Growth and nutrient uptake rates of duckweed cultivated on anaerobically digested dairy manure

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Abstract: Nutrient cycling from flushed dairy manure systems to croplands is a national research priority for sustainable dairy operations, resource utilization, and environmental protection. Cultivating aquatic plants on dairy wastewaters has been considered an effective approach for wastewater treatment/reuse and nutrient recycling. This study aimed to investigate nutrient uptake and biomass production of duckweed strains on dairy wastewater. Three duckweed strains, namely Landoltia punctata 0128, Lemna gibba 7589, and Lemna minuta 9517, were cultivated on anaerobically digested (AD) dairy manure wastewater over a period of 28 days. The highest reduction rate of total nitrogen (TN) was achieved by L. punctata from the AD dairy manure with a dilution ratio of 1:18 (83.1 mg·L⁻¹ TN) at 11.6% (±1.6%). The highest reduction rate of total phosphorus (TP) was achieved by L. punctata from the AD dairy manure with a dilution ratio of 1:27 (6.7 mg·L⁻¹ TP) at 15.4% (±4.4%). The corresponding fresh weight-based growth rate constants of L. punctata were 0.11 g·d⁻¹ and 0.17 g·d⁻¹ for the dilution ratios of 1:18 and 1:27, respectively. It has been shown that, among the three duckweed strains tested in this study, L. punctata has the greatest potential to be cultivated on the medium of diluted AD dairy manure for best N and P reduction and biomass production.

Keywords: Flushed dairy manure; Nutrient recovery; Wastewater treatment; Anaerobic digestion; Dotted duckweed (Landoltia punctata); Fat duckweed (Lemna gibba); Least duckweed (Lemna minuta)

As of 2018, Idaho was the fourth-largest producer of milk in the United States with 490 dairy farms and about 600 000 cows producing over 14.6 billion pounds of milk (USDA, 2018). About 59% of the Ida-
ho dairies are in the Magic Valley Region of Southern Idaho (IDA, 2018) which generate large volumes of wastewater stored in lagoon systems (Chen et al., 2014). The dairy wastewater contains large quantities of nitrogen (N) and phosphorus (P) which risk being transferred into the environment via run-off from over-irrigation and ammonia volatilization (Adhikari et al., 2015; Zhao et al., 2014). Concerns of the environmental impacts caused by concentrated animal productions have resulted in reporting requirements, mandatory management plans, and other regulations (Chen et al., 2014; Leytem et al., 2013; IDEQ, 2018). Identifying potential nutrient management technologies that utilize N and P contained in flushed dairy manure is essential to the sustainability of the Idaho dairy industry.

Duckweed is a family of tiny, flowering plants which belong to the Lemnaceae family and adapt to a variety of weather environments worldwide. There are five different genera including Lemna, Spirodela, Wolffia, Wolffiiella and Landoltia with 37 different species (Cheng and Stomp, 2009; Ziegler et al., 2015). Many species grow in various locations throughout the world and form distinct populations called strains. Each duckweed strain has a specific set of nutrient and growth characteristics under a range of environmental conditions. Duckweed has been studied as a means of domestic and agricultural wastewater treatment in the past 30 years (Mohedano et al., 2012). The interest in duckweed has stemmed from its ability to 1) double its biomass weight in as little as 2 to 3 days under ideal conditions, 2) uptake N and P from nutrient rich sources, and 3) be harvested and stocked without many challenges (Rusoff et al., 1980; Sooknah et al., 2004; Adhikari et al., 2015).

Previous studies have utilized duckweed for the removal of nutrients from swine and cattle wastewaters in large-scale applications such as constructed wetlands, wastewater stabilization lagoons, and storage ponds (Al-Nozaily, 2001; Mohedano et al., 2012; Adhikari et al., 2015). These studies have shown that the cultivation of duckweed is a promising technology to recycle nutrients from wastewaters. However, there are few studies on N and P uptake from anaerobically digested (AD) dairy manure by duckweed cultivation. To address nutrient management challenges, the aim of this study was to investigate the nutrient recovery and biomass production through duckweed cultivation on AD dairy manure for waste management and resource utilization in a value-added manner, resulting in a sustainable dairy industry in Idaho.

The specific objectives of this study were to 1) investigate the feasibility of cultivating duckweed on AD Idaho dairy manure, 2) identify the achievable N and P reduction rates by cultivation of the selected duckweed stains, and 3) determine the duckweed biomass growth rate constants for further research of scale-up cultivation.

1 Materials and methods

1.1 Pre-culturing duckweed strains

Three duckweed strains, i.e., Landoltia punctata 0128 (L. punctata), Lemna gibba 7589 (L. gibba), and Lemna minuta 9517 (L. minuta), were obtained from the Rutgers Duckweed Stock Cooperative and aseptically pre-cultured in 114 mm × 86 mm × 102 mm cultivation containers on 200 mg·L⁻¹ of Hoagland’s No. 2 Basal Salt Mixture for seven days (Zhao et al., 2014). The Hoagland’s No. 2 Basal Salt Mixture (hereafter named as Hoagland E-Medium) contains 115.03 mg·L⁻¹ of ammonium phosphate monobasic (NH₄H₂PO₄), 2.86 mg·L⁻¹ of boric acid (H₃BO₃), 656.40 mg·L⁻¹ of calcium nitrate tetrahydrate [Ca(NO₃)₂·4H₂O], 0.08 mg·L⁻¹ of cupric sulfate pentahydrate (CuSO₄·5H₂O), 5.32 mg·L⁻¹ of ferric tartrate (C₃Fe(H₂O)₆), 240.76 mg·L⁻¹ of magnesium sulfate anhydrous (MgSO₄), 1.81 mg·L⁻¹ of manganese chloride tetrahydrate (MnCl₂·4H₂O), 0.02 mg·L⁻¹ of molybdenum trioxide (MoO₃), 606.6 mg·L⁻¹ of potassium nitrate (KNO₃), and 0.22 mg·L⁻¹ of zinc nitrate hexahydrate [Zn(NO₃)₂·6H₂O]. Once a double-layer of duckweed growth was observed, the duckweed strains were transferred to larger rectangular PET cultivation containers (0.267 m²) for 30 days to allow for further growth as inoculum samples for testing.

1.2 Batch testing

1.2.1 Nutrient analysis

AD dairy manure was collected from an anaerobic digester operated on a local dairy in Southern Idaho. Chemical oxygen demand (COD), total nitrogen (TN), total Kjeldahl nitrogen (TKN), nitrates and nitrates (NO₃⁻+NO₂⁻), ammonia nitrogen (NH₃-N), ortho-phosphate-phosphorus (o-PO₄-P), total phosphorus (TP), and suspended solids (SS) of the collected AD manure were analyzed using a spectrophotometer (DR5000, Hach, USA) based on Method 8000, Method 10242, Method 10031, Method 10214, Method 10127, and Method 8006, respectively (Hatch Company, 2005). Total and volatile solids of the collected AD manure were analyzed by the standard methods of APHA (2015). Analyses of pH, dissolved oxygen (DO) and electric conductivity (EC) were conducted using the Sper Scientific 850049 water meter kit (Scottsdale, AZ). Table 1 summarizes the nutrient characteristics of the AD dairy manure used in this study.

1.2.2 Growth media preparation

Growth media were prepared to establish the desired nutrient concentrations by mixing AD dairy manure and deionized water. These corresponded to TN concentrations of 114, 84 and 57 mg·L⁻¹, respectively, and the initial pH values of the media were 8.11±0.03, 8.12±0.04, and 8.19±0.02, respectively, for the media with the dilution ratios of 1: 13, 1: 18 and 1: 27.

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The pH of the media was then adjusted to 6.7 every 2 days with 5% (v/v) acetic acid to keep an ideal growth environment and reduce the loss of ammonia nitrogen (NH$_3$-N) from the cultivation containers. After the pH of the media was adjusted to 6.7±0.01, the initial EC, DO and temperature of the media were measured at 1660±12 µS·cm$^{-1}$, 4.7±0.1 mg·L$^{-1}$, and 21.4±0.1 °C at the dilution ratio of 1: 13, 1.331±16 µS·cm$^{-1}$, 3.8±0.2 mg·L$^{-1}$, and 23.4±0.1 °C at the dilution ratio of 1: 18, and 1.066±10 µS·cm$^{-1}$, 4.5±0.1 mg·L$^{-1}$, and 23.6±0.1 °C at the dilution ratio of 1: 27, respectively. The media were then transferred to cultivation containers with transparent lids to prevent evapotranspiration. The cultivation containers have dimensions of 114 mm × 86 mm × 102 mm, a surface area of 0.011 6 m$^2$, and a total volume of 300 mL. An average of 0.46, 0.57 and 0.61 g of L. punctata, L. gibba, and L. minuta were added to the cultivation containers to cover 25% of the surface area of each container.

### 1.2.3 Experimental setup

Batch tests were conducted inside an environmental chamber (Environmental Growth Chambers, Chagrin Falls, Ohio) at an average light intensity of 10 000 lux and a photoperiod of 16: 8 (light: dark). The experiments consisted of a total 36 cultivation containers having a working volume of 200 mL in each container. Of the 36 cultivation containers, the three duckweed strains, i.e., L. punctata, L. gibba, and L. minuta, were applied in triplicates to 27 cultivation containers containing the media as described above. Nine control cultivation containers (without duckweed inoculation) consisted of the same media with three dilution ratios (1: 13, 1: 18 and 1: 27) in triplicates.

### 2 Sampling and analysis

Liquid medium samples were extracted from each of the cultivation containers with the duckweed treatments and the control for nutrient analysis on day 0, 4, 8, 12, 16, 20, 24, and 28, and measured for temperature, TN, TKN, NO$_3$-N+NO$_2$-N, NH$_3$-N TP, o-PO$_4$-P, pH, DO, and EC based on the methods mentioned previously. The duckweed samples were carefully filtered from the cultivation containers and separated from the medium using course tulle. Residual liquid on duckweed surfaces was blotted with absorbent paper and fresh weight of duckweed was measured with an analytical balance (Mettler AE 260 Delta Range). After weighting, the duckweed biomass samples were carefully re-inoculated back to their containers immediately.

The rate constant of the fresh weight-based growth (hereafter named growth rate constant, g·d$^{-1}$) and its standard deviation were established using a linear regression equation (Eq.1) (Cheng et al., 2009; Xu et al., 2010).

$$C_t = k \cdot t + C_0$$  \hspace{1cm} (1)

where $C_0$ is the initial duckweed fresh weight (g), $C_t$ (g) is the duckweed fresh weight (g) at time $t$ (days), and $k$ is the duckweed growth rate constant (g·d$^{-1}$).

The initial and final nutrient concentrations of the media were used to find the percent TN and TP reduction rates (Sooknah et al., 2004; Adhikari et al., 2015). The differences in TN and TP reduction rates between the media with the duckweed (treatments) and without the duckweed (control) were used to calculate the percentages of TN and TP uptake rates by the duckweed strains.

The fresh weights of duckweed biomass samples and the nutrient concentrations in the media were analyzed statistically using a two-way analysis of variance (ANOVA) to determine the validity of the experimental results between the treatments and controls by the statistical software of SPSS (SPSS Inc., Chicago, IL, USA). Tukey simultaneous tests were conducted to determine the statistical differences between treatments.

### 3 Results and discussion

#### 3.1 Nitrogen and phosphorus uptake by duckweed

Total nitrogen (TN) and total phosphorus (TP) concentrations within the media for the 28-day cultivation period are shown in Figure 1. The growth media was modelled to maintain conditions that would apply to the cultivation environment of a dairy lagoon where bacteria, algae, and duckweed would thrive symbiotically among the batch samples. The three duckweed strains grew well on the cultivation media from diluted AD dairy manure with the dilution ratios of 1: 18 (84 mg·L$^{-1}$ of TN) and 1: 27 (57 mg·L$^{-1}$ of TN). However, the three duckweed strains perished after 4 days of cultivation on the dilution ratio of 1: 13 (114 mg·L$^{-1}$ of TN) which indicated that the nutrient concentration in the medium with the dilution ratio of 1: 13 was too high for the duckweed strains. In the media with dilution ratios of
Kevin Kruger, et al: Growth and nutrient uptake rates of duckweed cultivated on anaerobically digested dairy manure

1: 18 and 1: 27 AD dairy manure, TN and TP were removed quickly in the first 12 days of cultivation, and the concentrations of TN and TP slowly decreased in the next 16 days of cultivation. This is in some disagreement with the study by Xu et al. (2010) in which the greatest ammonium nitrogen (NH₄-N) recovery occurred in the first 20 days of cultivation. Therefore, it is important to identify the period of fast nutrient reduction rates which in turn helps determine the timing and types of nutrients that need to be replenished to keep a healthy duckweed production system.

The changes of nutrient concentrations along with the reduction rates of TN and TP in the cultivation media are shown in Table 2. The highest TN reduction rate of 92.5%±0.3% was observed from the media with the dilution ratio of 1: 18 treated by L. punctata. Generally, TP reductions of 66.0%±0.6% to 85.6%±4.4% were achieved, and the highest TP reduction rate of 85.6%±4.4% came from the media with the dilution ratio of 1: 27 treated by L. punctata, which are in agreement with the previously reported N and P reduction ranges of 60%–98% from pretreated livestock wastewaters (Adhikari et al., 2015; Mohedano et al., 2012; Xu et al., 2010). In addition to the uptake by duckweed, reduction in N and P in the batch experiments could occur due to other means such as ammonia volatilization, nitrification/denitrification, bacterial transformation, sedimentation and algal uptake (Mohedano et al., 2012; Xu et al., 2010). The controls in this study were used to explore the possibility of N and P reduction via these alternative pathways to determine the uptake by the duckweed strains.

Figure 2 shows the results of TN and TP uptake by the duckweed strains from the cultivation media after a 28-day batch growth. These results were determined by the difference between the N and P reductions from the batch system (Table 2) of the duckweed treatments and the controls. At the dilution ratio of 1: 18, the duckweed achieved TN and TP reduction rates of 11.6%±1.6% and 10.6%±1.1% for L. punctata, 7.9%±1.6% and 9.3%±2.7% for L. gibba, and 4.1%±1.2% and 3.2%±0.8% for L. minuta. At the dilution ratio of 1: 27, reduction rates of 7.8%±2.9% of TN and 15.4%±4.4% of TP, 3.3%±0.9% of TN and 12.9%±2.4% of TP, and 1.6%±0.8% of TN and 5.6%±1.7% of TP for L. punctata, L. gibba, and L. minuta, respectively, were observed. In our batch systems, the combined alternative pathways (i.e., ammonia volatilization, nitrification/denitrification, bacterial transformation, sedimentation and algal uptake) contributed to a N reduction in the range of 65.0% to 80.8% in the growth media with the dilution ratio of 1: 18 and 53.8% to 61.6% in the growth media with the dilution ratio 1: 27. The main pathways for P reduction are bacterial transformations, biomass absorption either by duckweed or algae, and sedimentation. Mohedano et al. (2012) reported that P was most strongly reduced in the anaerobic stage of a lagoon system probably due to...

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Table 2  Total nitrogen (TN) and total phosphorus (TP) reductions in diluted anaerobically digested dairy manure with the dilution ratios of 1 : 18 and 1 : 27 treated with L. punctata, L. gibba and L. minuta at the end of the 28-day test

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Dilution ratio</th>
<th>TN</th>
<th>TP</th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Initial (mg·L⁻¹)</td>
<td>Final (mg·L⁻¹)</td>
<td>Reduction (%)</td>
<td>Initial (mg·L⁻¹)</td>
<td>Final (mg·L⁻¹)</td>
</tr>
<tr>
<td>L. punctata</td>
<td>1:18</td>
<td>83.1±1.4</td>
<td>6.3±0.2</td>
<td>92.5±0.3</td>
<td>9.9±0.0</td>
<td>2.5±0.1</td>
</tr>
<tr>
<td>L. gibba</td>
<td>83.1±1.4</td>
<td>9.3±0.1</td>
<td>88.8±0.3</td>
<td>9.9±0.0</td>
<td>3.3±0.0</td>
<td>66.0±0.6</td>
</tr>
<tr>
<td>L. minuta</td>
<td>83.1±1.4</td>
<td>12.5±0.3</td>
<td>85.0±1.5</td>
<td>9.9±0.0</td>
<td>3.6±0.0</td>
<td>63.8±0.3</td>
</tr>
<tr>
<td>Control</td>
<td></td>
<td>83.1±1.4</td>
<td>15.9±1.3</td>
<td>84.7±2.3</td>
<td>6.7±0.0</td>
<td>1.0±0.3</td>
</tr>
<tr>
<td>L. punctata</td>
<td>1:27</td>
<td>55.8±1.4</td>
<td>8.6±1.4</td>
<td>84.7±2.3</td>
<td>6.7±0.1</td>
<td>1.4±0.6</td>
</tr>
<tr>
<td>L. gibba</td>
<td>55.8±1.4</td>
<td>11.0±0.6</td>
<td>80.2±1.5</td>
<td>6.7±0.0</td>
<td>1.6±0.1</td>
<td>75.7±1.7</td>
</tr>
<tr>
<td>L. minuta</td>
<td>55.8±1.4</td>
<td>12.0±0.3</td>
<td>78.5±0.4</td>
<td>6.7±0.0</td>
<td>2.3±0.6</td>
<td>70.2±0.0</td>
</tr>
<tr>
<td>Control</td>
<td></td>
<td>55.8±1.4</td>
<td>12.9±0.1</td>
<td>76.9±0.8</td>
<td>6.7±0.0</td>
<td>2.3±0.6</td>
</tr>
</tbody>
</table>

Data in the table are mean ± S.D. (the standard deviation of triplicate samples).

3.2 Duckweed growth rate

It was observed that the first eight days of cultivation were most productive with L. punctata and L. gibba covering over three quarters of the surface area. L. minuta did not double its biomass in the first eight days of cultivation and covered less than half of the surface area. Throughout the entire cultivation period of 28-days for both the dilution ratios of 1:18 and 1:27, L. punctata and L. minuta maintained a healthy complexion on the surface of the AD dairy manure and the Hoagland E-medium. However, at day 16 the fronds on the first layer of L. gibba started turning white but maintained buoyancy to float on the surface of the PET containers until day 28 for the dilution ratio of 1:18. A double layer of L. punctata and L. gibba formed at the surface of the dilution ratio of 1:27 by the 28th day. Filamentous algae growth was visible on the surface of the controls as well as on the sides of the cultivation containers. Algae growth was not visible on the surface of the duckweed treatments but was visible on the sides of the containers by the end of the cultivation period.

The growth of duckweed biomass was quantitatively analyzed by the growth rate constants which were calculated from the measured fresh weights of duckweed biomass samples cultivated with the dilution ratios of 1:18 and 1:27 through a linear model (Tables 3 and 4). As shown in Figure 3, no lag phases were observed due to the pre-culturing of the duckweed strains on the Hoagland E-Medium. The zero-order regressions fit the experimental data well with $R^2$ values in the range of 0.79 and 0.97. Growth rate constants of the three duckweed strains were higher at the dilution ratio of 1:27 with the growth rate constants of 0.17±0.02, 0.19±0.06 and 0.04±0.00 g·d⁻¹ (10.0, 8.0, 1.6 mg·d⁻¹ dry weight) for L. punctata, L. gibba, and L. minuta, respectively. In this study, the differences between the growth rate constants of L. punctata and L. gibba at the dilution ratios of 1:18 and 1:27 were significant ($P \leq 0.05$) different from that of L. minuta.
In a study conducted by Toyama et al. (2018), the growth rate constants for duckweed strains of *L. punctata* and *L. gibba* were approximately determined as 30 and 4.5 mg·d⁻¹, respectively, when cultivated on AD domestic wastewater at 30.1 mg·L⁻¹ of NH₄-N and 3.3 mg·L⁻¹ of nitrate nitrogen (NO₃-N). It is possible that the higher nitrogen concentrations in this study inhibited growth of the duckweed strains as compared to the values reported by Toyama et al. (2018).

Table 3  Linear equations describing relationships between the duckweed biomass fresh weight and cultivation time. The regression equations were found to significantly fit a linear model (*P* ≤ 0.05).

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Dilution ratio of AD manure</th>
<th>Regression equation</th>
<th><em>R</em>²</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>L. punctata</em></td>
<td>1: 18</td>
<td>y=0.11x+0.47</td>
<td>0.89</td>
</tr>
<tr>
<td><em>L. gibba</em></td>
<td>1: 18</td>
<td>y=0.11x+0.57</td>
<td>0.87</td>
</tr>
<tr>
<td><em>L. minuta</em></td>
<td>1: 18</td>
<td>y=0.04x+0.62</td>
<td>0.79</td>
</tr>
<tr>
<td><em>L. punctata</em></td>
<td>1: 27</td>
<td>y=0.17x+0.47</td>
<td>0.95</td>
</tr>
<tr>
<td><em>L. gibba</em></td>
<td>1: 27</td>
<td>y=0.19x+0.57</td>
<td>0.86</td>
</tr>
<tr>
<td><em>L. minuta</em></td>
<td>1: 27</td>
<td>y=0.03x+0.62</td>
<td>0.83</td>
</tr>
</tbody>
</table>

Further research should include the verification of the N and P contents in the duckweed biomass and an analysis of the mechanisms of N and P assimilation by bacteria, algae, and duckweed strains when cultivated on flushed AD dairy manure. These factors will potentially maximize biomass production and nutrient reduction such that the cultivation of duckweed on dairy farms could be utilized as a protein supplement for dairy cattle.

### 4 Conclusions

The results obtained from this study have shown that *L. punctata*, *L. gibba* and *L. minuta* could be cultivated on diluted AD dairy manure for nutrient recovery and biomass production. On average, the TN and TP reduction rates were 11.6% and 10.6% for *L. punctata*, 7.9% and 9.3% for *L. gibba*, and 4.1% and 3.2% for *L. minuta* from the AD dairy manure with the dilution ratio of 1: 18. The TN and TP reduction rates with the dilution ratio of 1: 27 were 7.8% and 15.4% for *L. punctata*, 3.3% and 12.9% for *L. gibba*, and 1.6% and 5.6% for *L. minuta*. It is concluded that the duckweed strains recovered approximately the same percentages of TN and TP from the AD dairy manure at the dilution ratio of 1: 18. The duckweed strains had a higher TP recovery than TN recovery when cultivated on the AD dairy manure with the dilution ratio of 1: 27. At the dilution ratios of 1: 18 and 1: 27 the growth rate constants were 0.11 g·d⁻¹ and 0.05 g·d⁻¹ for *L. punctata*, 0.11 g·d⁻¹ and 0.19 g·d⁻¹ for *L. gibba*, and 0.04 g·d⁻¹ and 0.03 g·d⁻¹ for *L. minuta*. Both *L. punctata* and *L. gibba* had significantly higher growth rate constants when compared to *L. minuta*. It has been shown that, among the three duckweed strains tested in this study, *L. punctata* has the greatest potential to be cultivated on the medium of diluted AD dairy manure for the best N and P reduction and biomass production.

**Fig. 3**  Fresh biomass of *L. punctata*, *L. gibba*, and *L. minuta* cultivated on the anaerobically digested dairy manure with the dilution ratios of 1: 18 and 1: 27. The growth rate constants of both *L. punctata* and *L. gibba* were significantly higher than that of *L. minuta* for both the dilution ratios of 1: 18 and 1: 27 (*P* ≤ 0.05).

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