INTRODUCTION

Sperm competition, that is the competition between the ejaculates of two or more males for the fertilization of a given set of ova (Parker, 1970), and cryptic female choice, in which females influence the outcome of sperm competition (Eberhard, 1996), are important evolutionary forces across a diverse range of taxa in which females mate multiply. Variation in paternity success is therefore determined by differences in the various traits that might affect sperm competitive ability, which itself depends on both genetic and environmental factors, as well as on genotype-by-environment interactions (GEI). Seminal fluid is a major component of the male ejaculate that is often expected to mediate sperm competition, where different genotypes can differ in their seminal fluid expression as a response to different levels of sperm competition (i.e. exhibit GEI). We therefore here focussed on testing for GEI in expression of two recently identified seminal fluid transcripts, suckless-1 and suckless-2, which potentially modulate sperm competitive ability in the simultaneously hermaphroditic flatworm Macrostomum lignano via their effects on manipulating post-mating partner behaviour and ultimately the fate of transferred ejaculates. In addition, we sought to test for GEI in sperm competitive ability in a standardized sperm competition ($P_1$ and $P_2$) assay, to investigate the relationship between natural variation in the expression of these seminal fluid transcripts generated through GEI and relative paternity success. We found GEI for the expression level of suckless-1 and suckless-2, as well as for sperm competitive ability. Moreover, we found a positive relation between the expression of suckless-1 and relative paternity success ($P_1$). This suggests that natural variation in the expression of this seminal fluid transcript indeed can influence sperm competition outcomes in M. lignano.

KEYWORDS

gene/transcript expression, Macrostomum lignano, paternity success, post-mating sexual selection, simultaneous hermaphrodites, social group size, sperm competition
by many factors affecting sperm competitive ability of the ejaculate (Birkhead & Møller, 1998; Gage & Morrow, 2003; Lewis & Austad, 1990; Parker, 1970; Pizzari & Parker, 2009; Radwan, 1996; Simmons & Parker, 2006; Snook, 2005). Numerous adaptations related to the amount and quality of sperm such as sperm number, size, velocity, mobility and storage capacity affect relative paternity success under sperm competition (Birkhead & Møller, 1998; Bjork & Pitnick, 2006; Godwin et al., 2017; Parker & Pizzari, 2010; Pitnick, Hosken, & Birkhead, 2009; Pizzari & Parker, 2009; Snook, 2005; Wedell, Gage, & Parker, 2002). Alongside sperm, males typically also transfer large number of seminal fluid proteins/peptides (SFPs) during mating (reviewed by Avila, Sirot, LaFlamme, Rubinstein, & Wolfner, 2011; Hopkins, Sepil, & Wigby, 2017; Poiani, 2006) and these too are theoretically expected to be strongly shaped by sperm competition, favouring the evolution of SFP functions that confer a male fitness benefit through increased competitive fertilization success (Cameron, Day, & Rowe, 2007; Dapper & Wade, 2016; Dhole & Servedio, 2014).

SFPs can either decrease the risk of sperm competition, for example by preventing female re-mating, or increase the chance of sperm to fertilize eggs by serving as offensive or defensive tools against rival sperm in female genital tracts. Their well-known functions decreasing sperm competition risk include manipulation of female propensity to re-mate and/or attractiveness (e.g. Chapman et al., 2003; LaFlamme, Ravi Ram, & Wolfner, 2012; Lung & Wolfner, 2001) and blocking her genital tract by forming plugs to prevent additional successful copulations, as commonly occurs in many taxa (Barker, 1994; Jia, Duan, Jiang, & Wang, 2002; Mangels, Tsung, Kwan, & Dean, 2016; Sutter & Lindholm, 2016). On the other hand, the role of SFPs in sperm displacement has been reported for example in the fruit fly *Drosophila melanogaster* (Harshman & Prout, 1994) and in the seed beetle *Callosobruchus maculatus* (Yamane, Goenaga, Rönn, & Arnvist, 2015), in which seminal fluid of the second male to mate with a female causes a reduction in the number of sperm from the previous mating and correlations have been found between allelic variation at SFP loci and levels of sperm displacement, as well as resisting ability to being displaced (Clark, Aguade, Prout, Harshman, & Langley, 1995; Flumera, Dumont, & Clark, 2005). Moreover, in polyandrous ants and bees, seminal fluid enhances the survival of own sperm, while preferentially eliminating sperm of rival males (Den Boer, Baer, & Boomsma, 2010; Den Boer, Boomsma, & Baer, 2008).

So far, the literature on varying functions of seminal fluid modulating sperm competitive ability focuses mainly on separate-sexed organisms, but similar functions could also have evolved in hermaphrodites. Indeed, post-mating sexual selection has been suggested as a major evolutionary force shaping reproductive traits especially in simultaneous hermaphrodites (i.e. organisms with both male and female reproductive functions) (Charnov, 1979; Charnov & City, 1996; Marie-Orleach, Janicke, Vizoso, David, & Schärer, 2016; Michiels, 1998; Schärer, Janicke, & Ramm, 2015). Because frequent multiple mating is common in many reciprocally copulating simultaneous hermaphrodites, and individuals are capable of storing sperm from multiple ejaculate donors, there is an opportunity for selection to operate on differential fertilization success and resulting sperm competition among ejaculate donors (Anthes, 2010; Baur, 1998; Domínguez & Velando, 2013; Koene, 2005; Leonard, 2006; Michiels, 1998).

Our study organism, the flatworm *Macrostomum lignano*, is a reciprocally copulating simultaneous hermaphrodite in which self-fertilization does not occur (Ladurner, Schärer, Salvenmoser, & Rieger, 2005; Schärer & Ladurner, 2003), and is an emerging model organism to study ejaculate adaptations driven by sperm competition and sexual conflict (e.g. Janicke & Schärer, 2009a; Marie-Orleach et al., 2016; Patlar, Weber, & Ramm, 2019; Schärer, Littlewood, Waeschenbach, Yoshida, & Vizoso, 2011). *Macrostomum lignano* can adjust its sex allocation, that is the strategic investment to produce eggs and sperm, in response to sperm competition level. It has been clearly shown that in larger social groups, which predict larger mating group sizes (Janicke & Schärer, 2009a)—and where sperm competition intensity is high compared to small groups—worms invest more in their male sex function, as captured by traits such as testis size (Schärer & Ladurner, 2003; Janicke et al., 2013), testicular activity (Schärer, Ladurner, & Rieger, 2004), sperm production rate (Schärer & Vizoso, 2007) and spermatogenesis speed (Giannakara, Schärer, & Ramm, 2016). Moreover, it has been established that increasing investment in testis size and sperm production increases relative paternity success (Sekii et al., 2013; Vellnow, Marie-Orleach, Zadesenets, & Schärer, 2018). *M. lignano* also exhibits high mating rates, potentially due to the fact that individuals are motivated to donate sperm more than to receive it, just as in many other simultaneous hermaphrodites (Greeff & Michiels, 2017; Michiels, 1998; Schärer et al., 2015). However, the high motivation to mate more, in general, also increases the risk of receiving (excess) sperm and/or seminal fluid, and concomitantly increases risks of polyspermy, sexually transmitted pathogens and/or receipt of manipulative SFPs (Charnov, 1979; Schärer et al., 2015). Thus, it is likely that adaptations to gain control over the received ejaculate evolve in many simultaneous hermaphrodites, such as the sperm digestion common in gastropods (Baur, 1998; Greeff & Michiels, 2017; Michiels, 1998) and counter-adaptations to take control over own ejaculate such as bypassing the normal way of transferring sperm by hypodermic insemination in flatworms (Ramm, 2016; Ramm, Schlatter, Poirier, & Schärer, 2015; Schärer et al., 2011).

In *M. lignano*, there is a post-mating “suck behaviour” that often occurs immediately after mating. The precise function of this behaviour has not been definitively established, but it is proposed to be an adaptation to remove received ejaculate by sucking it out, that is potentially to gain control over the received ejaculate (Marie-Orleach, Janicke, & Schärer, 2013; Schärer, Joss, & Sandner, 2004; Schärer et al., 2011; Vizoso, Rieger, & Schärer, 2010). For example, Schärer et al. (2011) showed clearly some sperm shafts sticking out of the female genital opening after the suck has been performed, suggesting either sperm are removed out of the female storage organ or their position within it is altered, with potential implications for fertilization success. Supporting this
hypothesis about the function of the suck behaviour, sperm have morphological adaptations to resist being removed by the recipient; each sperm possesses a frontal feeler and two stiff lateral bristles to anchor itself in the antrum (Schärer, Joss, et al., 2004; Schärer et al., 2011; Vizoso et al., 2010). Moreover, a recently identified novel function of two seminal fluid transcripts, suckless-1 (Mlig-pro31) and suckless-2 (Mlig-pro32), which potentially mediate sperm competitive ability in *M. lignano*, occurs not by directly influencing sperm interactions between rivals but instead by manipulating this suck behaviour of the partner and presumably thereby affecting the fate of transferred ejaculates (Patlar, Weber, Temizyurek & Ramm., in press). We showed that the RNAi knockdown of these two transcripts in ejaculate donors increases the occurrence of the suck behaviour of their mating partner. This implies that the normal expression of these genes manipulates the partner to suck less often, meaning more sperm can likely be retained in the partner’s sperm storage organ, potentially enhancing paternity success (Patlar et al., 2019; Patlar et al., in press). We further showed substantial genetic variation in the expression of these transcripts (Patlar et al., 2019), indicating that this variation could be linked to variation in sperm competitive ability in *M. lignano*.

Although variation in sperm competitive ability can be expected to be depleted through strong directional selection, ejaculate traits often exhibit persistent genetic variation (Pitnick et al., 2009; Simmons & Kotiaho, 2002; Simmons & Moore, 2009). One potential explanation for this paradox is the existence of genotype-by-environment interactions (GEIs) (Danielson-François, Kelly, & Greenfield, 2006; Hunt & Hosken, 2014; Kokko & Heubel, 2008). GEIs create the potential to maintain genetic variation within populations exposed to conditions that vary in time and space (Gillespie & Turelli, 1989; Via & Lande, 1985) and could be especially important for understanding variation in sexually selected traits (Hunt & Hosken, 2014; Kokko & Heubel, 2008). So far, however, only a small number of studies have demonstrated GEI for sperm competitive ability itself (Engqvist, 2008; Lewis, Tigireros, Fedina, & Ming, 2012), although several others have shown substantial GEI for sperm traits (Evans, Rahman, & Gasparini, 2015; Marie-Orleach et al., 2017; Nystrand, Dowling, & Simmons, 2011; Simmons & Kotiaho, 2002; Snook, Bacigalupe, & Moore, 2010; Ward, 1998, 2000) and one showed GEI for expression of seminal fluid transcripts (Patlar et al., 2019), suggesting that variation through GEI can be widespread for ejaculate traits and potentially that it is related to, and could thereby help explain the maintenance of, variation in relative paternity success as an outcome of differential sperm competitive ability.

In this study, we therefore primarily aimed to investigate GEI for the expression of the *suckless-1* and *suckless-2* transcripts that potentially mediate sperm competition by manipulating partner suck behaviour, as well as for relative paternity success measured as first individual to mate (defensive sperm competitive ability; *Pd* 1) or second individual to mate (offensive sperm competitive ability; *Pc* 2) under sperm competition in *M. lignano*. We then sought correlative evidence of a potential link between these two traits, testing whether variation in seminal fluid expression generated through GEI (and/or other traits correlated with this variation) could predict relative paternity success under sperm competition.

### 2 MATERIALS AND METHODS

#### 2.1 Study organism

Cultures of the free-living marine flatworm *Macrostomum lignano* (Ladurner, Schärer, et al., 2005; Schärer & Ladurner, 2003) are kept in the laboratory at 20°C, 60% relative humidity, 14:10 light:dark cycle in six-well tissue culture plates (Techno Plastic Products AG, Trasadingen, Switzerland) containing artificial seawater (ASW) with 32‰ salinity, and fed ad libitum with the diatom *Nitzschia curvilineata*. Under these conditions, worms frequently copulate, up to around six times per hour, with the average copulation duration being relatively short, up to about 16 s, and they lay about 1–2 eggs per day (Schärer, Joss, et al., 2004; Schärer & Ladurner, 2003). During reciprocal copulations, both individuals transfer sperm and seminal fluid to each other via the stylet (male copulatory organ), and received sperm are stored in their female antrum (sperm storage organ) (Schärer, Joss, et al., 2004; Vizoso et al., 2010). Seminal fluid is produced by prostate gland cells located around the stylet (Ladurner, Pfister, et al., 2005; Vizoso et al., 2010) and includes a complex mixture of proteins (Weber et al., 2018).

In this study, we conducted experiments to investigate GEI for seminal fluid expression and relative paternity success using four different inbred lines: DV8, DV13, DV28 and DV71 (Patlar et al., 2019; Sekii et al., 2013). The origin and maintenance of these inbred lines are explained elsewhere (Patlar et al., 2019; Vellnow, Vizoso, Viktorin, & Schärer, 2017). In a previous study, these chosen inbred lines (hereafter also referred to as genotypes) were found to be slightly different in their overall seminal fluid investment and their plastic response to different group size manipulations, suggested promising potential for GEI in seminal fluid transcript expression, but GEI was not clearly established, likely due to low statistical power (Patlar et al., 2019). Another inbred line, DV1, which is the line commonly used in *M. lignano* studies (Janicke et al., 2013), was used to generate standardized recipient worms and a green fluorescent protein (GFP)-expressing outbred culture to generate sperm competitors in paternity assays. Note that the GFP marker is a dominant allele expressed in all somatic and gametic cell types allowing us to easily and reliably genotype the offspring following a double mating trial between a wild-type, non-GFP worm and a GFP-expressing worm in order to score paternity when in competition to fertilize the eggs of a wild-type non-GFP-expressing recipient (Marie-Orleach, Janicke, Vizoso, Eichmann, & Schärer, 2014; Vellnow et al., 2017).

It has been shown that GFP-expressing worms are not affected by carrying the GFP marker in their reproductive traits compared to wild-type outbred populations, which makes them reliable and powerful tools (Marie-Orleach et al., 2014). The GFP-expressing worms used in our experiment were from the outbred transgenic BAS1 culture (Marie-Orleach et al., 2014; Vellnow et al., 2017).
2.2 Assessing sperm competitive ability of genotypes

We followed a three-step experiment, since the aim was to estimate GEI effects on seminal fluid expression and sperm competitive ability and then their potential relation as a response to sperm competition level in the environment of chosen genotypes (Figure S1). Therefore, firstly we manipulated the sperm competition environment of genotypes by raising them from hatchlings in different social group sizes, namely pairs (group of two worms) and octets (group of eight worms). Second, once they were mature, we assessed the sperm competitive ability of these individuals (focals) originating from either a pair or an octet by conducting double mating trials in which virgin standardized mating partners (recipients) were mated sequentially with a focal worm followed by a (GFP-expressing) competitor, that is testing for the defensive sperm competitive ability of focals, \( P_1 \), or else a competitor followed by a focal, that is their offensive sperm competitive ability, \( P_2 \). Third, we measured seminal fluid gene expression of focals immediately after mating trials.

2.2.1 Social group size manipulation

We initially collected ca. 2- to 3-day-old juveniles from main stock cultures of each genotype (ca. 150 per line—\( F_0 \)) and divided them into two glass Petri dishes with ad libitum food to let them grow and lay eggs. Once they started to reproduce, we collected their 2- to 3-day-old offspring (\( F_1 \)) into one Petri dish (for randomization of juveniles collected from two Petri dishes of \( F_0 \)) and then we randomly distributed these \( F_1 \) offspring into 24-well tissue culture plates, including 1 ml of ASW and ad libitum food in each well, to form groups of pairs and octets. These offspring were raised for up to eight weeks in their given groups by transferring them to freshly prepared 24-well plates every week to prevent accumulation of their newly hatched offspring. We assumed that 8-week-old worms represent mature young adults considering their median life span is about 205 days (Mouton, Willems, Back, Braeckman, & Borgonie, 2009). Social groups were distributed on plates in a way that balanced for any potential plate position and genotype effects. In total, we formed 81 replicate pairs and octets. After eight weeks in social groups, one focal worm was chosen randomly from each group to compete against a rival GFP-expressing worm (either in the \( P_1 \) or \( P_2 \) assay) in double mating trials. Therefore, in order to avoid pseudo-replication for both the focal and GFP competitors, one individual worm was picked randomly from each group and immediately used for the sperm competition assays. In total, 40 replicates of pairs/octets of each genotype were tested in the \( P_1 \) assay and the remaining 41 of the original 81 replicates were tested in the \( P_2 \) assay. The \( P_1 \) and \( P_2 \) assays were run simultaneously, and for logistical reasons, we divided the assays into blocks performed over 13 days, with identical procedures on each day, and ensuring that recipients, genotypes and competitors used on each day were similarly aged (ranging from 55 to 62 days old) and randomly assigned. We paired each focal and its competitor with two recipients sequentially to simply increase the total number of offspring and thereby the precision of our paternity estimates, considering that individual worms lay relatively few eggs.

Each day, we paired a group of standardized virgin recipients (recipient one) with a focal worm (\( P_1 \) assays) or a GFP competitor worm (\( P_2 \) assays) for the first mating period of two hours. At the end of this two-hour period, the given focal or competitor from this first pairing was transferred to be immediately paired with a second recipient (recipient two) for a further two hours, whereas the recipient one was paired with the second individual to mate (either the GFP competitor—\( P_1 \) assay or a focal—\( P_2 \) assay, i.e. the opposite to the first period). At the end of these two hours, we then removed the second focal/competitor from recipient one and the first focal/competitor from recipient two, after which we immediately paired the second focal/competitor with recipient two. Immediately after their total four-hour
mating period, each focal worm was individually transferred to a 1-mL tube containing 25 μL RNA Later®, whereas GFP competitors were paired one by one with a separate group of virgin worms (DV1) in 24-well tissue plates, to verify their fertility. If a GFP worm did not sire any offspring with its partner after ca. seven–eight days, we paired it with at least three others to disentangle whether the GFP worm or its partner was the cause of the infertility. We later excluded data where GFP competitors did not achieve any reproductive success when paired with their additional partners.

Paternity assessment was done by counting the offspring of recipients one and two, which were isolated after the double mating trials to let them lay eggs. We sorted GFP-expressing or non-GFP-expressing offspring under a stereomicroscope equipped with epifluorescence (Nikon SMZ-18 stereomicroscope with a Nikon C-HGFI Intensilight fluorescence lightsource and GFP filter cube; Nikon GmbH, Düsseldorf, Germany) over a period of two weeks ensuring all produced offspring were observed. In total, 234 of 320 mating trials for the P₁ assay (for each line, the number of pairs and octets ranges between 15 – 25 and 23 – 30, respectively) and 248 of 328 mating trials for the P₂ assay (for each line, the number of pairs and octets ranges between 15 – 27 and 23 – 32, respectively) were successfully measured for paternity success. The reduction in targeted sample size for P₁ and P₂ mating trials was due to lost worms during social group size treatment or due to excluded recipients if both (recipients one and two) did not produce offspring. In total, paternities for 2,416 and 2,291 offspring were assigned in the P₁ and P₂ assays, respectively.

2.2.3 | Seminal fluid transcript expression

In order to evaluate the relationship between gene expression and paternity success, we measured the expression of suckless-1 and suckless-2 from eight randomly selected samples of each genotype/social group size combination from the worms used to assess P₁. We performed RNA extraction of the P₁ samples using the ReliaPrep™ RNA Tissue Miniprep System (Promega, USA, #Z6112,) following the manufacturer’s instructions. Afterwards, reverse transcription was performed using 4.0 μL of 10.0 μL total RNA solution with the cDNA-Synthesis Kit H Plus (VWR Peqlab, Germany, 732–3273). CFX Connect™ Real-Time PCR Detection System (Bio-Rad, CA, USA) was used for gene expression measurements. Reaction volumes were set at 10 μL, comprising 3 μL 2X SsoFast EvaGreen Supermix with low ROX (Bio-Rad, CA, USA), 150 nM of each primer pair (1.5 μL per pair), 3 μL nuclease-free water and 1 μL cDNA. Initial thermal cycling conditions were 1 cycle at 95°C for 5 min, followed by 39 cycles of denaturation at 95°C for 15 s and annealing/polymerization with a temperature of 59°C for 30 s. Raw C_t values (tripllicated technical measurements for each sample and transcript) were extracted from the CFX Software version 3.0. One sample, out of a total of 64, could not be quantified for gene expression because of possibly failed RNA extraction of this sample. We first evaluated the absolute expression values of technical replicates for consistency between them. To do so, we calculated the average value between all possible combinations of two replicates (average value of technical replicate one–two, one–three and two–three). If one technical replicate deviated by one C_t value or more from the other two (i.e. one C_t value increase is equal to ca. two times of mRNA/gene copies), we excluded this technical replicate (in total approximately 6% across all measurements) and based the expression measurement on the remaining two. Relative transcript expression values were calculated as ΔC_t (C_t of gene of interest – C_t of internal control) after averaging technical replicates, and results are reported as −ΔC_t values (Schmittgen & Livak, 2008). The gene macpiwi (Pfister et al., 2007) was used as an internal control, which is a stable gene in terms of expression at different group sizes (Patlar et al., 2019).

2.3 | Statistical analyses

All statistical analyses were performed with the R statistical software, version 3.2.3 (R Development Core Team, ). For transcript expression analyses, a two-way ANOVA approach was used to examine the main effects of genotype and social group size and the genotype-by-environment (group size) interaction, with relative transcript expression (−ΔC_t) as the dependent variable. To examine the GEI for paternity success, we fitted generalized linear models with a binomial error distribution with logit link function (P₁ and P₂ analysed in separate models) and models to examine three-way interaction of genotype, group size and mating order (with the two recipients). All models for paternity success analyses included the response variable as a matrix where the first column is the number of focal and the second column is the number of GFP offspring, and a random effect of focal ID (due to the fact that each focal was measured for up to two recipients). Significance tests of interactions were based on chi-square tests examining changes in deviance when the interaction term was dropped from the full model. To examine whether the number of sperm received/used may have been affected by social group size of foci’s, we further evaluated average total offspring number of recipients depending on environment using a two-way ANOVA approach testing recipient and social group size interaction for P₁ and P₂ assays. Finally, we fitted general linear models with a binomial error distribution with logit link function to examine main effects of the expression of transcripts on P₁ (i.e. the ratio of focal offspring to total offspring in the P₁ assay), for recipient one and recipient two separately. The rationale of fitting separate models for recipients one and two was, first, to be able to evaluate gene expression effects on each recipient separately considering that transcript expression measurements were performed only following matings with recipient two, and second, because we had found an effect of the order of mating with the two recipients on P₁ (see Results).

3 | RESULTS

As predicted based on our previous results, seminal fluid transcript expression differs significantly between genotypes for both
suckless-1 and suckless-2 and both exhibit significant GEI (Table 1). In addition, overall relative expression levels (−ΔCt) were similar between social groups (−3.79 ± 2.60 in pairs and −3.13 ± 2.65 in octets for suckless-1 and 0.31 ± 1.83 in pairs and 0.73 ± 1.40 in octets for suckless-2). Although the overall mean relative expression level of the two transcripts was apparently quite different (−3.46 for suckless-1 and 0.52 for suckless-2), the reaction norms showing differential expression pattern of transcripts across social groups were strikingly similar for all genotypes (Figure 1), suggesting GEI for expression of these transcripts manifests in a quite coordinated manner.

For the sperm competition assay, the initial model comparisons including mating order with recipient one (or two) as a discrete factor, plus genotype, social group size and their two-way interaction, showed that paternity success (scored as P1, plus genotype, social group size and their two-way interaction, showed that paternity success (scored as P1, plus genotype, social group size and their two-way interaction, meaning genotypes differ in their relative paternity success depending on both the social group size and mating order (Tables 2 and S2). Note that when we fitted models for GEI for each recipient separately, GEI was highly significant for both recipients in the P1 assay (p = .002 for recipient one, p = .005 for recipient two) and for the first recipient in the P2 assay (p < .001 for recipient one, p = .17 for recipient two) (Table S3).

The total offspring number of recipients did not differ between social groups, averaging 6.65 in pairs and 6.75 in octets for recipient one and 7.09 in pairs and 7.07 in octets for recipient two in the P1 assay, with no significant interaction found (F1,347 = 0.02, p = .88). The pattern was the same for the P2 assay (average offspring number was 5.88 in pairs and 6.39 in octets for recipient one and 6.18 in pairs and 6.31 in octets for recipient two; no significant interaction: F1,365 = 0.27, p = .61).

Finally, to determine how variation in seminal fluid transcript expression generated through GEIs might be associated with variation in paternity success, we regressed seminal fluid transcript expression data of individuals with their P1 scores. According to the generalized linear model fitted including additive main effects of suckless-1 and suckless-2, only the expression of suckless-1 had an effect on paternity success, but this was highly significant and consistent across both recipient one (p = .006) and recipient two (p < .001) (Table S4). We also calculated the correlation between the expression of the two genes, which was marginally significant but weak (r = 0.27, p = .037). Based on this, we then dropped suckless-2 from the models and evaluated and visualized only the effect of suckless-1 (Figure 3) for each recipient (GLM for recipient one: z = 3.14, p < .001, recipient two: z = 3.96, p < .001). To be precise, the average coefficient for suckless-1 = 0.24 (slope of log odds for recipient one = 0.24398 and recipient two = 0.2302), which is interpreted as the expected change in log odds for a one-unit increase in the expression level of suckless-1. The odds ratio can be calculated by exponentiating this value to get 1.27 which means we expect to see about a 27% increase in the odds of paternity success of genotypes overall, for a one-unit increase in the expression level of suckless-1.

| Effect          | df | SS     | F     | p    |
|-----------------|----|--------|-------|------|
| Group size      | 1  | 6.20   | 1.87  | .18  |
| Genotype        | 3  | 132.01 | 13.24 | <.001|
| Interaction     | 3  | 87.58  | 8.78  | <.001|
| Residuals       |    | 166.20 | 106.56|      |

Note: The significant p-values are written bold. (SS: sum of squares).

**FIGURE 1** Seminal fluid transcript expression of genotypes across social group size treatments. The mean relative expression (−ΔCt) of suckless-1 (left) and suckless-2 (right) of genotypes (inbred lines DV8, DV13, DV28 and DV71) in pairs and octets (social group sizes). Error bars show the standard errors of the mean.

**TABLE 1** Two-way ANOVA results for the seminal fluid transcript expression variation

| Effect          | df | SS     | F     | p    |
|-----------------|----|--------|-------|------|
| suckless-1      |    | 2.63   | 1.31  | .26  |
| suckless-2      |    | 30.30  | 5.02  | <.01 |

| Genotype        | df | SS     | F     | p    |
|-----------------|----|--------|-------|------|
| suckless-1      | 3  | 132.01 | 13.24 | <.001|
| suckless-2      | 3  | 87.58  | 8.78  | <.001|

**DISCUSSION**

We found evidence of GEI for the expression level of two seminal fluid transcripts that were recently functionally characterized, suckless-1 and suckless-2 (Patlar et al., in press), and for relative paternity success, as well as evidence linking suckless-1 expression to sperm competitive ability.
TABLE 2 Model comparisons to evaluate interaction effects for paternity success

|                          | \( P_1 \) | \( P_2 \) |
|--------------------------|-----------|-----------|
|                         | df | Chi | p>|Chi | df | Chi | p>|Chi |
| Model comparison for two-way interactions | 10 | 3.99 | .26 | 10 | 6.51 | .09 |
| Model without two-way interaction | 7 | 7 |
| Model comparison for three-way interactions | 17 | 11.15 | .01 | 17 | 10.48 | .02 |
| Model without three-way interaction | 14 | 14 |

Note: Generalized linear model comparisons (\( P_1 \) and \( P_2 \) as in separate models) based on likelihood ratio tests to evaluate the effects of two-way interaction (genotype-by-group size interaction) and three-way interaction (genotype-by-group size-by-mating order with the recipients). The full model for two-way interaction comparison incudes mating order, genotype and group size plus genotype-by-group size interaction as fixed factors, and focal ID as a random factor. The full model for three-way interaction comparison incudes mating order, genotype and group size plus all possible interactions as fixed factors, and focal ID as a random factor. The outcomes of the full models were added as supplementary tables (Tables S1 and S2).

Research on GEI in sexually selected traits is an important area of study in evolutionary biology and has been proposed to explain the maintenance of standing genetic variation in sexually selected traits that are often under strong selection (Clark, 2002; Hunt & Hosken, 2014; Ingleby, Hunt, & Hosken, 2010; Kokko & Heubel, 2008; Pomiankowski & Müller, 1995). The majority of studies have focused on either sexually selected traits involved in the premating episode of sexual selection (reviewed by Ingleby et al., 2010) or on sperm traits that were proposed as possible adaptations to sperm competition (reviewed by Reinhardt, Dobler, & Abbott, 2015). These studies have shown GEI on sperm characteristics such as sperm length (Morrow, Leijon, & Meerupati, 2008), sperm transfer rate (Engqvist, 2008), sperm velocity (Evans et al., 2015) and sperm motility (Purchase, Butts, Alonso-Fernández, & Trippel, 2010), whereas others have focused on testis size as a predictor of sperm production (Marie-Orleach et al., 2017; Nystrand et al., 2011) or on mating duration that is likely related to male ejaculate allocation in response to sperm competition (Breitman, Fricke, Hetherington, Stone, & Chapman, 2010; Breitman, Lízé, Walling, & Price, 2014). To our knowledge, there is only one previous study that directly tested GEIs for seminal fluid transcript expression, our own previous investigation also in M. lignano (Patlar et al., 2019; but see also Mangels et al., 2015). Here, we therefore provide novel evidence of GEI for another major aspect of the male ejaculate of adaptive significance under sperm competition.

First of all, we supported our previous results regarding significant genotypic variation for the relative expression of suckless-1 and suckless-2, beyond which we extended our findings by demonstrating also significant GEI for these transcripts. We already had some evidence of GEI for the expression of different seminal fluid transcripts from our previous study, where in total we showed significant GEI for 14 of 58 seminal fluid transcripts (Patlar et al., 2019), although this did not include suckless-1 or suckless-2, and where we also showed a lack of significant social group size effect for all 58 transcripts (Patlar et al., 2019; but see Ramm et al., 2019). In contrast to several studies that found high degrees of difference in expression of SFPs between manipulated sperm competition levels, the lack of group size and thus sperm competition effect in M. lignano may therefore be explained by the existence of GEI. Overall, these results suggest that GEI could be widespread for seminal fluid production traits, potentially helping to explain the maintenance of standing genetic variation for seminal fluid proteins that was shown in some previous studies (for gene expression: Smith, Hosken, French-Constant, & Wedell, 2009; Patlar et al., 2019; for protein abundance: Baer, Zareie, Paynter, Poland, & Millar, 2012; Goenaga, Yamane, Rönn, & Arnqvist, 2015; Mangels et al., 2015). In fact, one can expect some genetic variation to be maintained for some SFPs because of the lack of fitness relation; thus, selection does not decrease genetic variation. However, considering their varying functions are often tightly related to successful reproduction, one can expect the lack of selection is unlikely a general explanation for genetic variation in SFPs. Yet, phenotypic studies that have manipulated social environments to manipulate sperm competition have revealed considerable plasticity for seminal fluid expression according to the presence/absence of rival males (e.g., Fedorka, Winterhalter, & Ware, 2011; Harris & Moore, 2005; Mohorianu et al., 2017; Ramm, Edward, et al., 2015; Sloan, Lovegrove, & Simmons, 2018). Here, we argue that variation in social environment and thus the extent of GEIs may arise from heterogeneity in social conditions and could maintain genetic variation in seminal fluid expression. Further studies are needed to explain how often GEI occurs among organisms for seminal fluid expression, and another interesting question will be to what extent seminal fluid gene expression is controlled by genetic/epigenetic mechanisms to imprint social environmental heterogeneity to gene expression (Perry & Mank, 2014).

Second, we found some evidence for GEI for both \( P_1 \) and \( P_2 \), revealing that GEIs occur for relative paternity success and implying geno-type-by-environment interaction effects on donor sperm competitive...
ability. Notably, however, the differing reaction norms of different genotypes depended not just on the social group size, but also on mating order (i.e. whether we assessed paternity success in matings with the first or second recipient). To the best of our knowledge, only a few studies have so far examined GEI for sperm competitive ability, and one study provides very similar evidence to ours of GEI for $P_1$, but in that case depending on nutritional conditions affecting ejaculate contents in the flour beetle *Tribolium castaneum* (Lewis et al., 2012). Another study showed GEI for sperm transfer rate also depending on nutritional conditions, indirectly suggesting a GEI effect on paternity success depending on the amount of sperm transferred in the scorpion fly *Panorpa cognata* (Engqvist, 2008). Other studies mainly focused on genotype-by-genotype interactions for sperm competitive ability in other organisms; they provide strong evidence for female–male interactions and male–male interactions explaining variation in paternity success measured as $P_1$ and/or $P_2$ (Bjork, Starmer, Higginson, Rhodes, & Pitnick, 2007; Clark, Begun, & Prout, 1999; Clark, Dermitzakis, & Civetta, 2000; Dowling, Friberg, & Arnqvist, 2007; Firman, 2014).

Both GEI and individual genotype interactions for sperm competitive ability may well be taxonomically widespread, since many factors affect paternity success in sperm competition. For example, our results also indicate that the pattern of GEI depended crucially on the mating order. Our experimental design involved a focal worm being taken out from its respective group (pair or octet) and at once paired with a virgin recipient (one) for two hours, and thereafter immediately with a second virgin recipient (two) in the subsequent two hours. It has been shown that group size manipulation affects mating rate and average sperm transfer rate in *M. lignano* (Janicke & Schärer, 2009a, 2009b), and worms raised in pairs have more stored sperm in their seminal vesicle than worms raised in octets as a result of high mating rate in octets (Janicke & Schärer, 2009b; Marie-Orleach et al., 2014). It is possible that only few sperm are left in the seminal vesicles in worms raised in octets, and the genotypes which were used in this study may vary considerably among themselves for sexual traits. Therefore, genotypes might well differ in their testis size, sperm production rate or the amount of sperm stored in their seminal vesicles, and this may differentially affect ejaculate size and composition transferred to first versus second recipients—and thereby relative paternity success—depending on the group size and genotype. For example, after mating with recipient one, some genotypes might be faster or slower to replenish and restore sperm and/or seminal fluid proteins when paired with the second recipient, or genotypes might differ in their motivation to re-mate with novel partners depending on the group size from which they originated. However, the total offspring success did not differ overall between recipients depending on environment in both assays, suggesting that average number of sperm that successfully fertilized eggs were not affected by social environment manipulation.

A further interesting finding from our study is that variation in expression level of *suckless-1* generated through GEIs robustly predicted defensive sperm competitive ability ($P_1$), across both first and second recipients, potentially suggesting adaptive GEI for seminal fluid gene expression. Note that we did not evaluate offensive sperm competitive ability ($P_2$) in terms of its relation with seminal fluid expression because sperm displacement occurs in *M. lignano* (Marie-Orleach et
al., 2014) and it has been proposed that the shape of the copulatory organ could be important in sperm competition to outcompete sperm of the previous donor by mediating sperm displacement (Janicke & Schärer, 2009a): we therefore considered that paternity measurements of P2 assays might also depend strongly on variation in stylet morphology of the chosen genotypes. Moreover, when the worm mates first a previously unmated recipient, then only its sperm/seminal fluid affects whether the partner sucks or not, whereas if it mates second, then potentially both its and the first donor’s sperm/seminal fluid can affect recipient behaviour, confounding our test.

We expected that an increase in the expression of seminal fluid transcripts, assuming their expression is positively correlated with the amount of protein transferred, may result in an increased number of sperm being retained in the recipient’s sperm storage organ by manipulating the suck behaviour. Therefore, relative paternity success may be linked with the expression of these transcripts, and indeed that appears to be the case for one of them, *suckless-1*. Earlier work in *M. lignano* had found fewer sucks occur in virgin compared to sexually experienced worms depending on their partners mating status but not on the mating status of the individual that sucks, suggesting that individuals suck less often after copulating with a virgin individual that may transfer a larger ejaculate (Marie-Orleach et al., 2013). Our results suggest this could be because larger ejaculates transferred by virgin individuals contain more *suckless-1* proteins. However, our conclusion about the underlying mechanism should be treated with caution, because we did not directly manipulate transcript expression and it is likely that several other seminal fluid transcripts, as well as the other aspects of male allocation, sperm production, will to some extent co-vari with *suckless-1* expression (Patlar et al., 2019). Indeed, we previously demonstrated positive genetic correlation among seminal fluid transcripts and between testis size and seminal fluid investment (Patlar et al., in press), and recent studies clearly showed that increases in testis size have a positive impact on paternity success in *M. lignano* (Sekii et al., 2013; Vellnow et al., 2018). Therefore, the effect we found might be due to the collinearity between sperm and seminal fluid production, or between different seminal fluid components. If more than just *suckless-1* is involved in such a response, this could help explain why the natural variation in *suckless-1* expression investigated here was clearly linked to relative paternity success, but in a previous manipulative experiment where we knocked down the expression of (only) *suckless-1* we observed no such clear impact on paternity success (Patlar et al., in press). In the current study, it was striking that *P1* of some recipients (from both recipient one and two groups) scored as one hundred per cent (see Figure 3), particularly linked with higher expression level of *suckless-1* suggesting that an increase in seminal fluid expression, and especially *suckless-1* or transcripts that are highly positively genetically correlated in their expression with *suckless-1* (Patlar et al., 2019), might have an antagonistic effect on re-mating rate of the partner. Nevertheless, we note that we did not see any evidence for a link between paternity success and the expression level of a second seminal fluid gene, *suckless-2*. Further studies are needed to test for the actual effect of *suckless-1* on paternity success in competitive environments, especially by controlling the other aspects of male allocation. In fact, there are promising tools such as manipulation of sperm production in a dose-dependent manner in *M. lignano* (Sekii et al., 2013) that could be a very useful approach to control for the effect of variation in sperm production/transfer and thereby help disentangle the independent effects of seminal fluid proteins on sperm competitive ability.

To conclude, our study demonstrates that GEI occurs for seminal fluid transcript expression, depending on the social group size, and thus the level of sperm competition, and additionally, GEI also occurs for sperm competitive ability but depending on both group size and potentially on other traits, for example mating rate or average sperm transfer rate which themselves also depend on the level of sperm competition and individual genotypes. Further, we found evidence that natural variation in expression level of *suckless-1* generated through GEIs could predict relative paternity success and so influences the outcome of sperm competition.

**ACKNOWLEDGMENTS**

We thank A. Giannakara and M. Weber (Bielefeld University, Germany) for help in maintaining the laboratory cultures and T. Schmoll and P. Korsten (Bielefeld University, Germany) for useful statistical feedback. This work was funded by the Deutsche Forschungsgemeinschaft (DFG) (grant no. RA 2468/1-1).

**CONFLICT OF INTEREST**

SAR is a Reviewing Editor for *Journal of Evolutionary Biology*. The other author declares that there is no conflict of interest.

**ORCID**

Bahar Patlar https://orcid.org/0000-0002-0442-9061
Steven A. Ramm https://orcid.org/0000-0001-7786-7364

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