The origin and evolution of the angiosperms is one of the great terrestrial radiations and has had manifold effects on the global biota. Today, flowering plants generate the vast majority of human food, either directly or indirectly as animal feed, and account for a huge proportion of land-based photosynthesis and carbon sequestration. With a fossil record that extends back to just over 130 million years ago, flowering plants have diversified to include 250,000 to possibly 400,000 species occupying nearly every habitable terrestrial environment, and many aquatic ones. Understanding how angiosperms have accomplished this feat over a relatively short span of evolutionary time will elucidate many of the key processes underlying the assembly of Earth’s plant/animal associations and entire ecosystems.

Many scientists have understood the importance of broad, comparative genome sequencing since the beginning of the Arabidopsis thaliana and rice (Oryza sativa) genome sequencing projects [1-4]. Arabidopsis, a relative of cabbage, had already become the premier model for plant genetics, and half the world’s dependence on rice for food makes that crop plant an important model for the genetic architecture of traits important to humanity. More recently, poplar (Populus trichocarpa), grapevine (Vitis vinifera) and papaya (Carica papaya) have been sequenced as genomic models for woody crop plants [5-12]. These advances have been motivated by the realization that understanding the structure and evolution of plant genomes would contribute to society through enhancements to agriculture and forestry [13]. However, the few angiosperm nuclear genomes that have been sequenced so far reside on just two limbs within the angiosperm branch of the Tree of Life [14,15] and, therefore, aid us little in understanding the characteristics of the last common ancestor of all angiosperms (Figure 1). Many key angiosperm innovations, such as the origin of the flower and fruit, diverse pollination systems and double fertilization, large water-conducting vessel elements, diverse biochemical pathways, and many of the specific genes that regulate
key growth and developmental processes, appeared first among the basal angiosperm lineages [16-20]. A thorough understanding of processes that shape genes and genomic features, and of the many similarities and differences between model monocots (for example, *Oryza*) and eudicots (for example, *Arabidopsis*), requires a perspective based on evolutionary lineages. Such perspectives can be obtained only through analysis of an appropriately broad sampling of genomes, including lineages branching from the most basal node on the angiosperm tree [21]. But which basal angiosperm(s) should be given the highest priority for sequencing in the near future?

Recent phylogenetic analyses [14,15,17, 22] have identified *Amborella trichopoda*, a large shrub known only from the island of New Caledonia, as the single 'sister species' to all other living flowering plants. *Amborella* therefore offers the unparalleled potential to 'root' analyses of all angiosperm features, from gene families to genome structure, and from physiology to morphology. Furthermore, as the branching-point for *Amborella* is situated 'between' gymnosperms and all other angiosperms, a genome sequence for *Amborella* would help characterize processes that distinguish these two lineages of extant seed plants. The nuclear genome sequence of *Amborella* would contribute uniquely to efforts to reconstruct characteristics of the 'ancestral angiosperm'. The importance of *Amborella* in this regard is already widely appreciated [19,23]. Two recent papers, in fact, point specifically to basal angiosperms, including *Amborella*, as obvious choices for future nuclear genome sequencing efforts [24,25].

The genome structure of the ancestral angiosperm is currently much debated: did a whole-genome duplication pre-date or coincide with the origin of angiosperms (perhaps catalyzing innovation) or did the whole-genome duplication reported for several lineages of basal angiosperms [26] occur after the
Figure 2
Sequencing the nuclear genome for Amborella will root comparisons of monocot and eudicot genome sequences. (a,b) Sequence-based comparisons of the Amborella sequence (highlighted in yellow) with (a) Arabidopsis and (b) rice (Oryza) sequences for homologous genome segments (1, 1’, 2 and 2’) identify homologous genomic regions and genes (shown by colored arrows) that have undergone duplications and presumed gene loss in different segments. (c) From such comparisons investigators can identify the timings of segmental duplications and inversions, gene gains and losses, and whole-genome duplications (WGDs) in these three lineages. The large black circle indicates the monocot-eudicot split. The Amborella sequence resolves the timing of an inversion and a tandem duplication (versus loss of a duplicate) that distinguish homologous Arabidopsis and rice segments. Taken together, the map comparisons imply that the orientation of the green, blue and red genes in the Amborella sequence matches that in the common ancestor of monocots and eudicots. We can also infer that the purple gene was present in the common ancestor of monocots and eudicots. However, the homologous region would have to be sequenced in a gymnosperm to determine whether this gene was gained on the lineage leading to monocots and eudicots, or was present in the common ancestor of eudicots, monocots and Amborella and lost in the lineage leading to Amborella.
Figure 3
Synteny of the Amborella genome with other plant genomes. Illustrated here is a physical map of a 0.65 Mb region of the Amborella nuclear genome (highlighted in yellow) showing synteny with segments in each of the Arabidopsis, poplar, grapevine, and rice genomes. Two homologous segments are shown in each case: one above and one below the Amborella map. The physical map is based on high information content fingerprinting of an Amborella BAC library. Synteny was inferred over 5 Mb tracts of sequenced genomes on the basis of BAC-end sequences matching the reference genomes with TBLASTX bit scores of greater than 80. Red and green ovals depict BAC-end Amborella sequences with significant hits to known transposable elements and protein-coding genes, respectively.
divergence of Amborella? Was the common ancestor of Vitis, Populus, and Arabidopsis an ancient hexaploid that arose after the monocot-eudicot split? Did a separate genome-wide duplication occur early in monocot evolutionary history [8,11]? The answers to these questions are crucial for understanding angiosperm genome evolution and the diversification of flowering plants themselves. The Amborella Genome Project will address fundamental questions relating to the early evolution of gene content and genome structure in angiosperms (Figure 2), while providing comprehensive genomic resources for researchers studying all aspects of angiosperm biology [27].

In addition, two features of Amborella’s truly extraordinary mitochondrial genome raise compelling questions that warrant the sequencing of the Amborella nuclear genome. First, the Amborella mitochondrial genome is extraordinarily rich in ‘foreign’ genes acquired by horizontal gene transfer, far richer than any other plant mitochondrial genome [28]. These foreign genes were acquired from a wide range of donors. These findings raise important questions that can best be addressed with a complete nuclear genome sequence. For instance, is the Amborella nuclear genome also exceptionally rich in foreign sequences, and were these sequences acquired from the same donors as the foreign mitochondrial sequences? The Amborella nuclear genome sequence will enable subsequent experiments to determine what roles, if any, foreign nuclear genes play in Amborella. Second, the Amborella mitochondrial genome is exceptionally large, and much of the extra DNA is of unknown origin (Rice DW, Richardson AO, Young GJ, Sanchez-Puerta MV, Zhang Y, CWD, Knox EB, Munzinger J, Boone J, JDP, unpublished observations). We suspect that much of this unknown DNA was probably acquired from Amborella’s nuclear genome, a hypothesis that can only be tested once a complete nuclear sequence is available.

Ongoing deep transcriptome sequencing and physical mapping [26,29,30] form the foundation for this important project. Amborella cDNA sequences have already rooted gene trees and illuminated the timing of gene diversification relative to the origin of the angiosperms for many gene families [(31-34) and Duarte JR, Wall PK, Barakat A, Zhang J, Cui I, Landherr LL, Leebens-Mack J, Ma H, CWD, Kim S, et al., unpublished observations], and the potential for further evolutionary orientation of other gene families is great. The generation and analysis of a bacterial artificial chromosomes (BAC) fingerprint/end sequence physical map of the relatively small, 870 Mb Amborella genome [26] is already yielding new and exciting information about the genome structure of the earliest angiosperms and the retention of some syntenic blocks throughout angiosperm history (Figure 3). The physical map will also serve as a framework for assembling the sequence of the Amborella genome.

Given the available genomic infrastructure, the importance of Amborella as the sister to all other extant angiosperms, the large community of plant biologists who require a universal evolutionary reference for their studies, and the availability of cost-effective, ultra-high-throughput DNA sequencing technologies, it is our opinion that the Amborella genome is in an extremely strong position to warrant complete sequencing in the near future. Thus, the stage is set for a large-scale international Amborella genome sequencing initiative in support of fundamental and applied plant sciences, and we enthusiastically advocate such an endeavor.

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