1 Title: Methane Reduction Potential of Two Pacific Coast Macroalgae During in-vitro 2 **Ruminant Fermentation.** Charles G. Brooke^a (cgbrooke@ucdavis.edu), Breanna M. Roque^a (bmroque@ucdavis.edu), 3 Negeen Najafi^a (nnajafi@ucdavis.edu), Maria Gonzalez^a (mdcgonzalez@ucdavis.edu), Abigail 4 5 Pfefferlen^a (apfefferlen@ucdavis.edu), Vannesa DeAnda^a (vndeanda@ucdavis.edu), David W. Ginsburg^b (dginsbur@usc.edu); Maddelyn C. Harden^c (mharden@usc.edu), Sergey V. Nuzhdin^c 6 (snuzhdin@usc.edu), Joan King Salwen^d (jsalwen@standford.edu), Ermias Kebreab^a 7 (ekebreab@ucdavis.edu), and Matthias Hess^a (mhess@ucdavis.edu). 8 9 10 ^aDepartment of Animal Science, University of California, One Shields Avenue, Davis, CA, 11 95616, USA 12 ^bDornsife College of Letters, Arts and Sciences, University of Southern California, 3454 13 Trousdale Pkwy, CAS 116, Los Angeles, CA 90089, USA 14 ^cDornsife College of Letters, Arts and Sciences, University of Southern California, 3551 Trousdale Pkwy, Los Angeles, CA 90089, USA 15 ^dDepartment of Earth System Science, Stanford University, 450 Serra Mall, Stanford, CA, 94305 16 17 USA 18 Corresponding author: Matthias Hess 19 2251 Meyer Hall 20 Department of Animal Science 21 University of California, Davis 22 Davis, CA 95616, USA

P (530) 530-752-8809

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24 F (530) 752-0175 25 mhess@ucdavis.edu 26 27 **Conflict of interest:** The authors declare to have no conflict of interest. 28 **Funding:** Any opinions, findings, conclusions, or recommendations expressed in this publication 29 are those of the author(s). This work was supported through funds from the College of 30 Agricultural and Environmental Sciences at the University of California, Davis, by Elm 31 Innovations and by the Hellman Foundation. Maddelyn C. Harden was supported by ARPA-E 32 (Contract# 1726-1513) and Sergey V. Nuzhdin was supported by a Wyatt Foundation gift. 33 34 **Abstract:** 35 With increasing interest in feed based methane mitigation strategies, fueled by local legal 36 directives aimed at methane production from the agricultural sector in California, identifying 37 local sources of biological feed additives will be critical in keeping the implementation of these 38 strategies affordable. In a recent study, the red alga Asparagopsis taxiformis stood out as the most effective species of seaweed to reduce methane production from enteric fermentation. Due 39 to the potential differences in effectiveness based on the location from where A. taxiformis is 40 41 collected and the financial burden of collection and transport, we tested the potential of A. 42 taxiformis, as well as the brown seaweed Zonaria farlowii collected in the nearshore waters off 43 Santa Catalina Island, CA, USA, for their ability to mitigate methane production during *in-vitro* 44 rumen fermentation. At a dose rate of 5% dry matter (DM), A. taxiformis reduced methane

production by 74% ($p \le 0.01$) and Z. farlowii reduced methane production by 11% ($p \le 0.04$)

after 48 hours and 24 hours of *in-vitro* rumen fermentation respectively. The methane reducing

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effect of A. taxiformis and Z. farlowii described here make these local macroalgae promising candidates for biotic methane mitigation strategies in the largest milk producing state in the US. To determine their real potential as methane mitigating feed supplements in the dairy industry, their effect *in-vivo* requires investigation. **Key Words:** Asparagopsis taxiformis, Zonaria farlowii, feed supplementation, greenhouse gas mitigation, in-vitro rumen fermentation, macroalgae 1. Introduction Methane (CH₄) accounts for more than 10% of the greenhouse gas (GHG) emissions from the US (Myhre, 2013) and enteric fermentation from ruminant animals accounts for approximately 25% of the total anthropogenically produced methane (NASEM, 2018). Thus, efficient strategies to lower enteric CH₄ production could result in a significantly reduced carbon footprint from agriculture and animal production more specifically. In-vitro studies have demonstrated that some brown and red macroalgae can inhibit microbial methanogenesis (Machado, 2014) and they have been suggested as feed supplements to reduce methanogenesis during enteric fermentation (Machado, 2016; Dubois, 2013; Wang, 2008). In addition to its methane reducing affect, utilization of these macroalgae could promote higher growth rates and feed conversion efficiencies in ruminants via the potential net energy yield from the redistribution of energy from the microbial methanogenesis pathway, into more favorable pathways (i.e volatile fatty acids) (Hansen, 2003; Marín, 2009). Therefore, macroalgae

feed supplementation may be an effective strategy to simultaneously improve profitability and sustainability of beef and dairy operations.

In a recent study (Machado et al. 2014), the red alga *Asparagopsis taxiformis* stood out as the most effective species of seaweed to reduce methane production. In this work, the effect of a large variety of macroalgal species including freshwater, green, red and brown algae on CH₄ production during *in-vitro* incubation were compared and the obtained results showed that *A. taxiformis* amendment yielded the most significant reduction (~98.9%) of CH₄ production.

A major barrier to the implementation of an *A. taxiformis* based methane mitigation strategy is the availability of the seaweed, which has led to the exploration of alternative seaweed species. Previous investigations have collected *A. taxiformis* during diving excursions off the coast of Australia. Due to the potential differences in effectiveness based on the location and growing conditions from which the seaweed is collected and the financial burden of transport, we tested the potential of two different species of subtidal macroalgae (*A. taxiformis* and the brown alga, *Zonaria farlowii*) from Southern California for their ability to mitigate methane production during *in-vitro* rumen fermentation.

2. Materials and Methods

2.1 Experimental Design

To determine the effect of two locally sourced macroalgae species on methane production during in-vitro rumen fermentation, *Asparagopsis taxiformis* and *Zonaria farlowii* were supplemented to an *in-vitro* gas production system at a dose rate of 5% DM. Rumen fluid was diluted 3-fold with artificial saliva buffer (Oeztuerk et al., 2015). After homogenization, 200 ml of the mixture was allocated to 300 ml vessels fitted with Ankom head units (Ankom Technology RF Gas

Production System, Macedon, NY, USA). Each vessel recieved 2 g of rumen solids, and 2 g of a basic ration (Super basic ration — SBR, Table 1) commonly used in the dairy industry in California. Rumen solids and SBR were sealed in separate Ankom feed bags and seaweed was included in the respective SBR feed bags (Ankom, Macedon, NY). Vessels were placed in a shaking water bath (39°C) and incubated while mixed at 40 rpm. Foil gas bags (Restek, USA) were connected to the Ankom head units to collect gas at 24 and 48 hours respectively.

2.2 Pacific Coast Seaweed Collection and Preparation

Asparagopsis taxiformis and Z. farlowii were collected from Little Fisherman's Cove on the leeward side of Santa Catalina Island, ~35 km off the coast of Southern California, USA (Figure 1). The seaweed was shipped on ice to the University of California, Davis, where it was dried at 55°C for 72 hours and ground through a 2 mm Wiley Mill (Thomas Scientific, Swedesboro, NJ).

2.3 Rumen Fluid Collection

All animal procedures were performed in accordance with the Institution of Animal Care and Use Committee (IACUC) at University of California, Davis under protocol number 19263. Rumen content was collected from a rumen fistulated Holstein cow, housed at the UC Davis Dairy Research and Teaching Facility Unit. The rumen fluid donor was fed a dry cow total mixed ration (50% wheat hay, 25% alfalfa hay/manger cleanings, 21.4% almond hulls, and 3.6% mineral pellet, Table 1). Two liters of rumen fluid and 30 g of rumen solids were collected 90 min after morning feeding. Rumen content was collected via transphonation using a perforated PVC pipe, 500 mL syringe, and Tygon tubing (Saint-Gobain North America, PA, USA). Fluid

was strained through a colander and 4 layers of cheesecloth into a 4 L pre-warmed, vacuum insulated container and transported to the laboratory.

2.4 Greenhouse Gas Analysis

Methane and CO₂ were measured from gas bags using an SRI Gas Chromatograph (8610C, SRI, Torrance, CA) fitted with a 3'x1/8" stainless steel Haysep D column and a flame ionization detector (FID) with methanizer. The oven temperature was held at 90°C for 5 minutes. Carrier gas was high purity hydrogen at a flow rate of 30 ml/min. The FID was held at 300°C. A 1 mL sample was injected directly onto the column. Calibration curves were developed with Airgas certified CH₄ and CO₂ standard (Airgas, USA).

2.5 Statistical Analysis

Differences in CH₄ and CO₂ production were determined using unpaired parametric t-tests with Welch's correction conducted in Graphpad Prism 7 (Graphpad software Inc, La Jolla, CA). Significant differences among treatments were declared at $p \le 0.05$.

Table 1. Composition of dry cow diet and super basic ration (SBR).

	Dry Cow Diet		SBR	
Ingredient				
	Alfalfa	25%	Alfalfa	70%
	Wheat	50%	Dried distillers grain	15%

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Almond hulls Rolled corn 21.40% 15% Mineral pellets 3.60% 3. Results 3.1 Gas production profile of in vitro fermentation of rumen fluid amended with 5% A. taxiformis At a dose rate of 5% DM, A. taxiformis reduced methane production by 74% after 48 hours of invitro rumen fermentation ($p \le 0.01$, Figure 2B) and daily methane production remained nearly identical in the presence of A. taxiformis on both days (7.1±1.9 ml (g DM)^-1 and 6.6±2.5 ml (g DM)^-1 after 24 and 48 hours respectively). Methane production in the control vessels increased by 76% after 48 hours of incubation (20.3±11 ml (g DM)^-1 and 35.5±8.5 ml (g DM)^-1 at 24 and 48 hours respectively). While methane production varied with 5% DM inclusion of A. taxiformis, CO₂ production remained similar between treatment (41.9±6.2 ml (g DM)^-1 and 65.23±9.1 ml (g DM)^-1 at 24 and 48 hours respectively) and control vessels (47.4±13.4 ml (g DM)^-1 and 69.0 ± 15.9 ml (g DM) $^{-1}$ at 24 and 48 hours respectively). 3.2 Gas production profile of in vitro fermentation of rumen fluid amended with 5% Z. At a dose rate of 5% DM, Z. farlowii reduced methane production by 11% after 24 hours of in vitro rumen fermentation ($p \le 0.04$, Figure 3A). Daily methane production decreased slightly at 48 hours compared to 24 hours of incubation for both the control and treatment vessels (Control

151 $= 62.5\pm3.3$ ml (g DM) $^{-1}$ and 51.4 ± 2.9 ml (g DM) $^{-1}$ CH₄, at 24 and 48 hours respectively; 152 treatment = 55.3 ± 2.7 and 45.9 ± 3.7 ml (g DM)^-1 CH₄, at 24 and 48 hours respectively). 153 While methane production decreased slightly for all vessels at 48 hours, CO₂ production 154 nearly doubled (Control = 74.1 ± 7.7 ml (g DM)^-1 and 117.9 ± 14.6 ml (g DM)^-1 CO₂, at 24 155 and 48 hours respectively; treatment = 67.6 ± 4.1 ml (g DM) $^{-1}$ and 114.2 ± 6.0 ml (g DM) $^{-1}$ 156 CO₂, at 24 and 48 hours respectively). Carbon dioxide production from vessels amended with 157 5% DM of Z. farlowii did not differ from the control vessels at 24 or 48 hours ($p \le 0.27$ and $p \le$ 158 0.70 respectively). 159 160 4. Discussion 161 With increasing interest in feed-based biotic methane mitigation strategies fueled by legal 162 directives aimed at reducing methane production from the agricultural sector, identification of 163 local biotic feed-supplements will be critical to render large-scale methane mitigation strategies 164 economical. 165 The data presented here suggest that subtidal macroalgae from Santa Catalina Island, 166 Southern California reduced the *in-vitro* production of CH₄ when added to rumen content from 167 California dairy cattle, suggesting that California seaweed might represent a viable option for use 168 in feed based methane mitigation strategies. In addition to demonstrating the potential of the 169 local A. taxiformis for methane mitigation during enteric fermentation, we also demonstrated 170 significant methane reduction in the brown alga Z. farlowii, a species of seaweed commonly 171 found along the Southern California Bight, without obvious impact on CO₂ production (Figures 2

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and 3, panels A and B).

The effectiveness of a macroalgae in reducing methane production during rumen incubation has been linked to the concentration of halogenated bioactives including bromoform and di-bromochloromethane (Machado, 2016). However, in contrast to A. taxiformis, which has been shown to produce several halomethane compounds, Z. farlowii amendment only reduced methane on a short time scale. These findings suggest that either the bioactives in Z. farlowii are more bioavailable but less effective or concentrated, or methane reduction is occurring via a different compound or a different mode of action. Previous studies have identified multiple phenolic lipids produced by Z. farlowii from Southern California waters as possessing antimicrobial activity (Gerwick and Fenical, 1981). However, the reduction of methane in vessels amended with Z. farlowii was modest compared to those amended with A. taxiformis. Zonaria farlowii is commonly found along the Southern California Bight, which makes it a potential candidate for non-terrestrial farming operations along the Southern California Coast. A more in-depth nutrient analysis of Z. farlowii along with in-vitro assays will be essential to help determine its value for future methane mitigation strategies and to determine its potential for use in dairy operations.

5. Conclusion

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Asparagopsis taxiformis and Z. farlowii collected off Santa Catalina Island were evaluated for their ability to reduce methane production from dairy cattle fed a mixed ration widely utilized in California. The methane reducing effect of the A. taxiformis and Z. farlowii described in this study makes these macroalgae promising candidates for biotic methane mitigation strategies in the largest milk producing state in the US. With expected growth in livestock production, it is

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necessary to investigate and confirm the effect of these macroalgae in-vivo, in order to ensure that farmers have sufficient incentive to implement such strategies. 6. References Myhre G, Shindell D, Bréon F-M, Collins W, Fuglestvedt J, Huang J, Koch D. Lamarque J.-F, Lee D, Mendoza B, Nakajima T, Robock A, Stephens G, Takemura T, Zhang H. Anthropogenic and Natural Radiative Forcing. 2013 In: Climate Change: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate. Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA. 2013. https://www.ipcc.ch/pdf/assessmentreport/ar5/wg1/WG1AR5_Chapter08_FINAL.pdf. Accessed 15 March 2018. Academies of Science Engineering and Medicine **National** (NASEM). Improving Characterization of Anthropogenic Methane Emissions in the United States. The National Academies Press, Washington, DC. 2018. https://www.nap.edu/read/24987. Accessed 15 March 2018 Machado L, Magnusson M, Paul NA, de Nys R, Tomkins N. Effects of Marine and Freshwater Macroalgae on In-Vitro Total Gas and Methane Production. PLOS ONE 2014;9:e85289 Machado L, Magnusson M, Paul NA, Kinley R, de Nys R, Tomkins N. Dose-response effects of Asparagopsis taxiformis and Oedogonium sp. on in-vitro fermentation and methane production. J Appl Phycol. 2016;28:1443-1452.

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Figure 1. Map showing the location of Santa Catalina Island relative to the Southern California mainland. Inset: The red alga Asparagopsis taxiformis (A) and the brown alga Zonaria farlowii (B) were collected (2-5 m depth) in Little Fisherman's Cove, located ~0.6 km from the USC Wrigley Marine Science Center. Figure 2. Methane and CO₂ production during *in-vitro* fermentation of rumen fluid amended with A. taxiformis. Production of CH₄ [ml (g DM)^-1] and CO₂ [ml (g DM)^-1] from vessels without (n=4) and with 5% (n=4) A. taxiformis as additive. Methane and CO₂ were measured at 24 h (A & B respectively) and 48 h (C & D respectively). "*" indicate significant difference (p value ≤ 0.05), "ns" indicates not significant. Error bars represent the standard error from the mean. Figure 3. Methane and CO₂ production during *in-vitro* fermentation of rumen fluid amended with Z. farlowii. Production of CH₄ [ml (g DM)^-1] and CO₂ [ml (g DM)^-1] from vessels without (n=3) and with 5% (n=3) Z. farlowii as additive. Methane and CO₂ were measured at 24 h (A & B respectively) and 48 h (C & D respectively). "*" indicate significant difference (p value ≤ 0.05), "ns" indicates not significant. Error bars represent the standard error from the mean.





