Increased Resistance of Bt Aspens to *Phratora vitellinae* (Coleoptera) Leads to Increased Plant Growth under Experimental Conditions

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Abstract

One main aim with genetic modification (GM) of trees is to produce plants that are resistant to various types of pests. The effectiveness of GM-introduced toxins against specific pest species on trees has been shown in the laboratory. However, few attempts have been made to determine if the production of these toxins and reduced herbivory will translate into increased tree productivity. We established an experiment with two lines of potted aspens (*Populus tremula* x *Populus tremuloides*) which express Bt (*Bacillus thuringiensis*) toxins and the isogenic wildtype (Wt) in the lab. The goal was to explore how experimentally controlled levels of a targeted leaf beetle *Phratora vitellinae* (Coleoptera; Chrysomelidae) influenced leaf damage severity, leaf beetle performance and the growth of aspen. Four patterns emerged. Firstly, we found clear evidence that Bt toxins reduce leaf damage. The damage on the Bt line was significantly lower than for the Wt line in high and low herbivory treatment, respectively. Secondly, Bt toxins had a significant negative effect on leaf beetle survival. Thirdly, the significant decrease in height of the Wt line with increasing herbivory and the relative increase in height of one of the Bt lines compared with the Wt line in the presence of herbivores suggest that this also might translate into increased biomass production of Bt trees. This realized benefit was context-dependent and is likely to be manifested only if herbivore pressure is sufficiently high. However, these herbivore induced patterns did not translate into significant affect on biomass, instead one Bt line overall produced less biomass than the Wt. Fourthly, compiled results suggest that the growth reduction in one Bt line as indicated here is likely due to events in the transformation process and that a hypothesized cost of producing Bt toxins is of subordinate significance.

Introduction

The future challenges for forestry are demanding due to changes in climate and intensified land-use [1]. Fossil fuels will need to be replaced with renewable energy sources, which will affect not only agriculture practices, but also silviculture (tree production). Forests can, in theory, become a major source of bioenergy in the future and have the potential to mitigate the anticipated rise in CO2 over the next 50 years. However, this calls for improvements in tree characteristics as well as changes to management practices and technology [1,2].

To facilitate such mitigation, genetic engineering is a useful compliment to other practices as it may partially alleviate some of the constraints on conventional tree breeding. Conventional tree breeding is based on natural variation in economically important traits. Forest tree breeders therefore focus on quantitative traits controlled by several genes [3]. These constraints are associated with the late flowering, slow maturation, long reproductive cycles, and complex mating systems (including self-incompatibility and a high degree of heterozygosity) in trees. Difficulties in identifying the best parents (and controlling their mating), maintaining genetic gain with high heterozygosity [4], and understanding the complex genomes of many tree species causes problems for tree breeders. Genetic modification (GM), on the other hand theoretically allows modification of most individual traits in selected genotypes. Hence, GM technology is much more specific than classical breeding and it can accelerate and allow new strategies for breeding [5].

One main aim with genetic engineering of trees is to produce plants that are resistant to various types of pests [6,7]. Tree pests can severely effect growth and survival of forest trees and thus inflict large economic losses [8]. A warmer climate in the future could increase these problems [8,9]. The most common transformations for pest resistance involve the use of *Bacillus thuringiensis* (Bt) genes, enabling the plant to produce Cry toxins lethal to certain targeted insect pests. However, there are considerable risks for the evolution of pest resistance in wild
populations that needs to be evaluated and minimized \cite{10,11}. The Bt toxin leads to cell damage in the insect mid-gut (for more information see \cite{12}). More than 150 different Cry proteins have been identified \cite{12}, with examples including Cry3Aa proteins targeting coleopteran insects and the cry1 and cry2 families effective against lepidopteran species \cite{13,14}. The effectiveness of these toxins against specific pest species on trees has been shown repeatedly in the laboratory \cite{6,15} and in the field \cite{13}. Still, it is not clear to what degree Bt resistance also will translate into increase tree productivity. Establishing if, and to what degree, plant benefits from the Bt gene with respect to production is essential for cost-benefit analyses of Bt trees, which is the focus of our study.

It has been shown that the production of natural plant defenses are often associated with costs, i.e. there may be a trade-offs between growth and defense \cite{16–20}. It has sometimes proved difficult to demonstrate the costs of defensive compounds and such trade-off might also be transient or context-dependent \cite{21–23}. If such trade-off also should apply to trees producing Bt toxins is not clear at this point but if such costs exist the realized benefits with Bt resistance is likely context-dependent, i.e. influenced by herbivore levels.

We established an experiment with potted Bt-expressing aspens (\textit{Populus tremula}×\textit{P. tremuloides}) in the greenhouse to explore how experimentally controlled levels of a presumably targeted leaf beetle \textit{Phratora vitellinae} (Coleoptera; Chrysomelidae) affected leaf damage severity and performance of the plants. We hypothesized that GM aspens producing Bt toxins should suffer less damage by the leaf beetle than the isogenic wildtype (Wt) and that survival of the leaf beetle, \textit{P. vitellinae}, would be reduced on Bt aspens compared to the wildtype. In line with the above predictions, we further hypothesized that reduced herbivory would translate into increased growth in Bt aspens compared to the wildtype in the presence of the \textit{P. vitellinae}.

Methods

Plant material

We used three isogenic lines of an aspen hybrid (\textit{P. tremula}×\textit{P. tremuloides}) (INRA # 333-38) in which two lines were genetically modified to express Bt toxins, and one was a unmodified line considered a wildtype control (Wt) line. The two genetically modified lines are the Bt17 and Bt27 lines previously described by Genissel et al. \cite{6} and are modified to express a \textit{cry}3Aa Bt- protein targeting Coleopteran species. Bt17 and Bt27 produces toxins in concentrations of approximately 0.05% and 0.0025% of total soluble proteins in the leaves, respectively, and both lines have high resistance to the leaf beetle \textit{Chrysomela tremulae} \cite{6}.

Plantlets of all lines were propagated in the lab and subsequently planted in 3 L pots in commercially available soil in the greenhouse. During the first 10 days of the establishment phase the plants were covered by individual micro-greenhouses using transparent plastic bags. After removal of the micro-greenhouses, the plants were left an additional 14 days before the experiments started. During the experiment the plants received a commercially available NP-fertilizer (Webulls “Rika S”) and water was added to the plants when required.

Experimental design

We used a randomized block design with three plants in each block and a total of 30 blocks. Plants within a block consisted of one individual from each line (Wt, Bt17 and Bt 27) and each block was randomly assigned to the different herbivore treatments (see details below). At the start of the experiment, individual plants (approximately 28.4 cm (SE=0.3) in height at the time), were covered with a tent of fibre cloth. The cloth was commercially available and is used in agricultural practice to mechanically reduce damage by insect pests. Tents were 1.3 m tall to allow maximum tree growth.

Adult \textit{Phratora vitellinae} (Coleoptera; Chrysomelidae) individuals were collected in the field and to minimize the variation in plant responses due to variations in beetle life history state (e.g. sex and age etc.) the beetles were randomly assigned to different plants and density treatments. Furthermore, the beetles were collected from the same site at the same time (i.e. they belonged to the same generation). Thus, although variation in sex and age among the beetles used in the experiments might have resulted in increased variation in damage levels, the randomization of beetles to different treatments and the large number of beetles used should have minimized this influence. This beetle species is a common herbivore on both willow and aspen species \cite{24} and converts salicyl glucosides from the host plant into a larval defensive secretion which consists mainly of salicylaldehyde \cite{25,26}. A related beetle species has been shown to be attracted to highly defended trees, where their sequestration of defenses makes them better defended against predaceous ants \cite{27}. Herbivore treatments consisted of no (0 adults), low (3 adults), and high (7 adults) herbivore loads.

Response measurements

After 5 weeks the experiment was terminated and we counted the number of live adult beetles and larvae (no larvae were introduced to the plants, but some adults were reproducitively successful during our experiment). We also measured leaf damage and height, stem, leaf and root biomass of the trees (see details below). For leaf damage, every leaf was assessed for percent damage using a scale with 5% intervals (i.e. 0 equals 0, 1–5 equals 5%, 6–10 equals 10%, and so forth). In addition, each individual plant was destructively harvested and divided into stem, leaf and root parts. To isolate root material the soil was gently removed with a hand shower. The plant fractions were dried to constant mass in a dryer at 40°C.

Statistical analyses

We used two-way analyses of variance (ANOVA) to determine the effect of herbivore treatment (\(n=3\)), aspen line (\(n=3\)), and their interaction on plant height, dry mass, beetle survival and leaf damage. Plant height at the start of the experiment was used as a covariate in all analyses. When significant effects were shown, subsequent pair-wise comparisons (Tukey’s HSD) were used to identify differences between herbivore treatment or aspen line. In addition, when significant interactions occurred, we examined the effect of each factor at each level of the other factor, using simple contrast which test relationships among cell means \cite{28}. In all of the analyses outlined above, the assumptions of ANOVA were tested with residual plots and in cases of heterogeneous variances the data were log(\(x+1\)) transformed prior to analysis. All statistical analyses were performed in SYSTAT 13 \cite{29}.

Results

Insect survival

Herbivore treatment, aspen line and the interaction between these two factors had a significant effect on the survival of \textit{P. vitellinae} adults (\(F_{3,90}=49.12, P<0.001, F_{3,90}=38.12, P<0.001, F_{3,90}=10.68, P<0.001\), respectively). Further analyses of the interaction term revealed that beetle survival was significantly higher on Wt plants compared to the Bt lines in both the high and
low herbivore treatment ($P<0.001$ in all cases, Fig. 1). Survival was also significantly lower on line Bt27 than on Bt17 in the high herbivore treatment ($P=0.035$). The initial height of the plant at the start of the experiment had no effect on beetle survival ($P=0.941$). Importantly, adult beetles successfully reproduced on Wt plants, but did not reproduce on Bt lines; on average, 3.1 ($\pm 2.5$) live larvae were found on Wt plants in the high herbivore treatment, but, no live larvae were found on any of the Bt lines.

Degree of damage

Leaf damage was significantly influenced by herbivore treatment, aspen line and the interaction between these two factors ($F_{3,80} = 41.57, P<0.001$, $F_{3,80} = 64.945, P<0.001$, $F_{3,80} = 19.911, P<0.001$, respectively). In both the low and high herbivore treatments, the damage was significantly higher on the Wt than on Bt17 or Bt27 lines ($P<0.001$, in all cases, Fig. 1). Leaf damage did not differ significantly between Bt lines in any of the herbivore treatments ($P=0.758$ and $P=0.904$). Finally, leaf damage was unaffected by the initial height of the plant at the start of the experiment ($F_{3,80} = 2.780, P=0.099$).

Plant height

Aspen line, herbivore treatment and their interaction all significantly affected plant height, although the effect of herbivory was marginal ($F_{3,80} = 4.490, P=0.014$, $F_{3,80} = 3.078, P<0.052$, $F_{3,80} = 2.597, P<0.042$, respectively). Within herbivore treatments, trees from the Wt line were significantly taller than the Bt27 line, but not the Bt17 line in the no herbivory treatment ($P=0.008$ and $P=0.216$). In contrast, the Wt line was shorter than the Bt17 line but not the Bt27 line in the high herbivore treatment ($P=0.006$ and $P=0.364$; Fig. 2). No significant differences in plant height were found in the low herbivory treatment or between the Bt lines regardless of herbivore treatment ($P>0.05$ in all cases). Between herbivore treatments, trees from the Wt line were significantly taller in the no herbivory treatment than in the low and high herbivore treatments ($P=0.018$ and $P<0.001$), but Wt plants in low and high herbivore treatments did not differ in height ($P=0.112$). Furthermore, the height of the Bt lines did not differ between herbivore treatments ($P>0.05$ in all cases). Initial height of the plant at the start of the experiment had a significant effect on final height ($P=0.001$), but these differences were used as a covariate in all other analyses and were thus accounted for.

Plant mass

The dry mass of stems and leaves differed significantly among the lines ($F_{3,80} = 4.526, P=0.014$; $F_{3,80} = 8.576, P<0.001$, respectively) but there were no significant herbivory treatment or interaction effects ($F_{3,80} = 2.036, P=0.137$; $F_{3,80} = 0.314$, respectively). Within herbivory treatments, trees from the Wt line were significantly taller than the Bt27 line, but not the Bt17 line in the no herbivory treatment ($P=0.008$ and $P=0.216$). In contrast, the Wt line was shorter than the Bt17 line but not the Bt27 line in the high herbivore treatment ($P=0.006$ and $P=0.364$; Fig. 2). No significant differences in plant height were found in the low herbivory treatment or between the Bt lines regardless of herbivore treatment ($P>0.05$ in all cases). Between herbivore treatments, trees from the Wt line were significantly taller in the no herbivory treatment than in the low and high herbivore treatments ($P=0.018$ and $P<0.001$), but Wt plants in low and high herbivore treatments did not differ in height ($P=0.112$). Furthermore, the height of the Bt lines did not differ between herbivore treatments ($P>0.05$ in all cases). Initial height of the plant at the start of the experiment had a significant effect on final height ($P=0.001$), but these differences were used as a covariate in all other analyses and were thus accounted for.

![Figure 1. Beetle survival and degree of leaf damage.](image1)

The mean number of live Phratora vitellinae adults per plant and the degree of leaf damage on leaves from Wt, Bt17 and Bt27 plants at the end of the trials in the high (initially 7 beetles plant) and low (initially 3 beetles per plant) herbivore density treatments. Bars with different letters indicate significant differences among lines ($P<0.05$). doi:10.1371/journal.pone.0030640.g001

![Figure 2. Changes in plant height and stem mass.](image2)

Mean height and stem mass (and $\pm SE$) of plants from Wt, Bt17 and Bt27 lines at the end of the experiment. Bars with different letters indicate significant differences ($P<0.05$) among lines within the same herbivory treatment and different numbers inside the bars denote significant differences within the same line but between treatments. Please note that the ANOVA analysis revealed no significant interaction between line and herbivory for stem mass. As a result, no pair-wise statistical comparisons were conducted for stem mass and the bars therefore lack letters. doi:10.1371/journal.pone.0030640.g002
In contrast to our second hypothesis, the herbivore inflicted differences in height did not translate into significant differences in dry mass production, although the trend was similar to that for plant height (Fig. 2). This could potentially be due to the very high growth potential of the aspens in the greenhouse environment. The plants increased in height from an average of 28 cm to 97 cm during the 5 weeks of the experiment. Good growing conditions (unlimited water, nutrients, and light), are known to increase the ability of plants to compensate for herbivore damage [31–33]; but see also [34]. The degree of leaf damage was also relatively low with an average of only 3.7% leaf area affected on Wt plants in the high herbivore treatment. This degree of damaged should be compared to estimated levels of insect damages in aspen plantations ranging between 3.8% and ca 50% [35–37] and recent field experiments under semi-natural conditions with the same aspen lines which resulted in ~3.5% leaf damage [38]. Thus, it seems likely that the damage levels, although they did effect height, were too low to have any serious impact on biomass (see also [39]). The plants in our study were also only subjected to herbivory during a relatively short period of time and only during one growing season. Stronger growth responses might have been observed if the plants had been subjected to herbivory for a longer period of time. For example, repeated herbivory is known to reduce the ability of woody plants to compensate for biomass loss due to herbivory [31,33]. Below ground competition can also reduce compensatory ability in plants [40] but our aspens were grown singly in pots and were therefore not affected by belowground competition. Thus, it is likely that increased herbivore density and repeated herbivory, similar to what can be found in commercial aspen plantations, would lead to detectable growth advantage for Bt aspens. The reduction of plant height of Wt plants but not Bt plants with increased herbivory supported our second hypothesis, but we only found a trend and no significant effects on biomass production. On balance we therefore must conclude that our results only provide partial support for our second hypothesis.

In the absence of herbivores, plants from the Bt27 line actually grow less well than Wt plants. This could suggest that there is a cost associated with the production of Bt toxins. However, line Bt27 produced much less Bt toxins than line Bt 17 (approximately 0.0025% and 0.03%, respectively; [6] and plants from the Bt17 line did not show any reduction in growth compared to the wild type. Thus, the reduction in growth is therefore most likely due to events in the transformation process.

It is well known in plant genetic engineering that many events in the transformation process may cause variability in gene expression or gene silencing and have secondary, unintended effects on plant physiology and fitness [41–43]. However, it is currently not possible to determine which of these events that is the most likely cause of the reduced growth of the Bt 27 line or the likelihood that these effects would be manifested under natural growing conditions. In this respect, it is also important to point out that our results should not be considered as representative for Bt plants in general or even all lines of Bt aspens. Our lines were selected due to good performance in the greenhouse but as pointed out above, various factors could influence GM trees physiology and performance. A product-by-product evaluation is always necessary to evaluate both the potential benefits and the potential risks with GM plants [44].

To conclude, in this study we found clear evidence that Bt toxins reduce leaf damage and survival of the target insect herbivore (P. vitellinae). The relative increase in height of the Bt17 line compared with the Wt line in the presence of herbivores suggests that this also might translate into increased growth for Bt

**Discussion**

We found support for our first hypothesis; the Bt aspens negatively affected survival, growth and reproduction of P. vitellinae. Further, the degree of leaf damage inflicted by the leaf beetles was clearly lower on Bt lines than on the Wt line. An earlier study by [6] showed that these trees are highly resistant to the leaf beetle, Crysomela tremulae, and we found a similar pattern for the related species, P. vitellinae. Field experiments with Bt aspens also suggest a high efficiency against target herbivores [13,30]. This is promising as it suggests that the Bt effects are consistent across a wider range of field or greenhouse habitats. In fact, the degree of leaf damage in our experiment was so low on Bt plants in both herbivore treatments that they did not differ significantly from the no herbivore control (P>0.165 in all cases).

Although earlier field experiments and lab experiment do suggest a high efficiency against target herbivores [13,30] these studies did not deal with realized benefits in tree growth. Our greenhouse experiment made it possible to address the question of realized benefits under controlled levels of herbivory and has the advantage that the variation in other confounding factors can be kept to a minimum. In support of our second hypothesis, we found indications that increased herbivore resistance also resulted in growth advantages. Intensified herbivory reduced the relative height of the Wt line compared to one Bt line in the high herbivory treatment. In addition, Wt plants were taller in the no herbivore treatment than in both the herbivore treatments. At the same time we showed that this benefit was context dependent, i.e. depended on the degree of herbivory. We failed to detect any significant differences among aspen lines at the low herbivory treatment and the Bt27 line was shorter than the Wt line in the no herbivore treatment.

In the absence of herbivores, plants from the Bt27 line actually grow less well than Wt plants. This could suggest that there is a cost associated with the production of Bt toxins. However, line Bt27 produced much less Bt toxins than line Bt 17 (approximately 0.0025% and 0.03%, respectively; [6] and plants from the Bt17 line did not show any reduction in growth compared to the wild type. Thus, the reduction in growth is therefore most likely due to events in the transformation process.

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To conclude, in this study we found clear evidence that Bt toxins reduce leaf damage and survival of the target insect herbivore (P. vitellinae). The relative increase in height of the Bt17 line compared with the Wt line in the presence of herbivores suggests that this also might translate into increased growth for Bt
trees if the herbivore pressure is sufficiently high. Although we were unable to detect significant differences, we found a similar trend for stem biomass as for plant height. We detected no growth response corresponding to the concentrations of Bt toxins produced, suggesting that the indicated growth reduction in one Bt line is more likely due to events in the transformation process and that a hypothesized cost of producing Bt toxins is of subordinate significance.

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Author Contributions

Conceived and designed the experiments: JH EPA TGW CJL RJT AW GP. Performed the experiments: JH EPA. Analyzed the data: JH EPA. Contributed reagents/materials/analysis tools: CJL RJT GP. Wrote the paper: JH EPA TGW CJL RJT AW GP.

References

1. Smeets EMW, Faaij APC (2007) Bioenergy potentials from forestry in 2050 - An assessment of the drivers that determine the potentials. Climatic Change 81: 353–390.
2. Fenning TM, Walter C, Garland KMA (2006) Forest biotech and climate change. Nature Biotechnology 24: 613–617.
3. Groover AT (2007) Will genomics guide a greener forest biotech? Trends in Plant Science 12: 234–238.
4. Cheliak WM, Rogers DL (1999) Integrating Biotechnology Into Tree Improvement Programs. Canadian Journal of Forest Research-Review Canadien De Recherche Forestiere 20: 452–463.
5. FAO (2004) Preliminary review of biotechnology in forestry, including genetic modification. Forest Genetic Resources Working Paper FGR/59/E. : Forest Resources Development Service, Forest Resources Division, Rome, Italy.
6. Genisell A, Leple JC, Millot N, Augustin S, Jouanin L, et al. (2003) High tolerance against Chrysomela tremulae of transgenic poplar plants expressing a synthetic cry1Aa gene from Bacillus thuringiensis ssp. thuringiensis. Molecular Breeding 11: 103–110.
7. Pena L, Seguin A (2001) Recent advances in the genetic transformation of trees. Trends in Biotechnology 19: 500–506.
8. Ayres MR, Lombardero MJ (2008) Assessing the consequences of global change for forest disturbance from herbivores and pathogens. Science of the Total Environment 353: 390–398.
9. Bale JS, Masters GJ, Hodkinson ID, Awmack C, Bezemer TM, et al. (2002) Insect herbivores and the role of nectar: a review. Biological Reviews 77: 41–71.
10. Wenes AL, Bourgert D, Andow DA, Courtin C, Carré G, et al. (2006) Frequency and fitness cost of resistance to Bacillus thuringiensis in Chrysomela tremulae (Coleoptera: Chrysomelidae). Heredity 97: 127–134.
11. Genisell A, Augustin S, Courtin C, Pilate G, Lorme P, et al. (2003) Initial frequency of alleles conferring resistance to Bacillus thuringiensis in a field population of Chrysomela tremulae. Proceedings of the Royal Society of London Series B-Biological Sciences 270: 791–797.
12. Schnepp E (1998) Bacillus thuringiensis and its pesticidal crystal proteins. Microbiology and Molecular Biology Reviews 62: 775–790.
13. Ha JH, Tian YC, Han YF, Li L, Zhang BE (2001) Field evaluation of insect-resistant transgenic Populus nigra trees. Euphytica 121: 123–127.
14. Hussein HM, Habustova O, Schnal F (2005) Beetle-specific Bacillus thuringiensis Cry1Aa toxin reduces larval growth and curbs reproduction in Spodoptera littoralis (Boisd.). Pest Management Science 61: 1186–1192.
15. Klein T, Scouras EV, McCown BH, Rau C (2003) Leaf ontogeny influences leaf phenolics and the efficacy of genetically expressed Bacillus thuringiensis cry1A(a) d-endotoxin in hybrid poplar against gypsy moth. Journal of Chemical Ecology 29: 2565–2602.
16. Stevens MT, Waller DM, Lindroth RL (2007) Resistance and tolerance in Populus tremuloides: genetic variation, costs, and environmental dependency. Environmental Evolutionary Biology 21: 829–847.
17. Paul-Victor C, Zut T, Rees M, Kliebenstein DJ, Turnbull LA (2010) A new method for measuring relative growth rate can uncover the costs of defensive compounds in Arabidopsis thaliana. New Phytologist 187: 1102–1111.
18. Coley PD (1986) Costs and benefits of defense by tannins in a neotropical tree. Oecologia 70: 238–241.
19. Simms EL, Rausher MD (1997) Cost and benefits of plant-resistance to herbivory. American Naturalist 130: 570–581.
20. Straus SY, Rudgers JA, Lau JA, Irwin RE (2002) Direct and ecological costs of resistance to herbivory. Trends in Ecology & Evolution 17: 278–285.
21.ｵsaka A, Hohler PG, Fritz RS, Snall T (2010) Growth and chemical defense in willow seedlings: trade-offs are transient. Oecologia 163: 283–290.
22. Bergelson J, Purrington CB (1996) Surveying patterns in the cost of resistance in plants. American Naturalist 148: 536–538.
23. Koricheva J (2000) Meta-analysis of sources of variation in fitness costs of plant-herbivore defences. Biology Letters 3: 176–190.
24. Rowell-Rahier M (1984) The food plant preferences of Phoronia illiniana (Coleoptera: Chrysomelidae). B. A laboratory comparison of geographically isolated populations and experiments on confounding. Oecologia 64: 373–380.
25. Rank NE, Køpf A, Julkunen-Tiitto R, Talvanainen J (1998) Host preference and larval performance of the salicale-feeding leaf beetle Phoronia illiniana. Ecology 79: 618–631.
26. Pasternak JM, Rowell-Rahier M, Braekman JG, Dupont A (1983) Salicin from host plants as a precursor of salicylaldehyde in defensive secretion of Chrysomelinae larvae. Physiological Entomology 8: 307–314.
27. Kearsley MJ, Whitham TG (1992) Guns and butter - a no cost defense against predation for Chrysomela-confluens. Oecologia 92: 556–562.
28. Zar JH (1996) Biostatistical analysis. New Jersey: Prentice Hall, Inc.
29. Systat, Software, Inc (2009) SYSTAT 12: Chicago, USA.
30. Balestrazzi A, Allegro G, Confalonieri M (2006) Genetically modified trees expressing genes for insect pest resistance. Tree Transgenesis: Recent Developments, pp 253–273.
31. Hjälte J, Danell K, Ericson L (1993) Effects of simulated herbivory and intraspecific competition on the compensatory ability of birches. Ecology 74: 1136–1142.
32. Maschinski J, Whitham TG (1989) The continuum of plant responses to herbivory - the influence of plant-association, nutrient availability, and timing. American Naturalist 134: 1–19.
33. Persson IL, Bergstrom R, Danell K (2007) Browse biomass production an growth capacity after transgenic Bt-Aspens and non-transgenic trees responses to moose browsing along a productivity gradient. Oikos 116: 1639–1650.
34. Wise MJ, Abrahamson WG (2007) Effects of resource availability on tolerance of herbivory: A review and assessment of three opposing models. American Naturalist 169: 443–454.
35. Gruppe A, Fasseder M, Schopf R (1999) Short rotation plantations of aspen and balsam poplar on former arable land in Germany: defoliating insects and leaf constituents. Forest Ecology and Management 121: 113–122.
36. Tomescu R, Nef L (2007) Leaf eating insect damage on different poplar clones and sites. Annals of Forest Science 64: 99–108.
37. Coyle DR, McMillin JD, Hall RB, Han ER (2002) Cottonwood leaf beetle (Coleoptera, Chrysomelidae) defoliation impact on Populus growth and above-ground volume in a short-rotation woody crop plant. Agricultural and Forest Entomology 4: 293–300.
38. Axelson EP, Hjälte J, LeRoy CJ, Julkunen-Tiitto R, Wemström A, et al. (2013) Performance of insects on transgenic poplars: (Bt)-expressing aspens under semi-natural field conditions including natural herbivory in Sweden. Forest Ecology and Management 264: 167–171.
39. Antonen S, Pisanen R, Ovaska J, Mustakainen P, Saranpää P, et al. (2002) Effects of defoliation on growth, biomass allocation, and wood properties of Betula pendula clones grown at different nutrient levels. Canadian Journal of Forest Research-Review Canadien De Recherche Forestiere 32: 498–508.
40. Hubbard AP, Hellstrom K, Rautio P, Tsoumi J (2000) A test of the compensatory continuum: fertilization increases and below-ground competition decreases the grazing tolerance of tall weusread mustard (Erysimum straminum). Evolutionary Ecology 14: 353–372.
41. Causer MD, Baxton DR, Vogel KP (2002) Genetic modification of lignin concentration affects fitness of perennial herbaceous plants. Theoretical and Applied Genetics 104: 127–131.
42. Kaldorf M, Flurchg M, Mush HJ, Buscot F (2002) Mycorrhizal colonization of transgenic aspen in a field trial. Planta 214: 653–660.
43. Brodeur-Campbell SE, Vucetich JA, Richter DL, Waite TA, Rosemier JN, et al. (2006) Insect herbivory on low-lignin transgenic aspen. Environmental Entomology 35: 1696–1701.
44. Hjälte J, Lindau A, Wemström A, Blomberg P, Witzell J, et al. (2007) Unintentional changes of defense traits in GM trees can influence plant-herbivore interactions. Basic and Applied Ecology 8: 434–443.