Interpopulational and seasonal variation in the chemical signals of the lizard *Gallotia galloti*

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**ABSTRACT**

Communicative traits are strikingly diverse and may vary among populations of the same species. Within a population, these traits may also display seasonal variation. Chemical signals play a key role in the communication of many taxa. However, we still know far too little about chemical communication in some vertebrate groups. In lizards, only a few studies have examined interpopulational variation in the composition of chemical cues and signals and only one study has explored the seasonal effects. Here we sampled three subspecies of the Tenerife lizards (*Gallotia galloti*) and analyze the lipophilic fraction of their femoral gland secretions to characterize the potential interpopulational variation in the chemical signals. In addition, we assessed whether composition of these secretions differed between the reproductive and the non-reproductive season. We analyzed variations in both the overall chemical profile and the abundance of the two main compounds (cholesterol and vitamin E). Our results show interpopulational and seasonal differences in *G. gallotia* chemical profiles. These findings are in accordance with the high interpopulational variability of compounds observed in lizard chemical signals and show that their composition is not only shaped by selective factors linked to reproductive season.

**INTRODUCTION**

Communication is one of the main challenges to be met by animals (Bradbury & Vehrencamp, 2011). The outstanding diversity of communicative traits lies in subsequent evolutionary processes of diversification that often trigger a wide repertoire of signals with a vast-range of functions (Espmark, Amundsen & Rosenqvist, 2000; Maynard Smith & Harper, 2003). Coloration (e.g., Keyser & Hill, 2000), movements (e.g., Peters, 2008), sounds (e.g., Llusia et al., 2013), vibrations (e.g., Hebets, 2004), electric stimuli (e.g., Dunlap, 2002) and chemicals (e.g., Bacquet et al., 2015) are some examples of signals that animals employ to interact. Multiple pieces of evidence show that signaling traits can differ among populations of the same species (Barquero, Peters & Whiting, 2015; Giery & Layman, 2015; Barbosa, Rebar & Greenfield, 2016) or even change within a population throughout the year.
as a result of seasonal effects because of the variability of climate or reproductive dynamics, among other factors. For example, a study in the sagebrush cricket (Cyphoderris strepitan) showed that nightly calling duration differed among seasons as a result of the cost associated to the sound emission during the reproductive season (Sakaluk & Snedden, 1990). Likewise, seasonality can alter visual signals, as was described in blue tits (Parus caeruleus), in which UV and blue ornaments differed between reproductive and non-reproductive periods (Örnborg et al., 2002).

Although Darwin Jr (1859) noted the importance of chemical communication in social and sexual interactions, the underlying factors of the staggering diversity of chemical signals used by organisms in social and sexual interactions are relatively unclear and have attracted considerable interest of the scientific community in the last decades (Mason, 1992; Johnston & Del Barco-Trillo, 2009; Wyatt, 2014; Apps, Weldon & Kramer, 2015; Wyatt, 2017). Environmental variables (e.g., Weber et al., 2016; Baekens et al., 2017), trophic resources (e.g., Henneken et al., 2017) or physiological changes and hormonal levels (e.g., Kent et al., 2008) are examples of biotic and abiotic factors that may modulate the production and expression of these signals (Symonds & Elgar, 2008; Steiger, Schmitt & Schaefer, 2010). In this context, and given that many of the above-mentioned factors operate differently throughout the year, chemical signaling might also change under the effect of seasonality. However, the few studies that have addressed this question are primarily focused on insects (e.g., McNeil, 1991) or mammals (Johnston & Del Barco-Trillo, 2009), whereas lizards have been almost neglected (but see Alberts et al., 1992).

Lizards release chemical signals through feces, skin and specialized follicular glands (Mason, 1992; Weldon, Flachsbarth & Schulz, 2008; Mayerl, Baekens & Van Damme, 2015; García-Roa et al., 2017a). Specifically, the lipophilic fraction of the chemical secretions produced by the femoral and precloacal follicular glands might play a key role in lizard recognition, hierarchy establishment, or mate choice (Cooper Jr & Steele, 1997; Martín & López, 2006; Carazo, Font & Desfilis, 2008; Font et al., 2012; Pruett et al., 2016). Their composition (i.e., the number and abundance of chemical compounds) can differ between species, and/or sexes (Khannoon et al., 2011; García-Roa et al., 2016a; García-Roa et al., 2016b; García-Roa et al., 2017b; Martín et al., 2017). Nevertheless, the interpopulational variation in these secretions remains only known for a few species (Runemark, Gabirot & Svensson, 2011; Gabirot, López & Martin, 2012; Martin et al., 2013; MacGregor et al., 2017) and examples of seasonal variation are scarce (but see Alberts et al., 1992). Indeed, the fact that many lizard species from temperate areas only produce secretions during the mating season (Martin & López, 2014; Martin & López, 2015) has promoted a huge bias in the characterization of their composition (Alberts, 1990; Escobar, Labra & Niemeyer, 2001; Khannoon et al., 2011; Khannoon, 2012; García-Roa et al., 2016b). Many regions around the world, however, have favorable climatic conditions for lizards during most of the year. In this scenario, it would not be surprising that lizards from these areas could produce chemical signals across different seasons.

In this work, we analyzed the femoral gland secretions of three subspecies of the insular Tenerife lizard (Gallotia galloti). This is a large lizard endemic to the Canary Islands (Spain) divided in four subspecies: G.g. eisentrauti, G. g. galloti, G. g. palmae and G.g. insulanagae.
These lizards inhabit different islands of the Canary Archipelago: Tenerife Island (G. g. eisentrauti and G. g. galloti), La Palma Island (G. g. palmae), and the small islets of Roque de Anaga (G. g. insulanagae). In addition, while G. g. eisentrauti and G. g. palmae inhabit cloudy and wet densely vegetated forest areas in northern Tenerife and La Palma respectively, G. g. galloti lives in dry and sunny semidesert areas in the south and the center of Tenerife (Thorpe & Brown, 1989; Bohórquez-Alonso & Molina-Borja, 2014). Hence, the divergent ecological conditions of each area might modulate the expression of chemical signals, as it has been described for visual ornaments (Thorpe & Brown, 1989).

We investigated potential differences in the composition of the femoral gland secretions among particular subspecies in the entire chemical profile and in the abundance of the two main compounds: cholesterol and vitamin E (α-tocopherol). It has been hypothesized that these two compounds might have protective properties under different environmental conditions; cholesterol might protect semiochemicals in dry conditions and vitamin E in wet environments (Escobar et al., 2003; Weldon, Flachsbarth & Schulz, 2008; Martín & López, 2014). Since the environment varies considerably among subspecies (Thorpe & Brown, 1989; Bohórquez-Alonso & Molina-Borja, 2014), this could translate into different abundances of both compounds. Moreover, as this lizard species is active during the whole year, we also examined whether the femoral gland secretions differed between the reproductive and the non-reproductive seasons.

**MATERIAL AND METHODS**

**Study sites**

We conducted our study in two islands (Tenerife and La Palma) of the Canary Archipelago, Spain (Fig. 1). Because G. galloti lizards are active during the whole year, we focused on two different periods, spring (reproductive season; R) and winter (non-reproductive season; NR) (Salvador, 2015). In Tenerife, we visited two populations of G. galloti, in March (R) and December (NR) 2013. The G. g. eisentrauti population was located at El Pris in the North of the island (28°30′46″N, 16°25′4″W) and the G. g. galloti population was in Malpaís de Güímar, in the Southeast of the island (28°18′3″N, 16°23′49″W). The two sampled populations do not overlap in their distribution. Further, we studied a population of G. g. palmae close to El Pedregal at La Palma Island (28°37′13″N, 17°54′24″W) in April 2014 (R). Due to logistic reasons we could not sample this population in the non-reproductive season.

**Sampling and analysis of the femoral gland secretions**

Field-work was carried out between 9:00 a.m. and 2:00 p.m. We captured a total of 30 males, G. g. eisentrauti (7), G. g. galloti (7) and G. g. palmae (7) in April, and G. g. eisentrauti (5) and G. g. galloti (4) in December, by using pitfall traps baited with banana and tomato (Oppliger, Vernet & Baez, 1999). All of the studied lizards were adults (snout-to-vent length range: G. g. eisentrauti, 92–130 mm; G. g. galloti, 92–125 mm; G. g. palmae, 100–120 mm). Traps were hidden next to bushes and rocks to avoid exposure to direct sunlight. We checked traps every 15 min to avoid overheating of captured lizards. Due to the high air temperatures in the islands, lizards were transported within individual cotton
Figure 1 The islands of La Palma and Tenerife (Canary Islands, Spain). Red dots represent the sampling localities of *Gallotia galloti palmae*, *G. g. galloti* and *G. g. eisentrauti* male lizards in (A) La Palma and (B) Tenerife.
bags to a cool room. Ethics clearance and permits for this research were obtained from Cabildo Insular de La Palma (sampling permit: 201302/030-A/OT-098/2013) and Tenerife (13796/JBFT/CPA/AFF 97/13, 2013-00484). We collected femoral glands secretions of males to analyze and describe their chemical composition. Secretions were extracted by gently pressing glands and were introduced in glass vials with glass inserts and teflon-lined stoppers. In order to obtain blank controls, we used the same procedure without collecting secretion. Samples were stored at $-20^\circ$C until analyses. Lizards were healthy and we released them within the next 24 h in the same place where they had been captured.

We analyzed secretion using a Finnigan-ThermoQuest Trace 2000 gas chromatograph (GC), fitted with a column (5% diphenyl/95% dimethylsiloxane; Supelco, Equity-5, 30 m length $\times$ 0.25 mm ID, 0.25-$\mu$m film thickness), and a Finnigan-ThermoQuest Trace mass spectrometer (MS) as a detector. We operated in splitless analysis mode. We analyzed 2 $\mu$l of each sample dissolved in 200 $\mu$l of n-hexane (99%; obtained from JT Baker, Deventer, The Netherlands). We used helium as the carrier gas during the injection ($270^\circ$C) and detection ($250^\circ$C) phases with a constant septum purge. The temperature analysis program began at 50 $^\circ$C isothermal for 10 min, increased to 280 $^\circ$C at a rate of 5 $^\circ$C/min, and then isothermal ($280^\circ$C) for 20 min. Data recording began 7 min after the separation initiated using the Software XcaliburTM 1.4 (Thermo Fischer Scientific Inc., San Jose, CA, USA). Previous to the analytical process, we examined the repeatability of the GC-MS process with five consecutive injections of a standard compound (heptadecane) made in different days, being the relative standard deviation (RSD) always below 1 for retention times and 10 for peak areas, which ensures the repeatability of the analytical method. Because of the small amount of secretions that we could extract from each individual, samples were analyzed only once, as it is usually done in similar studies (Alberts et al., 1992; Escobar, Labra & Niemeyer, 2001; Khannooun, 2012; Sáiz et al., 2017). Our analyses were focused on the lipophilic fraction of the femoral gland secretions and not in their proteins (Mangiacotti et al., 2016). Impurities identified in the control vial samples were not considered.

For the initial identification of compounds embodied in femoral gland secretions, we used the NIST/EPA/NIH 2002 computerized mass spectral library, through chemicals mass spectra comparison. When possible, identifications were confirmed by comparison of spectra and retention times with those of authentic standards (Sigma-Aldrich Chemical Co, St. Louis, MO, USA). Then, we calculated relative proportions of each compound determined as the percent of the total ion current (TIC) in each sample. For the comparison of overall chemical profiles (i.e., list of the identified compounds and their relative abundance per sample), we corrected the problem of non-independence of proportions using logit transformation of the proportion data by taking the natural logarithm of proportion / (1–proportion) (Aebischer, Robertson & Kenward, 1993).

**Statistical procedures**

To assess potential differences in the chemical profile among subspecies, we used the software PRIMER v6.1.13 with the PERMANOVA+ v1.0.3 add-on package. We calculated the euclidean distances between every pair of individual samples to produce a resemblance matrix and then used a single factor permutational multivariate analysis of variance test.
(PERMANOVA) (McArdle & Anderson, 2001) based on the Euclidean resemblance matrix using 999 permutations. All identified compounds were included in the analysis. Possible differences were also analyzed with a canonical analysis of principal coordinates (CAP) (Anderson & Willis, 2003).

In addition, we tested for statistical differences between the reproductive and the non-reproductive season in G. g. eisentrauti and G. g. galloti populations. For this, we used a two-way PERMANOVA test and GLM analyses taking into account the effects of population, season and their interaction (Anderson & Willis, 2003). We confirmed both normality of data (Shapiro-Wilk’s test) and homogeneity of variances (Levene’s test) in all cases. Pairwise comparisons were performed with Tukey’s tests. All the statistical analyses were performed with R 3.2.2, SPSS 20.0.0 and STATISTICA v8.0 Software.

RESULTS

Interpopulational differences in femoral gland secretions

The lipophilic fraction of the femoral gland secretions of G. gallotia lizards clearly differed among subspecies in the number, abundance and presence-absence of some compounds (Tables 1 and 2). Considering the three populations together, during the reproductive period, vitamin E (TIC = 34.91%) was the most abundant compound, closely followed by cholesterol (TIC = 24.32%).

There were significant differences among the chemical profiles of the three subspecies during the reproductive season (PERMANOVA; pseudo \( F_{2,18} = 48.99, P = 0.001 \)). The pairwise comparisons showed significant differences in all cases (permutation tests; \( 6.20 < t < 7.34, P = 0.002 \) for all). The CAP analysis classified 95.2% of individuals within the correct subspecies in accordance with their chemical profiles, using leave-one-out cross-validation and \( m = 2 \) axes (\( \delta_{1}^{2} = 0.98, P < 0.001 \)).

Focusing on cholesterol secreted in the reproductive season, we found significant differences among subspecies (GLM; \( F_{2,18} = 22.30, P < 0.001 \)). However, the pairwise comparisons showed that G. g. galloti differed significantly from G. g. eisentrauti (Tukey’s tests, \( P < 0.01 \)) and G. g. palmae (\( P < 0.001 \)), but the difference between G. g. eisentrauti and G. g. palmae was not significant (\( P = 0.07 \)) (Fig. 2). We observed a similar pattern in vitamin E, which differed significantly among subspecies (GLM; \( F_{2,18} = 8.51, P < 0.01 \)). However, while G. g. galloti showed significant differences in the abundance of vitamin E with G. g. eisentrauti (Tukey’s tests, \( P < 0.01 \)) and G. g. palmae (\( P < 0.01 \)), G. g. eisentrauti and G. g. palmae did not significantly differ (\( P = 0.98 \)).

Seasonal differences in femoral gland secretions

We found seasonal variation in the composition of the femoral gland secretions between G. g. eisentrauti and G. g. galloti populations (two-way PERMANOVA; population: pseudo \( F_{1,19} = 73.38, P < 0.001 \); season: pseudo \( F_{1,19} = 9.73, P < 0.01 \); interaction population × season: pseudo \( F_{1,19} = 1.99, P = 0.13 \)). The CAP analyses classified 100% of individuals within the correct subspecies (\( \delta_{1}^{2} = 0.99, P < 0.001 \)) or season (\( \delta_{1}^{2} = 0.90, P < 0.001 \)) according to their chemical profiles. In addition, we found significant differences in the abundance of cholesterol between these two subspecies (GLM; \( F_{1,19} = 7.60, P = 0.01 \)) and
Table 1  List of compounds identified in male femoral secretions of the lizard *Gallotia galloti*. Three subspecies were sampled: *G.g. eisentrauti*, *G.g. galloti* and *G.g. palmae* in the reproductive (R: april) and non-reproductive (NR: december) seasons. The relative proportion of each component was determined as the percent of the total ion current (TIC proportion) and reported as the mean (±1SD). Characteristic ions (m/z) are reported for unidentified compounds. RT: Retention time.

| RT (min) | Compound                          | Gallotia galloti eisentrauti | Gallotia galloti galloti | Gallotia galloti palmae |
|---------|-----------------------------------|------------------------------|--------------------------|-------------------------|
|         |                                   | NR season (n = 5)            | R season (n = 7)         | NR season (n = 4)       | R season (n = 7) |
|         |                                   | mean ± SE                    | mean ± SE                | mean ± SE               | mean ± SE          |
| 14.6    | Nonanol                            | 0.05 ± 0.03                  | 0.04 ± 0.02              | 0.16 ± 0.10             | 0.08 ± 0.08       | 0.08 ± 0.07       |
| 15.0    | Decanal                            | 0.01 ± 0.05                  | 0.01 ± 0.01              | 0.02 ± 0.02             | 0.02 ± 0.02       | –               |
| 17.4    | 2,4-Decadienal                     | 0.01 ± 0.01                  | 0.02 ± 0.02              | –                       | –               | 0.01 ± 0.02       |
| 19.6    | Decanoic acid                      | 0.01 ± 0.01                  | 0.02 ± 0.02              | –                       | –               | –               |
| 20.0    | Decanol                            | 0.06 ± 0.05                  | 0.10 ± 0.06              | 0.18 ± 0.17             | 0.09 ± 0.12       | 0.10 ± 0.07       |
| 22.5    | Undecanol                          | 0.01 ± 0.01                  | 0.03 ± 0.03              | –                       | –               | 0.01 ± 0.01       |
| 23.0    | Dodecanal                          | 0.01 ± 0.01                  | 0.01 ± 0.01              | –                       | –               | –               |
| 24.3    | Dodecanoic acid                    | –                            | –                         | –                       | –               | –               |
| 24.8    | Dodecanol                          | 0.05 ± 0.02                  | 0.04 ± 0.03              | 0.13 ± 0.13             | 0.08 ± 0.10       | 0.07 ± 0.03       |
| 25.3    | Tetradecanal                       | 0.20 ± 0.09                  | 0.10 ± 0.06              | 0.20 ± 0.06             | 0.12 ± 0.07       | 0.19 ± 0.07       |
| 27.0    | Tetradecanol                       | 0.34 ± 0.18                  | 0.29 ± 0.04              | 0.25 ± 0.15             | 0.13 ± 0.11       | 0.29 ± 0.22       |
| 27.9    | 9-Hexadecanal                      | –                            | –                         | –                       | –               | 0.35 ± 0.11       |
| 27.6    | Pentadecanal                       | 0.27 ± 0.14                  | 0.05 ± 0.03              | 0.11 ± 0.04             | 0.06 ± 0.04       | 0.47 ± 0.33       |
| 28.7    | Tetradecanoic acid                 | 0.01 ± 0.01                  | 0.19 ± 0.14              | –                       | –               | –               |
| 29.2    | Pentadecanol                       | 0.17 ± 0.03                  | 0.04 ± 0.02              | 0.13 ± 0.10             | 0.08 ± 0.12       | –               |
| 29.7    | Hexadecanol                        | 1.23 ± 0.61                  | 0.75 ± 0.38              | 0.62 ± 0.24             | 0.27 ± 0.23       | 1.28 ± 0.94       |
| 30.3    | 6,10,14-Trimethyl-2-pentadecanone  | 0.01 ± 0.01                  | 0.03 ± 0.02              | 0.01 ± 0.01             | 0.02 ± 0.03       | –               |
| 30.7    | Pentadecanoic acid                 | 0.03 ± 0.02                  | 0.08 ± 0.06              | 0.03 ± 0.04             | 0.02 ± 0.02       | –               |
| 32.0    | Hexadecanoic acid, methyl ester    | –                            | –                         | –                       | –               | 0.78 ± 0.40       |
| 31.2    | Hexadecanol                        | 1.24 ± 0.72                  | 0.39 ± 0.24              | 1.12 ± 0.80             | 0.44 ± 0.34       | –               |
| 31.4    | 2-Heptadecanone                    | 0.20 ± 0.12                  | 0.05 ± 0.03              | 0.08 ± 0.03             | 0.05 ± 0.03       | 0.17 ± 0.11       |
| 31.8    | Heptadecanol                       | 0.23 ± 0.08                  | 0.03 ± 0.03              | 0.11 ± 0.07             | 0.08 ± 0.15       | –               |
| 32.3    | 9-Hexadecenoic acid                | –                            | 0.23 ± 0.20              | 0.02 ± 0.03             | 0.11 ± 0.10       | –               |
| 32.8    | Hexadecanoic acid                  | 0.44 ± 0.78                  | 4.06 ± 1.20              | 0.79 ± 1.46             | 2.96 ± 2.04       | 0.89 ± 0.76       |
| 33.2    | Hexadecanoic acid, ethyl ester     | 0.08 ± 0.07                  | 0.11 ± 0.11              | 0.04 ± 0.04             | 0.77 ± 1.58       | 0.09 ± 0.07       |
| 33.5    | Heptadecanol                       | 0.01 ± 0.01                  | 0.07 ± 0.07              | –                       | –               | 0.02 ± 0.03       |
| 33.7    | Octadecanol                        | 4.41 ± 3.07                 | 0.38 ± 0.32              | 1.67 ± 1.82             | 0.65 ± 0.65       | 3.00 ± 1.67       |
| 33.8    | Hexadecanoic acid, 1-methylethyl ester | –          | 0.31 ± 0.31              | 0.07 ± 0.12             | 0.11 ± 0.13       | –               |
| 34.6    | Heptadecanoic acid                 | –                            | –                         | 0.08 ± 0.03             | –               | –               |

(continued on next page)
Table 1 (continued)

| RT (min) | Compound                                      | Gallotia galloti Eisenrautti | Gallotia galloti | Gallotia galloti palmae |
|---------|-----------------------------------------------|------------------------------|-----------------|-------------------------|
|         | mean ± SE                                     | mean ± SE                    | mean ± SE       | mean ± SE               |
| 35.0    | Octadecanol                                   | 0.14 ± 0.10                  | 0.35 ± 0.27     | 0.09 ± 0.07             | 0.46 ± 0.43 | 0.52 ± 0.61 |
| 35.3    | 2-Nonadecanone                                | 1.02 ± 0.65                  | 0.12 ± 0.07     | 0.42 ± 0.24             | 0.16 ± 0.09 | 0.39 ± 0.29 |
| 35.6    | Nonadecanal                                   | 0.44 ± 0.39                  | 0.02 ± 0.02     | 0.35 ± 0.25             | 0.08 ± 0.14 | 0.53 ± 0.27 |
| 35.9    | 9,12-Octadecadienoic acid                    | 0.01 ± 0.01                  | 1.49 ± 1.11     | 1.62 ± 1.38             |            | 0.02 ± 0.04 |
| 36.0    | 9-Octadecenoic acid                           | 0.10 ± 0.18                  | 2.33 ± 1.08     | 4.84 ± 7.91             |            | 0.19 ± 0.50 |
| 36.3    | 9,12-Octadecadienoic acid, ethyl ester       | –                            | –               | 0.27 ± 0.50             |            | –            |
| 36.4    | Octadecanoic acid                             | 0.29 ± 0.52                  | 2.00 ± 1.10     | 0.03 ± 0.05             | 2.29 ± 1.35 | 0.61 ± 1.12 |
| 36.8    | Octadecanoic acid, ethyl ester               | –                            | –               | 0.16 ± 0.31             | 0.15 ± 0.23 | –            |
| 37.4    | Eicosanal                                     | 0.18 ± 0.14                  | 0.30 ± 0.16     | 0.16 ± 0.13             | 0.10 ± 0.12 | 0.27 ± 0.23 |
| 38.7    | Eicosanol                                     | –                            | –               | 0.14 ± 0.16             | 0.34 ± 0.27 | –            |
| 39.1    | 5,8,11,14-Eicosatetraenoic acid, ethyl ester | –                            | –               | 0.19 ± 0.13             | 0.26 ± 0.33 | –            |
| 39.7    | Eicosanoic acid                               | –                            | –               | 1.45 ± 1.24             |            | 0.19 ± 0.22 |
| 40.3    | Eicosanoic acid, ethyl ester                 | –                            | –               | 0.65 ± 0.79             |            | –            |
| 46.7    | Squalene                                      | 0.62 ± 0.22                  | 3.99 ± 2.92     | 0.41 ± 0.40             | 1.54 ± 0.97 | 0.51 ± 0.23 |
| 46.8    | Cholesta-2,4-diene                            | 0.24 ± 0.25                  | 0.32 ± 0.36     | 0.16 ± 0.23             | 0.02 ± 0.03 | 0.06 ± 0.07 |
| 47.0    | Unidentified terpenoid                        | 0.04 ± 0.05                  | 0.08 ± 0.10     | –                       | –          | –            |
| 47.4    | Cholesta-4,6-dien-3-ol                        | 0.13 ± 0.02                  | 0.05 ± 0.05     | 0.23 ± 0.09             | 0.14 ± 0.13 | 0.23 ± 0.38 |
| 47.6    | Cholesta-3,5-diene                            | 0.34 ± 0.16                  | 0.49 ± 0.34     | 0.33 ± 0.20             | 0.31 ± 0.29 | 0.32 ± 0.27 |
| 49.5    | γ-Tocopherol                                  | –                            | –               | 0.01 ± 0.01             | –          | –            |
| 50.2    | Cholesterol                                  | 0.48 ± 0.34                  | 0.58 ± 0.26     | 0.69 ± 0.55             | 0.70 ± 0.65 | 4.45 ± 11.21 |
| 50.7    | Cholesterol                                  | 51.92 ± 14.07                | 19.35 ± 6.65    | 54.21 ± 6.25            | 41.98 ± 13.32 | 11.65 ± 4.39 |
| 50.8    | α-Tocopherol (Vitamin E)                      | 18.76 ± 9.32                 | 41.71 ± 11.77   | 24.59 ± 6.79            | 19.97 ± 9.66 | 43.07 ± 14.23 |
| 51.3    | Cholestan-3-one                               | 0.16 ± 0.15                  | 0.24 ± 0.16     | –                       | –          | –            |
| 51.4    | Ergosta-5,22-dien-3-ol                        | 0.33 ± 0.34                  | 0.33 ± 0.25     | 1.16 ± 0.93             | 0.74 ± 0.73 | 0.54 ± 0.55 |
| 51.9    | Ergosterol                                    | 0.16 ± 0.12                  | 0.19 ± 0.26     | –                       | –          | 1.10 ± 1.93 |
| 52.1    | Campesterol                                   | 7.71 ± 2.21                  | 4.58 ± 2.83     | 3.52 ± 1.63             | 4.84 ± 2.40 | 12.38 ± 17.31 |
| 52.3    | Cholest-4-en-3-one                            | 1.53 ± 0.64                  | 1.55 ± 0.77     | 1.56 ± 0.87             | 1.18 ± 0.95 | 2.25 ± 2.09 |
| 52.5    | Stigmasterol                                  | 0.26 ± 0.17                  | 0.33 ± 0.29     | 0.14 ± 0.17             | 0.29 ± 0.21 | 0.95 ± 1.28 |
| 52.7    | Cholesta-4,6-dien-3-one                       | 0.33 ± 0.06                  | 0.30 ± 0.21     | 0.57 ± 0.28             | 0.29 ± 0.22 | 0.36 ± 0.30 |
| 53.0    | Stigmasterol derivative?                     | 0.05 ± 0.10                  | 0.13 ± 0.22     | –                       | –          | 0.82 ± 1.59 |
| 53.2    | Sitosterol                                    | 3.68 ± 1.61                  | 4.40 ± 1.28     | 2.87 ± 1.96             | 3.36 ± 1.95 | 4.99 ± 3.19 |

(continued on next page)
| RT (min) | Compound | Gallotia galloti eisentrauti | Gallotia galloti galloti | Gallotia galloti palmae |
|---------|----------|-----------------------------|--------------------------|-------------------------|
|         |          | NR season ($n = 5$)         | R season ($n = 7$)       | NR season ($n = 4$)     | R season ($n = 7$) |
|         |          | mean ± SE                   | mean ± SE                | mean ± SE               | mean ± SE |
| 53.3    | Stigmastanol | 0.93 ± 0.50                 | 0.50 ± 0.49              | 0.54 ± 1.08             | 0.38 ± 0.37 | 0.34 ± 0.84 |
| 53.8    | Unid. steroid(143,157,211,253,353,380,412) | 0.01 ± 0.02                | 0.28 ± 0.52              | –                        | –          | –           |
| 53.8    | Cholest-5-en-3-one | –                        | –                        | –                        | –          | –           |
| 53.9    | Hexadecanoic acid, ethenyl ester | 0.13 ± 0.08                | 1.13 ± 0.30              | 0.16 ± 0.10             | 0.26 ± 0.24 | 0.15 ± 0.24 |
| 54.1    | Hexadecyl 9-hexadecenoate | 0.06 ± 0.07                | 0.28 ± 0.22              | –                        | –          | –           |
| 55.3    | Cholest-4-ene-3,6-dione | –                        | –                        | –                        | –          | –           |
| 55.9    | Octadecyl 9-hexadecenoate | –                        | 0.44 ± 0.37              | 0.02 ± 0.04             | 0.16 ± 0.15 | –           |
| 56.2    | Octadecyl hexadecanoate | 0.04 ± 0.57                | 1.12 ± 1.52              | 0.48 ± 0.48             | 2.05 ± 3.48 | –           |
| 56.4    | 9-Octadecenyl hexadecanoate | –                        | –                        | –                        | –          | –           |
| 56.9    | Octadecanoic acid, ethenyl ester | –                        | –                        | 0.23 ± 0.27             | 0.48 ± 0.40 | 0.55 ± 0.54 |
| 57.6    | Eicosyl hexadecenoate | –                        | –                        | 0.19 ± 0.19             | 0.37 ± 0.43 | 1.26 ± 1.14 |
| 59.1    | Unidentified waxy ester? | –                        | –                        | –                        | –          | –           |
| 59.4    | 9-Octadecenyl 9-hexadecenoate | 0.04 ± 0.08               | 0.31 ± 0.35              | –                        | –          | –           |
| 60.5    | 9-Octadecenyl 9-octadecenoate | –                        | –                        | 0.24 ± 0.28             | 1.16 ± 1.40 | –           |
| 60.7    | 9-Octadecenyl octadecanoate | –                        | –                        | –                        | –          | –           |
| 60.9    | Octadecyl octadecanoate | 0.01 ± 0.02               | 0.75 ± 1.07              | –                        | –          | –           |
| 61.2    | Unidentified waxy ester? | 0.35 ± 0.60               | 0.68 ± 0.81              | 0.39 ± 0.31             | 0.28 ± 0.35 | 0.31 ± 0.35 |
| 65.4    | Unid. ester of 9-hexadecenoic acid | 0.03 ± 0.04               | 0.39 ± 0.22              | 0.04 ± 0.09             | 0.04 ± 0.07 | 0.06 ± 0.09 |
| 65.9    | Octadecyl eicosanoate | 0.03 ± 0.05               | 0.81 ± 0.56              | –                        | 0.79 ± 0.91 | 0.36 ± 0.51 |
| 67.2    | Unidentified waxy ester? | 0.36 ± 0.14               | 0.07 ± 0.10              | –                        | –          | –           |

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also between the reproductive and the non-reproductive seasons (GLM; $F_{1,19} = 18.01$, $P < 0.0001$); the interaction was significant (population $\times$ season: $F_{1,19} = 6.69$, $P = 0.01$) (Fig. 3). Pairwise comparisons showed that there were seasonal differences in the abundance of cholesterol in G. g. eisentrauti (Tukey’s tests, $P < 0.001$), but not in G. g. galloti ($P = 0.37$). Moreover, differences in cholesterol abundance between G. g. eisentrauti and G. g. galloti were only significant during the reproductive season ($P < 0.01$), but not in the non-reproductive season ($P = 0.99$). Regarding vitamin E, we did not find significant differences between subspecies or seasons (GLM; population: $F_{1,19} = 1.95$, $P = 0.17$; season: $F_{1,19} = 3.80$, $P = 0.06$); the interaction was significant (population $\times$ season: $F_{1,19} = 11.15$, $P < 0.01$). We observed significant seasonal differences in the amount of vitamin E in G. g. eisentrauti (Tukey’s tests, $P < 0.01$) but not in G. g. galloti ($P = 0.77$). Moreover, differences between both subspecies were significant during the reproductive season ($P < 0.01$), but not in the non-reproductive season ($P = 0.60$) (Fig. 3).
DISCUSSION

This study describes clear differences among three subspecies of G. gallotia lizards in the overall composition of their femoral gland secretions. Cholesterol and vitamin E were the two main compounds and their abundance also differed among subspecies. Interestingly, we also observed seasonal effects in the composition of secretions.

Few studies have examined and reported interpopulational differences in lizard chemical signals. For example, Escobar et al. (2003) found differences in precloacal secretions between two populations of Liolaemus fabiani lizards. These authors hypothesized that the variation could be ascribed to the different environmental conditions found in the two populations. Two recent works analyzing femoral gland secretions in insular lizards also showed interpopulational variability. In these studies, the hypothesis that climatic conditions might be shaping the composition of femoral gland secretions was also suggested (Runemark, Gabirot & Svensson, 2011; Martín et al., 2013). Climatic conditions could be an important factor modulating the composition of chemical signals, as it has been suggested not only for lizards (Martin & López, 2013; Heathcote et al., 2014; Martín et al., 2017; Baackens et al., 2017), but also for other animals (Sentis et al., 2015; Boullis et al., 2016). The fact that G. g. gallotia inhabits dry habitats and secretes the highest level of cholesterol, whereas G. g. eisenrauti and G. g. palmae inhabit areas with higher humidity and their secretions bear the highest abundance of vitamin E, is in line with the hypothesis that cholesterol and vitamin E might protect the chemical secretions in dry and wet conditions, respectively (Escobar et al., 2003; Weldon, Flachsbarth & Schulz, 2008; Martin & López, 2014). However, other alternative selective factors influencing signal evolution have been posited, such as differences in predation pressure (e.g., Hughes, Kelley & Banks, 2012), trophic resources (e.g., Hennekens et al., 2017), intrasexual selection (e.g., Ord, Blumstein & Evans, 2001), alternative “sexual strategies” (e.g., Pellitteri-Rosa et al., 2014), population density (e.g., Fowler-Finn, Cruz & Rodríguez, 2017), use of alternative signaling modalities (Johnstone, 1996), the degree of interspecific competition (Losos, 2009), or habitat use (e.g., Alberts, 1992). We cannot therefore ensure that, in our study, the chemical profile variation among...
populations is totally driven by climatic variables. The use of a single population for each climatic region hampers an accurate correlation between chemical profiles and climate.

An intriguing and neglected area in the field of lizard chemical communication is to find out whether chemical signaling changes throughout the year, mainly because most of examined species inhabit temperate regions where lizards only produce chemical signals during the mating season. Alberts et al. (1992) offered the first evidence in this respect, showing that the femoral secretions of Iguana iguana have more lipids during the reproductive season and thus chemical signals changed between seasons. Several years later, Martins et al. (2006) showed that the amount of secretion produced by follicular glands decreased during the non-reproductive period in comparison with the reproductive one. In that study, however, the information regarding chemical composition was lacking. Our results show that the composition of femoral secretions of G. g. eisentrauti and G. g. galloti differed considerably among seasons. Nevertheless, the analyses of the two most abundant compounds (cholesterol and vitamin E) showed seasonal variation in G. g. eisentrauti, but not in G. g. galloti. The effect of seasonality in the production and expression of these compounds might be variable. For instance, some compounds must be acquired through the diet (Weldon, Flachsbarth & Schulz, 2008) and therefore, changes in dietary availability could produce a higher or lesser degree of their expression in femoral gland secretions (García-Roa et al., 2017c). Recent research hypothesize that several compounds may be also under a trade-off between being used in metabolism functions or in chemical signaling (Kopena et al., 2011; Kopena, López & Martín, 2014b; Kopena, López & Martín, 2014a; Martín & López, 2015). If dietary availability and metabolic requirements of these compounds differ between seasons, this might also affect to the secreted lipophilic fraction. This could happen with vitamin E which is used in metabolic processes. In addition, health state (e.g., parasites, immune condition) or the endocrine profile (e.g., testosterone or corticosterone levels) may be also determinant variables affecting chemical signals in different seasons (Martín & López, 2015).

In Summary, we showed quantitative and qualitative interpopulational and seasonal differences in the femoral gland secretions of the insular lizard G. galloti. Although we could not tease apart the specific factors underlying this variation, our study provides an encouraging scenario to explore whether climate plays a key role shaping the chemical signals of G. galloti subspecies. Future research should replicate the effects of climatic conditions on a widespread species over different climatic regions. Finally, given that Gallotia lizards may use other sensory channels to transfer information (e.g., visual or acoustic), integrative analyses that consider other modes of communication would provide a global view of how these lizards communicate in a multimodal context.

**ADDITIONAL INFORMATION AND DECLARATIONS**

**Funding**

This work was funded by projects of the Spanish Ministerio de Economía y Competitividad (CGL2011-24150/BOS, CGL2012-40026-C02-01 and CGL2014-53523-P). The funders had
no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

**Grant Disclosures**
The following grant information was disclosed by the authors:
Spanish Ministerio de Economía y Competitividad: CGL2011-24150/BOS, CGL2012-40026-C02-01, CGL2014-53523-P.

**Competing Interests**
The authors declare there are no competing interests.

**Author Contributions**
- Roberto García-Roa conceived and designed the experiments, performed the experiments, analyzed the data, wrote the paper, prepared figures and/or tables, reviewed drafts of the paper.
- Rodrigo Megía-Palma conceived and designed the experiments, performed the experiments, analyzed the data, wrote the paper, reviewed drafts of the paper.
- Jesús Ortega wrote the paper, reviewed drafts of the paper.
- Manuel Jara analyzed the data, wrote the paper, prepared figures and/or tables, reviewed drafts of the paper.
- Pilar López contributed reagents/materials/analysis tools, reviewed drafts of the paper.
- José Martín conceived and designed the experiments, analyzed the data, contributed reagents/materials/analysis tools, wrote the paper, reviewed drafts of the paper.

**Animal Ethics**
The following information was supplied relating to ethical approvals (i.e., approving body and any reference numbers):
Ethics clearance and permits for this research were obtained from Cabildo Insular de La Palma with the sampling permit 201302/030-A/OT-098/2013 and Tenerife, 13796/JBFT/CPA/AFF 97/13, 2013-00484.

**Data Availability**
The following information was supplied regarding data availability:
The raw data has been uploaded as Data S1.

**Supplemental Information**
Supplemental information for this article can be found online at http://dx.doi.org/10.7717/peerj.3992#supplemental-information.

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