Are traded forest tree seeds a potential source of nonnative pests?

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Abstract. The international seed trade is considered relatively safe from a phytosanitary point of view and is therefore less regulated than trade in other plants for planting. However, the pests carried by traded seeds are not well known. We assessed insects and fungi in 58 traded seed lots of 11 gymnosperm and angiosperm tree species from North America, Europe, and Asia. Insects were detected by X-raying and molecular methods. The fungal community was characterized using high-throughput sequencing (HTS) and by growing fungi on non-selective agar. About 30% of the seed lots contained insect larvae. Gymnosperms contained mostly hymenopteran (Megastigmus spp.) and dipteran (Cecidomyiidae) larvae, while angiosperms contained lepidopteran (Cydia latiferreana) and coleopteran (Curculio spp.) larvae. HTS indicated the presence of fungi in all seed lots and fungi grew on non-selective agar from 96% of the seed lots. Fungal abundance and diversity were much higher than insect diversity and abundance, especially in angiosperm seeds. Almost 50% of all fungal exact sequence variants (ESVs) found in angiosperms were potential pathogens, in comparison with around 30% of potentially pathogenic ESVs found in gymnosperms. The results of this study indicate that seeds may pose a greater risk of pest introduction than previously believed or accounted for. A rapid risk assessment suggests that only a small number of species identified in this study is of phytosanitary concern. However, more research is needed to enable better risk assessment, especially to increase knowledge about the potential for transmission of fungi to seedlings and the host range and impact of identified species.

Key words: alien pests; fungal pathogens; insect pests; pathways risk assessment; pest risk assessment; phytosanitary risk; seed trade.

INTRODUCTION

Nonnative insect pests and fungal pathogens (later referred to as pests; FAO 2016) represent one of the major threats to trees and forest ecosystems worldwide (Boyd et al. 2013). These pests can cause significant ecological changes (Kenis et al. 2009, Loo 2009) and economic losses (Holmes et al. 2009, Aukema et al. 2011) by severely damaging or killing their host plants. For example, the Asian emerald ash borer (Agrilus planipennis Fairmaire) has caused a large-scale death of ash trees (Fraxinus spp.) in North America and is threatening native ash species in Europe (Herms and McCullough 2014, Orlova-Bienkowskaja 2014). Similarly, the fungal pathogen Cryphonectria parasitica (Murrill) Barr virtually eliminated the native American chestnut (Castanea dentata (Marsh.) Borkh.) from North American forests in the early 20th century, which led to significant changes in the forest species composition (e.g., Shackleton et al. 2018).

The number of nonnative forest pests has been increasing rapidly, mainly due to the growing international trade in plants for planting (P4P; Liebhold et al. 2012, Santini et al. 2013, Eschen et al. 2014). The origin of nonnative pests in Europe reflects historical patterns of trade, with North America historically being the most important trading partner (Desprez-Loustau 2009). Imports of P4P from Asia into Europe have been...
increasing recently, for example, imports from China have increased sixfold between 2000 and 2018 and are now equal to imports from North America (Eschen et al. 2014). Almost 50% of the nonnative terrestrial arthropods in Europe are of Asian (26.7%) or North American (21.9%) origin (Roques 2010) and nonnative pathogens in Europe follow a similar trend (35% from North America and 19% from Asia; Desprez-Loustau 2009). The importance of the recent shift in trade partners should not be underestimated because Eastern Asia has high plant diversity (Kreft and Jetz 2007) and exported plants might contain a high number of pests.

Some recent studies suggest that plant diversity is positively related to the diversity of associated insects (O’Brien et al. 2017, Liebhold et al. 2018), which could be due to the larger number of niches occupied by specialist herbivorous insects, which then could increase the diversity on higher trophic levels, such as of predatory insects (i.e., diversity–trophic-structure hypothesis; Hutchinson 1959). The relationship with plant diversity is much less explored for fungi. Fungal communities are normally composed of only small fractions of specialist fungi that show unclear levels of host affinity (Arnold and Lutzoni 2007, Higgins et al. 2007, Vincent et al. 2016). Therefore, a similar relationship between plant and fungal diversity could be expected. Similarly, geographic areas with higher plant diversity or more diverse plant taxa could have a higher associated insect or fungal diversity. This however does not necessarily imply a higher pest or pathogen diversity, since associated organisms may also be commensalists or mutualists (Rodriguez et al. 2009). Yet, the taxonomy of traded plant material may affect pest risk: angiosperms harbor more nonnative pests than gymnosperms (Santini et al. 2013, Eschen et al. 2015) perhaps because they are a more species-rich group. It has previously been suggested that gymnosperms and angiosperms recruit distinctive communities of associated organisms (Ishida et al. 2006, Arnold 2007, Zhou and Dai 2011), but we are unaware of studies of the differences in phytosanitary risk between angiosperms and gymnosperms (i.e., probability of pest transport).

Among the most damaging seed pests of angiosperms are larvae of weevils (Coleoptera: Curculionidae) and moths (Lepidoptera: Tortricidae), which can significantly affect crop yield (Myckzo et al. 2017). However, these larvae usually leave the seeds to pupate in the ground and are, thus, less likely to be moved with seeds. In gymnosperms, the dominant group of spermatophagous insects (i.e., insects that spend the whole life cycle in seeds) are chalcid wasps of the genus *Megastigmus* (Hymenoptera: Chalcidoidea) (Turgeon et al. 1994). Hence, spermatophagous insects are considered to be more relevant from a phytosanitary point of view, also because they are likely to go unnoticed during phytosanitary inspections (Auger-Rozenberg and Roques 2012). The world checklist of fungi includes over 550 seed-borne taxa with most records originating from North America and Europe (Mittal et al. 1990). However, interactions of seed-borne fungi with the host seem to be more complex than the interactions between spermatophagous insects and their hosts, with only a fraction of the fungi being pathogenic (Benitez et al. 2013, Cleary et al. 2019, Liu and He 2019). Moreover, only pathogens that are transmitted to the seedling (vertical transmission) may represent a phytosanitary threat (Burgess and Wingfield 2002).

Plant health regulations and phytosanitary measures are designed to reduce the risk of introducing pests. However, regulatory measures in most countries target only known pest species, which may overlook many potential nonnative pests as most were not known to be harmful prior to their introduction, or were not even known to science (Eschen et al. 2015). Furthermore, due to the increasing volume of international trade in P4P, only a small fraction of traded goods can be inspected (Brasier 2008), often only visually, which is not sufficient to detect latent, rare, or hidden pests (Venette et al. 2002, Slippers and Wingfield 2007). In addition, different types of P4P may vary in the number of potential pests that they carry. As a result, regulations for subcategories of P4P vary significantly, with trade in living plants, with soil, roots, or leaves, being significantly more restricted than trade in seeds (Anonymous 2000). In particular, trade in seeds is rarely considered to be a major pathway of pest introduction (Kenis et al. 2007, Liebhold et al. 2012, Santini et al. 2013) and seeds are, therefore, considered to represent a relatively low phytosanitary risk. This is best illustrated by the fact that among the 233 regulated organisms associated with P4P in the EU, *Fusarium circinatum* Nirenberg & O’Donnell, the causal agent of pitch canker, is the only seed-borne pathogen. This means that the European Union stipulates that only pine (*Pinus* spp.) and Douglas-Fir (*Pseudotsuga menziesii* (Mirb.) Franco) seeds are subjected to specific phytosanitary requirements for import and internal movement (Vetraino et al. 2018). In other parts of the world, the phytosanitary import requirements for seeds are often stricter to specifically prevent the introduction of seed-borne diseases (e.g., Canada, Australia). The value and volume of the international trade in forest tree seeds are very difficult to assess. Since seeds are considered to be part of the P4P pathway they are not listed separately in trade statistics and certainly not at the species level.

We assessed seed-borne insects and fungi associated with commercially traded seeds of 11 tree species. In particular, we explored the differences in overall insect and pathogen infestation and diversity, as well as in infestation by potentially harmful pests, between seeds of different taxonomic groups (angiosperms and gymnosperms) and origins (Europe, Asia, and North America). We tested the following hypotheses: (1) angiosperms have higher insect and fungal diversity than gymnosperms, (2) Asian seeds contain higher insect and fungal diversity than European and North American...
seeds, because Asia is one of the most biodiverse regions of the world, and (3) Europe and North America have more similar seed-borne pest communities, in comparison with Asia, as a result of the historically more intense trade in P4P between Europe and North America.

**METHODS**

**Seed origin and assessment of seed size**

Seeds and nuts (later referred to as seeds) of 11 tree species, belonging to three families within the angiosperms and gymnosperms, and native to Europe, Asia, or North America, were bought from three to eight localities from commercial seed suppliers in Europe and North America in autumn/winter 2016 (Table 1; Appendix S1: Table S1). A total of 58 seed lots (SL), of 100 seeds each, were obtained from the native continent of a tree species from locations with similar, temperate climates (i.e., Western North America, Europe, and Eastern Asia). Seeds of Asian species and of Asian origin were purchased from suppliers in Europe and North America. Hence, these seeds and their associated organisms originated from the native range of their host tree. We were able to obtain detailed information about the origin of most European and North America seeds, but could not obtain this information for the Asian seeds (Appendix S1: Table S1). From the 58 lots, five originated from a region that was slightly outside the native range of the tree species (but still from the continent or origin): two seed lots of *Picea abies* (L.) H. Karst. from Central France and Hungary (SL 24, SL 59), two seed lots of *Pinus sylvestris* L. from United Kingdom (SL 56, SL 57), and one seed lot of *Acer pseudoplatanus* L. from United Kingdom (SL 63; Appendix S1: Table S1).

**Insect and fungal assessments** were done on 53–58 seed lots (Appendix S1: Table S1), on 100 seeds per seed lot.

**Insect assessment**

For 56 out of 58 obtained seed lots, the seeds were X-rayed to reveal the presence of insects (Appendix S1: Table S1), as described by Roques and Skrzypczyńska (2003). Two seed lots of *Fagus sylvatica* L. (SL 72 and SL 73) were not assessed for insects because of late delivery. Infested seeds were dissected with a dissecting knife, and insects were collected and individually stored in collection tubes with 95–99% ethanol. All larvae were examined using a microscope and were identified to order level. All seeds were also visually inspected for exit holes of insects.

Genomic DNA was extracted from individual insect specimens with DNeasy Blood & Tissue Kit (Qiagen, Hilden, Germany) according to the manufacturer’s instructions. A fragment of the mitochondrial gene (mtDNA) cytochrome oxidase c subunit 1 (COI) was subsequently amplified by PCR in 25 μL reaction volumes containing 2 μL of DNA template, 0.2 μmol/L of each of the primers LCO1490 and HCO2198 (Folmer

<table>
<thead>
<tr>
<th>Tree species</th>
<th>Common names</th>
<th>Family</th>
<th>Group</th>
<th>Origin</th>
<th>No. seed lots</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Acer macrophyllum</em> Pursh</td>
<td>bigleaf maple, Oregon maple</td>
<td>Sapindaceae</td>
<td>Angiosperm</td>
<td>North America</td>
<td>6</td>
</tr>
<tr>
<td><em>Quercus garryana</em> Douglas ex Hook</td>
<td>Garry oak, Oregon white oak, Oregon oak</td>
<td>Fagaceae</td>
<td>Angiosperm</td>
<td>North America</td>
<td>6</td>
</tr>
<tr>
<td><em>Tsuga heterophylla</em> (Raf.) Sarg.</td>
<td>western hemlock, western hemlock–spruce</td>
<td>Pinaceae</td>
<td>Gymnosperm</td>
<td>North America</td>
<td>7</td>
</tr>
<tr>
<td><em>Acer pseudoplatanus</em> L.</td>
<td>sycamore</td>
<td>Sapindaceae</td>
<td>Angiosperm</td>
<td>Europe</td>
<td>3</td>
</tr>
<tr>
<td><em>Fagus sylvatica</em> L.</td>
<td>common beech</td>
<td>Fagaceae</td>
<td>Angiosperm</td>
<td>Europe</td>
<td>6</td>
</tr>
<tr>
<td><em>Pinus sylvestris</em> L.</td>
<td>Scots pine</td>
<td>Pinaceae</td>
<td>Gymnosperm</td>
<td>Europe</td>
<td>5</td>
</tr>
<tr>
<td><em>Picea abies</em> (L.) H. Karst.</td>
<td>Norway spruce, European spruce</td>
<td>Pinaceae</td>
<td>Gymnosperm</td>
<td>Europe</td>
<td>5</td>
</tr>
<tr>
<td><em>Acer palmatum</em> Thunb.</td>
<td>palmate maple, Japanese maple, smooth Japanese maple</td>
<td>Sapindaceae</td>
<td>Angiosperm</td>
<td>Asia</td>
<td>5</td>
</tr>
<tr>
<td><em>Pinus tabuliformis</em> Carrière</td>
<td>Manchurian red pine, southern Chinese pine, Chinese red pine</td>
<td>Pinaceae</td>
<td>Gymnosperm</td>
<td>Asia</td>
<td>4</td>
</tr>
<tr>
<td><em>Larix gmelinii</em> (Rupr.) Kuzen.</td>
<td>Dahurian larch</td>
<td>Pinaceae</td>
<td>Gymnosperm</td>
<td>Asia</td>
<td>3</td>
</tr>
</tbody>
</table>

Note: The number of seed lots of each species that were analysed is also indicated.
et al. 1994) and 1 × JumpStart REDTaq ReadyMix Reaction Mix (Sigma-Aldrich, Steinheim, Germany). PCR was performed using a Veriti 96-Well Thermal Cycler (Applied Biosystems, Foster City, California, USA) with the following settings: 2 min at 94°C, five cycles of 30 s at 94°C, 40 s at 54°C, and 1 min at 72°C, 35 cycles of 30 s at 94°C, 50 s at 51°C, and 1 min at 72°C, and a final extension of 10 min at 72°C. Successful amplification was confirmed by electrophoresis of the PCR products in 1.5% (w/v) agarose at 90 V for 30 min with ethidium bromide staining. Sanger sequencing was performed using the same primers and BigDye Terminator v3.1 Cyto Sequencing Kit (Applied Biosystems). Sequences were assembled and edited with CLC DNA Workbench (Qiagen) and compared against reference sequences in the BOLDSYSTEMS (Ratnasingham and Hebert 2007) and National Centre for Biotechnology Information (NCBI) GenBank databases (Geer et al. 2010). The specimens’ queries were matched to its best matching identified reference sequence for identification.

**Fungal assessment by plating**

Seed-borne fungal communities were determined by isolating fungal cultures from 53 out of 58 seed lots (Appendix S1: Table S1). For two seed lots of *Pinus ponderosa* Douglas ex Hook (SL 49, SL 50) and *Tsuga heterophylla* (Raf.) Sarg. (SL 51, SL 66) each and one seed lot of *F. sylvatica* (SL 73), fungal community was not assessed by plating because of late delivery. Seeds were first surface sterilized in 0.5% sodium hypochlorite (NaClO) for 5 min, followed by rinsing in sterile water for 2 × 5 min (Gamboa et al. 2003) to eliminate environmental contamination, and were then air dried in a laminar flow cabinet. The surface sterilized seeds were then cut in half using a dissecting knife. One half of each seed was set aside and kept in a freezer for high-throughput sequencing (HTS), and the other half was put on DNeasy PowerPlant Pro Kit (Qiagen) following the manufacturer’s instructions. This kit was used because it contains Phenolic Separation Solution, which improves the DNA yield from samples rich in phenols, such as some tree seeds. We used 40 μL of the solution for each sample, as recommended in the manufacturer’s instructions. DNA concentrations were quantified using the Qubit dsDNA BR Assay Kit (Thermo Fisher Scientific, Waltham, USA) on a Qubit 3.0 Fluorometer (Thermo Fisher Scientific), and DNA was diluted to 10 ng/μL. Samples that yielded <10 ng/μL were not diluted. The internal transcribed spacer region 2 (ITS2) was amplified with the 5.8S-Fung and ITS4-Fung primers (Taylor et al. 2016). PCR amplifications were performed in 20 μL reaction volumes containing 50 ng of DNA template, 1 mg/mL BSA, 1 mmol/L of MgCl₂, 0.4 μmol/L of each primer, and 0.76 × JumpStart REDTaq ReadyMix Reaction Mix (Sigma-Aldrich). PCR was carried out using Veriti 96-Well Thermal Cycler (Applied Biosystems) with the following settings: 2 min at 94°C; followed by 35 cycles of 30 s at 94°C, 30 s at 55°C, 1 min at 72°C; and a final extension of 10 min at 72°C. Successful amplification was confirmed by agarose gel electrophoresis, and PCR products were sent to Macrogen (Amsterdam, Netherlands) for purification and Sanger sequencing. The resulting sequences were aligned and edited using CLC DNA Workbench (Qiagen). Sequences were taxonomically classified based on the ITS2 region with a naive Bayesian classifier implemented in MOTHUR against the UNITE database (Nilsson et al. 2019) with a minimum bootstrap support of 80%.

**Fungal assessment by high-throughput sequencing (HTS)**

Seed-borne fungal communities of all 58 seed lots were also assessed using an HTS approach (Appendix S1: Table S1). The set-aside halves of the surface sterilized seeds from each seed lot were pooled and ground using a batch mill Tube Mill (IKA-Werke GmbH & Co. KG, Staufen, Germany) or with a pestle and mortar, under liquid nitrogen, if the seeds were small. DNA was extracted from 50 mg of ground seeds from each seed lot using DNeasy PowerPlant Pro Kit (Qiagen) following the manufacturer’s instructions. This kit was used because it contains Phenolic Separation Solution, which improves the DNA yield from samples rich in phenols, such as some tree seeds. We used 40 μL of the solution for each sample, as recommended in the manufacturer’s instructions. DNA concentrations were quantified using the Qubit dsDNA BR Assay Kit (Thermo Fisher Scientific, Waltham, USA) on a Qubit 3.0 Fluorometer (Thermo Fisher Scientific), and DNA was diluted to 10 ng/μL. Samples that yielded <10 ng/μL were not diluted. The internal transcribed spacer region 2 (ITS2) was amplified with the 5.8S-Fung and ITS4-Fung primers (Taylor et al. 2016). PCR amplifications were performed in 20 μL reaction volumes containing 50 ng of DNA template, 1 mg/mL BSA, 1 mmol/L of MgCl₂, 0.4 μmol/L of each primer, and 0.76 × JumpStart REDTaq ReadyMix Reaction Mix (Sigma-Aldrich). PCR was carried out using Veriti 96-Well Thermal Cycler (Applied Biosystems) with the following settings: 2 min at 94°C; followed by 35 cycles of 30 s at 94°C, 30 s at 55°C, 1 min at 72°C; and a final extension of 10 min at 72°C. Each sample was amplified in triplicates and successful PCR amplification confirmed by agarose gel electrophoresis before and after pooling the triplicates. Pooled amplicons were sent to the Génome Québec Innovation Center at McGill University (Montréal, Quebec, Canada) for barcoding using Fluidigm Access Array technology (Fluidigm, South San Francisco, California, USA) and paired-end sequencing on the Illumina MiSeq v3 platform (Illumina, San Diego, California, USA).
Sequence quality filtering, Exact Sequence Variant (ESV) clustering and taxonomic assignments

Sequence quality filtering, ESV clustering and taxonomic assignments were done on a bigger data set that consisted of the sequences obtained from the traded seed material (58 seed lots), and seeds collected in botanical gardens in Europe and North America (85 seed lots; Franić et al., unpublished manuscript), as described below. For the purposes of this manuscript, the final sample x ESV matrix was filtered so it contained only samples (seed lots) and corresponding ESV abundances obtained from traded seed material.

Quality filtering and clustering into ESVs (Callahan et al. 2017) was done with a customized pipeline largely based on UPARSE (Edgar 2013) implemented in USEARCH v.8 (Edgar 2010). Paired-end reads were merged using the USEARCH fastq_mergepairs algorithm (Edgar and Flyvbjerg 2015) with a required overlap of at least 30 base pairs and a minimum length of the merged sequence of 150 bp. PhiX fragments were removed using the nophix function built into USEARCH. PCR primers were detected and removed using Cutadapt (Martin 2011) allowing for one mismatch. Reads not matching the primers were discarded. The USEARCH fastq_filter (Edgar and Flyvbjerg 2015) function was used for quality filtering based on error probabilities, with a maximum expected error threshold of one. Sequences were de-replicated and all reads below an abundance threshold of four and chimeric sequences were discarded using USEARCH fastx_uniques. Sequences were clustered into ESVs using the UNOISE2 algorithm built into USEARCH with an alpha value of 2 (Edgar 2016). The final ESV sequences were checked with ITSx (Bengtsson-Palme et al. 2013), and sequences not verified as ITS2 regions were discarded. The final sample x ESV matrix was obtained by mapping all quality filtered reads against the ESV sequences using the usearch_global algorithm built into USEARCH with the search criteria of maxrejects 0, maxaccepts 0 and top_hit_tonly. Taxonomical classification of ESVs was performed using the naïve Bayesian classifier (Wang et al. 2007) implemented in MOthur (Schloss et al. 2009), with a minimum bootstrap support of 80%. ESV sequences were first compared against a customized ITS2 reference database collected from NCBI GenBank to exclude non-fungal sequences. Taxonomic affiliation of the fungal sequences was then obtained by querying against the UNITE database (Nilsson et al. 2019).

Comparison of plating and HTS

To determine if the fungal community detected by plating represents a subset of the fungal community captured by HTS, we mapped the ITS2 region of sequenced fungal cultures against the ITS2 region of the ESV sequences using the usearch_global algorithm built into USEARCH.

Identification of potential fungal pathogens

Parsing of fungal ESVs into guilds was done based on their taxonomic assignments by using the FUNGuild tool (Nguyen et al. 2016). FunGuild consists of a community-annotated data base and a bioinformatic script that assigns taxonomic names to functional guilds. Since we aimed at detecting all plant pathogens, the output table (Appendix S2: Table S1) was filtered by selecting all combinations of guilds in which plant pathogens were mentioned. For all ESVs that could be identified to species and that were recognized as potential plant pathogens, a literature search was done for reports or studies on their pathogenicity, known woody plant hosts and distribution range. For this, the Fungal Databases, U.S. National Fungus Collections, ARS, USDA (Farr and Rossman 2019), were used. Fungal cultures were parsed into guilds based on the guild of the fungal ESV they mapped with.

Statistical analyses

All statistical tests were performed using R (R Core Team 2018) and were considered significant at \( P < 0.05 \). Seed lots were the unit of analysis for all statistics. Comparisons were made between different groups (angiosperms and gymnosperms) and origins (Europe, Asia, and North America) of seed lots. The means and standard errors for the number of seeds infested with fungi per seed lot, and fungal infestation and diversity at the level of tree species are given in Appendix S3: Table S1.

We tested the effect of seed origin or taxonomic group on infestation rates by insects using Hurdle models (glmTMB function from the glmmTMB package (Brooks et al. 2017)), because the data were zero inflated. We tested for differences in overall infestation (seeds with larvae and/or exit holes), as well as in presence and number of larvae and exit holes separately. Binomial generalized linear mixed models were first used to analyze binary responses (presence or absence of larvae and exit holes). Seed lots of Asian origin were excluded from the analysis of exit holes because they did not show any exit holes. Truncated Poisson generalized linear mixed models were then fitted to the number of larvae and exit holes for those seed lots with insects or exit holes. In all models, taxonomic group (angiosperms and gymnosperms) and origin (Asia, Europe, North America) were fixed effects and tree species a random effect. Interactions were not included in these analyses.

The differences in the number of fungal cultures, morphotypes and ESVs among seed lots were analysed with the same fixed and random effect variables as mentioned above, but including interactions. We assumed a negative binomial distribution for the errors of these response variables, because Poisson models were overdispersed. The differences in percentages of potentially pathogenic ESVs and ESVs not identified to genus or species, as a
percentage of total ESVs per seed lot, were tested with models containing the same fixed and random effects, including interactions. For these models, we assumed a beta-binomial distribution for the errors because binomial models were overdispersed. A Tukey comparison of least-squares means was performed with the function lsmeans from the lsmeans package (Lenth 2016). The same function was used to calculate mean values and standard errors for fixed effects in the models.

To test the differences between fungal communities associated with seeds of different taxonomical groups and origins multivariate generalized linear models (multivariate GLMs) were fitted to the ESV abundance matrix. This approach was used because, unlike distance-based approaches such as PERMANOVA, multivariate GLMs account for the mean-variance relationship (Warton et al. 2012). The matrix was previously standardized across samples by randomly subsampling each sample to the lowest number of read counts obtained by any sample (here 7,038 read counts) to account for different sequencing depth across samples. Generalized linear models were fitted with the function manymglm from the mvabund package (Wang et al. 2012). Negative binomial distribution of sequence counts was assumed.

The differences in seed size among different groups and origins were assessed with the function lmer from the lme4 package (Bates et al. 2015) with log-transformed seed size as the response variable and the same fixed and random effect variable as above, including interactions. Angiosperm seeds were significantly larger than gymnosperm seeds (43.51 ± 20.72 and 10.13 ± 3.54 mm², respectively [mean ± SE]; $\chi^2 = 5.78$, df = 1, $P = 0.016$) but no differences in seed size were found among the different origins ($\chi^2 = 0.43$, df = 2, $P = 0.808$). The relationship between the average size of the seeds in a seed lot and the number of fungal cultures per seed lot, number of morphotypes per seed lot and the number of fungal ESVs per seed lot and was tested using glmmTMB, again with tree species as a random effect. A negative binomial distribution for the errors was assumed because Poisson models were overdispersed.

The relationship between the number of fungal morphotypes and ESVs per seed lot was tested using Spearman correlations. All seed lots in which fungi were assessed with both methods, plating and HTC (Appendix S1: Table S1), were included in this analysis.

### RESULTS

#### Insect assessment

**Infestation.**—Around 50% of the seed lots were infested, i.e., they contained larvae or had exit holes, but no differences in the infestation rate were found among seed lots of different groups or origins (50.58 ± 14.86% [overall mean ± SE]; Appendix S4: Table S1).

Nearly 30% of the seed lots contained insect larvae, with a total of 79 individuals detected. Insects were found in seed lots of only four tree species: *Larix gmelinii* (Rupr.) Kuzen., *P. abies*, *Q. garryana*, and *T. heterophylla* (Table 2). The number of larvae per seed lot corresponded to the number of infested seeds per seed lot (i.e., one larva per infested seed), except for *Q. garryana* (5.5 ± 0.9 infested seeds per seed lot) where sometimes more than one larva was found per seed. No differences in the presence or number of larvae were detected between angiosperm and gymnosperm seed lots, or between seeds of different origins (Appendix S4: Table S1).

Exit holes were recorded in around 30% of seed lots. A higher percentage of angiosperm than gymnosperm seed lots contained exit holes (73.06% ± 10.46% and 11.75% ± 6.43%, respectively; Appendix S4: Table S1) but the origins did not differ in the percentage of seed lots that contained exit holes (Appendix S4: Table S1). The number of exit holes per seed lot did not depend on taxonomic group or origin (Appendix S4: Table S1).

**Diversity.**—The 79 insect larvae belonged to four orders, namely Hymenoptera (48.1%), Lepidoptera (31.6%), Coleoptera (15.2%), and Diptera (2.5%), with two specimens (2.5%) unidentified. Hymenopteran and dipteran larvae were found only in gymnosperms, and lepidopteran and coleopteran larvae only in angiosperm seeds. Molecular analyses allowed the identification of 45 larvae to species, with a match of at least 99% in the barcode database (Table 3). Additionally, 12 specimens were identified to genus and two to the family level (Table 3). Two specimens could not be classified based on morphological features or by sequencing.

**Fungal assessment by plating**

**Infestation.**—Plating yielded fungal cultures from around 96% of the seed lots. No fungi developed on agar from

<table>
<thead>
<tr>
<th>Tree species</th>
<th>Origin</th>
<th>Group</th>
<th>SL with larvae (%)</th>
<th>Larvae density (no. larvae/SL)</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Larix gmelinii</em></td>
<td>Asia</td>
<td>Gymnosperm</td>
<td>100.0 ± 0.0</td>
<td>1.7 ± 0.3</td>
</tr>
<tr>
<td><em>Picea abies</em></td>
<td>Europe</td>
<td>Gymnosperm</td>
<td>80.0 ± 20.0</td>
<td>5.8 ± 2.3</td>
</tr>
<tr>
<td><em>Quercus garryana</em></td>
<td>North America</td>
<td>Angiosperm</td>
<td>83.3 ± 16.7</td>
<td>6.2 ± 2.4</td>
</tr>
<tr>
<td><em>Tsuga heterophylla</em></td>
<td>North America</td>
<td>Gymnosperm</td>
<td>57.1 ± 20.2</td>
<td>1.1 ± 0.6</td>
</tr>
</tbody>
</table>
two European seed lots (P. abies SL 24 and P. sylvestris SL 22). The number of fungal cultures varied with both origin and taxonomic group: overall, angiosperms yielded more fungal cultures per seed lot than gymnosperms and European seed lots had the most and North American seed lots the fewest fungal cultures. Differences between taxonomic groups also varied with origin: European angiosperms yielded more than 10 times as many fungal cultures as European gymnosperms (Appendix S4: Table S2), while Asian and North American angiosperms only had two to three times more fungal cultures than gymnosperms from the same continent (Fig. 1; Appendix S4: Table S2). The number of fungal cultures per seed lot was not affected by the size of the seeds ($\chi^2 = 0.36$, df = 1, $P = 0.548$) and this was consistent among angiosperms and gymnosperms (Group $\times$ Seed size: $\chi^2 = 1.75$, df = 1, $P = 0.185$).

**Diversity.**—A total of 4,202 fungal cultures were obtained from plated seeds. Overall, angiosperms had more fungal morphotypes per seed lot than gymnosperms. The origin of the seeds also had a significant effect, with European seed lots having the largest and North American seed lots having the lowest number of fungal morphotypes per seed lot. The number of fungal morphotypes per group also varied with origin: angiosperms had more morphotypes than gymnosperms in Europe but not in North America or Asia (Fig. 2; Appendix S4: Table S2). The number of fungal morphotypes per seed lot was not affected by the size of the seeds ($\chi^2 = 0.09$, df = 1, $P = 0.768$) and this was consistent among angiosperms and gymnosperms (Group $\times$ Seed size: $\chi^2 = 1.26$, df = 1, $P = 0.262$).

A total of 473 (11.3%) fungal cultures out of 4,202 recovered from seeds were sequenced. Good quality sequences were obtained from 441 cultures (93%). These sequences belonged to the classes Ascomycota (96%), Mucoromycota (2.5%), and Basidiomycota (1.5%). A total of 295 (66.9%) sequences were identified to one of 42 identified genera, and 142 (32.2%) to one of 53 identified species. In around 50% of the cases where two to five cultures of the same morphotype were sequenced, all sequences were mapped to the same ESV, around 70% were identified as the same genus and 75% to the same family.

**Potential fungal pathogens.**—A total of 107 out of 473 sequenced fungal cultures were assigned to one of 21 potentially plant pathogenic fungal genera, and of those, 51 fungal cultures to 18 potentially plant pathogenic fungal species (Table 4). Among these, two potentially plant pathogenic fungal species known to be associated with woody plants and absent from or with a limited distribution in Europe were identified, namely Diaporthe...
alleghaniensis R.H. Arnold (L. gmelinii SL 28) and Microascus cirrosus Curzi (P. abies SL 61).

Fungal assessment by high-throughput sequencing (HTS)

Infestation and diversity.—All 58 seed lots analysed using HTS were infected by fungi. The 2,376,400 high quality sequences (40,972 reads per seed lot) clustered into 1,905 Exact Sequence Variants (ESVs) (119.9 ESVs per seed lot). Overall, angiosperm seed lots had more fungal ESVs than gymnosperm seed lots and Asian seed lots had more fungal ESVs than seed lots from Europe and North America (Fig. 3; Appendix S4: Table S2). The number of fungal ESVs per seed lot increased with seed size ($\chi^2 = 3.93, df = 1, P = 0.047$), and angiosperms had larger seeds and more fungal ESVs than gymnosperms ($\chi^2 = 35.48, df = 1, P < 0.001$).

Around 72% (1,365 out of 1,905) of the ESVs were assigned to the Ascomycota, 18% (340) to the Basidiomycota, <1% (12) to the Mucoromycota, and 9% (188) ESVs remained unclassified. A total of 1,143 (60%) and 559 (30%) ESVs were assigned to 234 genera and 269 species, respectively.

Overall, gymnosperms, especially those from Europe, had a higher percentage of ESVs not identified to the genus level than angiosperms (44.15 ± 1.89 and 30.74 ± 1.84, respectively) (mean ± SE; Appendix S4: Table S2). However, the percentage of ESVs not identified to the species level was similar among groups and origins (Fig. 4; Appendix S4: Table S2).

Fungal communities differed significantly among seed lots of different taxonomic groups (df = 1, deviance = 5921, $P = 0.001$) and seed origins (df = 2,
deviance = 7216, P = 0.001) A total of 1,099 out of 1,905 ESVs (58%) were found exclusively in angiosperm seeds and 444 (23%) only in gymnosperm seeds. Only 362 (19%) ESVs were found to be associated with seeds of both groups. Around 65% of ESVs were unique to one continent and 12% of the ESVs were found in seed lots from all three continents. Europe and North America shared more than two times as many ESVs (14%) than either continent shared with Asia (4.8% and 4.3%, respectively).

Potential fungal pathogens.—A total of 532 out of 1,905 ESVs (28%) were not assigned to a guild, and 630 out of 1,905 ESVs were classified as potentially pathogenic (33%). A total of 508 out of 1,905 ESVs were assigned to one of 86 potentially plant pathogenic fungal genera and of those, 182 ESVs to 100 potentially plant pathogenic fungal species. No differences in the relative occurrence of potentially plant pathogenic ESVs were found among continents (37.67% ± 2.06%; Appendix S4: Table S2), but angiosperms had a higher percentage of potentially plant pathogenic ESVs than gymnosperms (48.53 ± 3.28 and 26.81 ± 2.50, respectively; Appendix S4: Table S2). Eight and three of the potentially pathogenic fungal species, which are known to be associated with woody plants, are absent or have a restricted distribution in Europe, respectively (Table 5).

Comparison of fungal assessment by plating and HTS

The number of fungal morphotypes per seed lot was positively correlated with the number of fungal ESVs per seed lot (r = 0.59, S = 10170, P < 0.001). All but one of the 441 sequences of good quality from fungal cultures mapped against 126 ESV centroids from HTS. A total of 350 out of 441 sequences had an identical match to ESV sequences, and only 15 had a match of <80%. Around 6% of fungal diversity assessed by HTS (1,905 ESVs) was captured by plating (126 ESVs).

Similar fractions of pathogenic ESVs were obtained by plating (59 out of 126 ESVs, 47%) and using HTS (630 out of 1,905 ESVs, 33%). Ten fungal species that are known to be associated with woody plant hosts and are absent or have limited distribution in Europe were identified by HTS only (Table 5), D. alleghaniensis was identified by HTS and plating and M. cirrosus was identified only by plating.

DISCUSSION

The trade in seeds of most tree species is not regulated because seeds are considered less likely to carry harmful organisms than other plants for planting (P4P). However, our results demonstrate that seeds contain many insects and fungi. The high infestation of commercially supplied seed lots is a clear indication of how often potential pests may be transported via traded seeds. Furthermore, the presence of organisms that are already known to be pests in these seeds indicates that measures should be taken to reduce pest risk.

Risk of pest introduction

The absence of significant differences in insect infestation levels between taxonomic groups and origins of the seeds suggests that movement of potential pests occurs in all directions and in both angiosperms and gymnosperms. Since all insect species identified in this study are highly host specific, the higher percentage of seed
Table 5. Fungal species known to be potential pathogens of woody plants, found in traded seed lots from different continents.

<table>
<thead>
<tr>
<th>Plant pathogenic species</th>
<th>Corresponding ESV</th>
<th>Origin of seed lots</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Absent from Europe</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Alternaria rosae</em> E.G. Simmons &amp; C.F. Hill</td>
<td>ESV0248</td>
<td><strong>Europe</strong> 57, 73, 11, 12, 50</td>
</tr>
<tr>
<td><em>Colletotrichum gigasporum</em> Rakotoni. &amp; Munaut</td>
<td>ESV2475</td>
<td><strong>North America</strong> 29, 75, 77</td>
</tr>
<tr>
<td><em>Diaporthe australafriicana</em> Crous &amp; Van Nieker</td>
<td>ESV0675</td>
<td><strong>Asia</strong> 57, 78, 5, 7, 10, 11, 14,15</td>
</tr>
<tr>
<td><em>Diaporthe siamensis</em> Udayanga, X. Z. Liu &amp; K.D. Hyde</td>
<td>ESV1811</td>
<td>77</td>
</tr>
<tr>
<td><em>Lasiodiplodia crassipora</em> T. Burgess &amp; Barber</td>
<td>ESV0546</td>
<td>27, 29, 74, 75, 76</td>
</tr>
<tr>
<td><em>Phaeoacremonium roseum</em> J.R. Urbez-Torres, P. Haag &amp; D.T. O’Gorman</td>
<td>ESV0482</td>
<td>16, 17, 20, 77</td>
</tr>
<tr>
<td><em>Pseudocercospora spaeroellae-eugeniae</em> (Rangel) Crous, Alfenas &amp; R.W. Barreto</td>
<td>ESV0479</td>
<td>75, 77</td>
</tr>
<tr>
<td><em>Eurytoma laricis</em> Yano, a common Eurasian larch seed chalcid</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Limited distribution in Europe</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Diplodia corticola</em> A.J.L. Phillips, A. Alves &amp; J. Luque</td>
<td>ESV2722</td>
<td><strong>Europe</strong> 19</td>
</tr>
<tr>
<td><em>Neonectria major</em> (Wollenw.) Castl. &amp; Rossman</td>
<td>ESV0139</td>
<td><strong>North America</strong> 63</td>
</tr>
<tr>
<td><em>Setomelanomma holmii</em> M. Morelet</td>
<td>ESV0362</td>
<td><strong>Asia</strong> 71, 73, 30</td>
</tr>
<tr>
<td><em>Diaporthe alleghaniensis</em> R.H. Arnold</td>
<td>ESV1258</td>
<td><strong>Europe</strong> 73</td>
</tr>
<tr>
<td><em>Phaeoacremonium roseum</em> J.R. Urbez-Torres, P. Haag &amp; D.T. O’Gorman</td>
<td>ESV2757</td>
<td><strong>North America</strong> 28, 30, 68, 74</td>
</tr>
</tbody>
</table>

Notes: New records for species-continent combinations are identified by boldface type. Numbers indicate the seed lots (Appendix S1: Table S1) in which the species were found. ESV, exact sequence variant.

Lots showing exit holes in angiosperm than gymnosperm seeds may be due to the biology of the insects. Specifically, larvae of *C. latiferreana* and of *Curculio* sp. were still found in *Q. garryana* acorns, whereas seeds of the other angiosperms only had exit holes. This indicates that larvae had been in the seeds, but left them to pupate before the seeds were traded. On the other hand, in gymnosperm seeds we detected mainly hymenopteran seed pests (*Megastigmus* species), which are known to spend their entire life cycle in the seeds and cannot be detected by external observation (Roques et al. 2003). These gymnosperm pests are therefore of greater phytosanitary risk than angiosperm insects, which usually leave the seeds to pupate in the soil.

Three insect species of phytosanitary concern have been identified in this study. The chalcid wasp *Megastigmus strobilobius* Ratzeburg is a native pest of northern European *Picea* species (*P. abies* and *Picea obovata* Ledeb.), but has now colonized plantations of its native hosts from Western Europe to the Lake Baikal (Roques and Skrzypczyńska 2003). Similarly, *Megastigmus tsugae* Crosby is known to infest seeds of *T. heterophylla* and *Tsuga mertensiana* (Bong.) Carr. in the North American range of these hemlocks, but has never been recorded in native or planted hemlocks in Asia or Europe (Roques and Skrzypczyńska 2003, Roques et al. 2003). The chalcid wasp *Megastigmus pictus* Förster is a specialist of the Eurasian larch species (*Larix* spp.) with a wide distribution in Europe and Asia (Roques and Skrzypczyńska 2003) but had not previously been documented in North America. This is probably also the case for the unidentified *Eurytoma* sp. found in Chinese seeds of *L. gmelinii*, the sequence of which is similar to *Eurytoma laricis* Yano, a common Eurasian larch seed chalcid. A specimen of the American *M. tsugae* was found in *Tsuga* seeds that were purchased from European suppliers, and a specimen each of the Eurasian *M. pictus* and *E. laricis* were found in *L. gmelinii* seeds purchased in North America. These results therefore provide evidence for movement of potential pests via the seed trade to the areas where they were previously not recorded.

Fungi were more frequent in seed lots than insects were, and more frequent in seeds of a seed lot than insects were (Appendix S3: Table S1), which indicates that the likelihood of introducing fungi is higher than the likelihood of introducing insects. However, the likelihood that an introduced insect is a pest appears higher than the likelihood of an introduced fungus being a pathogen. This is due to the majority of seed-borne fungi being commensal or beneficial species, while the seed-borne insects are all specialized herbivores. Moreover, while all insects that we extracted from the seeds were alive, we can only be sure that the fungi plated on agar were living. Since DNA is detectable for several weeks to months after the organisms’ death (Schena et al. 2004), we cannot be sure that Exact Sequence Variants (ESVs) detected by high-throughput sequencing (HTS) originated from living structures (e.g., spores, mycelia) and detection of DNA without evidence of viability of the pest would be insufficient reason to adopt specific restrictive measures in the phytosanitary context of international trade (Griffin 2012). Fungal cultures of *M. cirrosus* and *D. alleghaniensis*, which were isolated and
grown on agar, are therefore the only two fungal pathogens of woody plants that were certainly alive. HTS allowed the characterization of highly diverse fungal communities, including rare, slow growing, or unculturable fungi (e.g., biotrophs) and thus represents a step forward from traditional plating. However, only a small percentage of fungal ESVs could be identified to the species level due to incomplete reference sequences limiting the use of this method for phytosanitary purposes (Tedersoo et al. 2019). For illustration, in our study nearly 70% of all ESVs remained unidentified to the species level. Since pest risk assessment is usually done at the species level, these ESVs could not be used to assess phytosanitary risk.

Large differences in fungal diversity among seeds of different taxonomic groups and origins were found using HTS and plating. The hypothesis that angiosperm seeds harbor a higher fungal diversity than gymnosperm seeds was confirmed, both by plating and HTS analyses. The larger size of angiosperm seeds compared to gymnosperm seeds may account for this difference, as suggested by significant relationship between seed size and fungal diversity as assessed by HTS. The high variability of the number of fungal cultures and morphotypes and seed size among and within angiosperm and gymnosperm species is the most probable reason why we found no effect of seed size on the number of fungal cultures and morphotypes. The difference in fungal diversity might also be due to the fact that angiosperms are taxonomically more diverse than gymnosperms and thus could support a higher diversity of associated organisms. The use of HTS might have underestimated the diversity in angiosperm seeds, since a smaller fraction of the ground and mixed material was used for the assessment than for the smaller gymnosperm seeds, where sometimes all ground material was used. The differences in fungal diversity among angiosperms and gymnosperms could also be due to the different plant resistance strategies (Eyles et al. 2010, Zalamea et al. 2018), which still needs to be explored.

Angiosperm seeds harbored more unique and more potentially plant pathogenic ESVs than gymnosperm seeds, which indicates a relatively higher phytosanitary risk associated with their trade. This finding is supported by previous studies, where 58% of invasive forest pathogens in Europe were shown to be specialized on angiosperms and only 26% on gymnosperms (Santini et al. 2013). The relatively higher fungal diversity, as assessed by HTS, in Asian seeds than in North American and European seeds indicates potentially higher phytosanitary risk associated with them. The same pattern of higher biodiversity in Eastern Asia than in Europe and North America has already been shown for many other groups, such as plants (Kreft and Jetz 2007) and soil fungi (Tedersoo et al. 2014). The largest fraction of fungal ESVs was shared between European and North American seeds, which can be explained by the more intense history of trade in P4P between Europe and North America. Hence, the European invasion debt for Asian species seems to be larger than for North American species (Rouget et al. 2016). Consequently, we expect that new nonnative fungi will be imported to Europe and North America from Asia.

**Risk of pest establishment**

Only a small fraction of the insects and fungi detected in the seeds are likely to become established in the region of introduction. Successful establishment requires that the insect or fungus exits the seed at its destination and infects a new, susceptible host. Moreover, specific biological characteristics make some species better invaders than others. For instance, some species of chalcid wasps, such as *M. pictus*, found in *Larix* seeds from China, show thelytokous parthenogenesis and a single female may be enough to establish a new population (Boivin et al. 2014). In addition, many *Megastigmus* spp. have the possibility to extend their state of diapause to overcome periods of adverse conditions (Turgeon et al. 1994). *Megastigmus* chalcids are usually oligophagous but strictly related to a tree genus (Roques et al. 2003), however, suitable hosts for all identified *Megastigmus* species are probably available in the three regions. For example, *M. strobilobius* has been found to parasitize seeds of a number of Asian *Picea* species and North American *Picea pungens* Engelm. in arboreta in France and Denmark (da Ros et al., 1993; Roques and Skrzypczyńska 2003), which indicates that it might switch to new hosts in Asia and North America if introduced to these regions. However, *M. strobilobius* as well as *M. tsuga* have an arrhenotokous parthenogenesis, where unmated females only produce males (Boivin et al. 2014), which limits its invasive potential.

Seed borne pests of nut-bearing angiosperms leave the nuts in autumn to pupate in the soil (Roques et al. 2003). If larvae are present in traded seeds, appropriate conditions for pupation may be lacking. Moreover, the small number of insects per seed lot, and variation in their time of emergence, may limit the chance that emerging adults find a mate and can establish a population (Contarini et al. 2009).

While a wide variety of fungi was detected and fungi were found in almost every seed lot, it is unknown whether these seed-borne organisms are seed transmitted. In general, vertical transmission of fungi in trees seems to be rare compared with seed infestation from the surrounding environment (horizontal transmission; Arnold 2007). However, for some seed-borne fungi, such as *Siroccoccus conigenus* (Pers.) P.F. Cannon & Minter 1983, a pathogen that causes shoot blight in numerous conifer species in North America and Europe, and the above-mentioned *F. circinatum*, vertical transmission was confirmed (Sutherland et al. 1981, Storer et al. 1998, Evira-Recuenco et al. 2015). Thus, further studies would be needed to assess the potential for the detected fungi to be vertically transmitted and spread to other plants.
The presence of a susceptible host is also essential for successful establishment of fungal species. The probability of finding a host in the introduced region is likely to decrease with increasing phylogenetic distance between the host and the native species (Gilbert and Webb 2007, Gilbert et al. 2012). Tree species of the same genera on which some of the currently absent pathogens were found also occur in Europe, such as *Rosa* (Alternaria rosae E.G. Simmons & C.F. Hill), *Acacia* and *Hibiscus* (Colletotrichum gigasporum Rakotomin. & Munaut), *Corylus*, *Prunus*, and *Salix* (Diaporthe australafriicana Crous & Van Niekerk), and *Pistacia* (*M. cirrosus*), which makes establishment of these pathogens possible. Some of the other pathogen species not yet present in Europe are mostly associated with hosts within tropical tree genera and/or genera from the southern hemisphere (Farr and Rossman 2019) and may be less likely to establish in Europe. However, data about host range and ecology are unavailable for many fungal species, which makes it difficult to assess their likelihood of establishment, all the more since host shifts remain difficult to predict (Gilbert and Webb 2007).

**Risk of post-introduction damage**

Among the identified insect species, only the filbert-worm (*Cydia latiferreana* Walsingham) is known to cause significant damage in its current area of distribution. This moth is a major pest in commercial hazelnut (*Corylus* spp.) production in the northwestern United States, which relies mainly on cultivars based on European hazel (Chambers et al. 2011). Therefore, its introduction and establishment in Europe could result in serious damage of hazels. By contrast, the *Megastigmus* and *Eurytoma* infestation rates in the native range of spruce, larch and hemlock are low (Roques and Skrzypczyńska 1987, de Groot et al. 1994, Skrzypczyńska 2013) and the three identified *Megastigmus* species do not represent major pests of these host species. A low impact on native hosts is typical for pests in their native range (Keane and Crawley 2002). However, if an exotic pest is introduced together with its native host to an area where no congeneric hosts are present, considerable damage can occur because of the absence of natural enemies and competitors (Auger-Rozenberg and Roques 2012). For instance, the Douglas-fir seed chalcid, *Megastigmus spermotrophus* Wachtl, has become a major seed pest in European Douglas-fir seed orchards and plantations, whereas in its native North American range its impact is limited by its insect competitors that attack seeds earlier (Rappaport and Roques 1991). In a similar scenario, establishment of *M. tsugae* on *T. heterophylla* in Europe might considerably affect the regeneration of *T. heterophylla*.

Some of the fungal pathogens identified in this study can affect economically important woody plants occurring in Europe, and are thus of special phytosanitary concern. For example, *Lasiodiplodia crassispora* T. Burgess & Barber, which was found in seeds of *Pinus tabuliformis* Carrière and *Acer palmatum* Thunb. from China, causes stem necrosis and cankers in grapevine (*Vitis vinifera* L.) in South Africa and California (Urbez-Torres et al. 2010, van Niekerk et al. 2010). Fungus *D. australafriicana*, which was found in seeds of North American *Pinus, Tsuga*, and *Acer* species, is associated with stem cankers of blueberries (*Vaccinium* spp.) and European hazelnut (*Corylus avellane* L.) in Chile (Latorre et al. 2012, Elfar et al. 2013, Guererro and Pérez 2013). *Diplodia corticola* A.J.L. Phillips, A. Alves & J. Luque, identified in seeds of North American *Quercus* and Asian *Acer* species, is responsible for cankers and dieback in oak species in North America, Northern Africa, and Southern Europe (Farr and Rossmann 2019). Several of the fungal species detected are thought to be pathogens but little is known about their impact.

Current European regulations target known pest species that are absent from, or have a restricted distribution in, Europe. Yet, repeated introductions of relatively widespread species or genera can result in the formation and introduction of superior invaders (Lombaert et al. 2010, Dutech et al. 2012). Several fungal cultures of potentially pathogenic genera (e.g., *Botryosphaeria*, *Cytospora*, *Diaporthe*, *Fusarium*) and species that are already present in Europe (e.g., *Cytospora ribis* Ehrenb., *Neofusicocccum parvum* (Pennycook & Samuels) Crous, Slippers & A.J.L. Phillips, *Diaporthe cotoneastri* (Punith.) Udayanga, PW Crous and KD Hyde, *Diplodia seriata* De Not., *Petalopsis* spp.) were also recovered from non-European seeds. Similarly, a specimen of chalcid wasp *M. pictus*, which is known to be present in Europe, was detected in one Chinese seed lot that was imported to Europe. Admixture of naturally isolated populations of the same species can increase fitness and subsequently lead to larger and more productive populations with lower probabilities of extinction, and ultimately increase the chance of range expansion and impact (Turgeon et al. 2011, Riis and Darling 2014). Similarly, hybridization between congeneric species that are naturally isolated, might lead to formation of new, more harmful hybrids (Giordano et al. 2019) or hybrids with new or combined features obtained from their parental species (Brasier 2001). Hence, additional intra- and intrageneric introductions should also be avoided.

**Conclusions**

The results of this study, which is, to our knowledge, the first that simultaneously assessed insects and fungi associated with any type of traded PAP, strongly suggest that the trade of forest tree seeds is not as safe as previously believed. High infestation rates of seed lots of some tree species by insects and fungi, and the discovery of pests that could establish in new areas is alarming and stresses the need to reconsider phytosanitary measures in the seed trade. Measures for insects should potentially
focus on pests of gymnosperms, since they represent a higher phytosanitary risk in comparison with insects of angiosperm seeds: the risk associated with angiosperm seed-borne insects can be significantly reduced by ensuring that seeds are collected after insects emerge but it may be more difficult to avoid gymnosperm pests. Possible measures could include improvement of detection techniques used by phytosanitary inspectors at the border, and the development of production methods to reduce infestation rates and treatments to reduce the viability of seed-borne pests.

The situation seems to be much more complex for seed-borne fungi than for insects. The high fungal abundance and diversity, especially in angiosperms and Asian seeds, which also contained large fractions of pathogenic and unique fungi, is concerning. The lack of knowledge about the distribution, host ranges, and pathogenicity of most fungal species, as well as the limitations of detection and identification techniques, make it challenging to conduct pest or pathway risk assessments. Nevertheless, our study shows that potentially serious pathogens can be found in traded seeds, and that angiosperm seeds and seeds of Asian origin might represent a particularly great risk. Since fungal pathogens are very difficult to detect during border inspections, the most promising approach might be to work on developing effective treatments for reducing their viability in seeds or to regulate the trade in seeds of especially risky origins or species. Further studies of the biology and ecology are needed for better risk assessment and management of the fungal and insect pests identified in this study.

Acknowledgments

I. Franči, E. Allan, R. Eschen, M. Kenis, and S. Prospero designed the study. M.-A. Auger-Rozenberg and A. Roques supervised the insect assessment. S. Schneider optimized molecular laboratory protocols. M. Hartmann supervised the amplicon sequencing approach and performed the bioinformatic analyses of the HTS sequence data. I. Franči analysed the data and wrote the manuscript with an input from all co-authors. We thank Quirin Kupper (WSL) for his assistance with the fungal assessment by HTS, Hélène Blaumenstein, Stephanie Pfister and Sandrine Fattore (WSL) for their help with the fungal assessment by plating, Christopher Schefer (WSL) for his assistance with taking seed measures, Haley Smith (Dorena Genetic Resource Centre) and Philippe Lorme (INRA) for their help X-raying the seeds, and Valerie J. Fieland and Caroline M. Press (USDA Horticultural Crops Research Unit) for their assistance with fungal assessment of seeds obtained in North America. We are grateful to Maja Kuzman (University of Zagreb, Faculty of Science) for her advice about data manipulation in R. We thank the Genetic Diversity Center (GDC) at ETH Zurich for providing computational infrastructure and acknowledge the contribution of scientists at the McGill University and Génome Québec Innovation Center, Montréal, Canada, for the paired-end sequencing on Illumina MiSeq. This work was supported by COST Action “Global Warning” (FP1401). We also gratefully acknowledge the financial support of the Swiss National Science Foundation (Project C15.0081) and the Swiss Federal Office for the Environment. Contributions of I. Franči, R. Eschen, and M. Kenis were supported by CABI with core financial support from its member countries (see http://www.cabi.org/about-cabi/who-we-work-with/keydonors/ for details).

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Supporting Information
Additional supporting information may be found online at: http://onlinelibrary.wiley.com/doi/10.1002/eap.1971/full

Data Availability
Data are available from the Dryad Digital Repository at https://doi.org/10.5061/dryad.nr51124. Raw HTS sequences have been deposited in the Sequence Read Archive under accession number PRJNA550270. Assembled ITS and COI sequences obtained from fungal cultures and insects have been deposited in GenBank under accession numbers MN105153–MN105593 and MN122012–MN122067, respectively.