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9	Prevention not cure: a review of methods to avoid sea lice infestations in salmon
10	aquaculture
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28 ABSTRACT

The Atlantic salmon aquaculture industry still struggles with ectoparasitic sea lice despite 29 30 decades of research and development invested into louse removal methods. In contrast, methods to prevent infestations before they occur have received relatively little research 31 effort, yet may offer key benefits over treatment-focused methods. Here, we summarise the 32 range of potential and existing preventative methods, conduct a meta-analysis of studies 33 trialling the efficacy of existing preventative methods, and discuss the rationale for a shift to 34 the prevention-focused louse management paradigm. Barrier technologies that minimise host-35 36 parasite encounter rates provide the greatest protection against lice, with a weighted median 76% reduction in infestation density in cages with plankton mesh 'snorkels' or 'skirts', and 37 up to a 100% reduction for fully enclosed cages. Other methods such as geographic 38 spatiotemporal management, manipulation of swimming depth, functional feeds, repellents, 39 40 and host cue masking can drive smaller reductions that may be additive when used in combination with barrier technologies. Finally, ongoing development of louse-resistant 41 salmon lineages may lead to long term improvements if genetic gain is maintained, while the 42 development of an effective vaccine remains a key target. Preventative methods emphasise 43 host resistance traits while simultaneously reducing host-parasite encounters. Effective 44 implementation has the potential to dramatically reduce the need for delousing and thus 45 improve fish welfare, productivity and sustainability in louse-prone salmon farming regions. 46

47

48 INTRODUCTION

The global expansion of sea cage fish farming has driven considerable shifts in the population 49 dynamics of marine pathogens. For 40 years, ectoparasitic lice have been an intractable 50 problem for Atlantic salmon (Salmo salar) farming industries in Europe and the Americas 51 (Torrissen et al. 2013; Iversen et al. 2015). Louse infestations are almost ubiquitous on 52 salmon farms in these regions – primarily the salmon louse *Lepeophtheirus salmonis* but also 53 Caligus elongatus in the northern hemisphere, and Caligus rogercressevi in South America 54 (Hemmingsen et al. 2020). Lice are natural parasites of fish, but intensive salmon farming 55 amplifies louse densities, resulting in unnaturally high infestation pressure for both farmed 56 and wild salmonids. Lice feed on the skin, blood and mucus of host fish, and severe 57

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infestations can cause ulceration leading to stress, osmotic imbalance, anaemia and bacterial
infection (Grimnes and Jakobsen 1996; Øverli *et al.* 2014; González *et al.* 2016).
Accordingly, management of louse infestations on farmed fish is crucial to maintain
acceptable stock welfare, limit production losses and reduce impacts on adjacent wild
salmonid populations (Krkošek *et al.* 2013; Thorstad *et al.* 2015).

In most jurisdictions, the primary management approach is to monitor louse densities on 63 farmed fish, with mandatory delousing or other sanctions implemented when louse levels 64 exceed allowable limits. Regulations also cap the number of active sites or total biomass in 65 each management zone according to estimated infestation pressure on wild salmonids, and 66 may mandate coordinated fallowing or other measures (e.g. Norway: Ministry of Trade and 67 Fisheries, 2012). The introduction of chemotherapeutants in the 1970s allowed farms to treat 68 sea louse infestations without substantially reducing production (Aaen et al. 2015). However, 69 70 most chemotherapeutants are not environmentally benign, leading to concerns about 71 bioaccumulation and effects on non-target invertebrate species (Burridge et al. 2010). More 72 recently, treatment-resistant lice have emerged on farms in Europe and the Americas (Aaen et al. 2015) rendering many chemotherapeutants less effective. 73

The discovery of treatment-resistance has prompted a rapid and recent shift to mechanical 74 and thermal delousing methods in the Norwegian salmon farming industry (Overton et al. 75 76 2018), with these methods also gaining traction elsewhere (e.g. Canada, Chile, Scotland). Mechanical and thermal delousing are highly effective at removing mobile lice and have little 77 or no impact on non-target species. However, they are stressful for host fish and can lead to 78 elevated post-treatment mortality rates compared to the use of chemotherapeutants (Overton 79 et al. 2018). Low salinity or hydrogen peroxide baths are also effective in the right conditions 80 and do not accumulate, although the long-term prospects for these methods are uncertain 81 given the possibility of increasingly resistant lice (Treasurer et al. 2000, Helgesen et al. 2018, 82 Groner et al. 2019). Alternatively, around 50 million cleaner fish (lumpfish Cyclopterus 83 *lumpus* and several wrasse species) are deployed annually at Norwegian salmon farms to eat 84 lice directly off salmon (Norwegian Directorate of Fisheries 2018), with >1.5 million cleaner 85 86 fish also used in Scotland (Marine Scotland Directorate, 2017). However, it is unclear whether their efficacy (Overton et al. 2020; Barrett et al. 2020a) is sufficient to justify their 87 poor welfare in commercial sea cages (Nilsen et al. 2014; Hvas et al. 2018; Mo and Poppe 88 2018; Yuen et al. 2019; Stien et al. 2020). 89

Decades of innovation in louse control have allowed the salmon farming industry to continue 90 functioning in louse-prone regions, but not without significant environmental and ethical 91 concerns. Most research and development efforts so far have focused on treating at the post-92 infestation stage. This likely reflects the relatively rapid return on investment into new 93 delousing methods but may be a sub-optimal strategy if opportunities to invest in long term 94 solutions are missed (Brakstad et al. 2019). An alternative approach is to focus louse 95 management efforts on preventing infestation via proactive interventions ('preventative 96 methods' herein) that may significantly reduce the need for farms to delouse. Here, we 97 98 summarise the range of potential or existing preventative methods and conduct a metaanalysis of empirical estimates of sea louse removal efficacy for each method. Finally, we 99 discuss the rationale for a paradigm shift from reactive louse control to a proactive approach 100 that focuses on predicting and preventing infestations, and outline some possible strategies to 101 promote long term efficacy of preventative methods. 102

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104 WHAT PREVENTATIVE METHODS ARE AVAILABLE?

Preventative methods are deployed pre-emptively to reduce the rate of new infestations. 105 Within this classification, we include approaches that either: (1) reduce encounter rates 106 between salmon and infective copepodid stage lice; or (2) reduce the attachment success 107 and/or early post-settlement survival of copepodids via interventions that begin to act at the 108 moment of attachment or first feeding (Fig. 1). These approaches are distinct from control via 109 110 delousing treatments, which are generally implemented as a reaction to an existing infestation (i.e. 'immediate' control), or via cleaner fish, which may be deployed prior to infestation and 111 function on an ongoing basis (i.e. 'continuous' control) but are not typically effective against 112 newly attached lice (e.g. Imsland et al. 2015). 113

- 114 1. Reducing encounters
- 115 *1.1 Barrier technologies*

A growing understanding of louse physiology and host-finding behaviour has led to several important advances in louse prevention, and by using data on preferred swimming depths of infective copepodids in relation to environmental parameters (Heuch 1995; Heuch *et al.* 119 1995; Crosbie *et al.* 2019), farmers can now separate hosts from parasites using depthspecific louse barriers.

Barriers made from fluid-permeable plankton mesh or impermeable membranes can 121 dramatically reduce infestation rates by preventing infective copepodids from entering the 122 cage environment. 'Skirt' or 'snorkel' barriers prevent particles in the surface layers—where 123 most copepodids reside-from entering the cage while still allowing full water exchange 124 below the level of the barrier (Oppedal et al. 2017; Wright et al. 2017; Stien et al. 2018). 125 Salmon often choose to reside below the level of the skirt or snorkel, meaning that the barrier 126 functions by simultaneously (i) encouraging salmon to swim below the depth at which 127 infestation risk is highest, and (ii) protecting any individuals that use the surface layers, for 128 129 example, while feeding or refilling the swim bladder. In the most complete use of barrier technologies, fully-enclosed cages are supplied with louse-free water either filtered or 130 pumped from depths below the typical depth range of copepodids (e.g. 25 m: Nilsen et al. 131 2017). 132

Barrier technologies (particularly skirts) are already widely used by the industry, but specific 133 134 designs should be matched to local environmental conditions to avoid problems with low dissolved oxygen or net deformation (Stien et al. 2012; Frank et al. 2015; Nilsen et al. 2017). 135 For example, Nilsen et al. (2017) prevented deformation of impermeable tarpaulin barriers at 136 relatively sheltered sites by creating slight positive pressure within the cage (i.e. inside water 137 level 2-3 cm above sea level). At more exposed sites, it is preferable to use fluid-permeable 138 plankton mesh barriers (e.g. Grøntvedt et al. 2018). Brackish surface water can also reduce 139 the efficacy of skirts and snorkels by causing both lice and salmon to reside below the level 140 of the barrier (Oppedal et al. 2019), while there is evidence that barrier technology may 141 reduce the performance of cleaner fish when used in combination (Gentry et al. 2020). 142

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1.2 Manipulation of swimming depth

144 Salmon behaviour, primarily swimming depth, can also be manipulated in the absence of barrier technology to reduce spatial overlap (and therefore encounter rates) between hosts and 145 146 parasites, especially salmon lice. Typically, the aim is to reduce encounter rates by causing salmon to swim below the depths at which lice are most abundant. Deep swimming 147 behaviour can be promoted through the use of deep feeding and/or lighting (Hevrøy et al. 148 2003; Frenzl et al. 2014; Bui et al. 2020). Where surface feeding is conducted, reducing the 149 150 frequency or regularity of feeding (e.g. twice daily at varying times) can reduce the amount of time spent in the surface layers (Lyndon and Toovey 2000). Deep swimming can also be 151 152 forced by submerging cages to the desired depth (Dempster et al. 2008; Dempster et al. 2009), and there is evidence for reduced louse levels on salmon in submerged cages (Osland 153

et al. 2001; Hevrøy *et al.* 2003; Sievers *et al.* 2018; Glaropoulos *et al.* 2019). Long term
submergence can affect fish welfare as salmon lose buoyancy over time (Korsøen *et al.* 2009;
Macaulay *et al.* 2020), however recent research indicates most welfare concerns can be
addressed by allowing periodic surface access or fitting a submerged air-filled dome for swim
bladder refilling (Korsøen *et al.* 2012; Glaropoulos *et al.* 2019; Oppedal *et al.* In Press).

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1.3 Geographic spatiotemporal management

A range of spatiotemporal management approaches are applied at the landscape scale to 160 reduce infestation risk by controlling where and when salmon are farmed. Some farm sites 161 have consistently low louse abundances and rarely require delousing (www.barentswatch.no). 162 163 Locating farms to take advantage of beneficial oceanographic conditions and minimise connectivity with adjacent sites may reduce the number of host-parasite encounters over a 164 grow-out cycle (Bron et al. 1993; Samsing et al. 2017; Samsing et al. 2019). Fallowing 165 during periods of high propagule pressure may also delay first infestation after sea transfer of 166 smolts (Bron et al. 1993). 167

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1.4 Filtering and trapping

Filters and traps may be deployed in or around cages to remove infective copepodids from 169 the water column before they encounter salmon. Filter-feeding shellfish racks hung around 170 sea cages may reduce louse abundance if deployed at sufficient scale (Byrne et al. 2018; 171 Montory *et al.* 2020), while powered filters are effective in the context of preventing lice and 172 173 eggs from entering the environment during delousing (O'Donohoe and Mcdermott 2014). In 174 other fish farming systems, cleaner shrimp have been used to remove parasites or parasite eggs from fish and nets and reduce infestation or reinfestation risk (Vaughan et al. 2018a; 175 176 Vaughan et al. 2018b). However, this method may have limited application against sea lice because of the planktonic mode of dispersal and infestation (i.e. larvae do not develop within 177 178 the cage structure). Light traps have been tested in the field with mixed results (Pahl et al. 179 1999; Novales Flamarique et al. 2009), and increasing knowledge of host-locating behaviour in lice may present new possibilities for baiting traps with attractive chemosensory cues 180 (Devine et al. 2000; Ingvarsdóttir et al. 2002; Bailey et al. 2006; Mordue and Birkett 2009; 181 182 Fields et al. 2018). No preventative filtering or trapping methods have been widely deployed in the industry, but some systems have recently become commercially available (e.g. 183 'Strømmen-rør', Fjord Miljø; 'NS Collector', Vard Aqua). 184

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1.5 Repellents and host cue masking

Interventions may be used to repel lice or mask host cues, potentially reducing host-parasite 186 encounters even when parasites enter the sea cage. Repellents or masking compounds can 187 either be released into the water column or included in feed to alter the host's semiochemical 188 profile (Hastie et al. 2013; O'Shea et al. 2017). Indeed, some existing commercially available 189 functional feeds are claimed to reduce attraction of lice toward fish (e.g. Shield, Skretting; 190 Robust, EWOS/Cargill). Visual cues may also be important, and the effect of modified light 191 conditions on infestation rates have been trialled with mixed results. Browman et al. (2004) 192 concluded that ultraviolet-A and polarisation were not important for host detection at small 193 194 spatial scales. Light intensity interacted with salinity and host velocity to influence distribution of louse attachment in another study (Genna et al. 2005), while Hamoutene et al. 195 (2016) reported that 24-hour darkness affected the attachment location but not abundance of 196 salmon lice. 197

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1.6 Incapacitation

Several methods have been proposed for disabling or killing lice—from egg to adult stages in or around sea cages. These include ultrasonic cavitation (Alevy 2017; Skjelvareid *et al.* 2018; Svendsen *et al.* 2018), direct current electricity (Bredahl 2014) and irradiation with short wavelength light (Barrett *et al.* 2020b, Barrett *et al.* 2020c). Some have demonstrated efficacy at close range (Skjelvareid *et al.* 2018, Barrett *et al.* 2020b, Barrett *et al.* 2020c), but it is currently unclear whether any such methods can be effective at commercial scale.

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1.7 Louse population control

Interventions to suppress louse populations outside the cage environment would require careful consideration before deployment and must be specific to targeted louse species. Very little work has been done in this area, but possible avenues may include the release of parasites and pathogens that are specific to sea lice (Økland *et al.* 2014; Økland *et al.* 2018; Øvergård *et al.* 2018), or CRISPR-based 'gene drives' (McFarlane *et al.* 2018; Noble *et al.* 2019).

- 212 2. Reducing post-encounter infestation success
- 213 2.1 Functional feeds

Feeds that provide physiological benefits beyond basic nutritional requirements are termed functional feeds and are increasingly prevalent in industrial fish farming (Tacchi *et al.* 2011). Feed ingredients that modify the mucus layer or modulate skin immune responses may reduce initial attachment success or facilitate effective immune responses against newly-

attached lice (Martin and Krol 2017). Functional feeds may also include ingredients that are 218 toxic or repellent to attached lice - these are not necessarily distinct from in-feed 219 chemotherapeutants, except that they tend to be derived from 'natural' sources (e.g. plant-220 derived essential oils: Jensen et al. 2015). Functional feeds aimed at improving salmon louse 221 resistance are already commercially available (e.g. Shield, Skretting; Robust, EWOS/Cargill). 222 It will be important to test for any adverse effects of new functional feeds. For instance, 223 glucosinolates and beta-glucans have been shown to be effective for reducing louse 224 infestation (Refstie et al. 2010; Holm et al. 2016), but glucosinolates also have a range of 225 226 effects on liver, muscle and kidney function that would need to be investigated (Skugor et al. 2016). Hormonal treatments may also be effective at reducing louse infestation (Krasnov et 227 al. 2015), but preventative hormone treatments are likely to be perceived negatively by 228 consumers. 229

230

2.2 Vaccines

Vaccines against bacteria and viruses are increasingly widespread in fish farming. In Norway, 231 antibiotics have been almost entirely replaced by injectable multi-component oil-based 232 vaccines (Brudeseth et al. 2013), and there is increasing use of injected or orally administered 233 vaccines in North America and Chile (Brudeseth et al. 2013). However, to our knowledge 234 there is currently only one (partially effective) vaccine available for sea lice (C. 235 236 rogercressevi: Providean Aquatec Sea Lice, Tecnovax). While there are no in-principle barriers, the development of vaccines for ectoparasites is technically challenging; despite the 237 identification of numerous vaccine targets in a range of ectoparasites, the cattle tick 238 (Rhipicephalus microplus) remains the only ectoparasite with a highly effective vaccine 239 (Stutzer et al. 2018). 240

241 Successful development of a recombinant or DNA vaccine would allow cost-effective production and delivery (Raynard et al. 2002; Sommerset et al. 2005; Brudeseth et al. 2013). 242 243 Potential vaccines exist at various stages of development, from localisation of candidate antigens in lice (Roper et al. 1995), demonstration of antibody production in response to 244 inoculation with louse extracts (Reilly and Mulcahy 1993), and use of recombinant proteins 245 to vaccinate salmon in tank trials (Carpio et al. 2011; Carpio et al. 2013; Basabe et al. 2014; 246 Contreras et al. 2020). Recently, RNA interference has been used to knock down candidate 247 vaccine targets and assess potential efficacy through challenge experiments (Eichner et al. 248 249 2014; Eichner et al. 2015; Komisarczuk et al. 2017).

250 *2.3 Breeding for louse resistance*

Variation in louse resistance is considerable among Atlantic salmon and has a heritable 251 component (Glover et al. 2005; Kolstad et al. 2005; Gjerde et al. 2011; Tsai et al. 2016; 252 Holborn et al. 2019), indicating that there is sufficient additive genetic variation for selective 253 breeding. Observed variation in louse resistance is probably due to differences in expression 254 of both host cues and immune responses (Holm et al. 2015). Decades of selective breeding 255 has resulted in much higher growth rates for farmed salmonid strains (Gjedrem et al. 2012) 256 and increased resistance to some diseases (Leeds et al. 2010; Ødegård et al. 2018; Storset et 257 al. 2007; reviewed by Robinson et al. 2017). More recently, the development of high-258 throughput single nucleotide polymorphism (SNP) genotyping technology has enabled 259 relatively rapid and affordable genomic selection and fine mapping of quantitative trait loci 260 associated with disease resistance. 261

Quantitative trait loci explaining between 6-13% of the genetic variation in sea louse 262 resistance (louse density on fish) have been detected in North American and Chilean 263 populations of Atlantic salmon (Rochus et al. 2018; Robledo et al. 2019). Salmon families 264 with greater resistance to sea lice show upregulation of several immune pathway and pattern 265 recognition genes compared to more susceptible families (Robledo et al. 2018), and the two 266 major breeding companies in Norway (AquaGen and SalmoBreed) offer salmon lines that 267 268 have been selected using marker assisted section or genomic selection for sea louse resistance. Use of genomic selection has been shown to increase the accuracy of selection for 269 sea louse resistance by up to 22% (Tsai et al. 2016; Correa et al. 2017), and two generations 270 of genomic selection focused on just sea louse resistance led to a 40-45% reduced sea louse 271 infestation compared to unselected fish (Ødegård et al. 2018). 272

Other possible approaches for improving sea louse resistance in Atlantic salmon include 273 hybridisation of Atlantic salmon with more louse-resistant salmonid species (Fleming et al. 274 275 2014), genetic modification of Atlantic salmon with immune genes from other salmonids, or use of gene editing to modify protein function or regulate the expression of genes affecting 276 resistance. In the case of hybridisation or any genetic modification, the effect on other 277 production traits would need to be assessed before hybrids or edited fish are used by the 278 279 industry. Gene editing approaches have high potential (Gratacap et al. 2019), but successful implementation depends on knowing which genes to modify to have the desired effect, on 280 281 developing effective methods for implementing and spreading the gene edits through the

breeding population, and on the acceptability of the use of the technology by the generalpublic and government.

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285 EFFICACY OF PREVENTATIVE METHODS

286 To assess the state of knowledge on the efficacy of preventative methods, we conducted a systematic review and meta-analysis of published studies pertaining to preventative methods. 287 To find relevant studies, we searched ISI Web of Science, Scopus and Google Scholar in 288 February 2020 using the following search string: (aquacult* OR farm*) AND (salmon* or 289 Salmo) AND (lice OR louse OR salmonis OR Caligus). We also discovered additional studies 290 referenced within articles returned by the search string. Together, our searches returned 291 292 >1200 peer-reviewed articles, technical reports and patents relevant to lice and salmon aquaculture, of which 141 provided evidence on the efficacy of preventative methods and 293 were included in the review. 294

Studies that provided relevant response variables were included in a meta-analysis, allowing 295 the comparison of effect sizes across the range of preventative approaches. For inclusion, 296 studies were required to provide empirical measures of relative louse infestation densities for 297 treatment groups (preventative methods used) and control groups (no preventative methods 298 used). Studies that applied treatments to lice but did not directly test for effects on infestation 299 were not included. Effect sizes were standardised using the natural log of the response ratio: 300 301 $lnRR = ln(\mu_T/\mu_C)$, where μ_T is the treatment group response and μ_C is the control group 302 response. In most cases, response variables were either mean or median attached lice per fish. Where a study tested multiple qualitatively different treatments, each treatment was 303 304 considered a replicate comparison in the meta-analysis. Where there were several qualitatively similar treatments (e.g. a range of doses of the same substance) the strongest 305 treatment was included in the meta-analysis. Epidemiological studies typically did not have 306 clear control or treatment groups; in such cases, the area or condition with the highest louse 307 density was designated as the control group for the purposes of calculating a response ratio; 308 this practice may inflate average effect sizes. 309

A total of 41 articles provided 98 comparisons that met the criteria for inclusion in the metaanalysis. For each preventative approach, we calculated a median effect size. When calculating a median effect, weighting studies according to their sample size can reduce bias. However, this was difficult in practice due to inconsistent definition of units of replication and therefore sample size across studies. Given this, we applied weightings to studies within each preventative approach (except vaccination, breeding and functional feed approaches, which are usually challenge tested in tanks) according to the scale or level of evidence of the experiment (in descending order of relative weights, level A: multiple farm experiment – 1.0; level B: experiment in full size sea cages at a single site – 0.8; level C: experiment in small sea cages at a single site – 0.6, level D: observational/epidemiology – 0.4; level E: experiment in tanks – 0.2).

To allow a visual assessment of potential publication bias, we produced a 'funnel plot' in which study effect sizes are fitted against the precision (1/SE) of the effect. This is based on sample size as defined by the study authors, or else the best available approximation. Precision is typically increased by sample size and/or experimental power, and typically, in a field without publication bias, the average direction and size of effect should not vary systematically with study precision (Hedges *et al.* 1999; Nakagawa *et al.* 2017).

327 Which preventative methods are most effective against sea lice?

328 Comparison of response ratios revealed high variability in effect sizes among trials of preventative methods (Fig. 2), but evidence from sea cage trials indicates that barrier 329 technologies can drive the largest and most consistent reductions in louse infestation levels 330 (weighted median 78% reduction, range 8% increase to 99% reduction, n = 13; Fig. 2). 331 Efficacy of specific barrier technologies appeared to be related to the extent of coverage: 332 skirts were moderately effective (median 55% reduction, range 30-81%, n = 2), snorkels were 333 highly effective (median 76% reduction, range 8% increase to 95% reduction, n = 9), and in 334 the sole closed containment study (Nilsen et al. 2017), infestations were almost entirely 335 avoided (98–99.7% reduction). 336

Approaches utilising manipulation of salmon swimming depth offered variable outcomes, but 337 with strong effects in certain situations (weighted median 26% reduction, range 72% increase 338 to 93% reduction, n = 11; Fig. 2). Geographic spatiotemporal management of farming effort 339 (or related variables such as simulated current speed: Samsing et al. 2015) had similarly 340 variable effects (weighted median 13% reduction, range 81% increase to 73% reduction, n =341 14; Fig. 2). Functional feeds tended to have small but beneficial effects on sea louse 342 infestations (median 24% reduction, range 108% increase to 67% reduction, n = 32: Fig. 2), 343 as do published vaccine trial results (median 4% reduction, range 20% increase to 57% 344 reduction). Notably, deployment of multiple preventative methods in combination with 345

cleaner fish had highly variable effects in three published studies using replicated modern commercial sea cages (weighted median 9% reduction, range 143% increase to 49% reduction, n = 5: Bui *et al.* 2019b; Bui *et al.* 2020; Gentry *et al.* 2020).

Several potential preventative approaches have seen little effort to test their effects on infestation rates. The use of repelling non-host cues was effective in one small-scale cage study (53-74% reduction, n = 3: Hastie *et al.* 2013), as was filtering of copepodids using oyster racks ((32% reduction: Byrne *et al.* 2018) or light traps (12% reduction: Pahl *et al.* 1999), and the incapacitation of lice using electric fences (78% reduction: Bredahl 2014) and ultrasonic cavitation (37% increase to 39% decrease: Skjelvareid *et al.* 2018).

355 Efficacy of selective breeding for louse resistance should be interpreted with a long-term view. Iterative improvements tend to be small-moderate but can lead to large genetic gain 356 over generations (Yanez et al. 2014; Gjedrem 2015), especially if genomic or marker assisted 357 selection for sea louse resistance is given a high weighting in the overall breeding index 358 (Ødegård et al. 2018). Estimates of heritability in louse resistance are moderate to high 359 depending on the method used (range 0.07-0.35: e.g. Gjerde et al. 2011; Glover et al. 2005; 360 361 Houston et al. 2014; Holborn et al. 2019), indicating that there is sufficient heritable variation available for genetic improvement. 362

363 Is the evidence base representative and robust?

Most preventative approaches have only been assessed a few times. Among the 41 articles 364 365 that met the criteria for inclusion in the meta-analysis, 7 provided data on efficacy of barrier 366 technologies, 6 on manipulation of swimming depth, 1 on breeding, 13 on functional feeds, 2 on incapacitation, 2 on repellents or cue-masking, 5 on geographic spatiotemporal 367 management, 2 on trapping and filtering, and 3 on candidate vaccines. Most articles (n = 38)368 were primarily concerned with salmon lice L. salmonis (i.e. those in Europe and North 369 370 America), while the remaining 3 articles targeted prevention of sea lice C. rogercressevi (i.e. those in Central or South America). All tested efficacy using Atlantic salmon. 371

Levels of evidence ranged widely: Barrier technologies had the most rigorous evidence base, with multiple studies with evidence levels from A-C (Fig. 2). Evidence levels should be considered when interpreting estimated efficacy, as preventative approaches may vary in their scalability to commercial sea cages (e.g. viability of methods to filter or trap copepodids are likely to be highly dependent on water volume). Units of replication also varied widely between studies, from individual fish to tanks, sea cages or farms. 51 out of 98 comparisons treated individual fish as replicates, in most cases resulting in a pseudoreplicated design as individuals were kept within a comparatively small number of tanks or cages (often <3 tanks or cages per group). We recommend that where fish are treated as replicates, the number of tanks or cages should also be reported, and mixed effects statistical methods employed to account for non-independence between fish held within the same tank or cage (Harrison *et al.* 2018).

- Finally, the meta-analysis revealed possible evidence for publication bias, with fewer studies 384 than expected present in the area of the plot corresponding to low precision and negative 385 findings (Fig. 3). In other words, the funnel plot indicates that among studies with small 386 sample sizes and/or highly variable data, those with positive results regarding efficacy of a 387 preventative method were more likely to be published. Not publishing negative findings can 388 (a) artificially inflate estimates of efficacy when averaging across studies, and (b) lead 389 390 researchers to waste resources testing methods that have already been found to be ineffective, perhaps multiple times. Accordingly, it is important that researchers and managers are aware 391 of the potential for publication bias when considering the evidence for novel louse 392 393 management strategies (whether preventative or otherwise). The prevalence of publication bias is likely to be influenced by the type of study and preventative method. For example, 394 tests of barrier technologies and swimming depth manipulation are generally conducted in sea 395 cages, and given the effort and cost involved, results are perhaps more likely to be published 396 in full. Other approaches may be inherently more susceptible to publication bias, for example 397 when a large range of substances or doses are tested in the early stages of a study and only 398 those that are reasonably successful are reported. 399
- 400

401 THE NEW PARADIGM: A FOCUS ON PREVENTATIVE METHODS AGAINST 402 SEA LICE

The evidence base demonstrates that effective implementation of preventative methods can
reduce infestation pressure within sea cages and therefore reduce the need for louse control.
A prevention-focused louse management paradigm may lead to several key benefits:

406 (1) Most preventative methods have small if any impacts on non-target organisms (like
407 mechanical and thermal delousing methods, but unlike some common chemotherapeutants:
408 Burridge *et al.* 2010; Taranger *et al.* 2015).

(2) Delousing treatments cause stress and injury to stock, leading to welfare concerns and 409 production losses from reduced growth, higher mortality and a lower quality product 410 (Overton et al. 2018). By focusing on avoiding encounters and reducing initial infestation 411 success, preventative methods may be targeted at infective louse stages without also 412 impacting host fish (Fig. 4). Conversely, some preventative methods can selectively target 413 host traits to improve innate resistance (Fig. 4), such as promoting parasite avoidance 414 behaviour via behavioural manipulation or immune function via functional feeds and 415 selective breeding. 416

(3) Multiple preventative methods can be deployed together and on a continuous basis, 417 although specific combinations should be trialled first (Bui et al. 2020; Gentry et al. 2020). 418 This contrasts with current louse control methods, which are less amenable to being used in 419 combination (for example, cleaner fish should not be subjected to mechanical delousing 420 along with the salmon). The technical ability already exists to place farms strategically to 421 minimise connectivity (Samsing et al. 2019), and salmon with higher louse resistance are 422 already being stocked by some farms in combination with barrier technologies (primarily 423 skirts) and/or functional feeds for louse resistance. Effective use of multiple preventative 424 425 methods in combination could reduce louse densities by orders of magnitude without negative effects on fish welfare, although as with any control strategy, potential welfare 426 concerns (e.g. those arising from holding salmon at depth) should be tested and mitigated 427 prior to widespread deployment. Vaccines may eventually result in even greater additive 428 reductions in louse densities. 429

430

431 MAINTAINING LONG-TERM EFFICACY

Host-parasite interactions are subject to a coevolutionary arms race in which organisms must 432 433 constantly evolve to keep up with the coevolution occurring in opposing organisms (i.e. the Red Queen hypothesis: Hamilton et al. 1990). Most lice never encounter a potential host, and 434 those that do will likely only have one opportunity to attach. This could precipitate strong 435 selective pressures, and because farmed salmon represent the majority of available hosts for 436 437 lice in some regions (especially in the north-east Atlantic), louse control interventions on farms are likely to exert directional selection pressure on louse populations wherever certain 438 genotypes are favoured over others. Evolution of resistance occurred relatively quickly in 439 response to chemical delousing (global reviews: Aaen et al. 2015; Gallardo-Escárate et al. 440

2019) and presently remains high (Helgesen *et al.* 2018), although in areas where wild
salmonids are abundant, flow of susceptible genes from lice on wild hosts may help to
maintain treatment efficacy (Kreitzman *et al.* 2017).

It is currently unclear whether preventative methods will be similarly vulnerable to the 444 445 evolution of resistance in lice, but some methods will likely create suitable conditions. For example, barrier technologies that span the surface layers (e.g. 0-10 m) may select for lice 446 that preferentially swim deeper. Potential for evolution will depend on many factors 447 including the heritability of the resistance to the preventative treatment in lice, the levels of 448 genetic variation existing in the louse population, the intensity of selection, treatment season, 449 frequency and geographic locations, prevailing currents and tides (louse dispersal) and the 450 biological complexity of the preventative mechanism. Nonetheless, the preventative 451 paradigm does have the advantage of a diversity of methods that may disrupt directional 452 selection for resistance to a given method. Research is needed to outline the best way 453 454 forward, but management strategies to slow the evolution of resistance to preventative methods should heed lessons from other systems (e.g. antibiotic resistance in human 455 medicine: Raymond 2019). Potential strategies to slow the evolution of resistance to 456 457 preventative methods may include:

(1) Continuing to delouse when necessary. Effective use of preventative methods will greatly
reduce the required frequency of delousing, but periodic delousing will hamper the genetic
proliferation of any lice that successfully infest stock.

(2) Deployment of multiple methods in combination to counteract directional selection. For
example, combining skirts or snorkels with non-depth-specific methods such as functional
feeds or spatial management may reduce directional selection for louse swimming depth.

(3) Planning of spatial 'firebreaks' whereby farms are removed or fallowed at strategic areas
to minimise louse population connectivity, thus reducing reinfestation rates and potentially
slowing the spread of resistant genotypes between farming areas (Besnier *et al.* 2014;
Samsing *et al.* 2017; Samsing *et al.* 2019).

(4) Ongoing selective breeding for louse-resistant salmon lineages to ensure that genetic gains are not lost through random genetic drift. Using current cohorts of wild sea lice when calibrating breeding value predictions for each generation will help to ensure that genetic gains continue to be relevant and account for any evolutionary developments in the louse population. Like other vertebrates, salmon have a complex immune system and biology, which should provide a range of potential defence options against parasites. Genomic selection probably affects a number of biological processes in the fish, and sea lice would therefore need to have sufficient genetic variability to be able to successfully adapt and counter the genomic selection. Development of multiple louse-resistant salmon strains may dampen directional selection for corresponding adaptation in louse populations.

Conversely, preventative methods could be utilised in a way that promotes evolution of certain resistant traits (such as deeper swimming) in order to increase specificity of louse populations to salmon in farming environments, and therefore reduce infestation pressure on wild salmon. Modelling is needed to determine whether such an approach could prove beneficial in decoupling encounters between farm-derived lice and wild salmonids.

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484 CONCLUSIONS

Effective use of barrier technologies such as skirts, snorkels, or closed containment, coupled 485 with supplementary preventative methods may make delousing treatments unnecessary at 486 many sites, while high-risk locations may require additional management and regulation. 487 Breeding of louse-resistant salmon has begun; heritable variation exists, and cumulative 488 improvements are reducing susceptibility to lice in some salmon lineages. The successful 489 development of an effective vaccine would also be an important advance. In general, 490 preventative methods are preferable to reactive delousing, and moving towards a prevention-491 492 focused paradigm on Atlantic salmon farms may yield significant improvements in fish 493 welfare and productivity, while avoiding significant environmental impacts.

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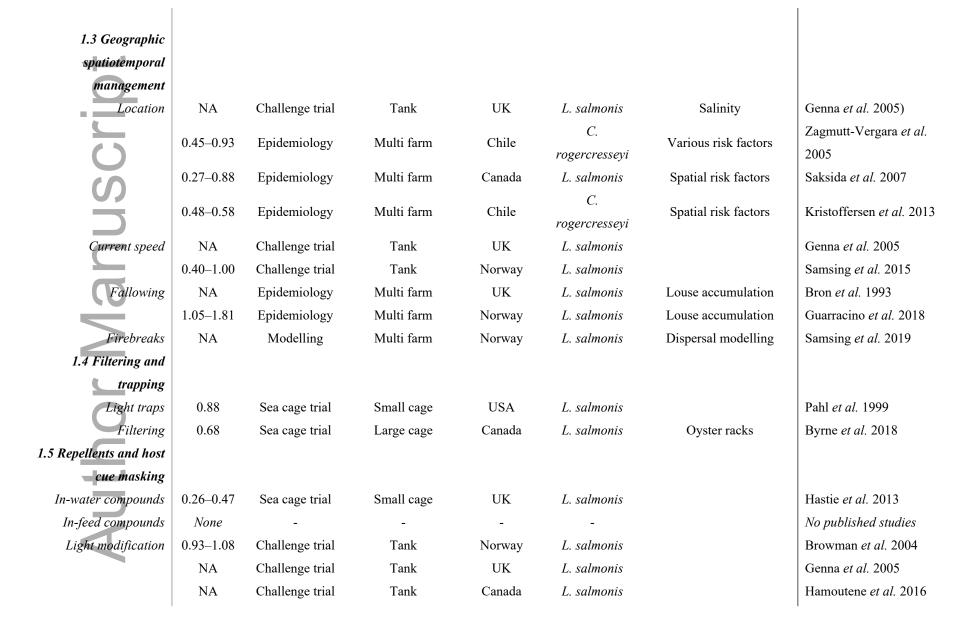
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TABLES

Table 1. Studies that assessed efficacy of preventative methods against louse infestation in Atlantic salmon. Effect sizes given are raw response ratios (treatment/control group) for louse infestation densities. Smaller values indicate more effective prevention. Where a study includes multiple treatment levels, the effect size range is given.

метнор	EFFECT SIZE (T/C)	STUDY TYPE	STUDY ENVIRONMENT	STUDY LOCATION	FOCAL LOUSE	NOTES	REFERENCE
1.1 Barrier technologies							
Snorkel cages	0.57	Sea cage trial	Small cage	Norway	L. salmonis		Stien et al. 2016
	0.05-0.37	Sea cage trial	Small cage	Norway	L. salmonis		Oppedal et al. 2017
Π	0.17	Sea cage trial	Large cage	Norway	L. salmonis		Wright et al. 2017
	0.24	Sea cage trial	Large cage	Norway	L. salmonis		Geitung et al. 2019
	0.36-1.08	Sea cage trial	Small cage	Norway	L. salmonis		Oppedal et al. 2019
Skirts	0.70	Sea cage trial	Multi farm	Norway	L. salmonis		Grøntvedt et al. 2018
L.	0.19	Sea cage trial	Large cage	Norway	L. salmonis		Stien <i>et al.</i> 2018
Closed containment	0.00-0.02	Sea cage trial	Multi farm	Norway	L. salmonis		Nilsen <i>et al.</i> 2017
1.2 Manipulation of							
swimming depth							
Forced submergence	0.08-1.72	Sea cage trial	Small cage	Norway	L. salmonis		Hevrøy et al. 2003
	0.31-0.45	Sea cage trial	Large cage	UK	L. salmonis		Frenzl et al. 2014
	1.09	Sea cage trial	Large cage	Norway	L. salmonis		Nilsson et al. 2017
	0.28	Sea cage trial	Small cage	Norway	L. salmonis		Sievers et al. 2018
	0.70	Sea cage trial	Small cage	Norway	L. salmonis		Glaropoulos et al. 2019
Deep lights/feeding	0.74	Sea cage trial	Large cage	UK	L. salmonis		Lyndon and Toovey 2000



1.6 Incapacitation							
Electricity	0.22	Sea cage trial	Small cage	Norway	L. salmonis	DC electric fence	Bredahl 2014
Ultrasound	0.61-1.37	Challenge trial	Tank	Norway	L. salmonis		Skjelvareid et al. 2018
Irradiation	None	-	-	-	-		No published studies
1.7 Louse population							
control							
Pathogens	None	-	-	-	-		No published studies
Gene drives	None	-	-	-	-		No published studies
2.1 Functional feeds							
Immunomodulation	0.56	Challenge trial	Tank	UK	L. salmonis	Nucleotides	Burrells et al. 2001
	0.61-1.09	Challenge trial	Tank	Canada	L. salmonis	Various additives	Covello et al. 2012
	0.48-1.31	Challenge trial	Small cage	Norway	L. salmonis	Various additives	Refstie et al. 2010
σ	0.70-0.81	Challenge trial	Tank	Canada	L. salmonis	Aquate, CpG	Poley et al. 2013
Ma	0.73–0.85	Challenge trial	Tank	Norway	L. salmonis	Various additives	Provan et al. 2013
	0.84	Challenge trial	Tank	Canada	L. salmonis	CpG	Purcell et al. 2013
	0.80	Challenge trial	Tank	UK	L. salmonis	Various additives	Jensen et al. 2015
<u> </u>	0.48-0.67	Cage trial	Small cage	Norway	L. salmonis	Sex hormones	Krasnov et al. 2015
0	0.78	Challenge trial	Tank	Chile	C. rogercresseyi	Various additives	Nunez-Acuna et al. 2015
	0.33-0.67	Challenge trial	Tank	Canada	L. salmonis	Peptidoglycan extract	Sutherland et al. 2017
uth	1.22	Sea cage trial	Large cage	Norway	L. salmonis	Skretting Shield (all cages had cleaner fish)	Bui <i>et al.</i> 2020
Ā	2.08	Sea cage trial	Large cage	Norway	L. salmonis	Skretting Shield (all cages had cleaner fish)	Gentry et al. 2020
Repellents/toxins	0.83	Challenge trial	Tank	Norway	L. salmonis	Phytochemicals	Holm <i>et al</i> . 2016
2.2 Vaccination							

Recombinant protein	0.43	Challenge trial	Tank	Chile	C. rogercresseyi	my32 protein	Carpio <i>et al.</i> 2011
	0.45–0.47	Challenge trial	Tank	Norway	L. salmonis	my32 protein	Kumari Swain <i>et al</i> . 2018
0	0.65–1	Challenge trial	Tank	Norway	L. salmonis	P33 protein offered strongest effect	Contreras et al. 2020
2.3 Breeding for louse							
resistance							
$\tilde{\mathbf{O}}$						Comparison of most	
Various	0.65	Sea cage trial	Small cages	Norway	L. salmonis	resistant and susceptible	Holm <i>et al.</i> 2015
						families	
Multiple methods	0.91	Sea cage trial	Multi farm	Norway	L. salmonis	All cages had cleaner fish	Bui et al. 2019b
						Functional feed + deep	
Ma	0.51	Sea cage trial	Large cage	Norway	L. salmonis	feeding and lighting (all	Bui et al. 2020
						cages had cleaner fish)	
						Functional feed + deep	
	0.79	Sea cage trial	Large cage	Norway	L. salmonis	feeding and lighting +	Bui <i>et al.</i> 2020
<u> </u>						skirt (all cages had	
						cleaner fish)	
						Functional feed + deep	
	1.91	Sea cage trial	Large cage	Norway	L. salmonis	feeding and lighting (all	Gentry et al. 2020
						cages had cleaner fish)	
						Functional feed + deep	
Authol	2.43 Sea	Sea cage trial	Larga ango	Norway	L. salmonis	feeding and lighting +	Gentry et al. 2020
		Sea cage unal	Large cage			skirt (all cages had	
						cleaner fish)	

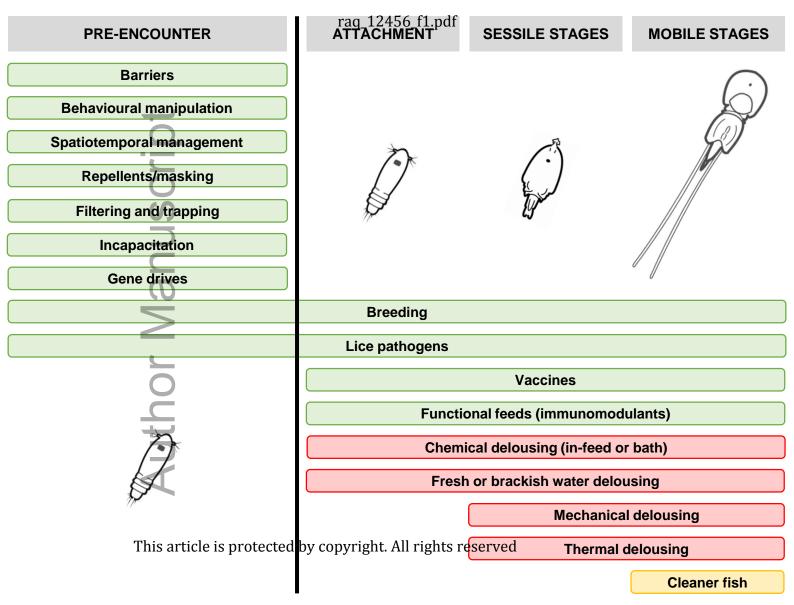
FIGURE CAPTIONS

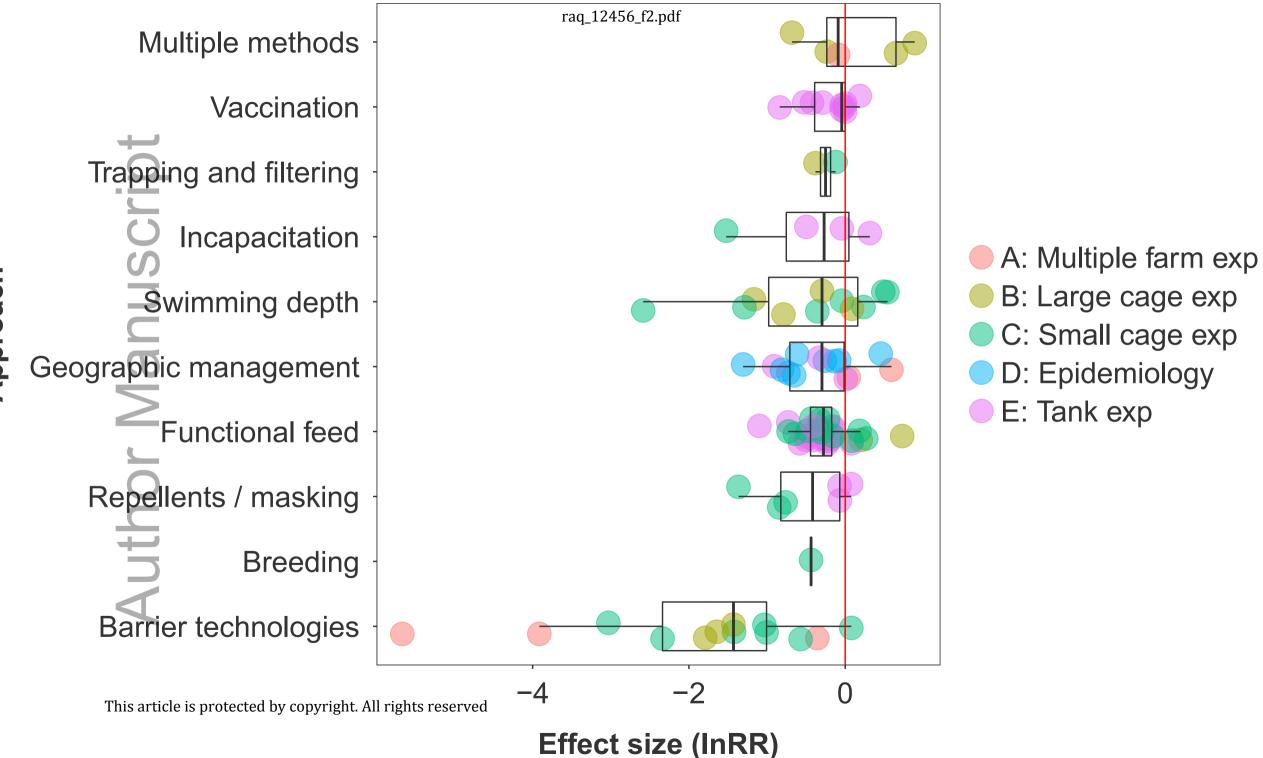
Figure 1. Sea louse infestation timepoints targeted by preventative methods and delousing treatments. Colours denote on-demand delousing (**red**), continuous delousing (**orange**) and preventative methods (**green**). Line drawings indicate the stage of louse predominantly affected by each method, L-R: larvae (nauplii and copepodids), sessile stages (chalimus I and II), and mobile stages (pre-adults and adults).

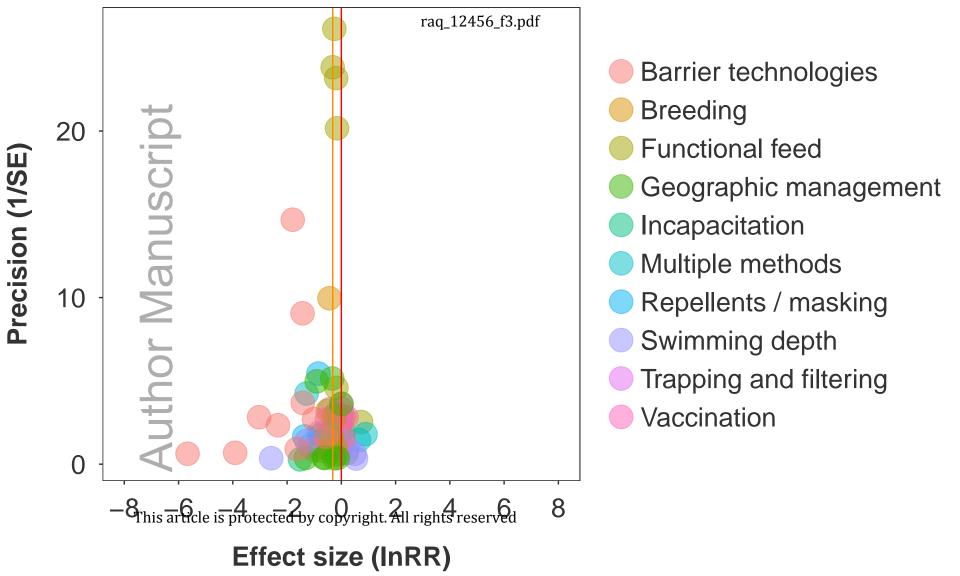
Figure 2. Distribution of effect sizes (natural log of the response ratio: lnRR) across studies testing preventive methods. Studies are grouped by the type of preventative method tested (Approach). Points denote the effect size of each study, coloured by the level of evidence provided. Negative values for lnRR indicate an effective approach. lnRR = 0 corresponds to no difference between control and treatment groups. Boxes indicate the median and 25-75% interquartile range of effect sizes from studies testing each approach.

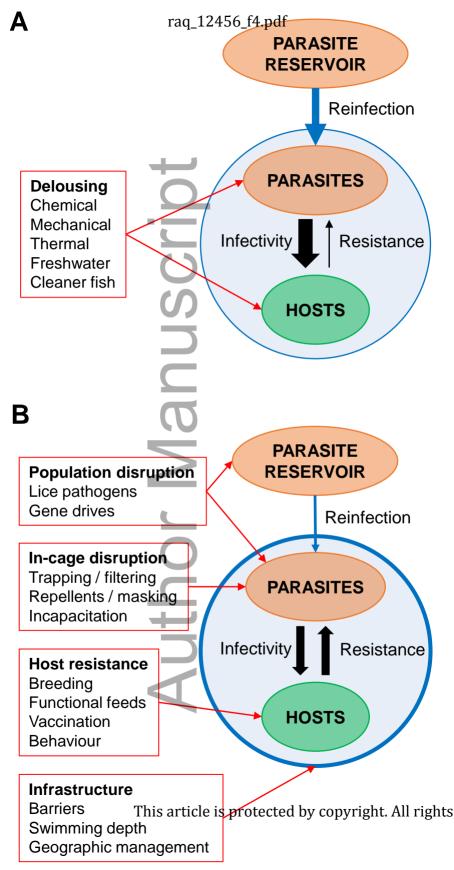
Figure 3. Funnel plot of published effect sizes (natural log of the response ratio) of preventative methods against sea louse infestations on Atlantic salmon. Effect sizes are plotted against the precision of the experiment (inverse of the standard error). The absence of studies on the right side of the plot is suggestive of publication bias against negative findings. **Red** line indicates zero effect (lnRR = 0), **orange** line indicates median effect size.

Figure 4. Conceptual diagram outlining: (A) the current delousing treatment-dominated paradigm for parasite control; (B) the new paradigm with a focus on prevention rather than treatment. Red arrows indicate management actions and how they are targeted (i.e. specificity, mediation). Blue arrows indicate supply of infective larvae (line thickness scales with number entering cages). Black arrows indicate host and parasite traits (line thickness scales scales with relative importance).









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