1	Full-fat black soldier fly larvae (Hermetia illucens) meal and paste in extruded diets for Atlantic salmon
2	(Salmo salar): Effect on physical pellet quality, nutrient digestibility, nutrient utilization and growth
3	performances
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23 Abstract

24 The present study investigated the effect of graded levels of black soldier fly larvae (BSFL) 25 (Hermetia illucens) meal and paste on physical pellet quality, digestibility and utilization of nutrients and growth performances in extruded diets for Atlantic salmon (Salmo salar). A total of 1260 Atlantic salmon 26 27 with 34 g of mean initial weight were randomly distributed into 21 fiberglass tanks and fed with one of 28 seven isonitrogenous, isolipidic and isoenergetic diets for seven weeks. The experimental diets consisted 29 of a control diet based on fishmeal, soy protein concentrate, corn gluten, faba bean and fish oil (Control-30 1); three diets with increased levels of full-fat BSFL meal, substituting 6.25% (6.25IM), 12.5% (12.5IM) 31 and 25% (25IM) of the protein content of Control-1; two diets with increased levels of full-fat BSFL 32 paste, substituting 3.7% (3.7IP) and 6.7% (6.7IP) of protein from Control-1 and an extra control diet with 33 0.88% of formic acid (Control-2). Pellet durability and hardness were overall high for all diets. However, 34 the expansion, sinking velocity and water stability of feed pellets were lower with increased inclusion of 35 BSFL meal and paste. Dietary inclusion of BSFL meal or paste did not affect the feed intake of fish. 36 Further, replacing the protein content of the control diet with up to 12.5% and 6.7% of BSFL meal and 37 paste, respectively, did not compromise fish growth rate or feed conversion ratio, although polynomial 38 contrast analysis showed that increasing BSFL meal level in the diet linearly (p<0.05) decreased these 39 parameters. However, apparent digestibility coefficient (ADC) of protein and lipid, protein efficiency 40 ratio and lipid retention were reduced linearly (p < 0.05) with increasing inclusion level of BSFL meal. 41 Further, increasing dietary levels of BSFL paste linearly (p<0.05) reduced ADC of protein, protein 42 efficiency ratio and phosphorous retention. Despite the decreased ADC of protein, protein retention was 43 not compromised by the inclusion of BSFL meal or paste. Replacement of 25% of dietary protein with 44 BSFL meal decreased (p<0.05) growth rate, accompanied by lower (p<0.05) ADC and utilization of lipids 45 and protein efficiency ratio. The present study showed that BSFL meal and paste could replace up to

46	12.5% and 6.7% of dietary protein, respectively, without compromising growth performance in Atlantic
47	salmon.
48	Keywords: black soldier fly larvae; Atlantic salmon; physical pellet quality; nutrient utilization; growth
49	performance
50	Abbreviations
51	FCR: feed conversion ratio; N: nitrogen; BSFL: black soldier fly larvae; AA: amino acids; Ca: calcium; DM:
52	dry matter; SPC: soy protein concentrate; ADC: apparent digestibility coefficient; WHC: water holding
53	capacity; FA: fatty acids; Y: yttrium; Mg: magnesium; K: potassium; Na: sodium; P: phosphorous; SME:
54	specific mechanical energy; SGR: specific growth rate; FBW: final body weight; PER: protein efficiency
55	ratio; LER: lipid efficiency ratio; SFA: saturated fatty acids
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70 1. Introduction

71 In recent years, insects have received growing attention as a sustainable ingredient for aquafeed 72 production (Henry et al., 2015; Makkar et al., 2014; Nesic and Zagon, 2019) although the production of 73 insects in sufficient volumes to compete with fishmeal and plant protein sources is yet to be achieved 74 (Sogari et al., 2019). The production of insects has environmental benefits such as lower greenhouse gas 75 and ammonia emissions (Oonincx et al., 2010), high land use efficiency (Alexander et al., 2017) and 76 efficient nutrient conversion (Oonincx et al., 2010; van Huis, 2013). The feed conversion ratio (FCR) of 77 insects fed food by-products ranged from 1.4-19.1 and nitrogen (N) conversion efficiency ranged from 78 22-87% depending on the insect species and growth media (Oonincx et al., 2015). The use of processed 79 insects in feed for aquaculture animals was recently allowed by the European Commission (Regulation 80 2017/893/EC, 2017), which promotes upscaling of this novel feed ingredient.

81 One of the most promising insect species for feed purposes is black soldier fly larvae (BSFL) 82 (Hermetia illucens) (van Huis, 2013). BSFL efficiently convert low-grade organic matter to high-quality 83 nutrients (Diener et al., 2009). As shown by Oonincx et al. (2015), the N conversion efficiency can reach 84 up to 43-55% in black soldier fly fed diets composed of food by-products. BSFL contain a moderate level 85 of protein (31-59%) and has an amino acid (AA) profile closer to fishmeal and superior to soybean meal 86 (Barroso et al., 2014; Makkar et al., 2014; Nogales-Mérida et al., 2019). In addition, BSFL is a good source 87 of lipid (11–49%) (Makkar et al., 2014; Nogales-Mérida et al., 2019) and minerals, particularly calcium 88 (Ca) (Finke, 2013; Makkar et al., 2014).

BSFL have successfully been used as a protein and lipid source in diet for Atlantic salmon (*Salmo salar*) reared in freshwater and seawater. Dietary inclusion of dried, defatted and chitin reduced BSFL
meal (60%) combined with BSFL oil (4.8%) (Belghit et al., 2018) and dried BSFL meal (10-20%) (Fisher et al., 2020) resulted in similar growth performance when compared with fishmeal and other protein
sources in Atlantic salmon pre-smolts. Similar findings have also been shown in Atlantic salmon post-

smolts when feeding partially defatted, dried BSFL meal (5-15%) (Belghit et al., 2019b; Lock et al., 2016).
Hence, previous research demonstrated that dried or partially defatted BSFL meal, has potential as an
alternative protein source in salmon feeds.

97 When considering commercial production, the processing of insect meal, particularly defatting is an additional cost. Thus, the use of whole BSFL meal is more cost-efficient. The use of full-fat insect 98 99 meals can be a challenge to the feed industry due to its high lipid content that can interfere with the 100 extrusion process (Lin et al., 1997), hence reducing the pellet quality (Sørensen et al., 2009). Thus, 101 processing of BSFL biomass into partially defatted protein-rich meal has become a common practice, 102 which allow high inclusion levels of insect meal in fish diets without reducing the technical quality of 103 extruded diets (Dumas et al., 2018). Several studies have used full-fat dried BSFL meal in pelleted diets 104 for rainbow trout (Sealey et al., 2011) and yellow catfish (Xiao et al., 2018), and extruded diets for 105 rainbow trout (Józefiak et al., 2019b) and Siberian sturgeon (Józefiak et al., 2019a). However, none of 106 them reported information on feed processing conditions or the impact of full-fat BSFL meal on extruder 107 parameters and physical pellet quality.

108 High-temperature processing can reduce the nutritional quality of protein feed resources 109 (Ljøkjel et al., 2000; Opstvedt et al., 1984; Opstvedt et al., 2003). Thus, processing BSFL at low 110 temperatures to produce a paste, maintaining the nutritional value and reducing the production cost, 111 could be beneficial for the aquaculture industry. As reported by Xu et al. (2020) feeding diets containing 112 undried BSFL pulp (4.4-17.5%) to mirror carp did not affect growth rate and FCR. To our knowledge, 113 there is no literature available regarding the use of undried full-fat BSFL ingredients in salmon diets. 114 Therefore, the present study used two types of BSFL; full-fat dried BSFL meal and BSFL paste 115 (ground frozen BSFL preserved with formic acid) and investigated their effect on nutrient digestibility, 116 nutrient utilization and growth performances when added in graded levels in extruded diets for Atlantic

117 salmon pre-smolts. In addition, the effect of increasing levels of BSFL meal or paste on extruder 118 parameters and physical pellet quality was also investigated.

119 2. Materials and Methods

120 2.1 Experimental Diets

121 BSFL meal and BSFL paste were produced at HiProMine S.A., Poznań, Poland. The BSFL feed was 122 normalized in terms of dry matter (DM) content by the addition of wheat middlings (17%) to fresh 123 vegetables and fruit mix, consisting of apples (15%), carrots (50%), potatoes (15%), and cabbage (20%) 124 and established at the level of 22% DM. Fresh vegetable and fruit pre-consumer waste was ground 125 (2000 rpm/1 min, (HPM milling system, 55 kw, Poland) to pass 2 mm screen and offered ad libitum to 126 BSFL. Substrates were not contaminated by any animal products in accordance with EC regulation (no 127 1069/09).

128 At the prepupal stage (10th day of rearing), larvae were harvested, sieved through 3 mm screen 129 and washed with water on drum separator at 90 °C for 10 minutes (HPM cleaning system, Poland). A 130 batch of BSFL was divided into two parts and frozen at -50 °C to produce BSFL paste or dried for meal. The BSFL were dried first at 130 °C for 1 hour, and then at 80 °C for 23 hours until a constant weight was 131 132 reached, using a chamber air flow dryer (HiProMine S.A., Poznań, Poland) to produce BSFL meal. In the 133 case of BSFL paste, BSFL were ground to pass 4 mm screen on continue flow homogenizer (HPM milling 134 system, 25 kw, Poland) and preserved with formic acid (2.5%). The analyzed chemical composition of 135 BSFL meal and paste is shown in Table 1.

136 Seven isonitrogenous, isolipidic and isoenergetic diets were formulated. The diets were 137 formulated to meet or exceed NRC (2011) requirements for all indispensable AA and other nutrients for 138 Atlantic salmon. The experimental diets consisted of a control diet based on fishmeal, soy protein 139 concentrate (SPC), corn gluten, faba bean and fish oil (Control-1); three diets with increasing levels of full-fat BSFL meal, substituting 6.25% (6.25IM), 12.5% (12.5IM) and 25% (25IM) of the protein content of 140

141 Control-1. In addition, two diets with increasing levels of full-fat BSFL paste, substituting 3.7% (3.7IP) and 142 6.7% (6.7IP) of the protein content Control-1 and an extra control with 0.88% of formic acid (Control-2) 143 were evaluated. The diet Control-2 was included as a control for BSFL paste diets, since the larvae used 144 to produce the paste were preserved with formic acid. Yttrium oxide was included in all the diets as an 145 internal marker for the determination of apparent digestibility coefficient (ADC) of nutrients. Crystalline 146 methionine was added to all the diets to ensure that the diets met or exceeded the methionine 147 requirement of Atlantic salmon pre-smolts (0.7%, DM basis (NRC, 2011)). The ingredient and analyzed 148 chemical composition of the experimental diets are shown in Table 2.

149 2.2 Production of experimental diets

150 The experimental diets were produced at the Norwegian University of Life Sciences (NMBU) Centre for Feed Technology (Fôrtek), Ås, Norway. The BSFL paste was ground frozen using a meat 151 152 grinder (Tripas-Wexiö, RK-82, Sweden). After thawing overnight at room temperature, water was added 153 to the paste (1 kg of water per 6 kg of paste) and ground using a pump grinder (Pedrollo TR 1.1, San 154 Bonifacio (VR), Italy). The macro ingredients including SPC, fishmeal, corn gluten meal, wheat bran, faba 155 beans, wheat flour, and BSFL meal were weighed and mixed with an ISDECA mixer (60-liter paddle-156 mixer, prototype, Fôrtek, Forberg, Norway) for 2.5 minutes. The material mix was then ground in a small 157 Hammer mill (Bill bliss, horizontal, 18.5 kW, USA) with a 1 mm sieve. The ground macro ingredients 158 mixture was mixed with the micro-ingredients. BSFL paste was also added using the ISDECA mixer when 159 producing the 3.7IP and 6.7IP diets. Formic acid was added to the Control-2 diet using the ISDECA mixer 160 equipped with spray nozzles (nozzle type: 11004, Spraying Systems Co., Norway). 161 The diets were extruded in a five-section Bühler twin-screw extruder (BCTG 62/20 D, Uzwil,

Switzerland) with reduced capacity, bypassing the conditioner, fitted with four 2.5 mm die holes. A small
K-tron feeder was used to feed the material directly into the first extruder section. A screw configuration

suitable for reduced extruder capacity was used for all the diets (Figure 1). The screw speed was

increased when the BSFL meal or paste content was increased in the diet. The pellets were dried at 60
 ^oC using fan heaters (15KW, Inelco heaters, Dania-heater 15kW, Fjerritslev, Denmark) for 1 hour and
 cooled at room temperature. The dried uncoated pellets were sieved (1.6 mm screen) and the
 percentage of dust/broken pieces was calculated. Dried extruded pellets were vacuum coated with fish
 oil in Gentle Vacuum Coater (GVC) – 80 prototype (Fôrtek, Amandus-Kahl).

170 2.3 Fish experiment and rearing facilities

The fish experiment was conducted at the Center for Fish Research, NMBU, Ås, Norway. The 171 172 experimental procedures were performed in accordance with the national guidelines for the care and 173 use of animals (The Norwegian Animal Welfare Act and the Norwegian Regulation and Animal 174 Experimentation). A total of 1260 Atlantic salmon (Aqua Gen Atlantic QLT-innOva SHIELD) with 34 g of 175 mean initial weight were distributed into 21 fiberglass tanks (300 L capacity) with 60 fish per tank. The 176 fish were kept under continuous light in recirculated freshwater with a water supply of 8.5 L min⁻¹. The 177 average water temperature was 14.8 °C during the experimental period. Dissolved oxygen levels were 178 kept above 7.0 mg L^{-1} in the outlet water. Triplicate tanks of salmon were fed one of seven 179 experimental diets over a period of seven weeks. Fish were fed ad libitum (i.e. 10% excess) with 180 electrically driven belt feeders twice a day for 2.5 hours. Daily feed intake in each tank was quantified 181 according to Helland et al. (1996), by collection of uneaten feed using wedge wire screens as explained 182 by Shomorin et al. (2019). Fish weight was measured at the start and end of seven-week experimental 183 period. Fifteen fish at the start of experiment and five fish from each tank at the end of the seven-week 184 experimental period were randomly sampled and euthanized by a sharp blow to the head and stored at 185 -20 °C. All sampled fish at the start of experiment and all sampled fish per tank at the end of the seven-186 week experimental period were pooled, homogenized and freeze dried prior to analysis of the chemical 187 composition. After the seven-week experimental period, fish were fed the experimental diets for 188 another two weeks for fecal collection. Fish were carefully stripped three times with seven days interval

189 (i.e. 0, 7 and 14 days after whole body sampling) for fecal collection from the posterior intestine

190 according to Austreng (1978). The feces were immediately weighed and stored at -20 °C prior to freeze

drying. Prior to weighing, sampling and stripping, fish were anesthetized with tricaine methanesulfonate

192 (MS-222) (80 mg L^{-1}) in small aerated tanks.

193 2.4 Physical pellet quality analysis

194 Physical quality parameters of oil-coated pellets were measured. Water stability of pellets 195 during 30, 60 and 120 minutes were measured according to Baeverfjord et al. (2006). The durability of 196 the pellets was estimated in triplicates using a Doris pellet tester (AKVAsmart, Bryne, Norway) according 197 to Hansen et al. (2010). Hardness was measured using 15 randomly picked pellets from each diet with a 198 Texture analyzer equipped with a 5 kg load cell (Tinius Olsen, H5KT, Salfords, England) according to 199 Øverland et al. (2009). The width of 30 randomly selected pellets per diet was recorded using the 200 Texture analyzer (Tinius Olsen, H5KT, Salfords, England) to determine expansion. The expansion (%) was 201 calculated as ((Pellet width-die diameter) \times die diameter⁻¹) \times 100. The sinking velocity was determined 202 by measuring the mean value of the time required for 10 randomly picked pellets to sink 1 m in 17 °C 203 tap water. In addition, the water holding capacity (WHC) was measured according to Nguyen et al. 204 (2015). The degree of starch gelatinization of feed was analyzed by the Nofima AS, Ås, Norway, 205 according to Kraugerud and Svihus (2011).

206 2.5 Chemical analysis

The feed and freeze-dried feces and fish were ground. DM content was measured by oven drying at 104 °C until a constant weight was reached. The N content of feed, feces and fish were analyzed by CHNS Elemental Analyzer (Vario El Cube elemental analyzer system GmbH, Hanau, Germany) and crude protein content was determined as N × 6.25. The crude protein content of BSFL meal and paste were estimated Kjeldahl N × 6.25 according to Commission Regulation (EC) No 152/ 2009. Samples were extracted with petroleum ether and acetone (70/30) and crude lipid content was

213 determined using an Accelerated Solvent Extractor (ASE200; Dionex Corp., Sunnyvale, CA, USA). The 214 starch content was determined as described by McCleary et al. (1994) with some modifications. Briefly, 215 the samples were treated with heat-stable α -amylase and amyl glucosidase-enzymes to degrade starch 216 into glucose and glucose content was measured by a spectrometer (RX4041 Randox Daytona+, England). 217 Gross energy was measured with PARR 1281 Adiabatic Bomb calorimeter (Parr Instruments, Moline, IL, 218 USA) according to ISO 9831. AA except tryptophan contents were analyzed according to Commission 219 Regulation (EC) No 152/2009 on a Biochrom 30 AA Analyzer (Biochrom Ltd., Cambridge, UK). Tryptophan 220 content was analyzed using a Dionex Ultimate 3000 HPLC system (Dionex Softron GmbH, Germering, 221 Germany) equipped with a Shimadzu RF-535 fluorescence detector (Shimadzu Corporation, Kyoto, 222 Japan) according to Commission Regulation (EC) No 152/2009. The fatty acid (FA) content was 223 determined using Trace GC Ultra gas chromatograph (Thermo Fisher Scientific, US) according to O'fallon 224 et al. (2007) by synthesizing the FA to FA methyl esters (FAME). Yttrium (Y), Ca, magnesium (Mg), 225 potassium (K) and sodium (Na) contents were measured using a microwave plasma atomic emission 226 spectrometer (MP-AES 4200, Agilent Technologies, USA) after acid decomposition in a microwave 227 digestion system (Start D, Milestone Srl, Italy). Total phosphorous (P) content was analyzed using a 228 commercial spectrophotometric kit (PH8328, Randox laboratories, County Antrim, UK) after combustion 229 and acid digestion according to Commission Regulation (EC) No 152/2009. The chitin content of BSFL 230 meal was measured according to Finke (2007). The formic acid content in BSFL paste, Control-2 diet and 231 diets containing BSFL paste was determined using HPLC-UV at Eurofins Agro Testing Norway AS. 232 2.6 Calculations

233 Specific mechanical energy (SME) (Wh kg⁻¹) was calculated as $(2 \times \pi \times 60^{-1}) \times (S_{rpm} \times Tk_{nm} \times T_{t/h}^{-1})$, 234 where S_{rpm} is screw speed, T_{knm} is Torque and $T_{t/h}^{-1}$ is throughput. Specific growth rate (SGR) (% body 235 weight day⁻¹) was calculated as [(In (Final body weight (FBW) (g fish⁻¹)) – In (Initial body weight (g fish⁻¹))/ Experimental period (days)] × 100%. Feed intake (g DM fish⁻¹) was calculated as Total feed intake (g

237 DM tank⁻¹)/Number of fish per tank. FCR (g g⁻¹) was calculated as Feed intake (g DM fish⁻¹)/ (FBW (g fish⁻¹)) 238 ¹) - Initial body weight (g fish⁻¹)). ADC of nutrients (%) was calculated as (1- [(Y concentration in diet/Y 239 concentration in feces) × (Nutrient concentration in feces/Nutrient concentration in diet)]) ×100. Fecal 240 excretion of minerals and N (%) was calculated as (100 – ADC of minerals or N). The dissolved N fraction 241 was calculated as (([Feed intake (g fish⁻¹) × N content in feed (%)/100] × ADC of N/100) – ([(FBW (g) × 242 Final N content in fish $(g g^{-1})) - ($ Initial body weight $(g) \times$ Initial N content in fish $(g g^{-1})))/($ [FBW $(g fish^{-1}) -$ Initial body weight (g fish⁻¹)]/1000). Protein and lipid efficiency ratios (g g^{-1}) were calculated as (FBW (g 243 244 $fish^{-1}$) - Initial body weight (g fish⁻¹))/ [Feed intake (g fish⁻¹) × Protein or lipid content in feed (%)/100]. 245 Apparent nutrient retention (% intake) was calculated as $[(FBW (g) \times Final nutrient content in fish (g g^{-1}))$ 246 - (Initial body weight (g)× Initial nutrient content in fish (g g^{-1}))]/[Feed intake (g fish⁻¹) × Nutrient content 247 in feed $(\%)/100] \times 100$.

248 2.7 Statistical analysis

The data were analyzed using one-way ANOVA, followed by a Tukey's test for comparison of means. Differences at p<0.05 were considered as significant. Linear and quadratic polynomial contrasts were used to evaluate the relationship between different parameters and dietary BSFL meal or BSFL paste levels as indicated in tables. The chosen level of significance was p<0.05 and threshold level of tendency was p<0.1. All the statistical analyses were performed using IBM SPSS Statistics 26 software. **3. Results**

255 *3.1 Feed production and pellet quality*

The extruder parameters used during the production of experimental diets are shown in Table 3. The full-fat BSFL meal increased the lipid content and the BSFL paste increased the moisture content of the feed mash prior to extrusion. To compensate this, the screw speed of the extruder was increased to obtain pellet with desirable expansion and physical quality. Despite the increased screw speed,

decreased die pressure and torque were observed, which resulted in a decreased SME. In addition to
 increased screw speed, the water added to the BSFL paste diets in the extruder was reduced.

Inclusion of BSFL meal and paste in the diets numerically increased the amount of fines before
coating. Pellet durability and hardness were overall high for all diets, but with a numerical reduction in
durability for the diet with 12.5% BSFL meal replacement (Table 4). The expansion, sinking velocity and
water stability of pellets after 30, 60 and 120 minutes were numerically lower with increased inclusion
of both BSFL meal and paste (Table 4 and Figure 2). Starch gelatinization varied among the experimental
diets, ranging from 54.4 to 95.2 %, but with high analytical variation within the treatments.

268 *3.2 Fish performance*

269 Only one fish died throughout the experimental period. Fish fed the BSFL meal diets had similar 270 feed intake compared to Control-1. According to the linear polynomial contrasts, FBW and SGR reduced 271 (p<0.05) with increasing dietary BSFL meal level. According to ANOVA, there were no differences in FBW 272 and SGR between the fish fed 6.25IM and 12.5IM diets and the fish fed Control-1 diet, while FBW and 273 SGR were lower (p<0.05) in fish fed 25IM diet than fish fed Control-1 diet. FCR of fish fed the BSFL meal 274 diets were not different from the fish fed Control-1, though it was lower (p<0.05) in 12.5IM than 25IM 275 diet. In addition, a linear relationship (p<0.05) was observed between FCR and BSFL meal level in the 276 diet (Table 5).

In contrast, 3.7IP and 6.7IP diets did not affect the growth performances of fish compared to
Control-1 or Control-2. A positive linear tendency (*p*=0.08) was, however, observed between dietary
BSFL paste level and FCR of fish. Growth performances did not differ between the two controls (Table 5). *3.3 Digestibility of nutrients, fecal excretion of minerals and dissolved fraction of N*

Although ADC of protein and energy in BSFL meal diets did not differ from Control-1, ADC of protein decreased linearly (*p*<0.05) with increasing dietary BSFL meal level. ADC of lipid was lower (*p*<0.01) for the 12.5IM and 25IM diets compared to the Control-1, whereas ADC of starch was highest

(*p*<0.01) for the 25IM diet. Furthermore, a negative liner relationship (*p*<0.001) was found between
 dietary BSFL meal level and ADC of lipid, and a positive linear relationship (*p*<0.01) was found in ADC of
 starch (Table 6).

ADC of AA, except tyrosine in the BSFL meal diets did not differ from the Control-1. Tyrosine digestibility was lower (p < 0.01) in 12.5IM and 25IM diets when compared with the Control-1. It was also observed linear reductions (p < 0.05) of the ADC of phenylalanine, histidine, lysine and tyrosine with increasing level of BSFL meal in the diet. The same trend was observed for ADC of arginine (p=0.05) and cysteine (p=0.07). However, the ADC of total AA was unaltered by the dietary inclusion of BSFL meal (Table7).

The ADCs of nutrients in salmon fed the BSFL paste diets did not differ from fish fed the Control-1 or the Control-2 diets. However, negative linear effects (*p*<0.05) were observed between dietary BSFL paste level and ADCs of protein and energy (Table 6).

Fecal excretion of P and Mg were not affected by dietary inclusion of BSFL meal. Fecal Ca and K
 excretions were lower (*p*<0.05) in 25IM diet and 12.5IM diet respectively, compared with Control-1.

298 Further, linear relationships (*p*<0.05) were observed between fecal Ca and K excretions and dietary BSFL

299 meal level (Table 6).

300 Fecal P excretion increased linearly (*p*<0.001) with increasing level of BSFL paste in the diet and

BSFL paste diets showed higher (*p*<0.01) P excretion than Control-2. Linear (*p*<0.05) and quadratic

302 (*p*<0.05) relationships were observed between fecal Ca and Mg excretions and dietary BSFL paste level.

303 6.7IP diet had higher (*p*<0.01) Ca excretion compared to Control-2 (Table 6).

304 The dietary inclusion of BSFL meal or paste did not affect the dissolved N discharges, whereas

fecal N excretion increased linearly (*p*<0.05) when increasing the level of BSFL meal and paste in the diet

306 (Table 6).

307 3.4 Nutrient utilization

The protein efficiency ratio (PER) linearly decreased (p<0.05) with increasing BSFL meal level and 25IM diet showed a lower (p<0.05) PER than Control-1, but dietary inclusion of BSFL meal did not affect the apparent protein retention in fish. Lipid efficiency ratio (LER) was lower (p<0.01) in all BSFL meal diets compared to Control-1. Further, linear (p<0.001) and quadratic (p<0.001) relationships were observed between the LER and dietary BSFL meal level. Apparent lipid retention and apparent energy retention decreased linearly (p<0.01) with increasing dietary BSFL meal level, where 25IM diet showed the lowest (p<0.05) retentions (Table 8).

PER decreased linearly (p<0.05) with increasing level of BSFL paste in the diet, and PER was lower (p<0.05) in the 6.7IP diet than in the Control-2 diet. However, apparent protein retention was not affected by dietary inclusion of BSFL paste. The 3.7IP diet showed higher LER (p<0.05) compared to the two controls and higher apparent lipid retention (p<0.01) compared to Control-2. Linear (p<0.05) and quadratic (p<0.01) relationships were also observed between LER and apparent lipid retention and the level of BSFL paste in the diet. Further, apparent P retention decreased linearly (p<0.05) when increasing dietary BSFL paste level (Table 8).

322 4. Discussion

323 *4.1 Feed production and physical pellet quality*

324 Adding the BSFL meal and paste to the diets led to an increased lipid and moisture content in 325 the mash prior to extrusion, respectively. Lipids act as lubricants, therefore a high lipid level in extrusion 326 increases the lubrication and reduces the friction in the extruder (De Pilli et al., 2015; llo et al., 2000; Lin 327 et al., 1997), resulting in a decreased dough temperature (Hansen et al., 2011; Lin et al., 1997). Similarly, 328 increased water content in the extruder can also act as a lubricant and decrease friction, leading to 329 reduced dough temperature (Huang et al., 1995; Lin et al., 1997). Lower dough temperature can reduce 330 starch gelatinization (Garber et al., 1997; Lin et al., 1997; Morken et al., 2012), which results in reduced 331 expansion (Garber et al., 1997) and physical pellet quality (Morken et al., 2012). Hence, to reduce the

adverse effect of high lipid and moisture contents in the mash during extrusion, the screw speed wasincreased with increasing BSFL meal and paste levels in the diet.

334 In general, a higher screw speed creates higher SME and leads to a higher dough temperature 335 during extrusion (Morken et al., 2012; Rolfe et al., 2001). A decreased dough temperature with 336 increased screw speed can also occur due to a decreased filling rate of the extruder (Huang et al., 1995) 337 or decreased residence time of the dough in the extruder, resulting in a less efficient heat transfer 338 between the extruder barrel and the dough (Della Valle et al., 1987; Huang et al., 1995; Lin et al., 1997). 339 A similar effect might have occurred in the present study, as indicated by the reduced SME, die pressure 340 and torque, because the resistance to screw rotation was proportional to the filling rate (Akdogan, 341 1996). In addition, the low barrel and die temperature, torque and SME in the high BSFL diets indicates 342 that increased screw speed was not optimal in compensating for the increased lipid level in the present 343 study. Further, modification of the screw configuration during extrusion of the high BSFL containing 344 diets could have given better results, as this has large effect on the extrusion parameters (Gogoi et al., 345 1996).

Starch gelatinization was reported to be reduced at high lipid (Hansen et al., 2010; Hansen et al., 2011; Lin et al., 1997) and increased screw speed (Lin et al., 1997) due to low dough temperature and hydrophobic properties of high lipid in the extruder. However, no clear relationship was found between starch gelatinization and increased inclusion of dietary BSFL meal in the present study. The decreased starch gelatinization observed with increased inclusion of BSFL paste was probably due to reduced dough temperature from the higher moisture content in feed as shown by Lin et al. (1997) and this is associated with reduction of WHC (Artz et al., 1990).

The reduced pellet expansion with increased dietary BSFL meal in the present study is in line with previous reports indicating that a decreased extrudate expansion was due to high lipid content in mash (Hansen et al., 2011; Ilo et al., 2000; Navale et al., 2015). The decreased pellet expansion in high

BSFL meal diets might also be related to lower barrel temperature, in particular the fifth barrel temperature (Bandyopadhyay and Rout, 2001; Kothakota et al., 2013; Pathania et al., 2013). The decreased expansion of BSFL paste diets might be associated with the higher screw speed and moisture content of the feed mash (Bandyopadhyay and Rout, 2001). This increased level of lipid content in the feed mash followed with reduced level of SME and pellet expansion is also, most probably, explaining the reduced pellet water stability in the present study as shown by Hansen et al. (2011).

Although Sørensen et al. (2009) showed that even a small increase in lipid content might adversely affect the physical quality of extruded diets, in the present study, pellet durability and hardness were not notably reduced with the inclusion of BSFL meal or paste in the diet. Similarly, BSFL and cricket meal did not affect the pellet durability in the extruded fish feed (Irungu et al., 2018). The presence of formic acid might also contribute to this in BSFL paste diets. As reported by the others, dietary supplementation of sodium diformate (Morken et al., 2011) and potassium diformate (Morken et al., 2012) increased durability/hardness of salmonid feed pellets.

369 4.2 Fish performance and nutrient digestibility and utilization

Dietary inclusion of both BSFL meal and paste did not affect the palatability of Atlantic salmon diets, as indicated by the similar feed intake among diets. Similar results were shown in Atlantic salmon pre-smolts (Belghit et al., 2018) and post-smolts (Belghit et al., 2019b) and rainbow trout (Dumas et al., 2018; Renna et al., 2017) fed BSFL meal diets.

As observed for both BSFL meal and paste in the present study, low dietary inclusion of BSFL meal, did not affect growth performance in Atlantic salmon pre-smolts (10-20%) (Fisher et al., 2020) or post-smolts (5-15%) (Belghit et al., 2019b) and rainbow trout (20%) (Józefiak et al., 2019b) compared to control diets based on fishmeal and other protein sources. Xu et al. (2020) also observed that dietary fishmeal replaced with BSFL pulp (4.4-17.5%), which contained crushed fresh larvae, did not influence the SGR or FCR of juvenile mirror carp. Yet, the highest BSFL meal inclusion level which replaced 25% of

dietary protein showed lower FBW and SGR in the present study. Similarly, FBW and/or SGR were also reported to be reduced in other studies where high levels of BSFL meal were included in the diets of Atlantic salmon pre-smolts (30%) (Fisher et al., 2020) and post-smolts (25%) (Lock et al., 2016) and rainbow trout (26.4%) (Dumas et al., 2018). In contrast, dietary inclusion of high levels of BSFL meal caused no adverse effect on growth performances in Atlantic salmon (60%) (Belghit et al., 2018) and rainbow trout (20-40%) (Renna et al., 2017).

386 The reduction of SGR in the present and previous studies may be attributed to the presence of 387 chitin in the BSFL. Chitin is a major component of insect cuticle (Chapman, 1998; Tharanathan 388 and Kittur, 2003). In the present study, whole BSFL meal including the cuticle was used. The chitin 389 content in the BSFL meal (i.e. 8% in DM basis) corresponded to a chitin level of 0.6,1.2 and 2.3% for the 390 meal diets and 0.4 and 0.6% for the paste diets. Previous studies reported reduced SGR in juvenile 391 turbot fed BSFL meal containing chitin (1.6-7.3%) (Kroeckel et al., 2012), Atlantic salmon fed chitin from 392 prawn shells (1-5%) (Karlsen et al., 2017) or chitin containing krill meal (2%) (Hansen et al., 2010) and 393 reduced weight gain in tilapia fed chitin (2-10%) (Shiau and Yu, 1999). Furthermore, feeding high levels 394 of chitin reduced BSFL meal had no adverse effect on growth performance of Atlantic salmon (Belghit et 395 al., 2018).

Chitin was reported to contain around 17.1 kJ/g of energy content, which could constitute a 396 397 substantial percentage of total energy intake (Gutowska et al., 2004), but, Atlantic salmon have been 398 reported to have a poor capacity to digest chitin (13–40%) (Olsen et al., 2006). This indicates chitin 399 function as a filler with low digestible energy content (Karlsen et al., 2017) that might limit growth rate 400 at high inclusion levels. In addition, the reduction in growth rate could also be a result of the reduced 401 ADC of nutrients. In accordance with the decreased ADC of protein with increasing dietary levels of BSFL 402 meal and paste in the present study, dietary inclusion of high levels of BSFL meal adversely affected ADC 403 of protein/AA in salmon pre-smolt (60%) (Belghit et al., 2018) and rainbow trout (40%) (Renna et al.,

404 2017). On the contrary, some research showed that ADCs of protein and most of the AA were not 405 affected by dietary BSFL meal inclusion in Atlantic salmon post-smolts (5-25%) (Belghit et al., 2019b; 406 Lock et al., 2016) and rainbow trout (20%) (Dumas et al., 2018). In agreement with present results for 407 BSFL meal diets, Belghit et al. (2018) and Belghit et al. (2019a) reported reduced ADCs of lipid and most 408 fatty acids with the inclusion of BSFL meal and oil in diets for Atlantic salmon. The lower ADC of 409 nutrients might also be attributed to chitin, because previous studies showed that feeding diets 410 containing chitin reduced ADC of nutrients in Atlantic salmon (Hansen et al., 2010; Karlsen et al., 2017) 411 and tilapia (Shiau and Yu, 1999). The chitin in insect cuticle exists in a matrix with proteins, lipids and 412 other compounds (Chapman, 1998; Kramer et al., 1995), which may reduce the access of digestive 413 enzymes, thus reducing ADCs of nutrients (Henry et al., 2015). In addition, chitin might further reduce 414 ADC of protein due to its capacity to bind proteins (Piccolo et al., 2017) and immobilize (Muzzarelli, 415 1980) or reduce the activity of proteolytic enzymes such as the brush border enzyme, leucine 416 aminopeptidase that break down peptides into AA (Belghit et al., 2018). It has also been suggested that 417 feeding chitin leads to decreased bile acid levels in the pylorus, and thereby reduce ADC of lipid as bile 418 acid is essential for activation of lipase and efficient lipid absorption (Hansen et al., 2010). In addition, 419 the FA composition of BSFL meal is presented in the supplementary table (Table A.1) showing that the 420 majority of the FA in BSFL meal were saturated fatty acids (SFA) (65% of total FA), which might increase 421 the SFA content in BSFL diets. High SFA dietary concentrations may also partially explain the decrease in 422 ADC of lipid in the present study, as the ADC of lipid decreases linearly with an increasing concentration 423 of dietary SFA. This has previously been demonstrated in salmonids (Hua and Bureau, 2009). 424 Based on the present results, acid detergent fiber fraction in BSFL meal contained 12% of AA,

424 Based on the present results, acid detergent fiber fraction in BSFL meal contained 12% of AA, 425 which was bound to chitin, and probably not available for digestion. The observed reduction of ADCs of 426 several AA in the present study might be because these AA were trapped in chitin that is concealed for 427 enzymatic digestion. Furthermore, in the analysis of AA, it appeared that the peak for tyrosine in the

HPLC chromatogram was overlapped with glucosamine, which is the building block of chitin (Ng et al.,
2001). Therefore, the reduced ADC of tyrosine was most likely linked to an overestimation of the
tyrosine content in the feces. However, the dietary inclusion of BSFL meal did not affect the ADC of total
AA although the ADC of protein was reduced. Thus, the observed reduction of ADC of protein might
partially be explained by the poorly digestible chitin.

433 Despite the decreased ADC of protein, apparent protein retention was not compromised by the 434 inclusion of BSFL meal and paste, probably indicating an increased utilization of digested proteins in the 435 fish fed the BSFL diets. The unaltered ADC of total AA and similar dissolved N level among the diets 436 might partially explain this. In addition, this might also partially be due to the content of nucleic acid in BSFL, which may have an N-sparing effect in salmon. As shown by other protein sources such as 437 438 bacterial meal, nucleic acids were suggested to have an N-sparing effect and increased N retention in 439 salmon, although the nutrient digestibility was slightly lower (Øverland et al., 2010). In line with the 440 present results, dietary replacement of fishmeal with BSFL meal did not affect protein retention in 441 Atlantic salmon post-smolts (Belghit et al., 2019b; Lock et al., 2016), gilthead seabream 442 (Karapanagiotidis et al., 2014) and yellow catfish (Xiao et al., 2018). However, dietary inclusion of BSFL 443 meal negatively affected PER and replacement of 25% of dietary protein with BSFL meal reduced PER in 444 the present study. In contrast, replacement of fishmeal and/or plant protein with BSFL meal did not 445 affect the PER in Atlantic salmon pre-smolts (Belghit et al., 2018) and rainbow trout (Józefiak et al., 446 2019b; Renna et al., 2017). Further, Fisher et al. (2020) reported even an increased PER at 30% inclusion 447 level in Atlantic salmon pre-smolts.

The apparent lipid retention values above 100% observed in several treatments of the present study indicated lipid synthesis outweighed lipid catabolism. Both LER and apparent lipid retention were negatively affected by the dietary inclusion of BSFL meal and the effect was worse with increasing level of BSFL meal in the diet. In agreement with this, lipid retention decreased at dietary inclusion of 33%

452 BSFL meal and higher in juvenile turbot (Kroeckel et al., 2012). In addition, two studies showed that 453 dietary BSFL meal negatively affected the whole-body lipid composition in rainbow trout (St-Hilaire et 454 al., 2007) and juvenile turbot (Kroeckel et al., 2012). The observed lower lipid utilization in salmon fed 455 diets containing BSFL meal was accompanied by low ADC of lipid in these diets and can be attributed to 456 the presence of chitin as discussed above. In addition, the most abundant SFA in BSFL was medium-457 chain lauric acid (40% of the total FA). Lauric acid is considered to be a good source of energy for 458 salmonids as it seems to be oxidized to a larger extent and used less for lipid deposition, resulting in low 459 tissue deposition (Belghit et al., 2019a; Renna et al., 2017) and subsequently reduce lipid retention and 460 LER. The increased energy production by lauric acid might also explain the observed comparable protein 461 retention of BSFL diets despite reduced ADC of protein due to a protein-sparing effect (Karalazos et al., 462 2011). Teo et al. (1989) also reported the potential protein-sparing effect of medium-chain triglycerides. 463 In agreement with this, previous studies have also shown that dietary inclusion of medium-chain 464 triglycerides improved N/protein retention in Atlantic salmon (Nordrum et al., 2000; Nordrum et al., 465 2003). In addition, protein synthesis is a highly energy requiring process (Nordrum et al., 2000) and the 466 high energy contribution by lauric acid might, therefore, have a positive effect on protein retention. 467 Nevertheless, the chitin and BSFL oil content in BSFL paste diets did not seem to be sufficient to cause a 468 negative impact on ADC of lipid. However, in contrast to BSFL meal, it was observed that 3.7% 469 replacement of dietary protein with BSFL paste improved both LER and apparent lipid retention. This 470 might be due to improved utilization of digested nutrients when the BSFL were subjected to low 471 temperature processing and preserved with formic acid or included in the diet at lower levels. 472 According to the results of present and previous studies (Finke, 2013; Fisher et al., 2020), BSFL 473 meal is more abundant in micronutrients (P, Ca, Mg, K) and BSFL have a mineralized cuticle in which Ca 474 is incorporated into the cuticle (Finke, 2013). In general P content in BSFL meal is lower than fishmeal 475 (Liland et al., 2017), which was reflected by a slight reduction of P level in BSFL meal diets. But the P in

476 insects is likely to be readily available, unlike plant-based phytate P (Finke, 2002). This might be the 477 reason for unaltered fecal P excretion and P retention of BSFL meal diets in the present study. Similarly, 478 whole fish P content was not altered by BSFL meal diets in Atlantic salmon pre-smolts (Belghit et al., 479 2018). The observed fecal Ca excretion values closer to or above 100% in the present study is most likely 480 due to Ca uptake by fish from water. The decreased fecal Ca excretion when increasing BSFL meal level 481 in the diet indicated that higher dietary inclusion of BSFL meal improved ADC of Ca. It has been reported 482 that the supplementation of diets with formic acid affects the intestinal pH of rainbow trout and 483 improves the ADC of P, Ca and Mg (Vielma and Lall, 1997). However, an opposite result was observed 484 for BSFL paste containing formic acid, where the increased dietary level of BSFL paste increased fecal 485 excretion of P indicated decreased ADC of P and accompanied by decreased P retention. Similarly, 486 increased fecal Ca excretion was observed with increasing inclusion level of BSFL paste in the diet, 487 indicating decreased ADC of Ca. The increased fecal N excretion of BSFL meal and paste diets and 488 increased fecal P excretion of BSFL paste diets indicate an increased environment impact of low 489 processed insect products as alternative protein sources, although the fecal P excretion of BSFL meal 490 diets and dissolved N fraction of the BSFL meal and paste diets were similar. Future work on further 491 processing such as defatting and dechitinization can help alleviate potential adverse environmental 492 effects of such insect ingredients.

493 **5.** Conclusions

The present study showed that BSFL meal and paste could replace up to 12.5% and 6.7% of dietary fishmeal and plant proteins, respectively, without compromising the growth performance or protein retention in Atlantic salmon. Nevertheless, protein and lipid digestibility, protein efficiency ratio and lipid retention decreased linearly with increasing dietary BSFL meal level, whereas increasing dietary BSFL paste level linearly decreased protein digestibility, protein efficiency ratio and phosphorous

- 499 retention. At higher replacement level of 25% BSFL meal, the growth rate was reduced, accompanied by
- 500 a reduction in digestibility and utilization of lipids and protein efficiency ratio.

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- 755 https://doi.org/10.1111/anu.13005.
- 756 Table captions
- 757 **Table 1:** Analyzed chemical composition (%, dry matter) of black soldier fly larvae (BSFL) meal and paste
- 758 **Table 2:** Ingredient and analyzed chemical composition of experimental diets with increased inclusion
- 759 level of black soldier fly larvae (BSFL) meal and paste¹
- 760 **Table 3:** Extruder parameters during the production of experimental diets with increased inclusion level
- 761 of black soldier fly larvae (BSFL) meal and paste¹
- 762 **Table 4:** Physical pellet quality of experimental diets with increased inclusion level of black soldier fly
- 763 larvae (BSFL) meal and paste¹
- **Table 5:** Performance of fish fed experimental diets with increased inclusion level of black soldier fly
- 765 larvae (BSFL) meal and paste¹

- 766 **Table 6:** Apparent digestibility coefficient (ADC) of nutrients (%), fecal excretion of minerals and nitrogen
- 767 (%) and dissolved fraction of nitrogen (g kg-1 fish body weight gain) of fish fed experimental diets with
- 768 increased inclusion level of black soldier fly larvae (BSFL) meal and paste¹
- 769 **Table 7:** Apparent digestibility coefficient (ADC) of amino acids (%) of fish fed experimental diets with
- increased inclusion level of black soldier fly larvae (BSFL) meal¹
- 771 **Table 8:** Nutrient retention parameters in fish fed experimental diets with increased inclusion level of
- 772 black soldier fly larvae (BSFL) meal and paste¹
- 773 **Supplementary Table A.1:** Analyzed fatty acid composition (%, dry matter) of black soldier fly larvae meal
- 774
- 775

776 Figure captions

- **Figure 1:** The screw configuration used during the extrusion of experimental diets. 20; 40; 60; 80; 100;
- 120: Length in cm of each screw element. R: Right. L: Left (Flow direction of each screw element). P:
- 779 Polygon. UC: Undercut conveying screw element (larger channel depth than the other conveying screw
- relements). The arrows indicate 5 mm spacer ring and 90° offset between the screw elements.
- 781 Figure 2: Water stability (dry matter retention %) of pellets with increased inclusion of black soldier fly
- 782 larvae (BSFL) meal and paste after 30, 60 and 120 minutes. Error bars indicate standard deviation.
- 783 Control-1: Control diet. 6.25IM, 12.5IM and 25IM: BSFL meal substituted 6.25%, 12.5% and 25% of
- protein content of Control-1. Control-2: Control diet with 0.88% of formic acid. 3.7IP and 6.7IP: BSFL
- paste substituted 3.7% and 6.7% of protein content of Control-1.