

letters Population ecology

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Long-term persistence of GM oilseed rape in the seedbank

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Coexistence between genetically modified (GM) and non-GM plants is a field of rapid development and considerable controversy. In crops, it is increasingly important to understand and Q5 predict the GM volunteer emergence in subsequent non-GM crops. Theoretical models suggest recruitment from the seedbank over extended periods, but empirical evidence matching these predictions has been scarce. Here, we provide evidence of long-term GM seed persistence in conventional agriculture. Ten years after a trial of GM herbicide-tolerant oilseed rape, emergent seedlings were collected and tested for herbicide tolerance. Seedlings that survived the glufosinate herbicide (15 out of 38 volunteers) tested positive for at least one GM insert. The resulting density was equivalent to 0.01 plants m^{-2} , despite complying with Q3 volunteer reduction recommendations. These results are important in relation to debating and regulating coexistence of GM and non-GM crops, particularly for planting non-GM crops after GM crops in the same field.

Keywords: volunteer; temporal gene flow; Brassica napus; seed; transgene

1. INTRODUCTION

Q4 Genetic modification (GM) technology makes it possible to engineer organisms with unique trait combinations, and it is currently a challenge to underpersistence of such organisms under field conditions (Pilson & Prendeville 2004; Snow et al. 2005). In particular, increasing effort is directed towards identifying and predicting the consequences of coexistence between GM and non-GM organisms (Ellstrand 2003; Pilson & Prendeville 2004). Part of the problem with coexistence is that the inserted transgenes disperse in the environment. The processes with which 55 this happens are analogous to non-GM escapees from agriculture (Ellstrand et al. 1999; Ellstrand 2003; Begg et al. 2006). The spread of GM organisms into non-GM populations may have implications by affecting the purity of non-GM crop and thus the consumer 60 willingness to buy such products of mixed origin (GM Science Review Panel 2003; Snow et al. 2005).

In agriculture, management strategies are adopted to reduce the GM volunteer plants (Pekrun et al. 1998; GM Science Review Panel 2003). Despite these measures, models now predict problems with volunteers from the seedbank, making it difficult to achieve GM contents below the 0.9% EU threshold (Lutman et al. 2005; Begg et al. 2006).

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In oilseed rape (OSR), Brassica napus L., experi-67 ments on gene flow between crop varieties are still 68 scarce (Légère 2005). Analysing temporal gene flow 69 through volunteer recruitment from the seedbank is 70 problematic in OSR because it is grown in a crop 71 cycle with a span of only a few years (but see Simard 72 et al. 2002; Lutman et al. 2005; Messean et al. 2007). 73 Long-term GM OSR seed persistence has instead 74 been investigated indirectly by, for example, sowing 75 non-GM seeds in cultivated or non-cultivated soil 76 (Pekrun et al. 1998; Lutman et al. 2003), seed burial 77 (Schlink 1998), adding seeds to semi-natural habitats 78 (Crawley et al. 1993, 2001) or molecular investi-79 gations of feral populations (Pessel et al. 2001). 80

In general, studies suggest that the majority of 81 seeds disappear from the seedbank within 2 years 82 (Crawley et al. 1993, 2001; Simard et al. 2002, 2005). 83 Recent models predict over 10-year OSR seed persist-84 ence in cultivated soil (Lutman et al. 2005; Begg et al. 85 2006), but empirical studies confirming this have not 86 been available (but see Messean et al. 2007). Here, 87 we investigated the long-term GM seed persistence in 88 a conventionally tilled system. Ten years after a GM 89 OSR trial in Sweden, a field was surveyed and 90 potential GM volunteers detected. Using a com-91 bination of crop use history, herbicide application and 92 molecular analysis, we investigated the presence of 93 descendents from the GM field trial. 94

2. MATERIAL AND METHODS

(a) Trial with GM OSR in 1995

98 In 1995, Plant Genetic Systems N.V. performed a field trial at 99 Lönnstorp Experimental Farm, Sweden (13°06' E, 55°40' N) with three transgenic OSR lines (OECD record number SWE95-005). 100 All these three lines were F_1 hybrids between a male sterile line and 101 a fertility restorer line (barnase and barstar transgenes, respectively; 102 table 1) and carried the transgene bar, which confers resistance to the herbicide glufosinate. Four 2×14 m subplots of each hybrid 103 line were sown in a 30×40 m trial plot. The remainder of the plot 104 and a 6-10 m border were sown with conventional OSR. The trial 105 was harvested in autumn 1995 (figure 1), with seed loss prevented 106 as much as possible. Shallow stubble tillage was performed twice to encourage germination before delayed ploughing in late November. 107 Rainfall was sufficient for OSR germination (figure 1). 108

(b) Field management 1996-2005

stand the fitness, competitiveness and long-term Q6 Between 1996 and 2005, wheat, barley and sugar beet were grown 110 (figure 1) in the trial plot. The field was ploughed every year and 111 harrowed before sowing. Volunteer occurrence during 1996-2005 112 was controlled by herbicides (a mixture of the herbicides tribenuron 113 (Express, DuPont Agro) and fluroxypyr (Starane, Dow Agro-114 Sciences)) and subsequent visual inspections. During the first 2 years after harvest, the field was controlled by the Swedish Board of 115 Agriculture, and for two additional years farm staff was under 116 obligation by the Swedish Board of Agriculture to control volunteer 117 rape in the field. During 1996-2005, farm staff controlled any volunteers observed with herbicides before flowering. Subsequent 118 inspections did not detect new volunteers. 119

(c) Analysis of OSR volunteers

121 Despite volunteer control, volunteers were still observed after 10 years. After harrowing in spring 2005, two persons searched the 122 trial field for 3 hours, collecting all detected volunteers. Volunteers 123 were planted in pots and kept outdoors. As controls we included 124 conventional (Vasaholm: 13°27' E, 55°38' N) and feral (Revinge: 13°25' E, 55°41' N) OSR plants. Unfortunately, at that time we did 125 not have access to glufosinate-tolerant control plants. After 19 days, 126 plants were hand sprayed with glufosinate herbicide (2% Basta). 127 Spraying was repeated after three weeks. Numbers of surviving 128 plants were recorded after spraying.

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Table 1. The plant genetic systems N.V. (PGS) transgenic lines grown in the trial of glufosinate herbicide (Basta)-resistant oilseed rape in 1995. (Identities of the transgene-carrying plasmids are given within parentheses.)



Figure 1. (*a*) Management of the GM Basta-tolerant OSR field trial in 1995, and (*b*) crops grown at the trial site from 1996 1 to 2005. Numbers in horizontal bar denote post-harvest monthly precipitation (mm) from August 30 to November 30.

Table 2. Survival of volunteer, crop and feral oilseed rape after glufosinate herbicide (Basta) application, and the frequencies of molecular markers for the transgenes *barnase* and/or *barstar* in the 15 surviving volunteers.

plant type	number of plants	number of plants surviving	plants positive for		
			barstar	barnase	barstar+barnase
volunteer	48	15	12	1	2
non-GM crop	67	0		_	_
non-GM feral oilseed rape	21	0		_	

Molecular analysis of surviving plants was performed to identify the occurrence of GM lines. One fresh leaf was collected from surviving plants, stored at -20°C and DNA was extracted using the DNeasy standard procedure (Qiagen). PCR analysis was performed with primers (23-24 mers) specific to the inserted constructs, *barstar* and *barnase*, which are genes for male sterility (*barnase*) or its restorer (*barstar*). Positive and negative control plants were included in the PCR analysis. The PCR temperature **Q7** cycle was 95°C per 4 min (1 cycle), 95°C per 1 min, 57°C per 1 min, 72°C per 2 min (5 cycles), 92°C per 30 s, 57°C per 30 s, 72°C per 2 min (25 cycles) and finally 72°C per 10 min. Products were visualized with ethidium bromide on agarose gels. Control plants with the *barnase* and *barstar* constructs were obtained from other sources and included in the analysis.

In January 2006, 40 soil samples (2.5 cm in diameter and approx. 25 cm deep) were randomly collected in the former trial field. Samples were stored at 2°C until sown in trays and placed in a greenhouse. The soil was watered whenever necessary and mixed several times to encourage germination (Lutman *et al.* 2003).

3. RESULTS

We found 38 volunteer OSR plants in the former ^{Q9} fields are rare. Available studies demonstrate that trial plot. Fifteen volunteers survived Basta application while none of the controls did (table 2). OSR persists for 5–6 or up to 8 years in agricultural fields (Simard *et al.* 2002; Lutman *et al.* 2005;

The difference in survival was highly significant (Fisher's exact probability test, p < 0.0001). All surviving OSR volunteers were positive for at least one, in two cases both, of the inserted genes (table 2, figure 2), thus clearly demonstrating a link between the GM OSR trial in 1995 and the volunteer population 10 years later. The density of GM OSR was 0.012 plants m⁻², and for all volunteers 0.04 plants m⁻². Seedlings from seven weed species germinated, but no OSR seedlings germinated.

4. DISCUSSION

Although temporal gene flow has been suggested to make the largest contribution to the mixing of OSR varieties in agricultural fields (Begg *et al.* 2006), data on long-term seed persistence in conventionally tilled fields are rare. Available studies demonstrate that OSR persists for 5–6 or up to 8 years in agricultural fields (Simard *et al.* 2002; Lutman *et al.* 2005; 256



Figure 2. OSR volunteers 25, 26, 28, 30, 36 and 41 tested for *barnase* (left part of the gel) and *barstar* (right part of the gel) **Q2** genes: *barnase* + denotes the *barnase* positive control and *barstar* + the *barstar* positive control. Only volunteers 26 and 30 were positive for *barnase*, but all volunteers (on this gel) were positive for *barstar*.

Gruber *et al.* 2007; Messean *et al.* 2007). Our finding of transgenic volunteers 10 years after cultivation contributes additional evidence that GM OSR can persist for considerable time in agricultural fields. The data appear to be consistent with theoretical Q11 predictions (Lutman *et al.* 2005; Begg *et al.* 2006). Our finding hybridized with each other and with the non-GM OSR plants in the trial plot. It is known that restoration of fertility in F_1 lines can be incomplete, and pollenproducing plants had either both the *barnase* and the *barstar* or only the *barstar* (Bisht *et al.* 2004). This could be why the majority of plants had only the

Seed loss at harvest, shallow cultivation and timing of ploughing have been identified as key factors to prevent Q10 incorporation of seeds into the seedbank (Begg et al. 2006). In the field trial, a protocol based on scientific advice was set up by the Swedish Board of Agriculture to prevent the occurrence of volunteer GM plants, and the trial was meticulously controlled. The shallow stubble tillage performed encourages OSR germination (Pekrun et al. 1998), and late ploughing eliminates seedlings (figure 1). Incorporation of seeds into the seedbank was therefore minimized as much as possible.

In the years after the GM field trial, OSR was not grown at the site and volunteers were controlled with herbicides and subsequent observations, so that sub-stantial seed return did not occur. Although every attempt was made to eliminate volunteers, there is a risk that low levels of seed return were possible due to overlooked volunteers. No other trials with GM OSR have been performed at Lönnstorp farm. The exten-sive control of volunteers makes it possible to conclude that the GM volunteers collected in 2005 most probably were recruited from 10-year old seeds, and provides evidence of long-term GM OSR seed persist-ence in conventional agriculture. In the year of cultiva-tion, the three hybrid lines probably self-pollinated,

hybridized with each other and with the non-GM OSR plants in the trial plot. It is known that restoration of fertility in F₁ lines can be incomplete, and pollenproducing plants had either both the barnase and the could be why the majority of plants had only the barstar gene. As only about one-quarter of the trial area $(336 \text{ m}^2/1200 \text{ m}^2)$ was sown with GM lines, volunteer density in commercial fields would probably be higher than the 0.01 plants m^{-2} reported here. Also, volunteer control in real fields would never be as strict as in this trial. This finding of volunteers, despite labour intensive control for 10 years, supports previous suggestions (Lutman et al. 2005; Begg et al. 2006; Messean et al. 2007) that volunteer OSR needs to be carefully managed in order for non-GM crops to be planted after GM crops.

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- Q2 We have edited the sentence 'Volunteers 26 and 30 were positive...' to 'Only volunteers 26 and 30 were positive...' Please check and confirm our edit.
- Q3 We have made a change to the sentence 'These results are important in relation...' Please review our edit.
- Q4 Both 'genetically modified' and 'Genetic modification' have the same abbreviation 'GM'. Please confirm which one to be followed.
- Q5 Reference Simard et al. (2005) has been cited in text but not provided in the list. Please supply reference details or delete the reference citation from the text.
- Q6 We have made a change to the sentence 'Between 1996 and 2005...' Please review our edit.
- 536 Q7 Please check and confirm the edit of the sentence 537 'The PCR temperature cycle was 95°C per 4 min...'
 - Q8 We have changed 'occurred' to 'germinated' in the sentence 'Seedlings from seven weed species germinated...' Please check and confirm our edit.
 - Q9 Please check and confirm the edit of the sentence 'Available studies demonstrate...'
 - Q10 The term 'seed bank' has been changed to 'seedbank'. Is it ok? Please confirm.
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