Signals of Selection in UK Bumblebees

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1 Keywords

² Genomics, bioinformatics, bumblebees, big data, HPC, selection.

2 Introduction

Bumblebees are critical in crop and wildflower pollination globally: these charismatic species are therefore vitally important economically and ecologically; for food security and ecosystem stability (Goulson et al., 2008; Cameron and Sadd, 2020; Sun et al., 2021). However, many species of bumblebee (*Bombus sp.*) face population declines caused by multifactorial stressors. Habitat loss, fragmentation and degradation; climate change; pathogens and pesticides have worked in combination to drive the decline of some species (Goulson et al., 2008; Cameron and Sadd, 2020). Understanding underlying genetic factors linked to these trends will be important in making inferences about population health and future trajectories, as well as implementing successful conservation action.

This study will focus on three bumblebee species which are resident in the United Kingdom: *Bombus terrestris*, *Bombus hortorum* and *Bombus ruderatus*. Whilst *B. terrestris* and *B. hortorum* are widespread across the UK, *B. ruderatus* has faced historic population decline and has a limited and fragmented geographical range (Ellis et al., 2005). Differences in the population dynamics of these three species could be explained by scrutiny of their genomes to understand differences in the selective events they face.

Using restriction-site associated DNA sequencing (RADSeq) data previously collected from these species at multiple sites in the UK, I aim to detect loci under positive selection. Selection in these species of bumblebee has not previously been investigated, so this novel study may be essential in understanding the diverging population trends and in targeting conservation plans. Specifically I aim to detect loci under selection and to identify differences in these findings between the species. Where possible, I will functionally annotate the selected sites by identifying genes associated with the significant loci. Finally I will attempt to identify the functional categories that these genes belong to in order to understand the underlying drivers and consequences of these selective forces.

3 Proposed Methods

In order to achieve the main goals outlined above I will undertake the following methods:

- 1. I will implement the *STACKS* pipeline (Catchen et al., 2013) for genome assembly and sample filtering of the raw RADSeq data (Rochette and Catchen, 2017).
- 2. Following this, I intend to execute multiple methods for identifying loci under positive selection in the *Bombus sp.* including the commonly-used software *BAYESCAN* (Foll and Gaggiotti, 2008; Ahrens et al., 2018) (e.g. Blanco-Bercial and Bucklin, 2016; Kang et al., 2017; Leiva et al., 2019; de Jong et al., 2021). Loci determined to be under positive selection will be compared to those identified using different methods, in order to isolate those that are consistently identified as outliers. This will improve robustness of inferences by reducing the impact of false positives. Through this analysis I will test the hypothesis that RADSeg data can be used to detect selection in non-model species.

- 3. A comparison of these significant loci between the species will be made to determine whether the species' genomes are similarly or differently affected by selection. I hypothesise that the species in decline (*B. ruderatus*) will display different signatures of selection to the other species.
- 4. Since reference genomes are available for *B. terrestris* and *B. hortorum*, I will place the significant loci within the physical context of their genomes to identify genes in linkage with these loci, which may therefore be affected by selection (Manel et al., 2016). I would expect to find signals of selection in areas of the genome which are functionally related to the stressors that these species face.

4 4 Anticipated Outputs and Outcomes

From the comparison of selection-detection methods I would hope to identify several loci consistently established to be under selection in the *Bombus sp.*' genomes. By comparing which loci were detected I hope to identify any common patterns between the species, which could be indicative of selection pressures shared by the genus; as well as any differences, which may offer insight into the differing population trends of the individual species. Where possible, I would like to identify any relevant functional roles associated with genes in linkage with the detected loci to put this study into some environmental context.

52 5 Timeline

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Table 1: Gantt chart of expected timeline. Numbered tasks indicate the broad methodological sections outlined above.

	April	May	June	July	Aug	Sept
Method						
Task 1: Filtering and assembly						
Task 2: Detecting selection						
Task 3: Species comparison						
Task 4: Functional categorisation						
Write-up						
Introduction						
Methods (simultaneous to tasks)						
Results						
Discussion						
Viva prep		<u> </u>				

53 6 Budget

 $_{54}$ 8Tb External hard-drive to backup data and run some analyses locally (£200) and 21" monitor to improve data exploration (£230).

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I have seen and approved the proposal and the budget.

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Dr Peter Graystock