal and shallow subtidal on a rocky shore in front of the DFFE Marine Research Aquarium in Sea Point, Cape Town (33° 55' 6.492'' S, 18° 22' 52.572'' E). This site is not privately owned or protected in any way, according to South African legislation (SAFLII, 2019). A fisheries research and development permit to collect, possess, transport, and engage in scientific investigation or practical experiment with the animals was issued successfully in terms of section 83 of the Marine Living Resources Act, 1998 (Act 18 of 1998) with permit reference number RES2023/65. All experimental procedures on animals followed the welfare guidelines of the DFFE. Daily biosecurity checks were performed for the duration of the study according to the biosecurity standard of the DFFE.

## Sea urchin collection

The sea urchins (*Parechinus angulosus*) were collected from the rock pools in front of the Marine Research Aquarium in Sea Point in May 2023. A total of 650 individuals of an average size of 4cm diameter were collected and immediately transported to plastic tanks with a flow-through system at the Marine Research Aquarium. Prior to the start of the experiment the urchins were starved for three weeks to reduce their gonad weight and ensure that all animals had a similar gonad state prior to the start of the trial. Thereafter, the urchins were stocked into oyster mesh baskets (L x W x D: 38 x 28 x 30 cm; mesh size: 6 mm) suspended in smaller plastic tanks (L x W x H: 42 x 36 x 30 cm) at 19 animals per basket (stocking density: ~ 0.6 kg m-2) and fed *Ecklonia maxima* for two weeks while they acclimatised to the experimental system. A similar test diameter size range of urchins (Table 1) were stocked in each basket across the various treatments to mitigate against growth rate differences due to different sized animals. *Parechinus angulosus* has a wide range of spinal colours (pink, light purple, dark purple, orange and red). Where possible, equal ratios of urchins with different test colours were selected for each basket.

Table 1. Initial size (test diameter in mm) and wet weight (g) of *P. angulosus* at the start of the trial

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Diet** | **Temperature** | **Treatment** | **Mean urchin test diameter (mm)**  **(mean ± se)** | **Mean urchin wet mass (g)**  **(mean ± se)** |
| formulated | ambient | fa | 32.91 ± 0.83 | 16.75 ± 0.77 |
| formulated | warm | fw | 35.61 ± 0.73 | 18.92 ± 0.75 |
| mixed | ambient | ma | 32.52 ± 0.85 | 15.00 ± 0.70 |
| mixed | warm | mw | 33.81 ± 0.90 | 16.91 ± 0.87 |
| ulva | ambient | ua | 33.71 ± 0.88 | 16.75 ± 0.83 |
| ulva | warm | uw | 33.80 ± 0.81 | 17.04 ± 0.79 |
| kelp | ambient | ka | 34.19 ± 0.96 | 17.83 ± 0.95 |
| kelp | warm | kw | 32.40 ± 0.72 | 15.70 ± 0.72 |

## Experimental setup

A flow-through experimental system was utilized for the trials, consisting of 32 rectangular plastic tanks (L x W x H: 42 x 36 x 30 cm). There were four tanks (replicates) for each treatment and each tank had a volume of 40 l; when accounting for the height of the outflow. Seawater for the ambient and the heated tanks was pumped from the kelp beds in front of the DFFE Marine Research Aquarium (MRA). Before entering the experimental systems, seawater passed through a drum filter and then a sand filter prior to entering a sump tank at the highest level of the MRA. For the ambient system, water from the sump was gravity fed into the experimental tanks at a rate of ~1000 L/min (~1.5 tank turnovers/hr). Conversely, for the heated system, water from the main sump at the MRA was gravity fed into two interconnected 2,500 L JoJo tanks where the water was constantly recirculated through a heat pump set at 19 oC before entering the experimental tanks, at a rate of ~ 1000 L/min (~1.5 tank turnovers/hr). This experimental system included two temperature treatments: ambient (A: ambient) incoming water and a consistent temperature (W: warm) of 18°C (temperature controlled using a heat pump).The exact temperature in the ambient and heated experimental tanks was continuously recorded at 30-minute intervals using a temperature probe (Star Oddi Starmon Mini temperature recorder) and the average temperature over the entire experimental period was (mean ± se) 15.36 ± 0.01 oC and 18.88 ± 0.01 oC for the ambient and heated systems, respectively. Constant aeration was provided in all tanks using airstones. Effluent water was returned directly to the ocean through the main effluent pipe of the MRA. The internal surfaces of tanks were manually cleaned of their sediments and fouling organisms twice a week, using a siphon and synthetic fibre brush.

Four feeding regimes were tested in quadruplicate: *Ulva lacinulata* (U), *Ecklonia maxima* kelp (K), a formulated feed containing 16% *Ulva* (F) (Cyrus et al., 2013), as well as a rotation of the forementioned diets (U, K, F) on a weekly basis to form a mixed diet (M), resulting in a total of 16 tanks (320 sea urchins). To ensure that feed was not a limiting factor for growth and development all feeding regimes were administered ad libitum three times a week ensuring that feed was constantly available, to avoid overfeeding, the amount of feed added at the start of the experiment was calculated as a percentage of the total body weight within each tank (U: 8%, F: 1.8%, K: 10%). Kelp fronds were cut into small segments (L x W: width of kelp frond x 5 cm) to increase surface area of feed available.

*(Add schematic)*

## Data collection

### *Survival rate, somatic growth and feed conversion ratio*

Somatic growth was measured in terms of urchin total wet weight (g) and diameter (cm) at six measurement timepoints (T0: initial, T1: 4 weeks, T2: 8 weeks, T3: 13 weeks, T4: 18 weeks, T5: 23 weeks) over the course of the study. At every timepoint all sea urchins were individually weighed and measured using standardised photographs taken with an iPhone 8. All images were processed using the “Urchin Vision” software developed by De Vos et al. (2023) and the average diameters and weights of animals in each tank were recorded.

Feed conversion ratios were measured for each tank by determining feed consumption over a 7-day period, divided by the corresponding wet weight increase over the 7 days using the following formulae:

All urchins were weighed at the start of the week (for week 0, 4, 8, 13, 18 and 23), these values were used for the somatic growth data. To allow the sea urchins to adapt to the treatments before feed consumption was measured, feed consumption measurements were recorded from week 8 onwards.

All urchins were measured again at the end of the week (for weeks 8, 13, 18 and 23), these values were used to calculate wet weight increase over the 7 days. The total feed consumed per tank over the 7 days was calculated as the difference between the feed introduced and the residual feed removed from the tank. The amount of feed added to the tank was weighed, in grams, at the start of the week.

Macroalgae tissues (Ulva and Ecklonia maxima) were still intact after 7 days when residual material was removed and therefore, no significant process of degradation and weight reduction within the experimental time was considered. These tissues were spun using a salad spinner to remove excess moisture before they were weighed. The formulated feed however, was significantly degraded, and the residual formulated feed in each tank was thus removed every 3 days and new formulated feed was weighed (g) and added for the experiment. The residual formulated feed was placed into foil weigh boats and dried to a constant weight in an oven at 60oC for 48 hours to remove excess moisture before being weighed. Feed consumption was calculated in milligrams of feed (dry weight) consumed per tank, per week. The feed consumption was then divided by the number of animals in the tank and by the number of days for the feed consumption experiment to calculate the average feed (dry weight in milligrams) consumed per animal, per day.

The somatic growth data collected in terms of weight (g) and size (cm diameter) was averaged per tank and used to calculate the specific growth rate (SGR). SGR (% growth/day) of each treatment was calculated for five intervals (between each respective timepoint) using mean weight (g) and size (mm) measurements per timepoint per tank per treatment:

Where mf = measurement final (g or cm); mi = measurement initial (g or cm); t = time (days) between final and initial measurements.

The intervals for the five specific growth rates calculated:

1. Week 4: week 0 - week 4 (t = 29 days)
2. Week 8: week 4 – week 8 (t = 29 days)
3. Week 13: week 8 – week 13 (t = 32 days)
4. Week 18: week 13 – week 18 (t = 36 days)
5. Week 23: week 18 – week 23 (t = 36 days)

### *Gonad quantity, quality and development*

Prior to dissection, the urchin was briefly placed in a mesh basket to drip dry, draining excess seawater from the sea urchin body. Whole animal wet weight (g) was recorded after which they were dissected in half using dissection scissors and tweezers to remove the gonad segments from the test. All other visceral tissues attached to the gonad were cleaned off before quantifying the total gonad wet weight (g) and calculating the gonadal somatic index (GSI) (%) using the following formulae:

Gonad quality was measured in terms of gonad colour, which was measured using a hand-held fibre-optic spectrophotometer (Lovibond® LC 100 spectrocolorimeter). The L\*a\*b\* colour space is used to quantifiably define colours (CIE reference) within the food industry (Suckling reference) and for sea urchin gonads (Suckling references). Three replicate measurements of CIE L\* (intensity or lightness), a\* (hue or redness) and b\* (chroma or yellowness) were recorded from each sampled gonad (Onomu et al., 2020) and averaged so that each urchin sampled had one set (CIE L\*, a\* and b\*) of colour measurements.

One gonad segment per urchin was kept for gonad development assessment, through histological analysis, to ascertain the gametogenic state of the urchin. The gonad segment was kept in a 50ml test tube and fixed in Davidson’s fixative (formula?) for a period of 48 hours. After 48 hr of immersion in the fixative, the samples were transferred to 70% ethanol for storage prior to paraffin histology (Bucke, 1989). Gonad maturity was analysed according to the method described by Cyrus, Bolton, and Macey (2015). Gonads were categorized into one of 6 different maturity stages of echinoid gametogenesis, namely (1) recovery, (2) growing, (3) premature, (4) mature, (5) partly spawned, and (6) spent (Vaïtilingon, Rasolofonirina, & Jangoux, 2005) (Table 2). Gonads having little or no gametogenic activity were considered as high-quality, that is gonads in the growing or premature stages (Onomu et al., 2020).

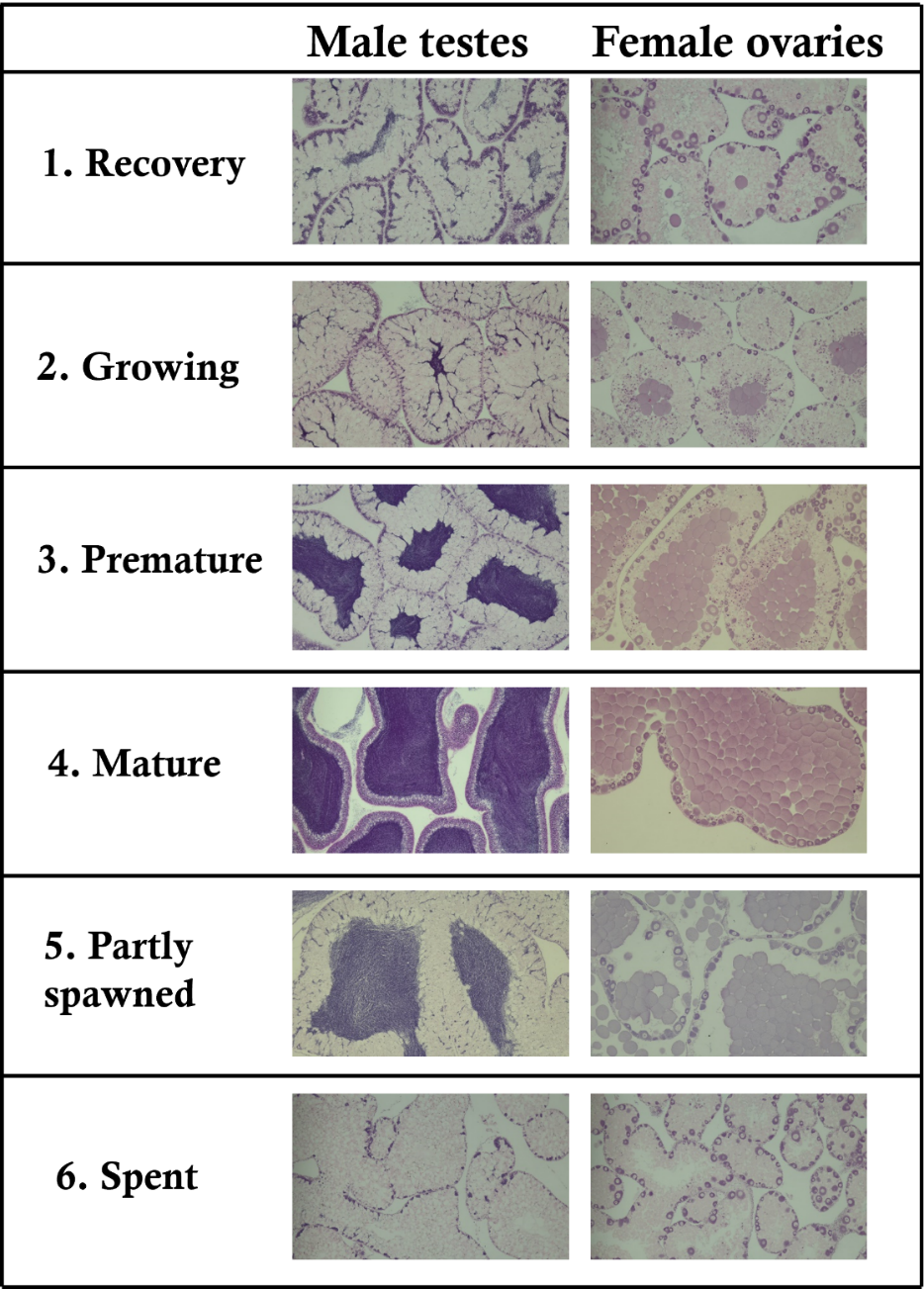


Figure 5: Gonad histology of *P. angulosus* male testes and female ovaries depicting the respective gonad maturity stages: (1) recovery, (2) growing, (3) premature, (4) mature, (5) partly spawned, and (6) spent

## Statistical Analyses

Data was stored in Microsoft Excel (V16.0.17126.20126) and data analyses and plots were performed using R statistical software (R Core Team, 2023). This study involved multiple datasets: survival rate, somatic growth rate (in terms of weight and size), gonad quantity (GSI), gonad colour (CIE L\*a\*b values) and gonad development (maturity).

Each dataset was tested for normality (Shapiro-Wilk) (Shapiro & Wilk, 1965) and homogeneity of variance (Levene’s test using *car* package; Fox & Weisberg, 2019) (Levene, 1960) and if these assumptions were met then a one-way analysis of variance (ANOVA) (Fisher, 1992) was conducted to assess the dietary treatment effect and a two sample t-test for the temperature treatment effect. Where significant results were found for dietary treatment effect, Tukey’s (Keselman & Rogan, 1977) post-hoc pairwise comparisons were conducted to determine dietary treatment group differences.

If data failed to meet the ANOVA assumption of normality following either a log or square root transformation, then a nonparametric Kruskal Wallis test (Kruskal & Wallis, 1952) was conducted for dietary treatment effects and a Mann-Whitney U test for temperature treatment effects. Where significant results were found for dietary treatment effect, Dunn’s (Dinno, 2015) post-hoc comparisons were conducted to determine dietary treatment group differences (using the *dunn.test* package). Due to multiple testing, a Bonferroni correction was applied to reduce the type I error rate.

If data had a normal distribution but failed to meet the assumption of homogeneity, then a Welch ANOVA (Welch, 1951) was conducted for dietary treatment effects and a two sample t-test with unequal variance for temperature treatment effect. Where significant results were found for dietary treatment effect, Games-Howell (Games & Howell, 1976) post-hoc comparisons were conducted to determine treatment group differences (using the *rstatix* package).

F test

# **Results**

## Temperature logs

Temperature was constantly monitored, due to seasonality and technical issues (power failures due to loadshedding and broken generators) there was a large degree of variability within both temperature treatments (Figure 6). From October to mid-November the DFFE MRA generator required repairs and was not working optimally. The most severe disruptions were near two data collection time points (week 18 and week 19). When the power was off there was no waterflow into the tanks and the heaters for the warm temperature treatment had no alternative power source. However, the temperature remained relatively constant for the heated tanks during this period.

(Note: I want to do an F test to compare the variances between the two temperature treatments but am struggling to normalise the data and cannot find information about a non-parametric equivalent for an F test)

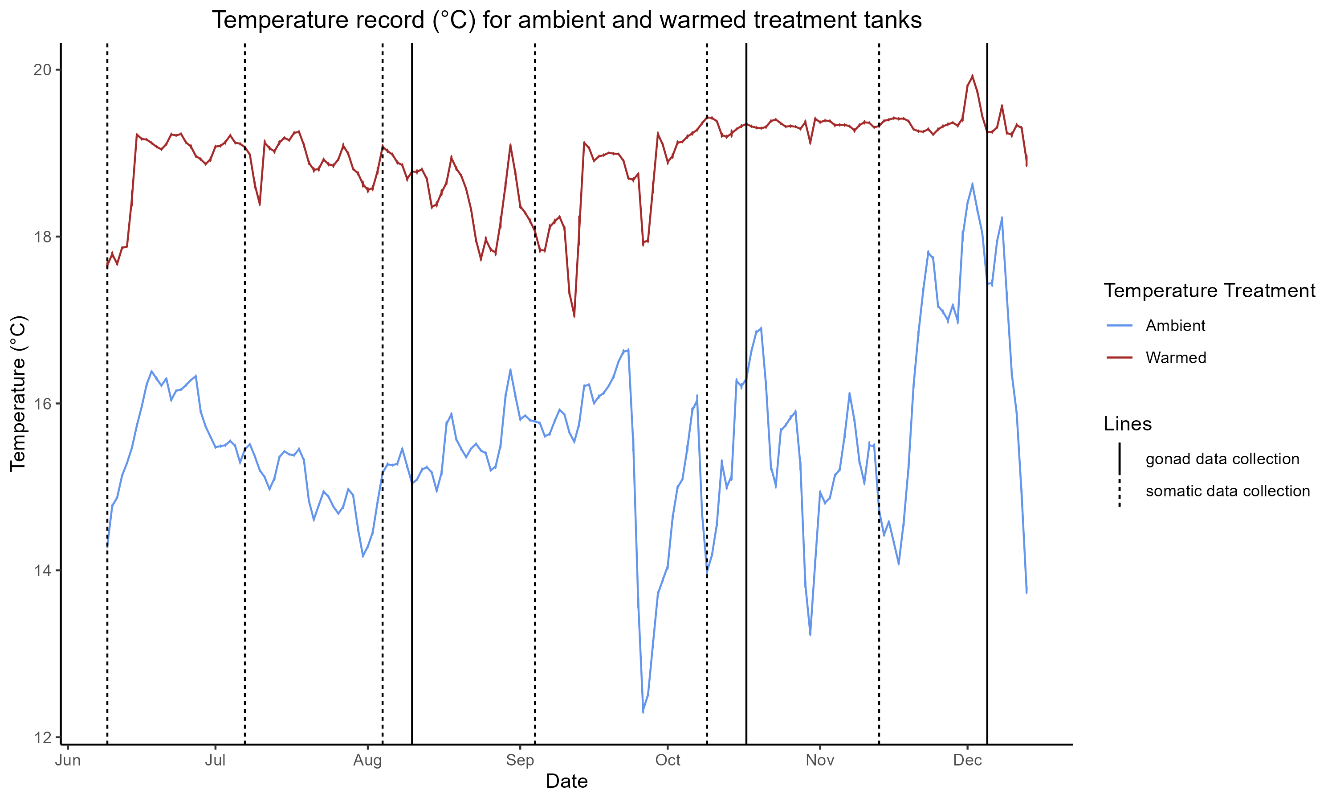


Figure . Temperature (oC) record for temperature treatments applied to tanks

## Survival

Diet had a significant effect on survival rate (%) (Kruskal-Wallis, p < 0.001) within the first nine weeks of the experiment (Figure 7). The kelp dietary treatment had a significantly lower (Dunn – Bonferroni test, p < 0.05) survival rate than all other dietary treatments in this period. Many urchins in the kelp dietary treatment group showed severe spine loss and were removed from the tanks for disease and water quality concerns (Appendix, Figure A1). Due to the consistently poor health (Appendix A2, morbidity graph) during the early stages of the experiment and concerns for the health of the animals, the kelp dietary treatment was suspended in week 9. Kelp dietary treatment urchins were ethically euthanized as per DFFE biosecurity regulation and kelp was removed from the mixed dietary treatment feeding regime. The mixed diet regime changed to *Ulva lacinulata* and a formulated feed containing 16% *Ulva* on a weekly basis from week 10 onwards.

Eliminating the kelp dietary treatment increased the significance of the temperature treatment effect on survival. The ambient temperature treatment group had a significantly lower survival rate than the warm temperature treatment group overall (Mann-Whitney U test, p <0.01). (Sig differences over time between week 10 and 24?)

A graph of different colored bars

Description automatically generated with medium confidence

Figure 7: Mean (± SE) of survival rate (%) for the Cape sea urchin, *P. angulosus*, fed different diet treatments (formulated feed with 16% ulva, *Ecklonia maxima* kelp, *Ulva lacinulata*, and a combination of the forementioned to form a mixed diet) under different temperature conditions (ambient ~ 15oC and warm ~18oC). Letters above bars represent dietary treatment groups that are significantly different to each other based on Dunn’s test with Bonferroni correction. Hashtags above bars indicate the level of significant difference between temperature treatment groups based on Mann-Whitney U tests (#: p<0.05, ##: p<0.01, ###: p<0.001). No letters or hastags indicate no significant differences.

## Somatic growth

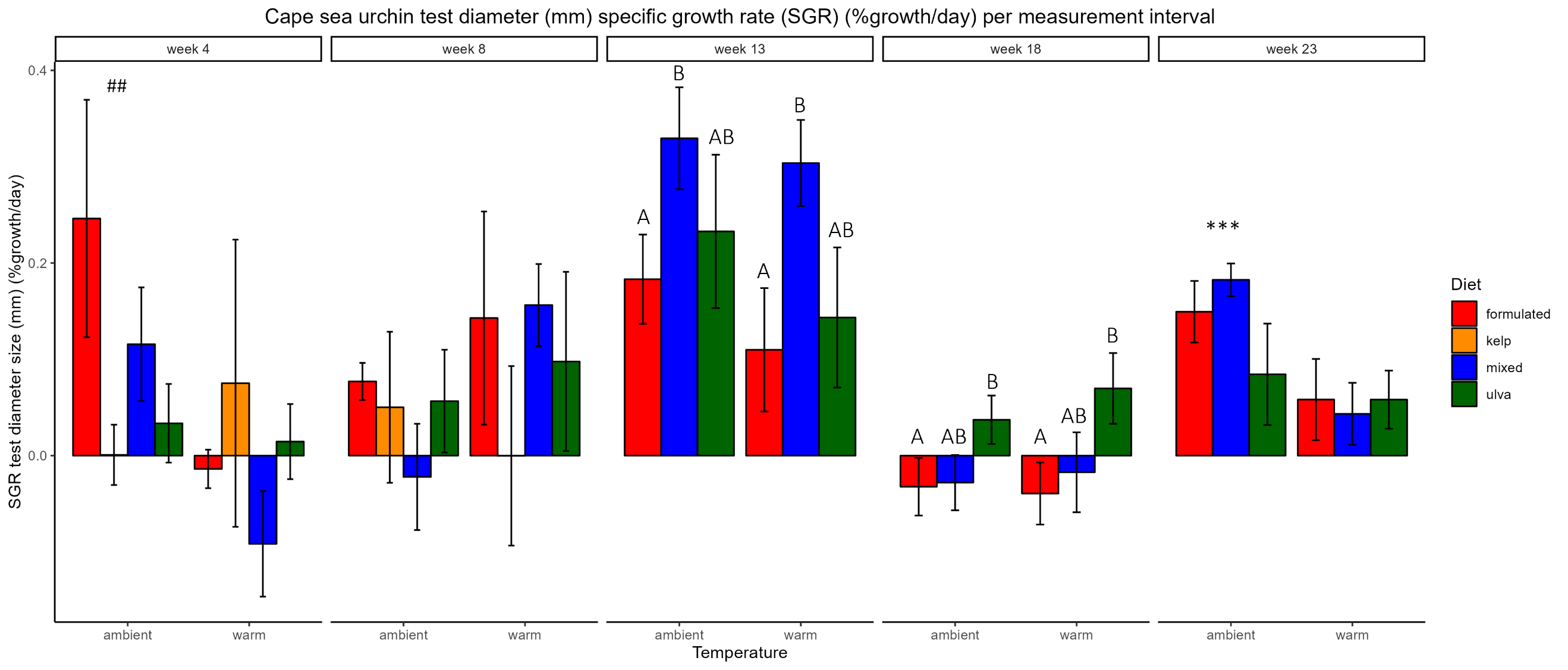
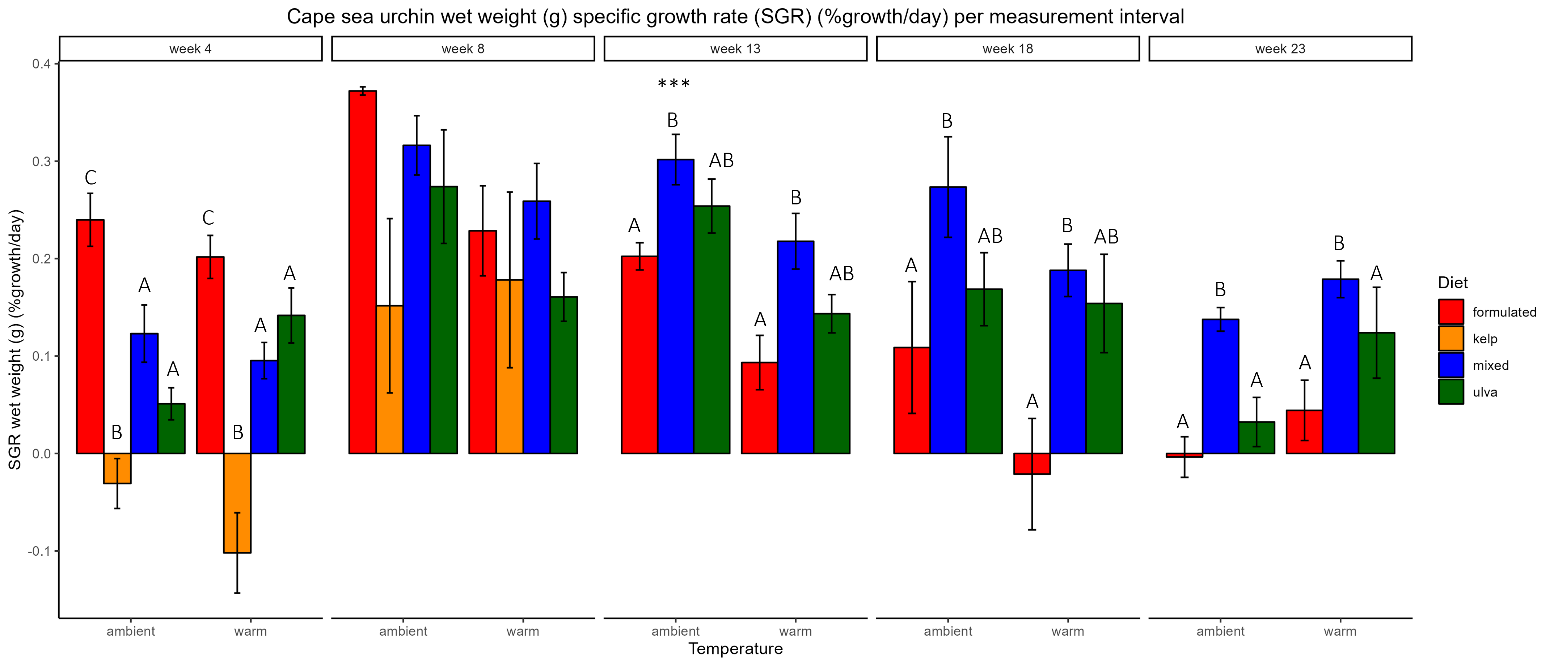
*Week 4-8*

Within the first four weeks of the experiment dietary treatment (ANOVA, p < 0.001) had a significant effect on the specific growth rates (SGR, % growth per day) of *P.angulosus* wet weight (g) (SGRweight). The formulated dietary treatment group gained weight significantly faster than all other dietary treatments (Tukey, p< 0.05) (Figure 8a). Temperature treatment (Mann-Whitney, p < 0.01) had a significant effect on the specific growth rate of *P.angulosus* test diameter (mm) (SGRsize) (Figure 8b). The test diameters of the ambient temperature treatment group grew significantly faster than the warm temperature treatment group (Mann-Whitney, p < 0.01). During this period, the kelp dietary treatment group lost weight as indicated by negative SGRweight values. There were no significant dietary or treatment effects on somatic growth in week 8.

*Week 13 - 23*

The ambient temperature treatment group gained weight significantly faster than the warm temperature treatment group in week 13 (T-test, p <0.001). Dietary treatment effects were significant for SGRweight in weeks 13, 18 and 23 (ANOVA, p < 0.05). The mixed dietary treatment had the fastest SGRweight during this period and differed significantly to the formulated dietary treatment group which had the slowest SGRweight during this period (Tukey, p < 0.05). At the final somatic measurement timepoint the mixed dietary treatment SGRweight was significantly faster than all other dietary treatments (Tukey, p < 0.05).

Dietary treatments had a significant effect on SGRsize in weeks 13 and 18 (ANOVA, p < 0.05). In week 13, the mixed dietary treatment had the fastest SGRsize during this period and differed significantly to the formulated dietary treatment group which had the slowest SGRsize during this period (Tukey, p < 0.05. In week 18 however, both the mixed (-0.02 ± 0.02) and formulated (-0.04 ± 0.02) dietary treatment groups had negative growth rates, indicating a lack of growth in test diameter. Temperature treatment had a significant effect on SGRsize at the final measuremeny timepoint, test diameters of the ambient temperature treatment group grew significantly faster than the warm temperature treatment group (T - test, p < 0.01).



**A**

**B**

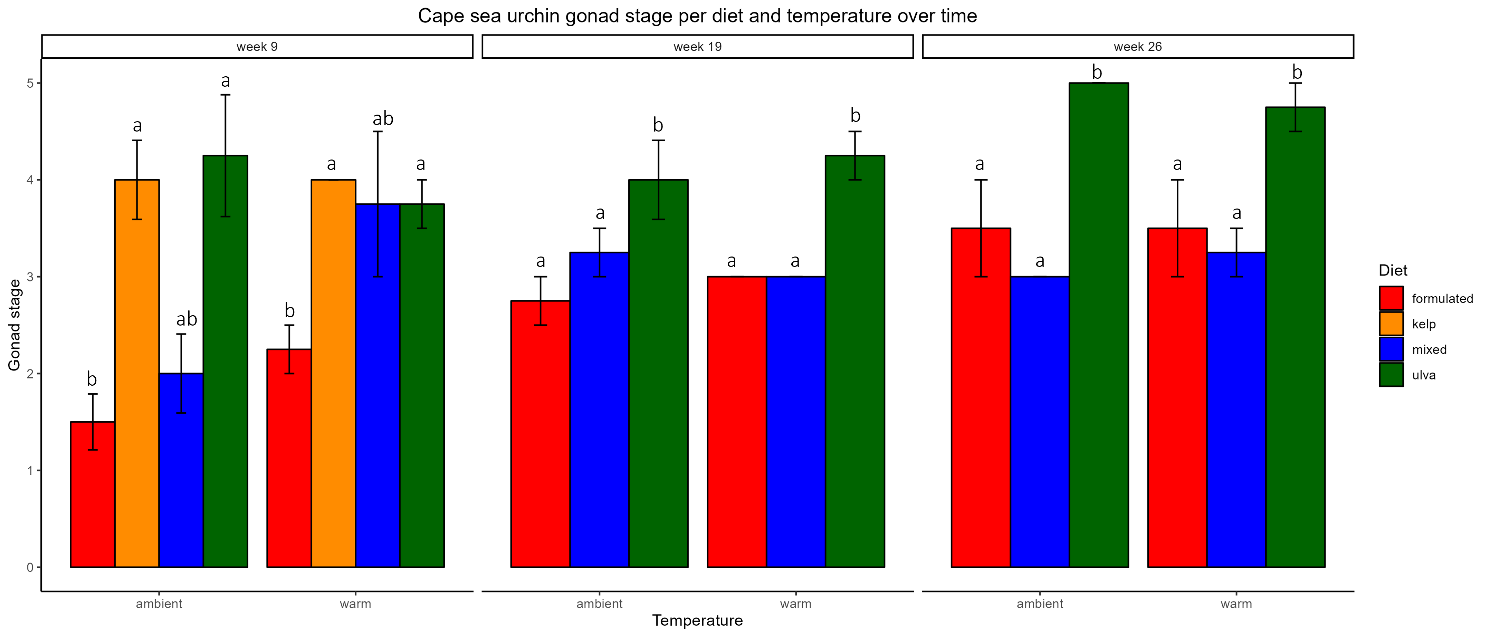
Figure 8: Mean (± SE) **A)** whole animal wet mass (g) and **B)** test diameter (mm) specific growth rate (SGR) (% growth day-1) for the Cape sea urchin, *P. angulosus*, fed different diet treatments (formulated feed with 16% ulva, *Ecklonia maxima* kelp, *Ulva lacinulata*, and a combination of the forementioned to form a mixed diet) under different temperature conditions (ambient and warm). Letters above bars represent dietary treatment groups that are significantly different to each other based on ANOVA (UPPERCASE), Kruskal-Wallis (lowercase) or Welch ANOVA (*lowercase italic*) followed by Tukey’s, Dunn’s with Bonferroni correction or Games-Howell post hoc pairwise comparison test respectively. Stars above bars indicate the level of significant difference between temperature treatment groups (\*:p<0.05, \*\*: p<0.01, \*\*\*:p<0.001). No letters or stars indicate no significant differences.

## Gonad quantity and quality

### Gonad development

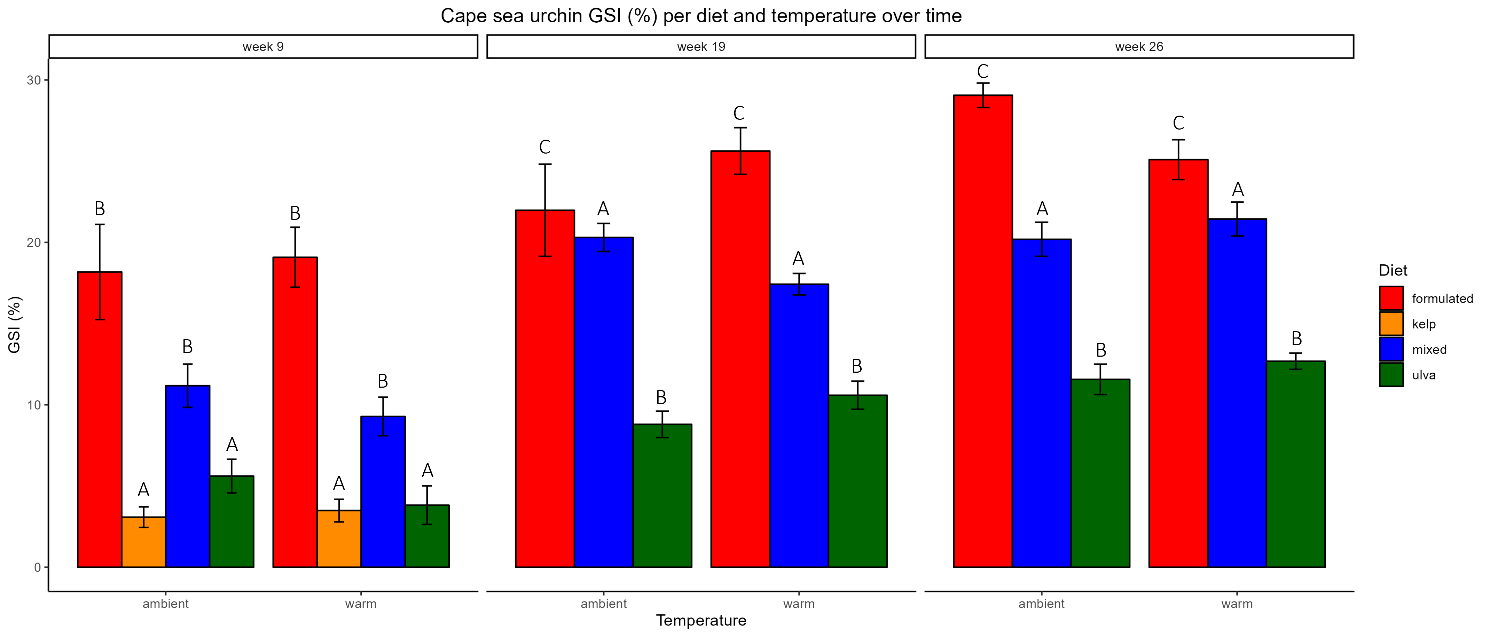
Dietary treatment had a significant effect on gonad development of *P. angulosus* throughout the experiment (Kruskal-Wallis, p < 0.001). Fro

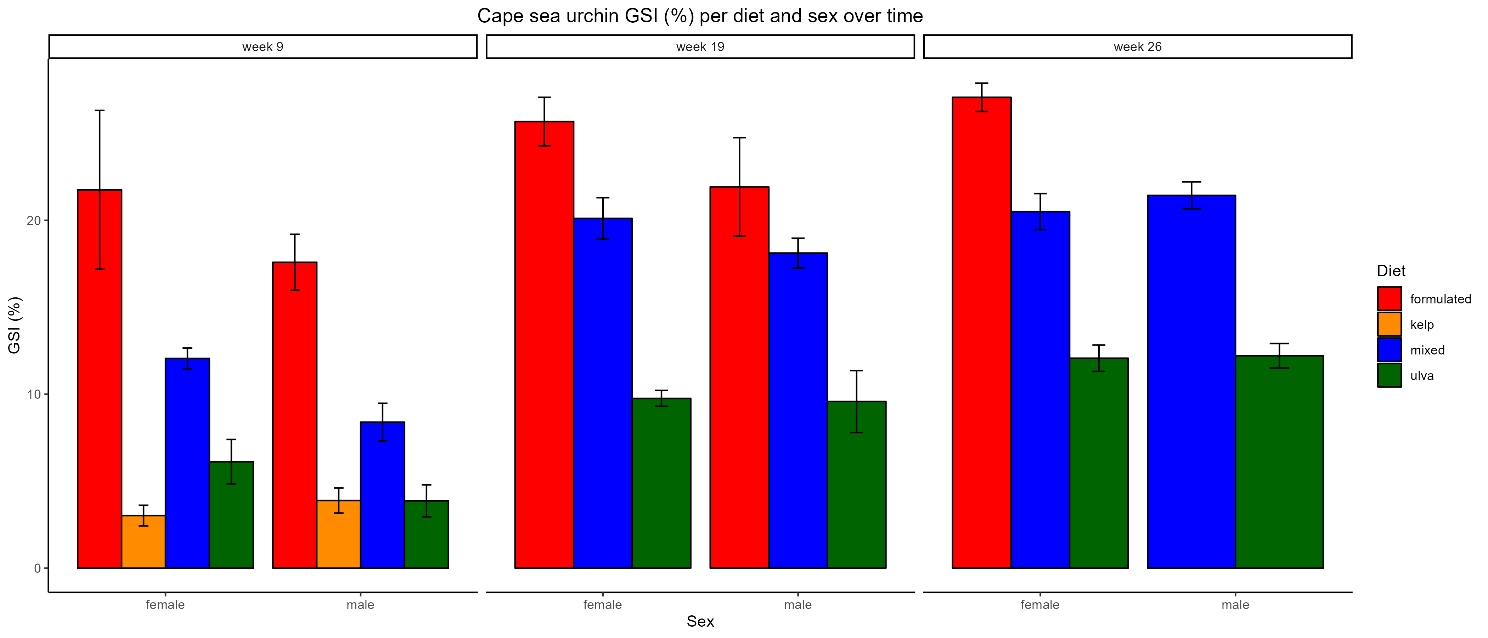
m the first gonad sample timepoint, the algal dietary treatment groups (kelp and ulva) were significantly more mature than the formulated treatment group (Dunn-Bonferroni, p < 0.01). In week 19 and 23, the ulva dietary treatment group continued to be significantly more mature than all other dietary treatment groups (Dunn-Bonferroni, p<0.05). Temperature treatment and sex did not have a significant effect on gonad development.



### Gonad quantity

(still busy)

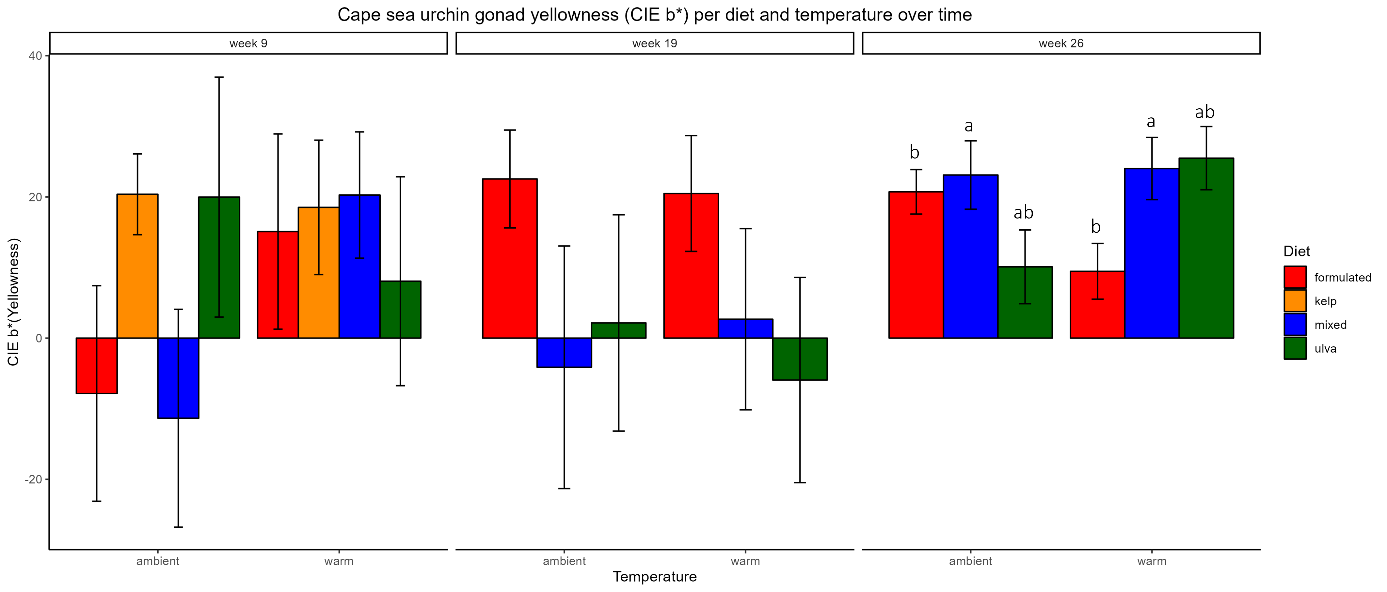
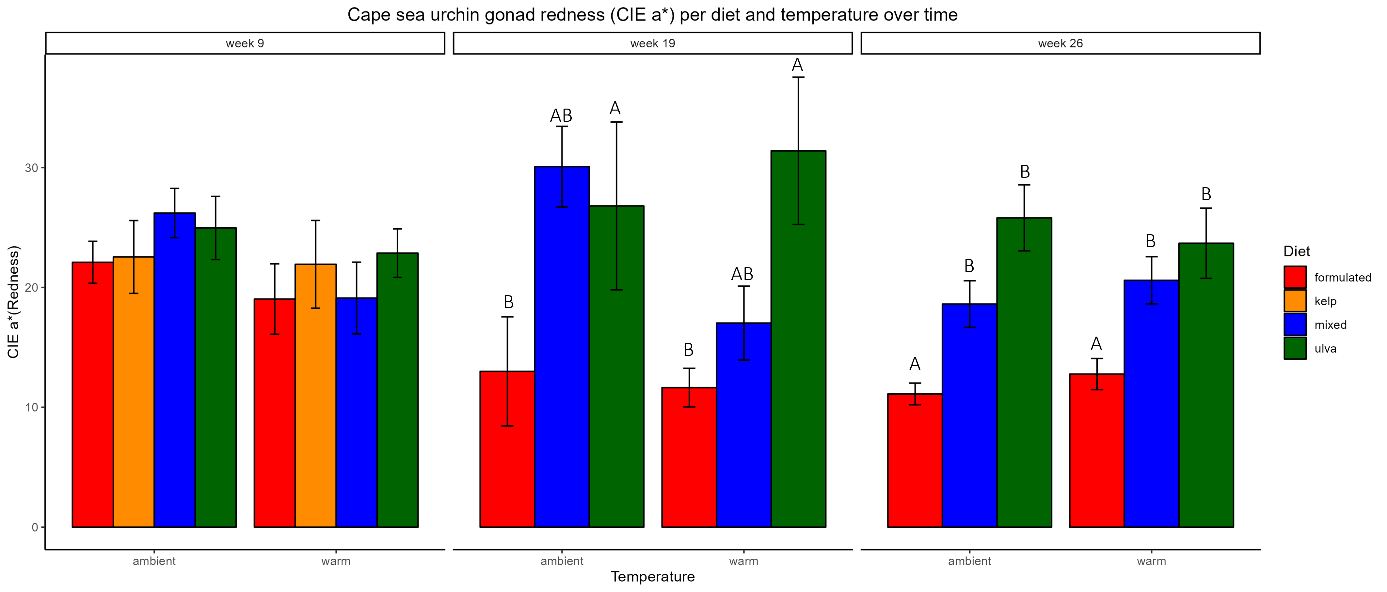
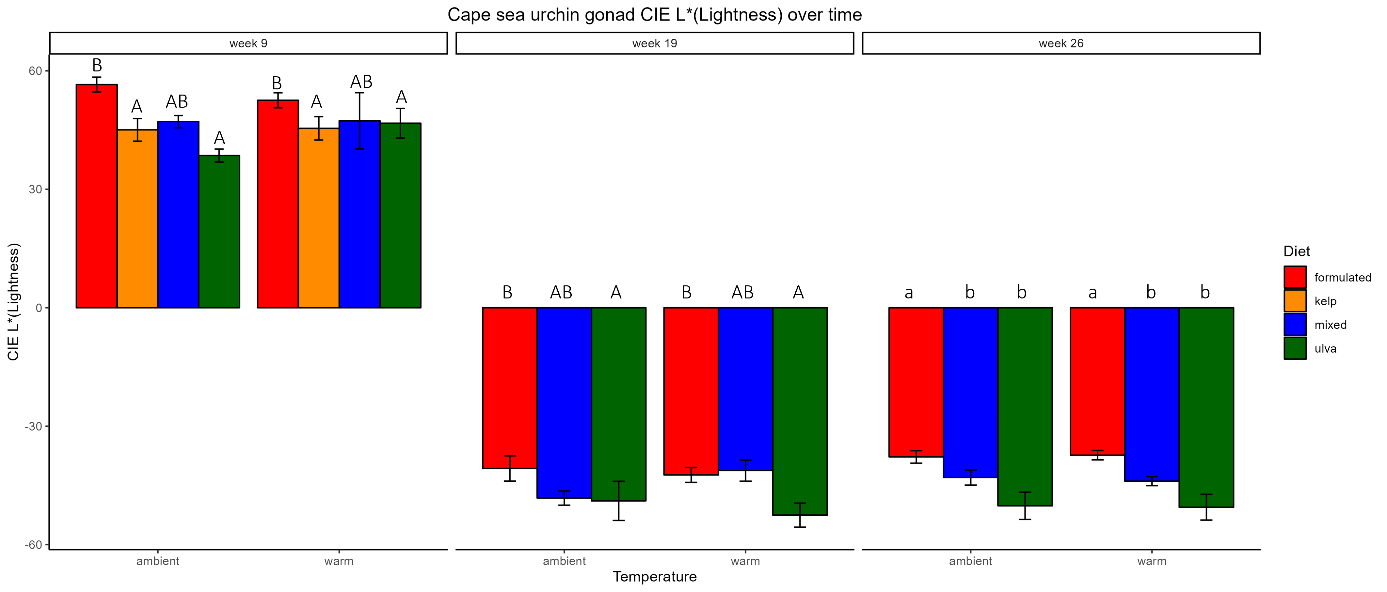




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### Gonad colour

(still busy)



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Description automatically generatedA graph of different colored rectangular shapes

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# **Discussion**

* Discuss most interesting results and possible reasons for unexpected results
* Did the project meet the project expectations & research objectives?
* Refer to other urchin work: how fast is their growth relative to other studies? How marketable is their gonad?

# **Conclusion**

* Is this feasible?
* What are the potential challenges?
* What were the research limitations?
* What are the opportunities for future research?

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# **Appendix**

## A close up of a plant Description automatically generatedSupplementary figures



Figure A1: Image of *P.angulosus* fed *Ecklonia maxima* dietary treatment taken in week 2 of the experiment.

Maybe a bar graph with morbidities?

GSI photos?

Gonad colour photos?

## Supplementary statistical tables

### Survival rate

Table A1.1. Kruskal-Wallis (H-values) test statistic dietary treatment effect and (two- sided) Mann-Whitney U test (Z-values) for temperature treatment effect on survival rate of *P.angulosus* fed one of four dietary treatments (formulated, *Ecklonia maxima*, mixed, or *Ulva lacinulata*) under one of two temperature treatments (ambient ~15 or warm ~18). Statistical significance is indicated as \*: p<0.05, \*\*: p<0.01, \*\*\*: p<0.001.

|  |  |  |
| --- | --- | --- |
| Week | Diet | Temperature |
| 9 (n=32) | H = 20.33 \*\*\* | Z = -1.42 |
| 10 (n=24) | H = 3.12 | Z = -2.4 \* |
| 18 (n=24) | H = 1.40 | Z = -2.79 \*\* |
| 24 (n=24) | H = 1.52 | Z = -2.50 \* |

Mean ± SE of *P. angulosus* survival rate (%) per diet, temperature and sex group. Statistically significant differences between dietary treatment groups per week are shown by different letters based on Kruskal-Wallis test followed by a Dunn’s (with Bonferroni correction) test pairwise comparisons respectively. Statistically significant differences between temperature groups are indicated with stars alongside the significantly larger group based on (one-sided) Mann-Whitney U test. Statistical significance is indicated as \*: p<0.05, \*\*: p<0.01, \*\*\*: p<0.001. No letters or stars indicate that there were no significant differences.

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
|  | Diet | | | | Temperature | |
| Week | F | K | M | U | A | W |
| 9 (n = 32) | 95.39 ± 2.20 b | 66.45 ± 4.21 a | 98.68 ± 1.32 b | 93.42 ± 2.77 b | 86.18 ± 3.56 | 90.79 ± 4.01 |
| 10 (n = 24) | 95.39 ± 2.10 | - | 98.68 ± 1.32 | 93.42 ± 2.77 | 92.54 ± 2.10 | 99.12 ± 0.59 \*\* |
| 18 (n = 24) | 93.42 ± 2.94 | - | 94.74 ± 3.85 | 89.47 ± 3.85 | 86.84 ± 3.21 | 98.25 ± 0.99 \*\* |
| 24 (n = 24) | 90.79 ± 3.55 | - | 94.08 ± 3.55 | 89.47 ± 3.85 | 85.52 ± 3.61 | 97.37 ± 1.21 \*\* |

### Somatic growth

Table A2.1. ANOVA (F-values) or Kruskal-Wallis (H-values) test statistic for dietary treatment effect and two sample t-test (t-values) or Mann-Whitney U test (Z-values) test statistic for temperature treatment effect on specific growth rate (SGR) (% growth per day) of the total wet weight (g) (SGR weight­) and test diameter (mm) (SGR size) for *P.angulosus* fed one of four dietary treatments (formulated, *Ecklonia maxima*, mixed, or *Ulva lacinulata*) under one of two temperature treatments (ambient ~15oC or warm ~18 oC). Statistical significance is indicated as \*: p<0.05, \*\*: p<0.01, \*\*\*: p<0.001.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  | SGR weight |  | SGR size |  |
| Week | Diet | Temperature | Diet | Temperature |
| 4 (n = 32) | F = 30.95 \*\*\* | t = 0.27 | H = 2.24 | Z = 2.60 \*\* |
| 8 (n = 32) | F = 2.42 | t = 1.71 | F = 0.45 | t = -1.17 |
| 13 (n= 24) | F = 5.00 \* | t = 3.78 \*\* | F = 4.40 \* | t = 1.14 |
| 18 (n = 24) | F = 6.49 \*\* | t = 1.51 | F = 4.88 \* | t = -0.41 |
| 23 (n = 24) | F = 9.91 \*\*\* | t = -1.87 | F = 0.54 | t = 2.91 \*\* |

Table A2.2. Mean ± SE for specific growth rate (SGR) (% growth per day) of the total wet weight (g) (SGRweight) for *P.angulosus* fed one of four dietary treatments (formulated, *Ecklonia maxima*, mixed, or *Ulva lacinulata*) under one of two temperature treatments (ambient ~15oC or warm ~18 oC). Statistically significant differences in dietary treatment shown by different letters based on ANOVA tests followed by a post hoc Tukey pairwise comparison test. Statistically significant differences between temperature groups are indicated as stars alongside the significantly larger group based on (one-sided) two sample t-test. Statistical significance is indicated as \*: p<0.05, \*\*: p<0.01, \*\*\*: p<0.001. No letters indicate that there were no significant differences.

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
|  | Diet | | | | Temperature | |
| Week | F | K | M | U | A | W |
| 4 (n = 32) | 0.22 ± 0.02 C | -0.07 ± 0.03 B | 0.11 ± 0.02 A | 0.10 ± 0.02 A | 0.10 ± 0.03 | 0.08 ± 0.03 |
| 8 (n = 32) | 0.30 ± 0.03 | 0.16 ± 0.06 | 0.29 ± 0.03 | 0.22 ± 0.04 | 0.28 ± 0.03 | 0.21 ± 0.03 |
| 13 (n = 24) | 0.15 ± 0.03 A | - | 0.26 ± 0.02 B | 0.20 ± 0.03 AB | 0.25 ± 0.02 \*\*\* | 0.15 ± 0.02 |
| 18 (n = 24) | 0.04 ± 0.05 A | - | 0.23 ± 0.03 B | 0.16 ± 0.03 AB | 0.18 ± 0.03 | 0.11 ± 0.04 |
| 23 (n = 24) | 0.02 ± 0.02 A | - | 0.16 ± 0.01 B | 0.08 ± 0.03 A | 0.06 ± 0.02 | 0.12 ± 0.02 |

Table A2.2. Mean ± SE for specific growth rate (SGR) (% growth per day) of the test diameter (mm) (SGRsize) for *P.angulosus* fed one of four dietary treatments (formulated, *Ecklonia maxima*, mixed, or *Ulva lacinulata*) under one of two temperature treatments (ambient ~15oC or warm ~18 oC). Statistically significant differences in dietary treatment indicated as letters based on ANOVA tests followed by a post hoc Tukey pairwise comparison test. Statistically significant differences between temperature groups are indicated as stars or hashtags alongside the significantly larger group based on (one-sided) two sample t-test (\*) or Mann-Whitney U test (#). Statistical significance is indicated \*/#: p<0.05, \*\*/##: p<0.01, \*\*\*/###: p<0.001. No letters or stars/hashtags indicate that there were no significant differences.

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
|  | Diet | | | | Temperature | |
| Week | F | K | M | U | A | W |
| 4 (n = 32) | 0.12 ± 0.08 | 0.04 ± 0.07 | 0.01 ± 0.05 | 0.03 ± 0.04 | 0.10 ± 0.04 ## | 0.00 ± 0.04 |
| 8 (n = 32) | 0.11 ± 0.05 | 0.03 ± 0.06 | 0.07 ± 0.05 | 0.08 ± 0.05 | 0.04 ± 0.03 | 0.10 ± 0.04 |
| 13 (n = 24) | 0.15 ± 0.04 A | - | 0.32 ± 0.03 B | 0.19 ± 0.05 AB | 0.25 ±0.04 | 0.19 ± 0.04 |
| 18 (n = 24) | -0.04 ± 0.02 A | - | -0.02 ± 0.02 AB | 0.05 ± 0.02 B | -0.01 ± 0.02 | 0.00 ± 0.02 |
| 23 (n = 24) | 0.10 ± 0.03 | - | 0.11 ± 0.03 | 0.07 ± 0.03 | 0.14 ± 0.02 \*\* | 0.05 ± 0.02 |

### Gonad quantity and development

Table A2.1. ANOVA (F-values), or Kruskal-Wallis (H-values) test statistics for dietary treatment effects and two sample t-test (t-values) or Mann-Whitney U test (Z-values) for temperature treatment and sex effects per week on gonad quantity (GSI) and development (gonad stage) for *P.angulosus* fed one of four dietary treatments (formulated, *Ecklonia maxima*, mixed, or *Ulva lacinulata*) under one of two temperature treatments (ambient ~15oC or warm ~18 oC). Where log transformations were performed, they are indicated as (log) prior to the test statistic. The last line of the table excludes the formulated dietary treatment data due to lack of data for females. Statistical significance is indicated as \*: p<0.05, \*\*: p<0.01, \*\*\*: p<0.001.

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
|  | GSI |  |  |  | Gonad stage |  |  |
| Week | Temperature | Diet | Sex | Week | Temperature | Diet | Sex |
| 9 (n = 43) | (log) t = 0.36 | (log) F = 33.11 \*\*\* | (log) t = -1.40 | 9 (n = 32) | Z = -1.13 | H = 18.89 \*\*\* | Z = -0.47 |
| 19 (n = 24) | t = -0.31 | F = 42.61 \*\*\* | t = 0.15 | 19 (n = 24) | Z = -0.27 | H = 15.07 \*\*\* | Z = 1.95 |
| 26 (n = 72) | t = 0.32 | F = 113.7 \*\*\* | t = 2.38 \* | 26 (n = 24) | Z = -0.65 | H = 14.66 \*\*\* | Z = 1.95 |
| 26 (n = 48) |  |  | t = -0.08 |  |  |  |  |

Table A2.2. Mean ± SE of *P. angulosus* gonad quantity (GSI) (%) values per diet, temperature and sex group. Statistically significant differences between dietary treatment groups per week are shown by different letters based on ANOVA test followed by a post hoc Tukey test pairwise comparisons. Statistically significant differences between sex groups per week are shown as stars alongside the significantly larger group based on one-sided two sample t-test. Statistical significance is indicated as \*: p<0.05, \*\*: p<0.01, \*\*\*: p<0.001. No letters or stars indicate that there were no significant differences. The last line of the table excludes the formulated dietary treatment data due to lack of data for females.

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | Diet | | | | Temperature | | Sex | |
| Week | F | K | M | U | A | W | F | M |
| 9 (n = 43) | 18.63 ± 1.61 B | 3.29 ± 0.46 A | 10.23 ± 0.90 B | 4.72 ± 0.80 A | 7.98 ± 1.43 | 7.44 ± 1.38 | 6.79 ± 1.36 | 8.66 ± 1.43 |
| 19 (n = 24) | 23.80 ± 1.63 C | - | 18.86 ± 0.74 A | 9.69 ± 0.65 B | 17.03 ± 1.99 | 17.88 ± 1.93 | 17.66 ± 2.17 | 17.25 ± 1.74 |
| 26 (n = 72) | 27.08 ± 0.82 C | - | 20.81 ± 0.73 A | 12.12 ± 0.53 B | 20.27 ± 1.31 | 19.74 ± 1.04 | 21.07 ± 0.98 \*\* | 16.55 ± 1.26 |
| 26 (n = 48) |  |  |  |  |  |  | 16.42 ± 1.00 | 16.55 ± 1.26 |

Table A2.3. Mean ± SE of *P. angulosus* gonad development (gonad stage) values per diet, temperature and sex group. Statistically significant differences between dietary treatment groups per week are shown by different letters based on Kruskal-Wallis test followed by a post hoc Dunn’s (with Bonferroni correction) test pairwise comparisons. Statistically significant differences between temperature treatment and sex groups per week are shown as stars alongside the significantly larger group. No letters or stars indicate that there were no significant differences.

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | Diet | | | | Temperature | | Sex | |
| Week | F | K | M | U | A | W | F | M |
| 9 (n = 32) | 1.88 ± 0.23 b | 4.00± 0.19 a | 2.88 ± 0.52 ab | 4.00 ± 0.33 a | 2.94 ± 0.37 | 3.44 ± 0.26 | 3.09 ± 0.41 | 3.24 ± 0.28 |
| 19 (n = 24) | 2.88 ± 0.13 a | - | 3.13 ± 0.13 a | 4.13 ± 0.23 b | 3.33 ± 0.23 | 3.42 ± 0.19 | 3.67 ± 0.22 | 3.08 ± 0.15 |
| 26 (n = 24) | 3.50 ± 0.33 a | - | 3.13 ± 0.13 a | 4.88 ± 0.13 b | 3.83 ± 0.30 | 3.83 ± 0.27 | 4.13 ± 0.26 | * 1. ± 0.24 |

### Gonad colour

Table A2.4. ANOVA (F-values), Kruskal-Wallis (H-values) or Welch ANOVA (*F-values)* test statistics for dietary treatment effects and two sample t-test (t-values) or Mann-Whitney U test (Z-values) for temperature treatment and sex effects on gonad colour in terms of lightness (CIE L\*), redness (CIE a\*) and yellowness (CIE b\*) for *P.angulosus* fed one of four dietary treatments (formulated, *Ecklonia maxima*, mixed, or *Ulva lacinulata*) under one of two temperature treatments (ambient ~15oC or warm ~18 oC). Where log transformations were performed, they are indicated as (log) prior to the test statistic. Statistical significance is indicated as \*: p<0.05, \*\*: p<0.01, \*\*\*: p<0.001. The last line of the table excludes the formulated dietary treatment data due to lack of data for females.

|  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | L\* (Lightness/Intensity) | |  | a\* (Redness/Hue) | |  | b\* (Yellowness/Chroma) | |  |  |
| Week | Temperature | Diet | Sex | Temperature | Diet | Sex | Temperature | Diet |  | Sex |
| 9 (n = 43) | t = -0.34 | F = 3.50 \* | t = -3.14 \*\* | t = 1.09 | *F = 0.66* | t = 2.07 \* | Z = -0.95 | H = 1.82 |  | Z = 1.92 |
| 19 (n = 24) | t = -0.19 | F = 4.43 \* | t = -3.38 \*\* | t = 0.68 | F = 6.27 \*\* | t (≠variance) = 3.69 \*\* | Z = 0.23 | H = 4.42 |  | Z = -0.40 |
| 26 (n = 72) | Z = 0.24 | H = 18.91 \*\*\* | Z = -1.77 | (log) t = -0.46 | (log) F = 18.85 \*\*\* | (log) t (≠variance) = 2.55 \* | Z = -0.42 | H = 6.31 \* |  | Z = -0.14 |
| 26 (n = 48) |  |  | t (≠variance) = -5.25 \*\*\* |  |  | t (≠variance) = 7.54 \*\*\* |  |  |  | Z = 1.09 |

Changes over time?

Table A2.5. Mean ± SE of *P. angulosus* gonad colour lightness (CIE L\*) values per diet, temperature and sex group. Statistically significant differences between dietary treatment groups per week are shown by different letters based on ANOVA or Kruskal-Wallis test followed by a post hoc Tukey (UPPERCASE) or Dunn’s (lowercase) (with Bonferroni correction) test pairwise comparisons respectively. Statistically significant differences between sex groups are shown as stars or hashtags alongside the significantly larger group based on one-sided two sample t-test (\*) or Mann-Whitney U test (#). Statistical significance is indicated as \*/#: p<0.05, \*\*/##: p<0.01, \*\*\*/###: p<0.001. No letters or stars indicate that there were no significant differences. The last line of the table excludes the formulated dietary treatment data due to lack of data for females.

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | Diet | | | | Temperature | | Sex | |
| Week | F | K | M | U | A | W | F | M |
| 9 (n = 43) | 54.53 ± 1.44 B | 45.25 ± 2.01 A | 47.24 ± 3.37 AB | 42.63 ± 2.44 A | 46.39 ± 1.81 | 47.31 ± 1.95 | 43.19 ± 1.47 | 50.70 ± 1.91\*\* |
| 19 (n = 24) | -41.53 ± 1.75 B | - | -44.73 ± 1.99 AB | -50.72 ± 2.78 A | -45.94 ± 2.17 | -45.38 ± 2.04 | -49.75 ± 1.80 | -41.57 ± 1.61 ## |
| 26 (n = 72) | -37.57 ± 0.97 a | - | -43.46 ± 1.11 b | -50.34 ± 2.32 b | -43.65 ± 1.62 | -43.93 ± 1.49 | -45.02 ± 1.34 | -39.82 ± 1.32 |
| 26 ( n = 48) |  |  |  |  |  |  | -50.78 ± 1.62 | -39.82 ± 1.32 \*\* |

Table A2.5. Mean ± SE of *P. angulosus* gonad colour redness (CIE a\*) values per diet, temperature and sex group. Statistically significant differences between dietary treatment groups per week are shown by different letters based on ANOVA or Kruskal-Wallis test followed by a post hoc Tukey (UPPERCASE) or Dunn’s (lowercase) (with Bonferroni correction) test pairwise comparisons respectively. Statistically significant differences between sex groups are shown as stars or hashtags alongside the significantly larger group based on one-sided two sample t-test (\*) or Mann-Whitney U test (#). Statistical significance is indicated as \*/#: p<0.05, \*\*/##: p<0.01, \*\*\*/###: p<0.001. No letters or stars indicate that there were no significant differences. The last line of the table excludes the formulated dietary treatment data due to lack of data for females.

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | Diet | | | | Temperature | | Sex | |
| Week | F | K | M | U | A | W | F | M |
| 9 (n = 43) | 20.57 ± 1.69 | 22.22 ± 2.34 | 22.66 ± 2.15 | 23.92 ± 1.59 | 23.62 ± 1.46 | 21.06 ± 1.81 | 24.59 ± 1.66 \* | 19.93 ± 1.53 |
| 19 (n = 24) | 12.32 ± 2.25 B | - | 23.55 ± 3.25 AB | 29.10 ± 4.40 A | 23.29 ± 3.51 | 20.02 ± 3.29 | 28.71 ± 3.34 \*\*\* | 14.60 ± 1.86 |
| 26 (n = 43) | 11.94 ± 0.79 A | - | 19.61 ± 1.37 B | 24.75 ± 1.98 B | 18.51 ± 1.52 | 19.02 ± 1.44 | 20.17 ± 1.28 \*\* | 14.25 ± 0.96 |
| 26 ( n = 48) |  |  |  |  |  |  | 26.52 ± 1.32 \*\*\* | 14.25 ± 0.96 |

Table A2.5. Mean ± SE of *P. angulosus* gonad colour yellowness (CIE b\*) values per diet, temperature and sex group. Statistically significant differences between dietary treatment groups per week are shown by different letters based on ANOVA or Kruskal-Wallis test followed by a post hoc Tukey (UPPERCASE) or Dunn’s (lowercase) (with Bonferroni correction) test pairwise comparisons respectively. Statistically significant differences between sex groups are shown as stars or hashtags alongside the significantly larger group based on one-sided two sample t-test (\*) or Mann-Whitney U test (#). Statistical significance is indicated as \*/#: p<0.05, \*\*/##: p<0.01, \*\*\*/###: p<0.001. No letters or stars indicate that there were no significant differences. The last line of the table excludes the formulated dietary treatment data due to lack of data for females.

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | Diet | | | | Temperature | | Sex | |
| Week | F | K | M | U | A | W | F | M |
| 9 (n = 32) | 3.63 ± 10.47 | 19.39 ± 5.56 | 4.46 ± 10.19 | 14.01 ± 10.67 | 8.88 ± 6.16 | 16.31 ± 5.59 | 18.71 ± 5.93 | 6.37 ± 5.58 |
| 19 (n = 24) | 21.51 ± 4.99 | - | -0.73 ± 10.00 | -1.90 ± 9.89 | 6.85 ± 8.02 | 5.74 ± 7.16 | 2.54 ± 8.53 | 10.05 ± 6.35 |
| 26 (n = 24) | 15.09 ± 2.74 b | - | 23.56 ± 3.20 a | 17.79 ± 3.73 ab | 17.97 ± 2.70 | 19.66 ± 2.69 | 18.13 ± 2.36 | 21.01 ± 2.46 |
| 26 ( n = 48) |  |  |  |  |  |  | 20.49 ± 3.60 | 21.01 ± 2.46 |

Table A2.5. Mean ± SE of feed consumed per week (g) for *P. angulosus* per diet and temperature group combinations. The feed given to the mixed diet group at the measurement timepoint, (U) ulva or (F) formulated, is shown in brackets alongside the values. Statistically significant differences between dietary treatment groups per week are shown by different letters based on ANOVA or Kruskal-Wallis test followed by a post hoc Tukey (UPPERCASE) or Dunn’s (lowercase) (with Bonferroni correction) test pairwise comparisons respectively. Statistically significant differences between temperature groups are shown as stars or hashtags alongside the significantly larger group based on one-sided two sample t-test (\*) or Mann-Whitney U test (#). Statistical significance is indicated as \*/#: p<0.05, \*\*/##: p<0.01, \*\*\*/###: p<0.001. No letters or stars indicate that there were no significant differences.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| All temperatures | Diet | | | |
| Week | F | K | M | U |
| 8 (n = 32) | 10.12 ± 0.45 | 31.50 ± 1.61 | 12.63 ± 0.34 (F) | 48.65 ± 4.39 |
| 13 (n = 24) | 10.12 ± 0.33 | - | 36.31 ± 5.14 (U) | 46.46 ± 5.63 |
| 18 (n = 24) | 8.98 ± 0.65 | - | 11.66 ± 0.58 (F) | 37.24 ± 3.07 |
| 23 (n = 24) | 7.10 ± 0.39 | - | 25.63 ± 3.57 (U) | 27.86 ± 5.47 |
| Ambient temperature |  |  |  |  |
| 8 (n = 32) |  |  |  |  |
| 13 (n = 24) |  |  |  |  |
| 18 (n = 24) |  |  |  |  |
| 23 (n = 24) |  |  |  |  |
| Warm temperature |  |  |  |  |
| 8 (n = 32) |  |  |  |  |
| 13 (n = 24) |  |  |  |  |
| 18 (n = 24) |  |  |  |  |
| 23 (n = 24) |  |  |  |  |

Table A2.5. Mean ± SE of feed consumed per animal per week (g) for *P. angulosus* per diet and temperature group combinations. The feed given to the mixed diet group at the measurement timepoint, (U) ulva or (F) formulated, is shown in brackets alongside the values. Statistically significant differences between dietary treatment groups per week are shown by different letters based on ANOVA or Kruskal-Wallis test followed by a post hoc Tukey (UPPERCASE) or Dunn’s (lowercase) (with Bonferroni correction) test pairwise comparisons respectively. Statistically significant differences between temperature groups are shown as stars or hashtags alongside the significantly larger group based on one-sided two sample t-test (\*) or Mann-Whitney U test (#). Statistical significance is indicated as \*/#: p<0.05, \*\*/##: p<0.01, \*\*\*/###: p<0.001. No letters or stars indicate that there were no significant differences.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| All temperatures | Diet | | | |
| Week | F | K | M | U |
| 8 (n = 32) | 0.56 ± 0.03 | 2.21 ± 0.16 | 0.67 ± 0.01 | 2.71 ± 0.19 |
| 13 (n = 24) | 0.60 ± 0.01 | - | 2.08 ± 0.27 | 2.79 ± 0.25 |
| 18 (n = 24) | 0.54 ± 0.03 | - | 0.69 ± 0.02 | 2.32 ± 0.14 |
| 23 (n = 24) | 0.47 ± 0.02 | - | 1.57 ± 0.19 | 1.80 ± 0.31 |
| Ambient temperature |  |  |  |  |
| 8 (n = 32) |  |  |  |  |
| 13 (n = 24) |  |  |  |  |
| 18 (n = 24) |  |  |  |  |
| 23 (n = 24) |  |  |  |  |
| Warm temperature |  |  |  |  |
| 8 (n = 32) |  |  |  |  |
| 13 (n = 24) |  |  |  |  |
| 18 (n = 24) |  |  |  |  |
| 23 (n = 24) |  |  |  |  |