

Forests and global change: what can genetics contribute to the major forest management and policy challenges of the twenty-first century?

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53 54

55 Abstract

56 57

The conservation and sustainable use of forests in the 21st century pose huge 58 challenges for forest management and policy. Society demands that forests provide a 59 wide range of ecosystem services, from timber products, raw materials and renewable 60 61 energy to socio-cultural amenities and habitats for nature conservation. Innovative management and policy approaches need to be developed to meet these often 62 conflicting demands in a context of environmental change of uncertain magnitude and 63 scale. Genetic diversity is a key component of resilience and adaptability. Overall, 64 forest tree populations are genetically very diverse, conferring them an enormous 65 potential for genetic adaptation via the processes of gene flow and natural selection. 66 Here, we review the main challenges facing our forests in the coming century and 67 focus on how recent progress in genetics can contribute to the development of 68 appropriate practical actions that forest managers and policy makers can adopt to 69 promote forest resilience to climate change. Emerging knowledge will inform and 70 clarify current controversies relating to the choice of appropriate genetic resources for 71 planting, the effect of silvicultural systems and stand tending on adaptive potential 72 and the best ways to harness genetic diversity in breeding and conservation programs. 73 74 Gaps in our knowledge remain and we identify where additional information is needed (e.g. the adaptive value of peripheral populations or the genetic determinism 75 of key adaptive traits) and the types of studies that are required to provide this key 76 understanding. 77

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Keywords: sustainable forestry; assisted migration; climate change; adaptation;
genetic diversity; gene conservation.

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84 Introduction

The development of appropriate and informed policy for the conservation and 85 sustainable use of forests worldwide, during the 21st century and beyond, presents 86 several challenges. The long life-span of trees makes them susceptible to both short 87 and long term environmental disturbances (Alberto et al. 2013). The natural cycles of 88 environmental disturbances are being modified by human activities and this is 89 predicted to intensify in the future with major effects on biodiversity (Alfaro et al. 90 2014, Sala et al. 2000). Thus, policies that guide forest management decisions must be 91 amenable to both short and long term assessment. Innovative forest management and 92 93 policy approaches are necessary to meet the often conflicting demands of a changing society and the need to deliver an array of provisioning, regulating and socio-cultural 94 ecosystem services (MEA 2005, TEEB 2012). To ensure effective implementation in 95 96 practice, these new approaches need to be more responsive to the future needs of society in general, more amenable to cross-sectorial cooperation and more evidence-97 based than is currently the case (Vanhanen et al. 2007). 98

99 Because of the incomplete scientific evidence on the effect of silviculture on long 100 term forest sustainability and because forests provide a wide range of goods and 101 services for society, forest-related societal challenges can result in controversy, 102 polarization of views, stalemates and deadlocks (van Eeten 1999). While some of the 103 controversies are philosophical in nature, others could be resolved by further scientific 104 research. For example, it is debatable whether management based on natural 105 regeneration is more sustainable than planting in reforestation programs (Espelta et al. 106 2003, Morrissey et al. 2010) because the outcome will depend on a number of factors, 107 including availability of sufficient appropriate seed sources for natural regeneration, 108 ecological traits of the planted trees species, etc. Similarly, there is lack of clarity 109 regarding the relative merits of gradual tree selection versus clear-felling as a 110 111 harvesting strategy for timber production in commercial forests (Lundqvist et al. 112 2013). Other controversies relate to the relative benefits of stand conversion compared to ecological restoration; the use of exotic species (or non-autochthonous parent 113 material of native species) instead of native material of local origin as a source of 114 115 forest reproductive material in plantation forests; the use of multiple versus single species forest plantations; the acceptance and application of clonal forestry and the 116 use of genetically-modified trees in intensive forestry; among others (MCPFE 2008). 117 118

In addition to these uncertainties and societal controversies, there is the recognition 119 that climate change will have an impact and should therefore be included in these 120 longstanding debates (Noss 2001). The realization that many local habitats will cease 121 to offer the same potential for forest cover and production as they have in the past 122 represents a serious paradigm shift for forestry. The uncertainty regarding the 123 magnitude of the change at different spatial scales presents yet another difficulty for 124 the development of appropriate forestry policy. For example, due to their location at 125 the rear-edge of species distributions, countries in the Mediterranean region are 126 generally expected to experience the consequences of climate change more abruptly 127 than the more northerly countries of Europe (Hampe & Petit 2005). Nevertheless, 128 habitat changes are also expected to be severe at high latitudes and generally greater 129 in the lowlands than at high elevations (Loarie et al. 2009). There is therefore, a need 130 for sound, informed policy development that is tailored to both local and regional 131 scales to ensure that the most appropriate forest management strategies are 132 implemented. The recognition that there is no single approach underlines the 133 134 importance of selecting appropriate practices that fit the conditions and objectives of each particular situation (Millar et al. 2007). 135

136

The term adaptation (see Annex 1 for definitions of this and other terms used 137 throughout the paper) is commonly used to cover a range of purposes in the context of 138 global, and particularly climate, change (Biesbroek et al. 2010). In forestry, the term 139 is generally used to encompass all aspects of forest management that need to be 140 modified to prepare forests so that they can continue to provide their usual ecosystem 141 services under (the uncertainties of) global change (Bernier & Schoene 2009), as in 142 "adapting forests to climate change" or "adaptive forestry". However, from a genetic 143 perspective, adaptation has a different, more restrictive meaning, i.e. the evolutionary 144 capacity of a population to cope with a changing environment through genetic 145 146 changes. The ability of forests to adapt in a genetic way, i.e. genetic adaptation, to the new environments that will result from climate change depends on: (i) the speed and 147 intensity of climate change across species ranges, (ii) the existence of genetic 148

diversity among individuals in natural populations and (iii) the ability of this diversity 149 to migrate between populations through pollen and seed movement, i.e. gene flow 150 (Savolainen et al. 2007; Kremer et al. 2012). Populations with high genetic diversity 151 for adaptive traits (i.e. rich in genetic resources) are more likely to be able to respond 152 via natural selection to the challenges posed by climate change (Jump et al. 2008). 153 The adaptive potential (Scotti 2010) of forests worldwide thus depends on the genetic 154 resources found both within populations and among forest stands that are connected 155 via effective gene flow. 156

157

In 1994, in recognition of the importance of genetic diversity for forest sustainability, 158 FOREST EUROPE (the pan-European political process for the sustainable 159 management of the continent's forests, http://www.foresteurope.org/) created 160 161 EUFORGEN (http://www.euforgen.org/), a collaborative network of European 162 countries whose remit is to promote the conservation and sustainable use of forest genetic resources. However, despite this high level recognition of the importance of 163 forest genetic diversity, the National and European forest policy agencies frequently 164 165 ignore this aspect when developing indicators and guidelines on how forest management should prepare for and react to climate change (Koskela et al. 2007). 166 Similarly, the role of forest genetic resources is often omitted from the 167 recommendations (Graudal et al. 2014) issued by habitat and species conservation 168 agencies and non-governmental organizations, such as those within the framework of 169 the European Habitat directive (Council Directive 92/43/EEC). This is despite the 170 consideration that forest genetic resources in forest management strategies are key in 171 ensuring that forests can adapt to climate change (Lefèvre et al. 2013). The overall 172 lack of attention given to the genetic dimension of forest adaptation is probably due to 173 a general lack of knowledge of the recent advances in forest genetics among forest 174 managers and policy makers. 175

176

In this article, we review the major challenges faced by forest managers and policy 177 makers in temperate Europe in the 21st century for which recent progress in genetics 178 can contribute to the development of appropriate practical solutions. We deliberately 179 focus on the practical issues that arise from these forest challenges, listing them as 180 they are perceived in a management context. Seven major challenges emerged from 181 discussions between the stakeholders of the Linktree project who had direct roles in 182 developing policy and appropriate management to prepare forests for climate change. 183 184 These range from natural regeneration to assisted migration and from conservation to breeding. We address naturally regenerated and plantation forests but exclude 185 genetically modified organisms from our report as they are not currently available for 186 most species of interest to European forestry. We present examples where uncertainty 187 adds complexity to management decisions. For these examples, we provide insights 188 into how science, and particularly an understanding of genetics, can help to inform 189 policy directions. In doing so, we aim at bridging the gaps between science, 190 management and policy which typically operate within different spheres of influence 191 and across different time frames and are not always driven by the same concerns. 192 193

We also aim to demonstrate that some key forest management and policy questions
can be resolved by genetic studies, in particular those using new generation
technology that is now increasingly becoming accessible and of practical use for
forest trees. These studies give insights into the current and future consequences of

and, in turn, can guide how management strategies should be adapted to foster the
 evolutionary potential of forests under changing conditions. Figure 1 shows a
 summary of the processes affecting the genetic diversity of forest trees and how
 management can use and modify these processes to prepare forests for climate
 change.

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205

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Figure 1. Processes affecting the genetic diversity of forest trees and how

management can harness these processes to prepare forests for climate change. The
pressure applied by forest management on the state of genetic diversity has impacts
that affects the forest as a whole. Climate change drives the system and the selection
pressure it imposes can be reduced by management practices.

212

213 214 Challenge 1: Will the genetic resources available in natural forests remain 215 appropriate and be sufficient under climate change?

216

This question relates to the adaptive potential of forest tree populations and their reaction to climate change. Depending on the management goal (e.g. production, protection or recreation), phenotypic traits of interest to managers will be wood quantity and quality or survival and health. Standing genetic variation and the phenotypic plasticity found in forest tree populations provide the means to cope with the new conditions.

223

Climate envelope modelling shows that species distribution ranges and the species 224 composition of forests are likely to change considerably over the coming decades (e.g. 225 Thuiller et al. 2005). However, this modelling approach, based on correlation of 226 geographic distribution with a range of climate variables, cannot accurately predict 227 the future distribution of species because it does not take into account important 228 ecological and evolutionary processes such as competition at the community level, 229 dispersal, phenotypic plasticity and adaptive potential. In the few studies in which 230 genetic diversity has been included in the modelling frameworks, it has been found to 231 modify the predicted range of forest tree species under climate change considerably 232 (Benito-Garzón et al. 2011, Oney et al. 2013). This highlights the urgent need for 233

appropriate genetic methods and data to determine the adaptive potential of existingmaterial.

236

Currently, the best available source of genetic data on phenotypic traits of importance 237 for local genetic adaptation comes from "common garden" trials which contain 238 material sourced from a range of origins, called provenances in the forestry literature. 239 Published results from such trials show that adaptive genetic diversity is large in most 240 forest tree species, mainly within but also among populations (Ducousso et al. 1996), 241 although there are remarkable exceptions (e.g. Pinus pinea, Mutke et al. 2010). These 242 trials also provide evidence of local genetic adaptation across the range of species and 243 demonstrate that most tested provenances display large phenotypic plasticity for key 244 adaptive traits when grown under different climatic conditions (Rehfeldt et al. 2002). 245 246 These findings are important because they indicate that, provided climate shifts are 247 not too steep and some degree of growth yield reductions are acceptable for management objectives, current genetic resources of many tree species are likely to 248 remain appropriate in many of their current habitats, particularly within the core of 249 250 geographic distribution ranges.

251

Unfortunately, common garden trials are only available for a restricted number of 252 253 commercially important forestry species, they rarely include representatives of all the ecologically relevant seed sources and in many cases the trials are insufficiently 254 replicated across the distribution range of the target species. Notably, peripheral 255 populations (i.e. coming from the ecological and geographical distribution margins) 256 are under-represented in existing field trials. There is therefore an urgent need for the 257 establishment of pan-European and range-wide species and provenance trials of novel 258 and native species (see for example the REINFFORCE initiative, 259

<u>http://reinfforce.iefc.net/).</u> These should be established using provenances from across
 the distribution range grown at multiple sites which include locations within the
 current distribution ranges as well as those that are beyond the current range but are
 predicted to become suitable in the future for the target species.

264

When field trials already exist, their analysis at a national or range-wide scale could provide immediate answers to some of the questions raised by managers regarding the potential of their populations to adapt to climate change (see for example <u>www.treedivnet.ugent.be</u>, an international network of common garden experiments where monocultures are compared with mixed species or mixed provenance treatments, and EU funded initiatives, such as Trees4Future,

271 <u>http://www.trees4future.eu/</u>, and the COST action FP1202, <u>http://map-fgr.entecra.it/</u>,).

272 The measurement of meaningful adaptive traits in these trials is recognized to be

challenging in terms of the time and expertise required to obtain them (e.g. drought

- tolerance, phenology, seed production, and other life history traits, Chambel et al.
- 2007) and rapid and cost-effective phenotyping methods of many hundreds of
 individuals need to be developed if the full benefit of these expensive trials is to be
- individuals need to be developed if the full benefit of these expensive trials is to be
 secured (Neale & Kremer, 2011). Progress in fast phenotyping approaches is being
- made with modern methods such as terrestrial Lidar scanning for measurement of tree
- size and form (Eysn et al. 2013), chlorophyll fluorescence for estimating stress via
- 280 photosynthetic efficiency (Salmela et al. 2011) and others (Alia & Majada 2013)
- becoming available. Greenhouse trials, in which treatments such as drought and pests
- and diseases can be imposed, represent a useful and sometimes alternative strategy to expose and assess standing genetic variation and phenotypic plasticity in key traits for

- seedlings and young trees (Sagnard et al. 2002, Chambel et al. 2007). Results from these trials will provide an understanding of variation in adaptive traits and how it is distributed across the landscape. This knowledge will form a framework with which to guide advice on the scale of current and future movement of planting material can be practiced without the fear of maladaptation. It will also contribute to models which aim to determine the evolutionary potential of our forests to cope with climate change.
- 290

291 Advances in molecular approaches are beginning to make it possible to link phenotypes to variation in genetic markers (González-Martínez et al. 2006). For 292 293 example, such approaches have already been used to explore cold and drought resistance in several conifers (Neale & Kremer 2011, Eveno et al. 2008, Grivet et al. 294 2011). By taking advantage of new cost-effective genotyping-by-sequencing (GbS) 295 296 methods it is now possible to adopt a genome scanning approach to generate millions of data points across the genome. In one of the first such studies in forest trees it was 297 shown that extensive variation exists within natural populations of pines for genes 298 linked to ecologically-relevant traits (Parchman et al. 2012). Ultimately, it is 299 300 anticipated that these approaches will provide managers with the practical tools to determine whether their targeted trees and stands have the capacity to adapt to climate 301 change, at low cost even in non-commercial species. 302

303

In conclusion, most genetic studies to date indicate that there is a large amount of 304 heritable adaptive variation available in tree species on which natural selection can 305 operate. Thus, the rather dramatic predictions made by climate envelope models of 306 large scale changes in forest distribution within the coming decades require 307 reassessment and refinement to include genetic processes. However, adaptive 308 potential depends on the species and local ecological conditions and demographic 309 constraints could severely limit the capacity of populations to evolve under the most 310 severe climate changes. Thus, although usually appropriate, management scenarios 311 based solely on locally existing forest material and genetic resources may be 312

313 *hazardous particularly at range and ecological edges.*

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Challenge 2: What should be monitored to determine whether the native species in natural woodlands are currently at risk under climate change?

318

Both a lack of genetic adaptation and demographic collapse can drive populations to extinction. Determining which genetic and demographic thresholds will lead to maladaptation is crucial to make informed management decisions, particularly in forests where, despite predictions, evidence of decline due to climate change is not yet strong. This can be achieved by genetic monitoring.

324

Although forest monitoring is often carried out by countries to assess the state of their forests and the social, economic and environmental benefits they derive from them, it rarely includes any assessment of genetic resources (Graudal et al. 2014).

Aravanopoulos (2011) defines genetic monitoring as "the quantification of temporal

changes in population genetic variation and structure". He suggests that genetic

330 monitoring should focus on ecologically and economically important forest tree

331 species and on four key genetic processes: (1) natural selection, (2) genetic drift and

- (3) gene flow and one demographic process linked to gene flow: the mating system.
- 333 Carried out at regular intervals, genetic monitoring will provide much needed

information on adaptive potential. Protected areas where there is less human mediated 334 interference and management practices are carried out with a lighter touch than in 335 other forest types constitute excellent sentinels of the effects of climate change and 336 are thus also appropriate candidates for genetic monitoring. 337 Neutral molecular markers can now be used with relative technical and financial ease 338 to monitor genetic drift, gene flow and the mating system. The relationship between 339 neutral genetic diversity and standing genetic variation at fitness related genes in large 340 populations is not straightforward (Le Corre & Kremer 2013) and genomic 341 approaches are not yet sufficiently generally accessible to be applied to clarify the 342 relationship. There is therefore a need to determine the most efficient ecological 343 proxies that could be used instead to assess the adaptive potential of forest 344 populations by forester. Aravanopoulos (2011) suggests that several demographic 345 346 indicators, such as: age class structure, filled seed set and seedling density can be used 347 as these proxies. A system of genetic monitoring for forest tree species broadly based

- on the concepts discussed here, was proposed and successfully tested in Germany for
 providing early warning signs of environmental changes affecting forests (Konnert et
 al. 2011).
- 351

Studies using novel approaches will be needed to assess standing genetic variation 352 and adaptive potential directly. General information can come from progeny tests in 353 common gardens (Houle 1992) or in situ experimentation (e.g. reciprocal 354 transplantation schemes), where heritability and genetic variance can be measured. 355 Sophisticated association and genome-wide ecological studies will increasingly 356 provide information on standing genetic variation available for adapting to changing 357 environments (see references above in Challenge 1). It is expected that genes involved 358 in genetic adaptation will vary along the geographical range of species (see Hancock 359 et al. 2011 for the model plant species Arabidopsis thaliana and Prunier et al. 2012 360 for an example in forest trees), thus the establishment of an inventory of such 361 adaptively important genes (and their allelic variants) across the full species range is a 362 priority for current and future applied genetics research. 363

364

In conclusion, monitoring the potential of forests to adapt genetically under climate is necessary over broad areas to inform management. Monitoring based on parameters used in classical forest management (such as adult age classes, seedling density and presence of pests) can now be coupled with molecular genetics assessment methods to provide early warning signs of maladaptation risks. Incorporating genetic monitoring as a component of the current pan-European forest monitoring efforts would provide key indicators of evolutionary and adaptive potential.

372 373

374 Challenge 3: Is assisted migration the answer for managing forests and their 375 habitats under climate change?

376

This question typically arises when sound evidence suggests that current genetic resources at a site will cease to offer an appropriate option under future climate (Challenge 1) and introduction of non-local resources is being considered. Assisted migration concerns the sourcing of seed and planting stock and can take two forms: the use of exotic species (i.e. species that do not naturally occur at the planting location or, more generally, within the country) or the use of non-local genetic resources (i.e. populations of naturally occurring species which originate from other parts of the species distribution range). In management plans, the definition of
assisted migration is often restricted to the use of exotic species and the option to use
conspecific material from a different origin is frequently overlooked.

388 The use of exotics has been practiced in forestry for centuries, often with great success in terms of sustaining wood production (e.g. Pseudotsuga menziesii, Pinus 389 radiata, Abies grandis, Picea sitchensis, etc introduced in Europe). Such successful 390 introductions are often the culmination of extensive trials and on the basis of their 391 results, the elimination of many species from the candidate list. Although there are 392 numerous examples of successful introductions (e.g. mountain habitat restoration in 393 France in the 19th century (Charry 1996, Bartoli & Musch 2003), there are also 394 notable instances of spectacular failures. A classic example is eastern white pine, 395 396 Pinus strobus L., a five-needle pine which was introduced into France from North America in the mid-16th century, and into England and Germany 150 and 300 years 397 later, respectively. In the mid-19th century the European plantations were colonized by 398 the pathogenic fungus Cronartium ribicola of Asian origin, with catastrophic results 399 400 due to lack of resistance to the disease (Karlman 2001). Consequently, this species is now rarely used in European forestry. Also, some of the successfully introduced 401 species in Europe have become invasive (e.g. Robinia pseudoacacia and Quercus 402 403 rubra), causing habitat management problems on a massive scale.

404

The translocation of genetic resources from one region of the species distribution to 405 another is also a form of assisted migration (sensu Richardson et al. 2009), i.e. within-406 species assisted migration or "assisted gene flow" sensu Aitken and Whitlock (2013). 407 Again, foresters have used this management option extensively at least since the 19th 408 century, very often as a strategy for planning large scale reforestation programs when 409 local seed supplies were insufficient or thought to be less productive than non-local 410 411 resources (Bartoli & Musch 2003) or to overcome budget limitations: for example 4000 tons of oak seed were brought into Germany for planting from south-eastern 412 Europe (Kleinschmit, 1993); cherry seedlings were sold in Europe from seeds 413 obtained in jam factories (Kleinschmit & Kleinschmit, 2009). Here again, despite 414 some general success, maladaptation of the planting stock has also resulted in severe 415 economic lost (see for example the massive dieback of Portuguese seed sources of 416 maritime pine planted in South West France after the 1984-1985 severe winter frost, 417 Timbal et al. 2005). 418

419

420 It is now well recognized that assisted gene flow is neither risk-free for the target species themselves nor for their associated communities (see for example Aitken & 421 Whitlock 2013). This understanding underpins the European Council Directive 422 1999/105/EC on the marketing of forest reproductive material which requires seed 423 companies to attach clear passport data to traded resources. This traceability allows 424 forest managers to make informed decisions regarding the likelihood of maladaptation 425 of purchased planting stock and the potential ecological consequences of using it. 426 Some managers have attempted to reduce the risk of planting failure under climate 427 change by mixing seed lots within and among regions of provenance and among years 428 (i.e. 'composite provenancing'). However, according to Ennos et al. (1998) the use of 429 assisted gene flow for such insurance purposes should only be considered if levels of 430 variation within indigenous populations are so low that they are unlikely to be able to 431 adapt to future conditions by themselves and if the imported material is considered to 432

be better adapted to future climatic conditions than the indigenous populations, whichare both difficult to ascertain (see Challenge 2).

435

Thus, assisted gene flow could offer a reasonable solution for situations in which local 436 resources are declining, adult tree mortality is above a set threshold and natural 437 regeneration is low or absent despite supplementation efforts using locally sourced 438 seed (Ledig & Kitzmiller 1992, Hubert & Cottrell 2007, CRGF 2008). This is a form 439 of genetic rescue where hybridizing local resources with genetically unrelated 440 resources could restore fitness (Hedrick et al. 2011). It requires seed material to come 441 from stands "preadapted" to a future (and uncertain) climate and it is not free of risk 442 as it could also result in outbreeding depression (i.e. lower overall fitness because of 443 disruption of co-adapted gene complexes). Ecological risks, such as within-species 444 445 introgression or modification of biotic interactions, although somehow attenuated compared to those from exotics, are also present with assisted gene flow (Aitken & 446 Whitlock 2013). Broad scale analyses such as those of Rehfeldt et al. (2002) for Pinus 447 sylvestris, where the fitness of exotic provenances is compared to that of the local 448 449 provenances at a trial site, enable the appropriateness of local resources to be evaluated and the need for transfer across seed zones to be assessed. 450

451

In conclusion, assisted migration programs can disrupt local genetic adaptation and affect the present and future dynamics of forest genetic resources. We can take advantage of past experiences in the introduction of species and provenances, but these are not real experiments to test the effectiveness of present-day assisted migration programs. Caution is needed in the use of extensive assisted migration as the responses will likely affect not only trees but also the whole forest community in which they are established.

459 460

Challenge 4: Is natural regeneration the most appropriate management technique for promoting the adaptation of natural forests to climate change?

463

464 Natural regeneration is being promoted in a European context of close-to-nature forest management for several reasons including those that relate to cost reduction, 465 decreased disturbance, better selection potential due to larger seedling density and 466 conservation and continuous evolution of the local gene pool. It is thus generally 467 468 considered to be the most appropriate management method in protected habitats. However, the success of natural regeneration is variable and factors such as soil type, 469 abundance of competing weeds, herbivore pressure, species light requirements and 470 availability of seed sources may force management to consider other regeneration 471 472 approaches.

473

Whether natural regeneration is a better strategy for regenerating forests than 474 plantation remains an open question (Espelta et al. 2003, Morrissey et al. 2010), but it 475 is clear that within this debate, genetic considerations are rarely given the importance 476 they deserve (except when the management goal is to use the local gene pool or 477 GMOs are involved). This is unfortunate as genetic diversity considerations can be an 478 important basis for preferring one strategy over the other (Ledig & Kitzmiller 1992), 479 particularly in small forest stands (Vranckx et al. 2014) and their relevance is central 480 to climate change related issues (Koskela et al. 2007, Alfaro et al. 2014). 481 482

The question geneticists must address to help management in this context is whether 483 the level of genetic diversity (and the amount of seedlings) available from natural 484 regeneration is sufficient (Vranckx et al. 2014). Decisions regarding whether natural 485 regeneration is more appropriate than planting for the maintenance of genetic 486 diversity (and ultimately adaptive potential) can be informed by some level of genetic 487 monitoring based on appropriate methods (Aravanopoulos 2011 and see Challenge 2). 488 If the stand to be regenerated is genetically of average or above average diversity for 489 the species, and if seedlings are abundant, then natural regeneration is likely to be the 490 most appropriate option. When using natural regeneration, managers should make 491 sure that: i) adult trees contributing to natural regeneration are plentiful and not too 492 similar phenotypically (see minimum requirements for conservation of genetic 493 diversity in Koskela et al. 2013), ii) varied silvicultural treatments are implemented at 494 495 the landscape scale (see the effect of structure and density on the spatial genetic 496 diversity of seedlings in Sagnard et al. 2011) and iii) enough time is left for natural regeneration to be thinned by varying levels and forms of natural or silviculture-based 497 selection (competition for light and water, resistance to herbivores, pests and 498 499 diseases).

500

If natural regeneration is scarce and/or a species in the stand has significantly lower 501 than average genetic diversity which may result in detrimental effects such as fixation 502 of deleterious alleles at adaptive loci (see how to monitor in Challenge 2), then a 503 strategy based on planting is likely to be beneficial. Managers who adopt this 504 approach should ensure that there is high genetic diversity in their planting stock by 505 insisting that seed lots used for plantation are based on equal contributions from at 506 least 30 to 50 seed trees (legal standards of most countries do not include this 507 requirement). They should also give careful consideration to the geographic origin of 508 the stock and chose the source that is most likely to be adapted to the conditions at 509 planting site. 510

511

512 Greater mortality (and thus potential for selecting advantageous phenotypic traits) is expected under a natural regeneration regime than one based on planting. A plantation 513 will typically install 1200 to 2500 seedlings/ha, whereas this number increases by one 514 or two orders of magnitude with natural regeneration. Amm et al. (2012), for 515 example, show that selection intensity could be as high as 0.0001 between the seed 516 production and the recruitment stage in Abies alba Mill. For any given heritable 517 518 phenotypic trait of importance to forestry, it would be necessary to increase initial densities of plantations dramatically to reach the same level of selection intensity as 519 that of natural regeneration, and these are unlikely to be acceptable under current 520 operational conditions. 521

522

In conclusion, to provide sufficient material on which natural selection can operate to bring about evolutionary change, the option of natural regeneration is likely to be most appropriate as it typically provides a much larger base population than is the case for plantations. Forestry will need to rethink its strategies for long rotation species to make it possible for selection to occur in those areas where climate change is expected to have its strongest impacts.

529

530

531 Challenge 5: Should/can the conservation of genetic diversity be included as a 532 component of habitat and species conservation strategies?

- 533 Managers tend to give much less consideration to the conservation of the genetic 534 resource (gene conservation) than to that of the species and habitat. Habitat 535 conservation usually aims to protect a set of particular ecological processes that 536 sustain a defined range of species, often rare or endangered, or are particular to an 537 ecosystem. In contrast, the in situ conservation of a genetic resource aims to maintain 538 the process of genetic adaptation under natural selection and demographic shifts so 539 that the individuals of a target species which are best adapted to current conditions 540 survive and breed to produce the next generation. Thus, a holistic biodiversity 541 conservation strategy consists of several different strands, the aims and approaches of 542 which are not necessarily compatible.
- 543 544

545 Management of protected areas is mostly directed towards the conservation of rare 546 and endangered species and habitats (Peterken 1977, Klein et al. 2009) and there is 547 little direct focus on gene conservation. In contrast, *in situ* gene conservation is generally practiced in large plots consisting of the major forest tree species in 548 549 common habitats where genetic processes can occur. The networks of gene conservation units for forest tree species in Europe follow a set of minimum 550 requirements, including range-wide genetic diversity representativeness and their 551 management procedures aim to shield conservation units from maladaptive gene flow 552 (see Koskela et al. 2013 for a full description). Thus, networks of gene conservation 553 units provide a level of protection of biodiversity that is seldom considered in 554 protected areas that, we believe, warrants worldwide recognition such as 555 categorization as IUCN protected areas. Gene conservation units are also useful for 556 demonstrating that the integration of genetic principles into management does not 557 hinder silviculture and forest planning and instead, can significantly contribute to 558 adaptive forest management by introducing evolutionary considerations into everyday 559 practice (Lefèvre et al. 2013, Koskela et al. 2013). 560

561

562 The application of the greenway or ecological corridor approach used by planners to establish a network of linked land and water habitats in the landscape (Ignatieva et al. 563 2011) may in some cases reconcile habitat, species and gene conservation strategies. 564 Corridors can facilitate both migration and gene flow which are key processes in the 565 maintenance of adaptive potential (Savolainen et al. 2007). Gene flow is generally 566 predicted to make a strong contribution to genetic adaptation (as it maintains the 567 568 overall genetic diversity, on which natural selection operates) in most forest tree species (Kremer et al. 2012). However, large gaps of knowledge still exist. For 569 example, the effects of different forest landscape components on the rates of long-570 distance gene flow are currently poorly understood. Although studies tend to 571 demonstrate that fragmentation indeed increases genetic drift and loss of fitness 572 (Vranckx et al. 2011) and the presence of woodland corridors and hedges generally 573 improve genetic connectivity, the distance that native forests have to be apart to 574 experience significantly reduced genetic connectivity is species specific and depends 575 on pollen and seed dispersal. Monitoring whether ecological corridors really do 576 promote gene flow, evolutionary processes and migration to more suitable habitats, as 577 well as identifying which species are most at risk by fragmentation, remain a 578 necessity. 579

580

The role of habitat connectivity is particularly controversial for the maintenance of evolutionary potential for populations from the periphery of the distribution range and

this is particularly relevant in the context of climate change. Models have shown that 583 pollen flow may in fact decrease genetic adaptation rate in some instances (Kuparinen 584 et al. 2010). Leading edge peripheral (as well as numerous small) populations in 585 northern Europe do not always demonstrate local genetic adaptation (Leimu & 586 Fischer 2008). For these, pollen flow from the core of the distribution area may be 587 beneficial, particularly from a climate change perspective as core populations are 588 better adapted to the future conditions that are predicted at the leading edge. In this 589 case, connectivity will reduce inbreeding and drift at the leading edge (Willi & 590 Fischer 2005). Conversely, disjunct populations at the rear edge, with a long history 591 of demographic stability (Hampe & Petit 2005), will suffer from maladaptive gene 592 flow from core populations (Lenormand 2002). Rear edge populations, geographically 593 or ecologically disjunct from species distribution ranges, may constitute hotspots of 594 595 evolutionary potential provided their census size is not too small (Lesica & Allendorf 1995). Plans to reconnect these to other populations within the main distribution range 596 of the species should therefore be resisted to maintain their genetic distinctiveness. 597 However, on the other hand, gene flow from these rear edge populations could be 598 599 helpful for the adaptive evolution of core and leading edge populations in of the to climate change. 600

601

In conclusion, gene conservation of forest tree species should be viewed as an 602 integral part of biodiversity conservation, alongside that for species and habitats. As 603 each conservation focus has discrete objectives that may require different 604 management strategies, it is important to run them in parallel and non-exclusively 605 over the landscape at national and regional levels. However, protected forest areas in 606 which there is little or no active management can sometimes directly contribute the 607 conservation of forest genetic resources and genetic diversity monitoring (Challenge 608 2) should become a priority concern there for the most relevant species. 609

610 611

612 Challenge 6: Do the current objectives of tree breeding programs address 613 climate change sufficiently? Does the current usage of forest reproductive 614 material represent the best practice in preparation for climate change?

615

Tree breeding involves selecting individuals that have particularly desirable traits 616 and crossing these individuals to improve the trait or traits of interest within 617 618 populations. Breeding programs can therefore produce genetic resources which are valuable in enabling forests to adapt to climate change and to provide more and/or 619 better ecosystem services. Historically, selection in these breeding programs has 620 targeted commercially important traits such as growth, wood density, stem 621 straightness and complete resistance to particular pathogens in clonal forestry (e.g. 622 poplars), with the goal of marketing the desired product within a minimum time span. 623 This breeding paradigm only makes sense if the environment remains constant during 624 the entire production rotation. 625

626

Despite risks related to climate change in the forest sector, many European countries
continue to subsidize and support the use of a limited set of clones or varieties
produced by breeding programs established long ago, and this for a limited set of
commercial tree species (Alia & Majada 2013). Many species of interest in the 1970s
have dropped out of financial support schemes and seed orchards (mostly for conifers)
have been neglected or even destroyed and are no longer used to provide seed.

Moreover, new breeding efforts are scarce outside a few dominant commercial 633 species. Both practice and breeding efforts are thus too limited and inadequate for the 634 challenges of the 21st century (which is recognized by the community and has led, for 635 example, to the EU initiative Trees4Future, see Challenge 1). A few breeding 636 programs, however, are now incorporating in their objectives functional traits of 637 relevance to adaptive potential (e.g. water use efficiency, phenology along with the 638 more traditional traits of growth and wood quality, e.g. the Maritime Pine breeding 639 programs in France and Spain, Bouffier et al. 2013), providing an example to follow 640 in other species. 641

642

Modern breeding methods based on genomic selection offer great potential for 643 multipurpose breeding programs. Genomic selection has the potential to accelerate 644 645 breeding in long-generation species, substantially increasing genetic gains while also providing a flexible framework to incorporate changes in breeding targets. It is 646 already widely used in animal and crop selection but has only recently started to be 647 applied to forest trees in species such as eucalypts. Advances are also being made 648 649 rapidly towards implementation in conifers and other broadleaves such as poplar (Plomion et al. 2015). By genotyping at regular intervals across the genome with a 650 large number of markers (classically over 10 markers/cM, i.e. over 15,000-30,000 651 markers in forest trees), genomic selection approaches are able to predict phenotypes 652 with variable degrees of accuracy, depending on the effective population size of the 653 base population, linkage disequilibrium, heritability and the genetic architecture of the 654 desired trait (Grattapaglia & Resende 2010). 655

656

There are calls from the research community for breeding programs to consider more 657 seriously genetic resources outside the current very few commercial species, to form 658 links with conservation programs and to address traits and uses that have not yet been 659 considered (Fins et al. 2006). For assisted gene flow and ecological restoration for 660 example, breeding for increased genetic diversity and evolvability alone might be a 661 reasonable goal in itself (Alia & Majada 2013, Lefèvre et al. 2013). Benefits from 662 state-of-the-art selection theory and practice can be down-scaled and used in so called 663 low-input breeding programs. Low-input breeding carries both the ideas of selection 664 towards rusticity and a highly cost-effective selection process in terms of direct 665 benefits to users (Dawson et al. 2008). Low input breeding is based on a breeding 666 cycle that ceases early in the selection process, uses information from both wild 667 668 populations and common gardens, and relies heavily on molecular markers to delineate meaningful geographic zones, track candidate genes and maintain a high 669 effective population size (Lindgren & Wei 2006, El-Kassaby & Lstiburek 2009). Such 670 breeding strategies represent flexible and viable alternatives to complicated and costly 671 (high-input) breeding programs that are designed for high yield situations (Wang et al. 672 2010). 673

674

675 In conclusion, policy-makers should be aware that breeding programs need to adapt quickly to the challenges of the 21st century while existing ex situ collections and past 676 selection efforts should be rescued and conserved. Whether based on a high or a low 677 input strategy, breeding programs should include the assessment of phenotypic traits 678 that are likely to be important in conferring genetic adaptation to climate change 679 (which may well be found outside usually investigated resources and will require 680 well-organized international collaboration). Low-input breeding strategies represent 681 an opportunity to do so for species that are traditionally under-represented in 682

breeding programs because of their low market value. This may be useful, as in the
future such species (e.g. Mediterranean trees) may become increasingly important
under climate change.

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Challenge 7: What are the socio-economic benefits of genetic conservation and breeding under global change scenarios?

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696

In addition to changing perceptions of the importance of genetic resources for better
coping with increased disturbances, the views of society are also shifting regarding
the role of genetic resources in meeting the demands for forest goods and services.
The two main current drivers of this shifting perception are climate change and
expected future demands by end-users.

Future demands by European end-users are anticipated to focus on an increase in 697 renewable energy and materials, both to mitigate climate change and to promote trade 698 699 and economic growth (Farizo et al. 2014, Soliño et al. 2012). This will be accompanied by a need to produce more product per unit of land area as the demands 700 for urbanization, recreation, nature conservation and food production should reduce 701 the amount of land that is available to forestry. There will be pressure to improve the 702 quality of timber (e.g. by increasing stiffness in solid wood or by lowering the lignin 703 content). According to Vanhanen et al. (2007), the maturing forest product markets in 704 705 Europe will place pressure on the forest sector to restructure through development and adoption of technical and social innovations. At the same time, the demand for 706 recreation and nature conservation is also increasing, and the desirability of a uniform 707 product (which can be more easily achieved via clonal forestry) for the bio-economy 708 has to be balanced against the need to maintain high levels of biodiversity. 709

710

Forest trees are characterized by a low level of domestication in comparison to other 711 organisms, and consequently, the high levels of intra-specific genetic variation 712 (among and within populations) that most tree species naturally contain has largely 713 been maintained. This variation is essential for the long term adaptability of the tree 714 species themselves as well as the associated species that depend on them. It is also 715 crucial because it serves as the reservoir for breeding resources in the future. 716 However, overall social perceptions of the benefits of biodiversity conservation to 717 718 forest ecosystem services are usually based on species richness and the importance of intra-specific genetic diversity is generally not accorded the consideration it deserves. 719 This is possibly because genetic diversity is a component of biodiversity that is not 720 easily visible or accessible to most people. Also, the perceived importance of 721 biodiversity varies with the socio-economic environment (Dominguez-Terrero and 722 Soliño, 2011). Intra-specific genetic diversity has an intrinsic economic value which, 723 despite its importance, is unfortunately poorly recognized in economic valuation 724 scenarios (Thorsen & Kjaer 2007). Highly-diverse forests are also better prepared to 725 response to societal demands for new specialized forest products and increased 726 productivity under climate change. 727

728

In conclusion, increasing the societal perception of the value of genetic diversity in managed forest ecosystems should be a priority. New methodological developments in forest and landscape genetics could help to assess the socio-economic importance of conservation or breeding activities. Raising awareness in forest managers, policy makers and conservationists of the essential role of genetic diversity on biodiversity
dynamics and adaptability of forests to future conditions is urgently needed.
Improvements in knowledge transfer beyond academia are required.

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739

738 Conclusion

740 Climate change will impact natural forests forcing trees, their keystone species, to move, adapt or disappear locally. Climate change is also likely to alter the health of 741 forest trees worldwide. The speed and extent to which these alterations are likely to 742 occur remain difficult to predict accurately. Ultimately, the decision to adopt one 743 management strategy over another is in the hands of managers and policy makers who 744 745 should be able to make "no regret" decisions. We have shown that genetic diversity 746 will play a vital role in determining the resilience and adaptability of forest tree populations to climate change whatever the management objectives chosen 747 (production, protection or recreation).. Genetic knowledge will help managers and 748 749 policy makers make informed, science-based decisions for prioritizing strategies and it is essential that practitioners keep abreast of scientific advances in this field. Of 750 course, much uncertainty remains and we expect that results from common gardens 751 and the rapid advances that are being made in genomics will soon contribute much 752 needed information for critical issues such as assisted migration and assisted gene 753 flow. This developing understanding will also help tree breeders to apply multi-trait 754 selection approaches to produce the most appropriate phenotypes needed under 755 changed climate conditions. Managing forests and other natural landscapes to 756 conserve and sustainably use genetic diversity is a sensible, practical and risk-757 reducing strategy under uncertain environmental conditions, that can be applied in 758 production, protection and recreation forests alike. 759

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