Invited Review

History and future perspectives of barley genomics

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Abstract

Barley (Hordeum vulgare), one of the most widely cultivated cereal crops, possesses a large genome of 5.1 Gbp. Through various international collaborations, the genome has recently been sequenced and assembled at the chromosome-scale by exploiting available genetic and genomic resources. Many wild and cultivated barley accessions have been collected and preserved around the world. These accessions are crucial to obtain diverse natural and induced barley variants. The barley bioresource project aims to investigate the diversity of this crop based on purified seed and DNA samples of a large number of collected accessions. The long-term goal of this project is to analyse the genome sequences of major barley accessions worldwide. In view of technical limitations, a strategy has been employed to establish the exome structure of a selected number of accessions and to perform high-quality chromosome-scale assembly of the genomes of several major representative accessions. For the future project, an efficient annotation pipeline is essential for establishing the function of genomes and genes as well as for using this information for sequence-based digital barley breeding. In this article, the author reviews the existing barley resources along with their applications and discuss possible future directions of research in barley genomics.

Key words: Hordeum vulgare, genome sequencing, genetic resources

1. Introduction

1.1. Origin of genomic diversity in barley

Before barley (Hordeum vulgare ssp. vulgare) was domesticated, hunter-gatherers used the ancestral wild form (H. vulgare ssp. spontaneum) of domesticated barley as a human food source. Both wild and domesticated barley were found in archaeological sites in the Fertile Crescent dating back about 10,000 years, which is believed to be the origin of barley domestication. Wild barley kernels can be distinguished from those of domesticated barley by their brittle and smooth rachis. The ancestral wild form of barley shares the same genome as domesticated barley and is classified as a barley subspecies. Millions of generations of this wild barley provide a source of diversity to the present-day cultivated form, although domestication partially narrowed the diversity. Soon after domestication, mutations responsible for agronomically valuable traits, such as the jump from two to six row spike, spring growth habit, and hulless caryopsis, were selected for and spread quickly to all cultivated barley within a few thousand years. Allelic diversity at these loci corresponds with ecological conditions and with different uses of the cereal (human food, animal feed, and malt production) and made barley landraces well suited for cultivation throughout the world in most conditions, except in the tropics. Naturally occurring diversity in locally adapted landraces was the only source of available diversity until barley cross-breeding and mutation induction started in the early 20th century.

2. Relationships of plant species within the Poaceae family

Barley ranks fourth among grain cereals (Poaceae species) after maize (Zea mays), wheat (Triticum aestivum), and rice (Oryza
3. Genetic and genomic resources

3.1. Seed collection for barley

Barley is the only cultivated species of the genus *Hordeum*, which includes about 32 species and about 45 taxa. More than 485,000 accessions for the genus *Hordeum* are preserved at more than 200 different institutions worldwide. These collections include 299,165 accessions of *H. vulgare* ssp. *vulgare* (primarily new and old cultivars and landraces), 32,385 accessions of *H. vulgare* ssp. *spontaneum*, 4,681 accessions of wild species, and a substantial representation of genetic stocks, breeding lines, and mapping populations. Many accessions are duplicated between gene banks for safety or to avoid quarantine problems. The world’s largest seed storage facility is the Svalbard Seed Vault, which preserves over 1 million crop-related accessions, including 92,075 *Hordeum* accessions as of July 2020 (https://www.nordgen.org/sv/).

As an East Asian Center of barley genetic resources, Okayama University maintains 10,980 cultivars and landraces, 2,498 genetic stocks, and 628 wild barleys. Their collection, preservation, and distribution are partly supported by the National Bioresource Project (nbrp.jp). These materials have been collected since the 1940s for crop evolutionary studies at Okayama University. About 5,300 cultivated barley accessions have been intensively characterized both by genomic markers and for some agronomically relevant traits, such as resistance to powdery mildew. The data sets are available online from Barley DB (http://www.shigen.nig.ac.jp/barley/).

These collections are part of the South and East Asian subset of the international barley core collection, which constitutes the entire *ex situ* (stored) barley genetic diversity. Okayama University is responsible for the distribution of the South and East Asian subset, comprising 380 accessions. Other materials can be requested from the United States Department of Agriculture (USDA) small grain collection (Americas), the International Center for Agricultural Research in the Dry Areas (ICARDA, West Asia and North Africa and ssp. *spontaneum*), Leibniz-Institut für Pflanzengenetik und Kulturpflanzenforschung (IPK, Germany), and the Nordic Genetic Resource Center in Sweden (for wild *Hordeum*).

Barley has been the subject of much mutation research and breeding. These mutants have been mainly collected at the USDA small grain collection, located in Aberdeen, Idaho, USA, and at the Nordic Genetic Resource Center, in Alnarp, Sweden. To enhance the utility and accessibility of these mutants, >400 mutant alleles have been introgressed into the cultivar Bowman. Information on the Bowman introgression lines is available in the Barley Genetics Newsletter, Vol. 26 (http://wheat.pw.usda.gov/ggpages/bgn/26/bgn26t.html). These historical mutants provide a rich resource for functional studies and gene cloning.

Okayama University also maintains mutants, tetraploid lines, linkage testers, and near isogenic lines for barley. These lines were largely developed through the research activities at Okayama University and used for the development of linkage maps and genetic analysis of mutant traits. Tetraploids helped to develop a series of trisomic lines that can be used to identify the chromosome location of a given locus of interest. Currently, three mapping populations have been deposited from the North American barley genome mapping project. Another mapping population (Haruna Nijo × H602) was developed at Okayama University. These populations were instrumental in developing molecular genetic maps for barley and are being exploited for high-throughput mapping by single-nucleotide polymorphism (SNP) arrays. Recombinant chromosome substitution lines (backcross introgression lines) are also useful resources to identify genes in specific genomic regions to precisely study and map a locus of interest. Several sets of populations are available for distribution.

3.2. Genomic resources

Multiple genomic resources have been developed to analyse partial or total genomic sequences and their associated functions in barley.

Since 2000, several large barley expressed sequence tag (EST) projects have generated large numbers of sequences from expressed genes (http://harvest.ucr.edu/). Eight different genotypes provided the material for these projects. The polymorphisms identified between genotypes contributed to promoting high-throughput SNP genotyping, genetic mapping, and marker generation. The first comprehensive barley full-length cDNA (FL-cDNA) sequences were collected by Sato et al. mRNA samples were isolated from 15 organs and treatments from the Haruna Nijo cultivar and later pooled to develop a FL-cDNA library using the Cap-trapper method. A total of 5,006 clones were sequenced (http://www.shigen.nig.ac.jp/barley/). Another set of about 25,000 clones was generated and sequenced from the same cultivar, from 40 different organs and treatments. These sequences have allowed the annotation of genes on contigs or chromosome-scale genome sequences.

Barley geneticists have developed high-quality genetic maps, based on mutant phenotypes, using classical three-point linkage tests. These efforts are being updated and complemented with genome-wide genetic maps generated from molecular markers. Stein et al. developed a consensus barley map with 1,032 EST-based loci assayed using a combination of marker assays. Sato et al. developed a high-resolution barley EST map with 2,890 loci using a single-mapping population. High-throughput and high-quality multiplex PCR-based genotyping assays based on Golden Gate technology (Illumina Inc., CA, USA) involve the allele-specific detection of SNPs. A consensus genetic linkage map containing >2,900 gene-based SNP markers has been developed (http://harvest.ucr.edu/).
Several bacterial artificial chromosome (BAC) libraries have been constructed. The first library was developed from the Morex cultivar, it has been joined by other five Morex BAC libraries that collectively cover >2.5 genome equivalents. A library from the cultivar Haruna Nijo consists of 294,912 clones with an average insert size of 115.2 kbp and a coverage of about 6.6 genome equivalents. Another BAC library from wild barley accession H602 was developed at Okayama University (unpublished). Libraries from Haruna Nijo (coded as HNB) and H602 are available from the National Bioresource Project (nbrp.jp).

4. Genome sequencing

4.1. The nuclear genome

The International Barley Genome Sequencing Consortium (IBSC) was established in 2006 to generate a high-quality barley genome sequence. Two general approaches have been undertaken for the analysis of genome structure in barley: (i) identifying a minimum tiling path of 87,075 genetically mapped BAC clones and sequencing these clones (BAC-by-BAC strategy) and (ii) performing whole-genome shotgun sequencing.

The development of a BAC-based physical map for genome sequencing is a large-scale and worldwide effort. Madishetty et al. developed a high-throughput approach to use overgo probes to identify gene-containing BAC clones. They used >10,000 overgo probes derived from EST sequences to identify 83,381 gene-containing clones from the initial Morex BAC library. Fingerprinting of these clones resulted in contigs that comprise roughly two-thirds of all barley genes. Four new Morex BAC libraries have been prepared to cover an estimated 25 haploid genome equivalents. From these libraries, approximately 350,000 clones (covering about 14 genome equivalents) have been fingerprinted and assembled in contigs. To complement these efforts, BAC-end sequencing will be conducted on 350,000 BACs. In addition, there are efforts to integrate the resulting BAC contigs with the SNP-based genetic maps. Thus, a robust BAC-based physical map was integrated with the genetic map.

Of the 5.1 Gbp of the barley genome, IBSC developed a physical map of 4.98 Gbp, with more than 3.9 Gbp anchored to a high-resolution genetic map. Projecting a deep whole-genome shotgun assembly, EST, FL-cDNA, and newly developed RNA-seq data onto this framework supports 79,379 transcript clusters, including 26,159 assembly, EST, FL-cDNA, and newly developed RNA-seq data onto this framework supports 79,379 transcript clusters, including 26,159 transcript sequences will also contribute to the annotation of pan-genome assemblies (Stein et al., unpublished). As it is essential to isolate intact mature transcripts (mRNA) to obtain FL-cDNA sequences, techniques are being developed to maintain transcript integrity.

4.3. Transcriptomes

Deep sequencing of the transcriptome (RNA-seq) from the cultivar Morex and FL-cDNAs from the cultivar Haruna Nijo helped to annotate the reference genome of the cultivar Morex. The recent development of a single-molecule sequencing technique may also support the sequencing of long transcripts. To this end, an international collaborative project on full transcript sequencing for major haplotypes is underway (Waugh et al., unpublished). These long transcript sequences will also contribute to the annotation of pan-genome assemblies (Stein et al., unpublished). As it is essential to isolate intact mature transcripts (mRNA) to obtain FL-cDNA sequences, techniques are being developed to maintain transcript integrity.

4.4. Genomic information and available databases

The barley research community maintains a diverse array of databases that house information pertaining to barley genetics and genomics that can be easily accessed. Table 1 lists such databases and their respective information type.

For genome assembly and annotation, EnsemblPlants provides easy access to the most updated barley genome assembly, including chromosome sequences, genes, transcripts, and predicted proteins. The same website also supports Basic Local Alignment Search Tool (BLAST) searches against the barley genome. Similarly, IPK allows the user to conduct BLAST searches against all sequence resources published by the worldwide barley community, including activities related to the International Barley Sequencing Consortium.
Table 1. Barley genome databases

| Name                  | URL                                      | Function                      |
|-----------------------|------------------------------------------|-------------------------------|
| EnsemblPlants         | http://plants.ensembl.org/Hordeum_vulgare | Browser, BLAST                |
| IPK (IBSC) barley BLAST server | https://webblast.ipk-gatersleben.de/barley_ibsc/ | BLAST                          |
| PLEXdb                | http://www.plantgdb.org/prj/PLEXdb/      | Gene expression analysis      |
| HarvEST               | http://harvest.ucr.edu/                  | cDNA sequence                 |
| barleyGenes           | https://scs.hutton.ac.uk/barleyGenes/    | cDNA-seq data                 |
| bex-db                | https://barleylc.dna.affrc.go.jp/bexdb/  | cDNA, gene expression         |
| GrainGenes            | http://www.graingenes.org                | Markers, maps, mutants, etc.  |
| Barley DB             | http://www.shigen.nig.ac.jp/barley/      | Seed collection, cDNA sequence|

Several databases are specifically related to transcripts. PLEXdb is a public resource for gene expression analysis of plants and plant pathogens. This website currently hosts microarray data sets from a range of species, including barley and wheat. HarvEST is principally an EST database viewing platform that emphasizes gene function and is geared towards comparative genomics and oligonucleotide design, in support of activities such as microarray content design, functional annotation, and physical and genetic mapping. The site also allows the display of consolidated maps of approximately 3,000 SNP markers from four barley mapping populations and offers a rice synteny viewer. A subset of these mapped SNPs is integrated with the minimum BAC clone tiling path. barleyGenes provides access to predicted genes from an assembly of whole-genome shotgun sequences from barley (cultivar Morex). These genes were predicted from the mapping of RNA-seq data to the genome assembly. Gene expression levels were also calculated from the RNA-seq data and are available in the form of FPKM values associated with the predicted genes. bex-db provides 5′- and 3′-end sequences for 175,000 FL-cDNA clones as well as their expression data in a searchable database. The database also provides a genome browser, showing the locations of cDNA sequences on the barley genome.

Several databases contain barley genomic resource information. GrainGenes, the database of choice for legacy and classical genetics data, gathers information such as genetic maps, genes, alleles, genetic markers, phenotypic data, quantitative trait loci studies, experimental protocols, and publications about Triticeae species. Barley DB includes information on barley germplasm and barley genome resources from Okayama University. The database contains 5,006 FL-cDNAs and 134,928 EST entries. The barley germplasm collection from Okayama University and associated information are also available on the site.

5.2. Sequence accessions to summarize natural sequence variation

The IPK crop gene bank has a major world collection of barley seed samples. To estimate the diversity in genome sequences in wild and cultivated barleys, a genotyping-by-sequencing platform was applied to single plants from 21,405 accessions in the IPK barley collection. All haplotype reads were aligned to the reference genome from the barley cultivar Morex, which allowed the detection of 171,263 bi-allelic SNPs. A principal component (PC) analysis of cultivated barleys indicated that PC2 separated Eastern and Western barleys, whereas PC1 set Ethiopian barleys apart. Figure 1 shows that the geographical origin of collection agreed well with the SNP analysis. Representative accessions based on PC analysis are being used to analyse the barley pan-genome (Stein et al., unpublished), which may present a full complement of sequence variation within the barley genome.

5.3. Exome sequencing

Plant biologists and breeders concentrate most of their efforts on deciphering the functions of genes. Exome sequencing specifically targets coding sequences from the genome, and hence alleviating the need to sequence entire genomes and their repeat content, while bringing the computational needs and cost down to a reasonable range. The barley exome capture system selectively enriches for 61.6 Mbp of coding sequences based on the Morex cultivar HC (high-confidence gene model) sequences, FL-cDNA sequences, and de novo assembled RNA-seq consensus sequence contigs by hybridizing the gene-related fragments (or exons) from genomic libraries. The platform provides a highly specific and targeted capture of barley exons and closely related species. Using this platform, Russel et al. sequenced the exomes of a collection of 267 geo-referenced landraces and wild accessions of barley. A combination of genome-wide analyses demonstrated that the patterns of variation in barley have been strongly shaped by geography and that variant-by-environment associations for individual genes are prominent. A high-density 50 K SNP array with the Illumina Infinium whole-genotyping array was also developed, based on SNP design on exome capture data from 170 lines from a barley diversity panel.
caused barley grains to remain on the inflorescence at maturity, genes underlying important phenotypes that were isolated by our homology-based approaches. Here, we provide two examples of cloning strategies. In other cases, several approaches were combined homology-based cloning, whereas others were isolated via positional nomic and industrial traits. Some such genes were isolated through natural diversity remains a major source of agronomically relevant genomic resources established

5.5. Identification of useful genes for the genetic and genomic resources established

The use of pooled sequencing approaches may accelerate genetic mapping and identification of causal mutations. Mascher et al. applied exome sequencing to pooled samples from a barley mapping population segregating for the phenotype caused by the mnd (many noded dwarf) mutant, which increases the rate of leaf initiation. The pool of mutant plants should display a local and specific peak in read depth corresponding to the mutant genomic background around the candidate locus, which can be confirmed by the analysis of independent mutant alleles exhibiting the same phenotype.

Another example of mapping by exome sequencing borrowed from an existing pipeline for QTL-seq, which was initially developed for genome sequencing in rice. This `exome QTL-seq` approach allowed the mapping of the causal locus underlying the black lemma and pericarp (Blp) phenotype from a segregating population derived from doubled haploid barley lines. Exome sequences were assembled into pseudo-contigs by first-ordering exomes based on the genomic coordinates of their respective genes and then limiting each locus to 200 bp in the pseudo-map (but including all relevant SNPs). Short reads generated by the sequencing of the exome capture library are then analysed through this QTL-seq pipeline. The causal loci responsible for the trait of interest are identified based on the relative enrichment in SNP allele frequencies from their original genomic background, as described above for QTL-seq.

5.6. Variations in the proteomes useful for trait analysis

A unique example of an application of transcript/protein sequence information in barley is related to its industrial product: malt and beer. Iimure et al. conducted a two-dimensional gel-based proteome analysis to identify proteins associated with quality traits related to malt and beer production. Several protein species were identified in malt, wort (the first extraction step after malt maceration and mashing), and beer by gel electrophoresis, followed by trypsin digestion and mass spectrometry analyses and/or liquid chromatography tandem mass spectrometry. In addition, low-molecular-weight polypeptides were isolated from beer by the combination of non-enzymatic digestion and mass analysis. Collectively, these data sets of polypeptides from barley proteomes provide a platform for analysing protein functions in beer. Several novel proteins related to beer enzyme digestion and mass analysis. Collectively, these data sets of polypeptides from barley proteomes provide a platform for analysing protein functions in beer. Several novel proteins related to beer.

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![Figure 1. Principal component analysis of 19,778 domesticated barleys based on 76,102 genotyping-by-sequencing markers from the IPK Bridge Web Portal (https://bridge.ipk-gatersleben.de/bridge). Accessions from Germany (left), Japan (middle), and Ethiopia (right) are plotted on PC1 (y axis) and PC2 (x axis). The proportion of variance explained by the PCs is indicated on the axes.](https://bridge.ipk-gatersleben.de/bridge)
have also been turned into efficient protein or DNA markers for trait selection in malting barley breeding.54

5.7. Transformation, genome editing, and functional validation of identified genes
As described above (see description of the brt1/brt2 and Qsd1 loci), introducing a gene of interest by stable transformation is the standard technique for validating the gene responsible for the target trait. Targeted genome modification technology (so-called genome-editing) offers tremendous promise as a technology that can efficiently produce mutations in desired genes, and several cases of barley genome-editing by clustered regularly interspersed short palindromic repeats (CRISPR) and the CRISPR-associated nuclease Cas9 have already been reported.55 However, transformation and genome-editing experiments may suffer from some limitations resulting from the low transformation potential of some accessions. Indeed, the old Scottish malting cultivar Golden Promise is one of the few reliable haplotypes for Agrobacterium (Agrobacterium tumefaciens)-mediated transformation. Hisano and Sato56 identified loci controlling transformation amenability in the regions of chromosomes 2H and 3H in an F2 population derived from a cross between the cultivars Golden Promise and Haruna Nijo. Introducing these genomic regions in target haplotypes may increase their transformation efficiency and genome-editing capabilities.

5.8. The digital bioresource project for digital breeding
Natural and induced variation provides opportunities to analyse traits of interest. However, how to combine sequence and trait information remains a challenging question that needs to be addressed. High-quality genome sequences are essential to establish a digital bioresource centre for world barley ex situ collections but are unfortunately also insufficient. To better understand the contribution of genomic sequence variation to various traits, automatic sequence annotation must become faster and more efficient, since this process currently relies on a slow and manual data curation step. Automated phenotyping is also emerging as an essential tool to diminish the bottleneck associated with the characterization of increasing numbers of accessions. Systematic gene inactivation or modification by genome editing may provide a functional picture of candidate genes of interest in the target haplotype. However, agronomically important traits are often controlled by multiple interacting genes, which demands a deep knowledge of trait-based genetics. Finally, the combined information collected from genomic sequences and the systematic functional analysis of genes may provide novel strategies for trait improvement, for example, the sequence-based digital breeding of barley.

6. Conclusion and future perspectives
The genomes of various organisms have been sequenced and analysed for some 40 years. During this period, development and application of new technologies including computer software frequently changed sequencing strategies. Initially, most plant genome sequencing projects had to rely on a single haplotype to establish a reference genome. However, it has become clear that to sequence multiple haplotypes in parallel would be essential for our understanding of the genetic and genomic features underlying the natural and induced variations. Newly developed technologies of various sorts have allowed us to make it quite efficient to analyse multiple accessions and lines simultaneously, even for species possessing a large genome size such as barley.

However, we realize at the same time that we need to establish genetically stable reference accessions to obtain consistent results for genomic and proteomic sequencings. In these regards, a digital bioresource including sequence and related information of all accessions appears to be quite challenging. In the meantime, however, sequencing of a limited number of accessions that cover most of the major sequence variations in a crop species may provide an alternative and efficient strategy to establish the ‘pan-genome’.

High-quality sequence assembly will certainly be an indispensable component of the pan-genome infrastructure. The ongoing attempt of a barley pan-genome project aims to construct chromosome-scale sequence assembly for 20 genotypes, consisting of landraces, cultivars, as well as a wild barley accession selected to represent the global barley diversity. The details of the project will be published in the near future which is expected to present an advanced view of barley genomics.

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Conflict of interest
The author declare that there is no competing interest.

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