Investigating the status of pyrethroid resistance in UK populations of the cabbage stem flea beetle (*Psylliodes chrysocephala*)

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Declaration of interest: none

HIGHLIGHTS

- UK populations of cabbage stem flea beetle exhibit high levels of resistance to pyrethroid insecticides
- Pyrethroid resistance in UK populations is largely the result of increased metabolism
- Resistance levels are highest in the South East of the UK
- Resistance is advancing to the North and West of England.

ABSTRACT

The cabbage stem flea beetle, *Psylliodes chrysocephala* L. is a major pest of winter oilseed rape in several European countries. Traditionally, pyrethroid insecticides have been widely used for control of *P. chrysocephala*, but in recent years control failures have occurred. In line with previous surveys, UK populations of *P. chrysocephala* were found to exhibit high levels of resistance to the pyrethroid lambda-cyhalothrin. This resistance was suppressed by pre-treatment with the cytochrome P450 inhibitor PBO, suggesting that the resistance remains

largely metabolic. The L1014F (kdr) mutation in the voltage-gated sodium channel, which confers relatively low levels (10-20 fold) of resistance to pyrethroids, was also found to be widespread across the UK regions sampled, whereas the L925I (super-kdr) mutation was also present but much less common. The current survey also suggests that higher levels of pyrethroid resistance are starting to spread to the North and West of England, and that resistance levels continue to remain high in the South East.

Keywords: cabbage stem flea beetle: oilseed rape; pyrethroid resistance

1. Introduction

The cabbage stem flea beetle, *Psylliodes chrysocephala* (Coleoptera: Chrysomelidae) is an established and key insect pest of winter oilseed rape, particularly in the UK (Graham and Alford, 1981) and Germany (Zimmer *et al.*, 2014), and is a significant pest of other *Brassica* species in several European countries (Bromand, 1990; Bartlet and Williams, 1991; Bartlet, Mithen and Clark, 1996). *P. chrysocephala* inflicts damage at both the larval and adult stage, with the tunnelling of the larvae into the leaf petioles and main stems causing the most damage through weakening of the upper section of the roots and lower parts of the stems (Williams, 2004).When infestation is high, the plant tips distort, the stems wilt and the infested plants become more susceptible to fungal infections such as *Phoma lingam* (Alford, 2003), the bacterial disease *Erwinia* sp. and frost damage (Højland *et al.*, 2015; Højland and Kristensen, 2018). Adult *P. chrysocephala* cause damage by feeding on stems, cotyledons and the first true leaves during crop emergence resulting in 'shot-holing' symptoms, leading to poor plant vigour or potential seedling death before emergence when fields are heavily infested (Zimmer *et al.*, 2014). Prior to 2014, *P. chrysocephala* affected approximately 67% of the area of oil seed rape grown in the UK causing an annual 1% yield loss (Clarke *et al*., 2009). However, in 2014, serious crop losses due to adult beetles (2.7% of the national crop) were recorded, the most serious losses (5–14%) being in eastern and southern England (Wynn, Ellis and Alves, 2014). In the autumn of 2015, a more extensive survey found that over 65% of crops had some damage, and that the damage was more widely distributed across the country than in 2014, although nationally only 1% of crops was lost (Alves, Wynn and Stopps, 2015). Subsequent surveys have confirmed that the average numbers of larvae per plant have risen substantially in all regions since 2014 (as summarised in Dewar, 2017).

Prior to December 2013, control of *P. chrysocephala* relied on the protection of oilseed rape seedlings by systemic neonicotinoid seed treatments containing either imidacloprid, thiamethoxam or clothianidin, followed by the application of foliar pyrethroid sprays later in the season if needed (Højland *et al.*, 2015; Højland and Kristensen, 2018). However, in December 2013, the European regulatory authorities (EU commission, 2013) banned the use of neonicotinoid seed treatments on all outdoor flowering crops, thus preventing their use in oil seed rape, leading to the increase in *P. chrysocephala* and the increased use of pyrethroid sprays. Today, pyrethroids (e.g. lambda-cyhalothrin) are the only class of insecticide that remain for chemical control of *P. chrysocephala* in the UK and other parts of mainland Europe.

The continuous use of pyrethroids to control *P. chrysocephala*, coupled with the lack of alternative insecticides with different modes of action, has led to a high selection pressure, driving the development and spread of resistance. Resistance to lambda-cyhalothrin was first reported in 2008, in north-western Mecklenburg, Western Pomerania, a major oilseed rape growing area in Northern Germany (Heimbach and Müller, 2013). Zimmer *et al*., (2014) reported the presence of the L1014F kdr mutation in the voltage-gated sodium channel, with high frequencies of the allele (90-100%) being found in populations collected from across Northern Germany, with the beetles exhibiting a low level resistance against a range of pyrethroids including lambda-cyhalothrin and tau-fluvalinate. More recently, studies by Højland *et al*., (2015) and Højland and Kristensen (2018) have shown that pyrethroid resistance resulting from the kdr mutation has continued to spread northwards, and is now present in populations from both Demark and the UK, whilst in Germany it has spread further south. Despite the presence of kdr in UK populations, Højland *et al*., (2015) found that the high pyrethroid resistance levels, with control failures being observed at the full field rate, did not completely correlate with the *kdr* genotype suggesting that another mechanism of resistance, such as metabolic resistance, is also present. Given the lack of alternative insecticides with

different modes of action, the presence and spread of pyrethroid resistance is concerning for the chemical control of *P. chrysocephala*.

The present study has determined the current status, extent and geographical spread of pyrethroid resistance in UK populations of *P. chrysocephala*. Bioassays, based on glass vial exposure of adult beetles to lambda-cyhalothrin, were carried out on samples collected in 2018 and 2019 to examine how resistance had changed over this time across the UK. The presence of the *kdr* and super-*kdr* target-site mutations in UK populations was also monitored, and the potential contribution of a metabolic resistance component in the beetles assessed by pre-treatment with the synergist PBO, which is a cytochrome P450 inhibitor.

2. Methods

2.1 Collection of field samples of *Psylliodes chrysocephala*

In July/August 2018 and 2019, live *P. chrysocephala* adults were collected from oil seed rape pods freshly harvested from the fields at Rothamsted Research, Harpenden, Hertfordshire, using a hand-held battery-powered pooter. Insects were maintained at 15±1°C, with 65% relative humidity in a light:dark photoperiod of 12:12h. Adults were kept in a mesh cage and fed continuously on a diet of Chinese cabbage *(Brassica rapa* spp*)*. Further samples were received by post from oilseed rape fields across the UK and were kept in sealed plastic bags or plastic containers containing Chinese cabbage or oilseed rape plant material and moist tissue paper, maintained in the same environmental conditions as the Rothamsted samples.

2.2 Bioassays to test the effect of pyrethroids on *Psylliodes chrysocephala*

P. chrysocephala samples were tested for resistance to the pyrethroid lambda-cyhalothrin (Syngenta, UK) using a glass vial bioassay based on IRAC (Insecticide Resistance Action Committee) Method 031 (www.irac-online.org/methods/weevils-and-flee-beetles/2014). Glass vials (14ml: 7cm tall/ 2cm diameter) (S Murray and Co, Surrey, UK) were prepared by coating

the inner surface with different concentrations of the insecticide. Initial stock solutions were prepared by diluting the technical grade insecticide in technical grade acetone. Three doses, equivalent to 4%, 20% and 100% of the recommended field application rate of lambda-cyhalothrin (7.5 g ai/ha) were used (Table 1). The controls were glass vials treated with acetone only. To coat vials, 500µl of solution was pipetted into the vials which were then placed horizontally without lids on a roller in a fume hood. Vials were rotated at room temperature for at least 2 hours until all the acetone had evaporated. Vials were then left vertically at 4°C overnight before attaching the screw tops the following day.

Table 1. Concentrations of lambda-cyhalothrin tested on *P. chrysocephala* via glass vial bioassay.

Concentration		% Field rate
(g/ha)	ppm	
0.3	0.204	4
1.5	1.02	20
7.5	5.1	100

The adult beetles (see 2.1) were used within a few days of collection and only those capable of walking or jumping when released onto a tray inside a three-sided Perspex cage were collected, using a hand-held battery-powered pooter. A minimum of ten beetles were transferred from the inverted pooter through a small funnel into each vial. The vials were then resealed and left at 18±1°C under a 16:8h light:dark photoperiod. After 24 hours, the beetles were transferred to untreated glass vials without lids under upturned 200ml plastic disposable cups (VWR International Ltd, Dublin, Ireland), to allow for a potential recovery which can occur in insects with metabolic resistance. After a further 24 hours, the beetles were released onto a tray and individuals scored using a fine paint brush according to three categories: ‘mobile’ (capable of jumping or walking in a coordinated way), ‘affected’ (incapable of jumping or coordinated movement) or ‘dead’ (no movement). Scoring of the beetles from each vial was done for 10 minutes to avoid adults that were simulating death, a behaviour shown by this species that has

probably evolved through predation pressure. Results were expressed as percentage mortalities. Following scoring, beetles in each category were transferred to Eppendorf tubes and snap frozen using liquid nitrogen before being stored in a freezer at -80°C.

2.3 TaqMan PCR assay to detect the presence of kdr/skdr in *Psylliodes chrysocephala*

TaqMan genotyping assays (Livak, 1999) were used to determine the presence of the mutations responsible for the kdr (L1014F) and super-kdr (L925I) sodium channel substitutions in individual adult beetles. Primer Express v.2.0 (Life Technologies) was used to design the primer and probe sequences for the assays (Table 2). In both assays, VIC reporter dye-labelled probes were used to detect the wild-type susceptible allele and 6-FAM reporter dye-labelled probes to detect the resistant allele. Each probe contained a 3' non-fluorescent quencher dye.

Table 2. Primer and probe sequences used for TaqMan assays to detect the L1014F (kdr) and L925I (skdr) mutations in *Psylliodes chrysocephala*.

Primer/Probe		**Sequence**
Primers	kdr-F	GGACTGTATGCTAGTCGGTGATGT
	kdr-R	GCAAAGCCAAGAAGAGATTCAGTA
	skdr-F	GCCAAGTCATGGCCAACTT
	skdr-R	TATAATGCACAGCACAAAGGTCA
Probes	kdr-VIC	TTACCACAAGATTACC
	kdr-FAM	TTACCACAAAATTACC
	skdr-VIC	TGGGTGCTTTAGGTAA
	skdr-FAM	TGGGTGCTATAGGTAA

PCR reactions (15µl) contained 1.5µl (50ng) genomic DNA, 7.5µl SensiFast probe mix (Bioline Reagents Ltd, UK), 0.375µl of kdr or skdr primer/probe mix (800nM of each primer and 200nM of each probe) and sterile water. Reactions were run on an Applied Biosystems 7900HT real-time PCR system, with initial incubations at 50ºC for 2 minutes and 95ºC for 10 minutes, followed by 40 cycles of 95ºC for 15 seconds and 60ºC for 45 seconds. The increase

155 in VIC and 6-FAM reporter dye fluorescence was monitored in real time and an allelic
156 discrimination analysis performed using the 7900HT Sequence Detection System software.

157 2.4 Use of a synergist to identify the presence of metabolic resistance in *Psylliodes*
158 *chrysocephala*

159 Pre-treatment with the insecticide synergist Piperonyl butoxide (PBO), obtained from Sigma-
160 Aldrich (Missouri, USA), was used to detect potential metabolic resistance mechanisms *in-*
161 *vivo*. PBO was diluted in technical grade acetone to give an equivalent concentration of 4g
162 $aiha^{-1}$. 500µl of solution was then used to coat glass vials (see 2.2). Ten beetles per replicate
163 were transferred to the PBO-coated vials for 1 hour before being transferred to either untreated
164 control vials or vials coated with lambda-cyhalothrin at the 100% field rate (7.5 g $aiha^{-1}$). The
165 beetles were then bio-assayed in parallel to beetles from the same sample not pre-exposed to
166 PBO.

167 **3. Results and Discussion**

168 3.1 Survey of pyrethroid resistance in *Psylliodes chrysocephala* across the UK

169 To determine the current extent and geographical spread of resistance to pyrethroid insecticides
170 in UK *P. chrysocephala* populations, and how this compares to previous reports (Højland *et*
171 *al*., 2015), bioassays with lambda-cyhalothrin were conducted on adult beetle samples from
172 Rothamsted Research's farm in Hertfordshire and oilseed rape fields located across England,
173 Scotland and Wales. The bioassays allowed the samples to be categorised as being either
174 completely susceptible, or to contain beetles that were 0-25%, 25-50%, 50-75%, 75-99% and
175 100% resistant, depending on the percentage of beetles per sample surviving treatment with
176 7.5 g ai ha^{-1} lambda-cyhalothrin. The bioassay is an approved test method (method 031) for
177 determining resistance in *P. chrysocephala* (IRAC, 2014) and was used by Zimmer *et al*.,
178 (2014) to monitor the emergence and geographic spread of pyrethroid resistance in *P.*

179 *chrysocephala* in Germany, by Højland *et al*., (2015) to determine the spread of pyrethroid
180 resistance in Danish, British and German samples and most recently by Højland and Kristensen
181 (2018) when investigating lambda-cyhalothrin resistance in Danish populations. Similar
182 bioassays have also been used to monitor the spread of pyrethroid resistance in European
183 populations of pollen beetle (*Meligethes aeneus*), another major pest of oilseed rape (Zimmer
184 and Nauen (2011), Slater *et al.,* (2011), Nauen *et al*., (2012)).

185 In 2018, a total of 42 *P. chrysocephala* samples, obtained from four different regions across
186 England, but primarily from counties in the East (Fig. 1), were tested. Of these only five
187 samples were found to contain no mobile beetles at 100% of the recommended field rate for
188 lambda-cyhalothrin, which would be expected if the sample was susceptibility. However, for
189 these five samples mortality was found to be <90% at 20% of the field rate, suggesting
190 resistance is present as judged by the IRACs 'susceptibility rating scheme' (IRAC, 2014). The
191 other 37 samples all showed some level of resistance with the highest resistance, at 89% being
192 the sample from Bishop Cannings (Wiltshire).

193 In 2019 a total of 145 *P. chrysocephala* samples were obtained from across England,
194 representing more of the country (Fig. 1), two samples were received from Wales and one from
195 Scotland. Only the Scottish sample was found to be truly susceptible to lambda-cyhalothrin,
196 displaying 100% mortality at 20% of the recommended field rate. Worryingly, populations
197 containing 100% resistant beetles were recorded for the first time in the UK. Overall, the
198 distribution maps for pyrethroid resistance in UK populations of *P. chrysocephala* (Fig. 1)
199 suggest that higher levels of resistance are starting to spread to the North and West of England
200 and that resistance levels continue to remain high in the South East.

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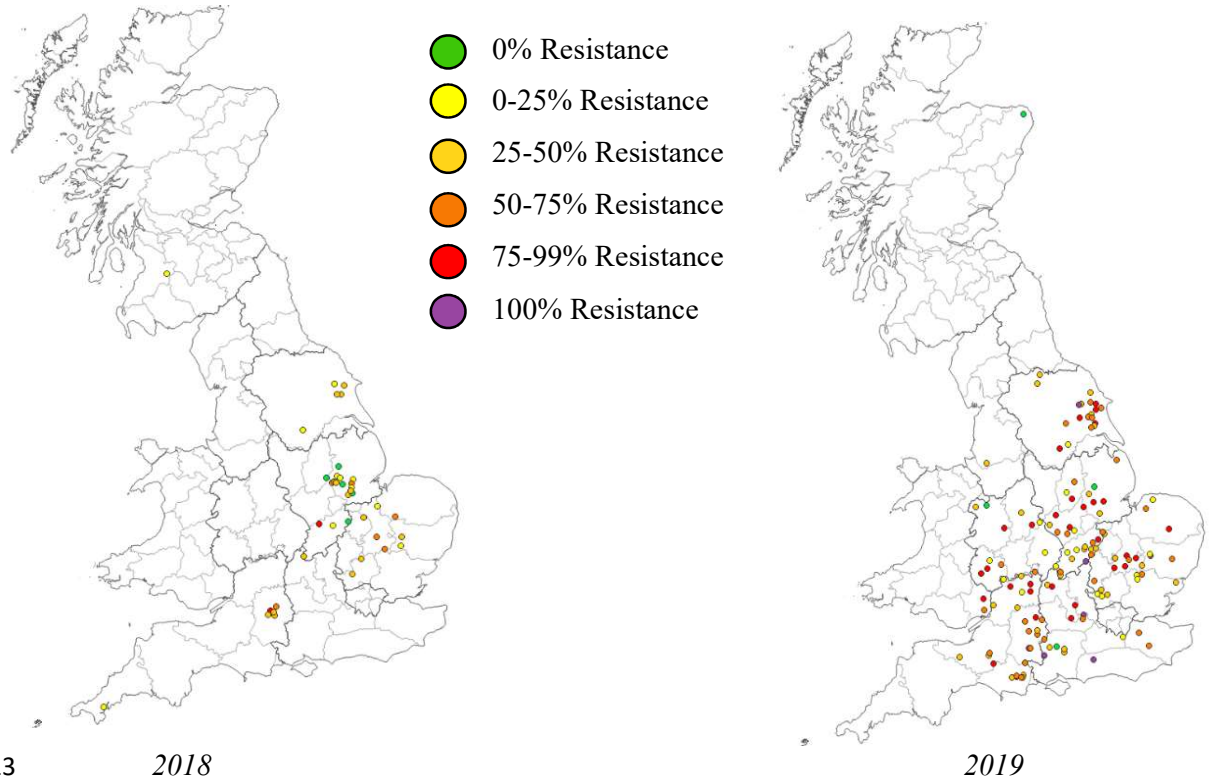

Figure 1. Pyrethroid resistance in *P. chrysocephala* in the UK for 2018 and 2019. The maps were created using QGIS (version 3.0.3) and use a 6-category colour scale to show the level of resistance. The map is divided into counties (light grey borders) and regions (dark grey borders).

Over the two years of monitoring, the percentage of pyrethroid-resistant beetles in each sample increased (Fig. 2). In 2018 the percentage in the 0-10% resistance category was approx. 16% whereas in 2019 this was down to 5%. In 2018 there were no samples in the 90-100% resistance category whereas in 2019 this increased to approx. 5%. The median resistance value was 28.5% in 2018 and 60% in 2019.

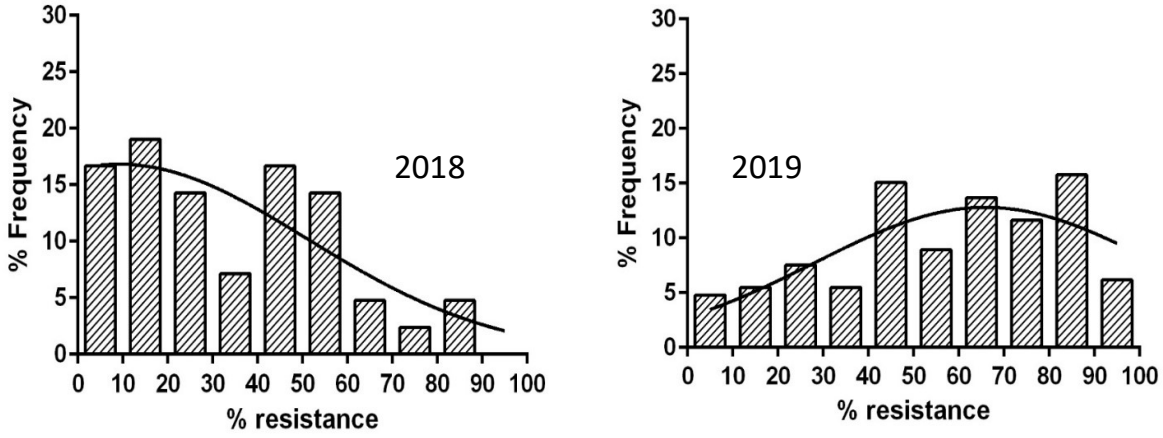

Figure 2. Histograms showing the shift in the relative % frequency of pyrethroid resistant *P. chrysocephala* in 2018 and 2019.

The median resistance levels recorded in 2019 were highest (approx. 60%) in the East Midlands, South East, South West and Yorkshire, and the Humber (Fig. 3). However, overall in each region the resistance levels were fairly evenly spread, with no statistically significant difference in resistance being found between regions.

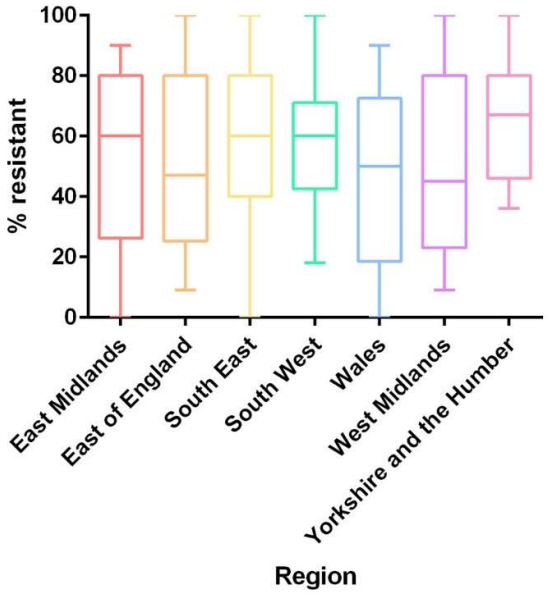

Figure 3. Boxplot of % resistance to pyrethroid in *P. chrysocephala* in each region of the UK surveyed in 2019.

When these 2019 data are broken down further to county level (Fig. 4), it is clear that there are also no significant differences in resistance across counties and that there are no resistance 'hotspots'. This suggests resistance is highly localised, almost on a farm-by-farm basis.

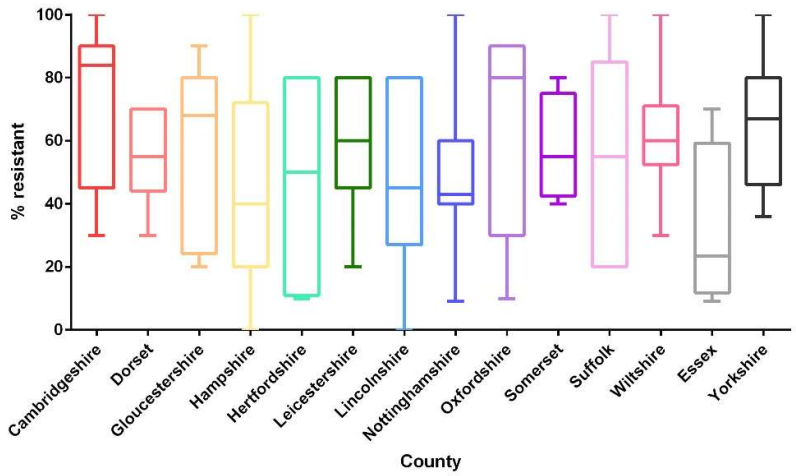

Figure 4. Boxplot of % resistance to pyrethroid in *P. chrysocephala* in each county of the UK surveyed in 2019.

3.2 Pyrethroid resistance mechanism(s) in *P. chrysocephala*

The TaqMan assays (see 2.3) were used to detect the presence of the L1014F (kdr) and L925I (s-kdr-like) substitutions in the 2018 and 2019 *P. chrysocephala* samples (Table 3). In 2018, 40 beetles from seven UK samples of *P. chrysocephala* were tested using only individuals that survived the 100% field rate of lambda-cyhalothrin, enabling the genotype associated with the resistant, mobile phenotype to be determined. The samples were from Great Saxham (Suffolk), Bishop Cannings (Wiltshire), Rothamsted (Hertfordshire), Linton (Cambridgeshire), Feltwell (Norfolk) and Horbling and Grantham (Lincolnshire). The L1014F mutation was present at all of the sites, with 47.5% of the beetles being homozygous for the resistant allele (RR), 37.5% heterozygous (SR) and the remaining 15% *kdr* SS, although this genotype was not present in the Suffolk or Wiltshire populations. In the Lincolnshire, Cambridge, Norfolk and Hertfordshire samples the *kdr* SS genotype represented 5%, 5%, 2.5% and 2.5% of the mobile beetles respectively. The detection of *kdr* SS genotypes in beetles that displayed the mobile phenotype after treatment with the label rate of lambda-cyhalothrin, suggests the presence of

another resistance mechanism in *P. chrysocephala*. In contrast to L1014F, the L925I mutation was much less common, with two samples (Norfolk and Rothamsted) being all SS and the overall percentage of beetles showing the homozygous L925I genotype (RR) being only 2.5%. This suggests that a potential fitness cost is associated with this mutation.

In 2019, *P. chrysocephala* individuals were screened for kdr from sites close to those sampled in 2018 (although a sample from Oxfordshire was also included) and again, only beetles that survived the 100% field rate of lambda-cyhalothrin were used. The L1014F mutation was present at all sites except the one from Scotland. The percentage of beetles homozygous for the *kdr* resistance allele (RR) increased in three of the samples, Suffolk, Norfolk and Hertfordshire but decreased overall from 47.5% to 36%. Given that the percentage of beetles resistant to lambda-cyhalothrin in each sample increased between 2018 and 2019, but there was an overall decrease in the homozygous and heterozygous L1014F mutation, this further suggests the presence of another resistance mechanism in *P. chrysocephala*. Whilst the L925I mutation was less common than the L1014F mutation, it was found to be present in the Wiltshire and Hertfordshire samples which contained only the wild-type metabolic genotype (SS) in 2018. In the Oxford sample 15% of the beetles tested for the s-kdr mutation were homozygous for the resistant allele (RR). Overall the percentage of beetles showing the homozygous genotype (RR) was 6.6%.

2018

Region	**County**	**Populations**	**No°**	***kdr status***			
				SS	**SR**	**RR**	**%RR**
East Midlands	Suffolk	1	4	0	3	1	25%
	Wiltshire	1	8	0	4	4	50%
	Lincolnshire	2	10	2	2	6	60%
East of England	Cambridgeshire	1	8	2	2	4	50%
	Norfolk	1	6	1	3	2	33%
	Hertfordshire	1	4	1	1	2	50%
	Total	7	40	6 (15%)	15 (37.5%)	19 (47.5%)	

Region	County	Populations	No°	*skdr status*			
				SS	**SR**	**RR**	**%RR**
East Midlands	Suffolk	1	4	3	1	0	0%
	Wiltshire	1	8	5	3	0	0%
	Lincolnshire	2	10	9	1	0	0%
East of England	Cambridgeshire	1	8	5	2	1	13%
	Norfolk	1	6	6	0	0	0%
	Hertfordshire	1	4	4	0	0	0%
	Total	7	40	32 (80%)	7 (17.5%)	1 (2.5%)	

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297 **2019**

Region	County	Populations	No°	*kdr status*			
				SS	**SR**	**RR**	**%RR**
East Midlands	Suffolk	1	4	0	1	3	75%
	Wiltshire	1	5	3	1	1	20%
	Lincolnshire	2	16	2	8	6	38%
East of England	Cambridgeshire	1	8	1	6	1	13%
	Norfolk	1	8	2	4	5	63%
	Hertfordshire	1	5	1	1	3	60%
SE England	Oxfordshire	1	19	7	4	8	42%
Scotland	Aberdeenshire	1	10	10	0	0	0%
	Total	9	75	26 (34.7%)	25 (33.3%)	27 (36%)	

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Region	County	Populations	No°	*skdr status*			
				SS	**SR**	**RR**	**%RR**
East Midlands	Suffolk	1	5	5	0	0	0%
	Wiltshire	1	3	1	1	1	33%
	Lincolnshire	2	16	11	5	0	0%
East of England	Cambridgeshire	1	8	5	3	0	0%
	Norfolk	1	8	7	1	0	0%
	Hertfordshire	1	6	4	1	1	17%
SE England	Oxfordshire	1	20	13	4	3	15%
Scotland	Aberdeenshire	1	10	10	0	0	0%
	Total	5	76	56 (73.7%)	15 (19.7%)	5 (6.6%)	

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300 **Table 3.** Detection of kdr/skdr alleles in *P. chrysocephala* using TaqMan assay

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304 3.3 Bioassays of *P. chrysocephala* using lambda-cyhalothrin and the synergist PBO

305 The insecticide synergist piperonyl butoxide has been shown to inhibit both P450
306 monooxygenases and esterases, thereby acting as a tool for the identification of metabolic
307 resistance in insect samples (Young, Gunning and Moores, 2006). To investigate the lack of
308 correlation between lambda-cyhalothrin resistance and *kdr* frequency, and to determine
309 whether P450 monooxygenases (and or esterases) may play a role in mediating pyrethroid
310 resistance in UK *P. chrysocephala* populations, synergist bioassays with PBO pre-treatments
311 were conducted on five *P. chrysocephala* samples.

312 When exposed to lambda-cyhalothrin at the recommended field rate, the percentage of beetles
313 affected was 47% (North Yorkshire), 75% (Wiltshire), 8% (Wiltshire), 40% (Leicestershire)
314 and 40% (Hertfordshire) (Fig. 5). However, all adults pre-treated with PBO, prior to exposure
315 to lambda-cyhalothrin at the same field rate were killed. This strongly suggests that a
316 metabolic-based mechanism for pyrethroid resistance is present in the *P. chrysocephala.*

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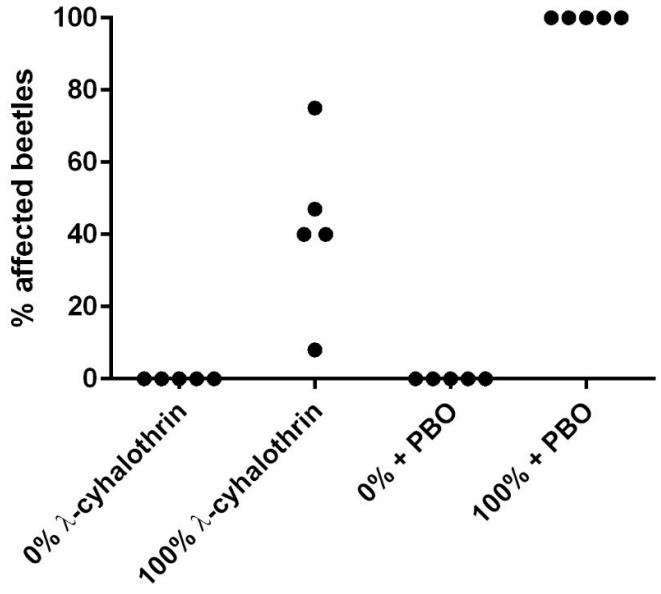

328 **Figure 5.** Restoration of insecticide (pyrethroid) susceptibility in *P. chrysocephala* following
329 pre-treatment with PBO. Samples tested were from North Yorkshire, Wiltshire (x2),
330 Leicestershire and Hertfordshire.

4. Conclusions

Since the EU-imposed ban on neonicotinoid seed treatments, the pyrethroid insecticide lambda-cyhalothrin has been widely used for chemical control of *P. chrysocephala* in the UK. This has resulted in a high selection pressure and led to the development of resistance, particularly in the South East of England. In the current study, populations of *P. chrysocephala* from around the UK were found to exhibit high levels of resistance to lambda-cyhalothrin, but some of this resistance was suppressed by the cytochrome P450 inhibitor PBO. This suggests that, as well as target site resistance, there is P450 mediated- detoxification of lambda-cyhalothrin, although further research is required to identify the specific P450(s) involved and elucidate the exact mechanism of resistance. This resistance to pyrethroids has resulted in widely-reported control problems for this pest in the farming press (e.g. Clark 2014; Casswell, 2014; FarmingUK team, 2015; Hill, 2017; FarmingUK team, 2017; Case, 2018; Allison, 2019; Dyer, 2019; Gillbard, 2019) since the introduction of the neonicotinoid ban.

The development of pyrethroid resistance in *P. chrysocephala* is a matter of growing concern, with the increasing spread of resistance threatening the production of oilseed rape both in the UK and mainland Europe (Heimbach and Müller, 2013). Thus, continuing to monitor the extent and geographical spread of pyrethroid resistance in this pest is particularly important at a time when synthetic pesticides are becoming less effective, through the evolution and selection of resistance, and less favoured, through EU legislation. Clearly there needs to be informed decision making on how to best deploy pesticides effectively in the future.

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