SURVEY AND SUMMARY

Sequence capture by hybridization to explore modern and ancient genomic diversity in model and nonmodel organisms

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ABSTRACT

The recent expansion of next-generation sequencing has significantly improved biological research. Nevertheless, deep exploration of genomes or metagenomic samples remains difficult because of the sequencing depth and the associated costs required. Therefore, different partitioning strategies have been developed to sequence informative subsets of studied genomes. Among these strategies, hybridization capture has proven to be an innovative and efficient tool for targeting and enriching specific biomarkers in complex DNA mixtures. It has been successfully applied in numerous areas of biology, such as exome resequencing for the identification of mutations underlying Mendelian or complex diseases and cancers, and its usefulness has been demonstrated in the agronomic field through the linking of genetic variants to agricultural phenotypic traits of interest. Moreover, hybridization capture has provided access to underexplored, but relevant fractions of genomes through its ability to enrich defined targets and their flanking regions. Finally, on the basis of restricted genomic information, this method has also allowed the expansion of knowledge of nonreference species and ancient genomes and provided a better understanding of metagenomic samples. In this review, we present the major advances and discoveries permitted by hybridization capture and highlight the potency of this approach in all areas of biology.

INTRODUCTION

The emergence of next-generation sequencing (NGS) technologies has radically revolutionized medical, biotechni-
capture approaches have some limitations. Indeed, their application is not suited for genomic regions that are several megabases in size because of the multiple primer pairs or probes required to cover large regions and the difficulty of designing them with the necessary high sequence specificity. Similarly, the number of target biomarkers is also limited because of primer design, which quickly causes the application of these strategies to become fastidious and expensive. Finally, neither of these approaches has been developed into commercially available products, making them more expensive and less amenable to a broad range of research applications.

Hybridization capture coupled with high-throughput sequencing has been developed to overcome these limitations, allowing the rapid and simultaneous detection of numerous and potentially large targeted DNA regions (Figure 1). The hybridization between the sequencing library and probes can be performed in solution or on a solid support. In solid-phase hybridization capture, herein referred to as array-based hybrid selection (AHS), DNA probes are bound to a solid support such as a glass microarray slide (11,12). DNA libraries are applied to the surface of the support, and targeted DNA fragments hybridize with the immobilized probes. Nonspecific unbound molecules are washed away, and the enriched DNA is eluted and amplified for downstream direct sequencing. In solution hybridization capture, also referred to as solution-phase hybrid selection (SHS), free DNA or RNA probes are biotinylated to enable the selection of targeted fragment-probe heteroduplexes using magnetic streptavidin beads (13). Non-targeted library fragments are removed from the liquid phase through repeated washes, and targeted fragments are then eluted from the beads and amplified for subsequent sequencing. Methods relying on hybridization are the most widely used, with numerous available cost-effective commercial platforms (14–17). Because these approaches employ longer probes than PCR methods, they are generally less specific and show a greater tolerance for the discovery of polymorphisms (13). Furthermore, hybridization capture is highly scalable to targets of different lengths, equally enriching short discontinuous targets or long genomic regions (18), and is highly multiplexable through the use of barcodes to pool many libraries together.

Hybridization capture methods allow the effective enrichment of a wide range of targets and offer a good compromise between specificity, multiplexing and scalability. Additionally, they match the scale of current high-throughput NGS and have become reference methods for resequencing efficiently contributes to the characterization of genetic variations in clinical genetics, agronomical traits and environmental adaptations over time and space. Targeted sequencing directed to the protein-coding portion of the genome (exome) or the captured transcriptome improves the resolution for rare variants detection. Overlooked off-targets sequences obtained during sequence capture by hybridization provide important source to study the unknown flanking regions (new variants, virus and mobile genetics elements integration sites, regulatory signals). These approaches are also very efficient even for degraded samples. Ancient DNA (aDNA) extracted from paleontological remains or museum specimens has been explored for understanding the evolutionary history of numerous lineages (humans, livestock, pets, wild animals, fish, insects, plants, microorganisms) of both extinct and extant species. Hybridization capture is also used to reveal new sequences for a broad range of organisms even distantly related and without reference genomes. Some erroneous relationships in the previously established phylogeny and identifying of new clades have been highlighted. Complex microbial ecosystems could also be efficiently explored allowing to access to complete phylogenetic or functional biomarker diversity as well as providing a first glimpse into the identification of metabolic capabilities through the association of biomarkers. In this review, we present the major hybridization capture experiments and results in the above-mentioned fields and illustrate how powerful this approach can be for revealing information that is inaccessible or very difficult to obtain when using other conventional molecular approaches.

RESEQUENCING OF LARGE GENOMIC REGIONS

Based on knowledge of reference genomes, the application of NGS to the genomes of particular individuals gives rise to an exhaustive list of genomic variations. Therefore, DNA resequencing can efficiently contribute to the characterization of genetic variations such as single-nucleotide polymorphisms (SNPs), insertions and deletions (indels), structural variants (inversions, translocations) and copy number variants (CNVs) (8). Nevertheless, the high sequencing depth required to distinguish real variants from sequencing errors, occurring at a rate of 0.1–1% (19), with high reliability, makes this approach difficult and expensive (2). Moreover, complete genome resequencing is not always necessary when only particular genes or genomic regions are of interest. Conventional resequencing approaches based on bidirectional sequencing of targeted PCR amplicons are particularly well suited for the detection of SNPs but are less well suited for the detection of other variations and are not sufficient because of the high levels of multiplexing required when large or numerous genomic regions have to be characterized (20). Consequently, resequencing has evolved toward the hybridization of DNA to resequencing microarrays containing tiling probes designed to detect interesting genomic variants (21). This approach is commonly applied in genome-wide association studies (GWASs) to simultaneously detect thousands of SNPs across a complete genome associated with a studied phenotype (22). However, GWASs are generally restricted to common genomic variants, excluding rare and undiscovered genetic modifications (8), and are principally relevant for the detection of SNPs. Thus, gene capture approaches by hybridization using tiling probes have been developed to specifically enrich and resequence numerous meaningful subsets of the genomes of multiple individuals with a high depth and to better associate genotypes and phenotypes in a cost-effective manner (Figure 2A).
Figure 1. Principle of hybridization capture. A sequencing library containing targeted biomarkers is constructed and hybridized against a set of specific probes. Hybridization can be performed either in solution (solution hybrid selection, SHS) with biotinylated probes captured by streptavidin-coated magnetic beads, or on a solid support (array-based hybrid selection, AHS) on which probes are spotted. After hybridization, nontarget sequences are washed away, and the enriched sample is eluted and sequenced.

Pioneering studies using hybridization capture were carried out by Okou et al. (12) and Albert et al. (11), who applied high-density microarrays with tiling probes for the enrichment and resequencing of large genomic regions on the order of ten to hundreds of kilobases. Okou et al. (12) studied the enrichment of coding and noncoding regions of human genes involved in fragile X syndrome. Similarly, Albert et al. (11) resequenced areas of 200 kb, 500 kb, 1 Mb, 2 Mb and 5 Mb surrounding the human BRCA1 gene involved in Burkitt’s lymphoma. Likewise, Gnirke et al. (13) developed an SHS method that simultaneously targeted four human genomic regions ranging from 220 to 750 kb in size. These approaches have been used by Beaudoin et al. (23) to resequence 55 genes known to be associated with ulcerative colitis, by Ruark et al. (24) to target 507 genes implicated in DNA repair in association with breast cancer and ovarian cancer, by Clarke et al. (25) to discover SNPs associated with traits of agronomical and nutritional importance through 47 Brassica napus specific genomic regions, or even by Schiessl et al. (26) to explain phenological variations in flowering in Brassica through deep resequencing of 29 regulatory genes. Similarly, Tennessen et al. (27) used hybridization capture to decipher origins and dynamics of strawberry octoploid subgenomes. Jupe et al. (28) discovered and annotated pathogen resistance gene family members, Fu et al. (29) evidenced SNPs in maize genome and Faucon et al. (30) detected variants involved in Aedes aegypti insecticide resistance in more than 760 candidate genes. More recently, the efficacy of capture coupled to third-generation sequencing has been demonstrated by delineating the breakpoint junctions of low copy repeat associated complex structural rearrangements in patients diagnosed with Potocki–Lupski syndrome (31).

The potential of hybridization capture to accurately detect genomic variants in large specific genomic targets, as demonstrated in the congruous studies mentioned above, opened the door for its application to a broader range of biomarkers, rather than a single gene or a limited number of restricted regions. Hybridization capture has therefore been
Figure 2. Applications of hybridization capture. (A) Capture for resequencing. Specific tiling probes are used to capture genomic regions of interest and to identify sequence polymorphisms. (B) Exome capture. Tiling probes targeting the whole exome are used to identify coding mutations in the genome. (C) Off-target capture. Tiling probes are used to enrich a biomarker whose indirectly captured flanking regions are of interest. (D) Nonreference species capture. Tiling probes designed based on divergent genomes are used to capture biomarkers or genomes of nonreference species. (E) Ancient DNA capture. Tiling probes designed based on modern species are used to enrich ancient DNA and reconstruct biomarkers or genomes. (F) Metagenomic sample capture. Nonoverlapping exploratory probes are used to enrich all variants of a biomarker from a complex sample to study its diversity and its genomic context.

EXOME CAPTURE

Exome capture, also known as whole exome sequencing (WES), is targeted sequencing of the protein-coding portion of the genome. This method employs capture by hybridization with exon-specific tiling probes to target the protein-coding variants in the best understood subset of the genome (Figure 2B) (32). Indeed, the exome represents approximately 1% of the human genome, corresponding to a total of approximately 30 Mb, but harbors 85% of the known genetic disease-causing mutations (33,34). Moreover, gene exons are more likely to contain pathogenic variants than introns or regions between genes (35–37), and they harbor fewer repetitive elements, making their analysis easier (38).

Exome capture has rapidly become a standard practice in clinical genetics for determining the basis of human dis-
In addition to the common variants associated with MI risk, APOA5 genes involved in myocardial infarction (MI), therefore particularly difficult (52). However, the detection of rare carcinogen genetic alterations in these complex samples through conventional molecular approaches is therefore particularly difficult (52). However, the detection of subclonal mutations can be readily achieved by exome capture, as illustrated by Agrawal et al. (53) in a survey of head and neck squamous cell carcinoma (HNSCC).

In addition to allowing the detection of causal variants involved in genetic disorders and opening new horizons for medical treatments, WES has been shown to be an effective tool for medical diagnosis (54). Worthey et al. (55) successfully used exome capture to diagnose a patient through the identification of gene variants for the first time. In these cases, despite a comprehensive clinical evaluation suggesting an immune defect, clinicians were unable to establish a definitive diagnosis, thereby limiting the medical management of the patient. The use of WES revealed a hemizygous missense mutation in the gene responsible for the X-linked inhibitor of apoptosis deficiency. In a different context, through the analysis of mutations detected using WES, Choi et al. (56) fortuitously diagnosed congenital chloride diarrhea in a patient thought to be suffering from Bartter syndrome, a renal salt-wasting disease. In addition, a major project emerged 5 years ago: the Deciphering Developmental Disorders (DDD) study (57). The purpose of this study is to determine genetic variants underlying severe undiagnosed developmental disorders in 12 000 children through multiple complementary genome-wide approaches, including exome capture. Thus far, the exomes of more than 1000 children have been sequenced, and more than 30% of the children have been diagnosed, demonstrating the potential of WES in this field (58). This approach benefits from large databases of pathogenic variants of human genes, such as that of the Exome Aggregation Consortium (ExAC), which combines exome sequencing data from more than 60 000 individuals with various diseases (http://exac.broadinstitute.org), and from bioinformatics tools linking mutations discovered through WES back to a medical diagnosis (59, 60).

Exome capture has also been used to sequence the messenger RNA (mRNA) fraction as complementary DNA (cDNA) in human medical studies to extend information obtained from DNA-based investigations and reveal information that is inaccessible based on analysis of DNA alone. This method provides an interesting alternative to RNA-sequencing (RNA-Seq) of mRNA, allowing targeted enrichment of the complete coding fraction of the transcriptome. Mercer et al. (61) and Halvardson et al. (62) demonstrated that, due to the deep coverage of exome capture, this method reveals rare and unannotated transcripts whose expression levels are below the detection limits of conventional sequencing approaches. Moreover, Cabanski et al. (63) and Cieslik et al. (64) showed that exome capture targeting coding transcripts improves performance compared with RNA-Seq for degraded samples for which either limited amounts of RNA are available or polyA selection is arduous, respectively. They demonstrated that genetic variants and gene fusions are accessible through exome capture and that the obtained absolute and differential gene expression levels are accurate for tumor samples, which has also been shown by Clark et al. (65) for long noncoding RNAs. Therefore, exome capture appears to be a complete tool for medical resequencing, providing exhaustive information necessary for determination of the genetic basis of various pathologies and their diagnosis.

Exome capture has also been used in many other model organisms for different purposes. For example, WES has
been employed for to variant identification in medical model species, such as mice (66), pigs (67) and dogs (68,69), providing valuable tools for disease association studies. Moreover, as not all protein-altering variants induce diseases (70), exome capture has also been used for the detection of variants responsible for modifications in phenotypic traits of economic importance in crops and livestock. Mascher et al. (71) and Saintenac et al. (72) explored natural variation and genetic diversity respectively in barley and in wheat. Muraya et al. (73) applied exome capture in maize to detect sequence variation affecting biomass production. Other studies have investigated the cattle exome to determine gene variants associated with milk production (74), fertility (75) and growth (76). Similarly, the *Eucalyptus* exome has been studied, to identify gene mutations responsible for wood property traits (77). This method has also been employed in black cottonwood (78), wheat (72,79), barley (71,80) and switchgrass (81) to elucidate genomic variation in these model species and to better link genotypes and phenotypes for further agronomic improvements. Another similar application of exome capture in this field is for the identification of variants generated through mutagenesis to investigate gene function. Indeed, random mutagenesis approaches induce various unknown mutations affecting phenotypes. WES has been effectively used in such agronomic studies to link phenotypes back to gene mutations, replacing comparative genomic hybridization microarrays (82) and bringing variant detection to the whole genome level as demonstrated by Henry et al. (83) and King et al. (84). Similarly, Bolon et al. (85) successfully used exome capture to detect deletion mutations induced in 120 000 soybean seeds through fast neutron radiation. After phenotypic screening based on traits of interest such as seed composition, maturity, morphology, pigmentation and nodulation, these authors identified candidate genes, allowing the determination of agronomically important markers for soybean crop breeding.

These applications of exome capture in a wide range of studies, from discovering the origins of genetic disorders to determining key agronomic biomarkers, and in a broad spectrum of species, illustrate the potential of hybridization capture for the resequencing of this genomic subset. Nevertheless, the information obtained from the exome remains incomplete. Therefore, many studies have taken advantage of the off-target sequences obtained from hybridization capture to increase the amount of information generated from experiments.

**OFF-TARGET CAPTURE**

While exome sequencing primarily targets exons, a significant portion of the captured fragments come from outside of these coding regions (86,87) (Figure 2C). Indeed, the hybridization process selects for fragments that contain at least a portion of targeted exons, resulting in enrichment of flanking regions such as introns, untranslated regions (UTRs), promoters and intergenic regions. In a typical exome sequencing experiment, these off-target sequences represent approximately 40–60% of the obtained reads (e.g. 31,32,37,78,79); this percentage can be related to many different factors, such as the exome capture platform, the size of DNA libraries and sequencing technologies (4,14). The off-target sequences generated are usually excluded from subsequent analyses. However, these sequences provide a potentially important source of unknown information that could greatly extend the utility of exome sequencing.

Following the study initiated by Guo et al. (88), numerous researcher have devoted attention to these overlooked off-target sequences. Indeed, these studies have demonstrated that reads and SNPs outside of targeted regions coming from different exome capture platforms are of good quality, suggesting that these data can be exploited. They have also shown that through the exploitation of complete sequencing data, it is possible to more than double the number of high-quality SNPs in comparison with those that are made available by focusing on exome regions. Indeed, they found that approximately 50% of the SNPs identified from exome sequencing were in target regions, while 27% were in the 200 bp flanking regions, and the remaining 23% were in regions >200 bp away from exons, illustrating that sequences falling outside of the targeted regions should not be ignored in data processing. A good illustration of a subsequent off-target investigation is a study on the adaptation of residents of the Tibetan Plateau to high altitude through exome capture (89). The authors identified SNPs located in different genes, and showed that the SNPs exhibiting the greatest difference in frequency compared with low-altitude residents were intronic, suggesting the significance of introns and supporting the importance of off-target regions.

In many studies, off-target WES reads have been shown to be a reliable and rich source of human mitochondrial DNA (mtDNA) because of the relative high abundance of the corresponding genome within the total DNA (90). Indeed, each mammalian cell contains approximately 100 mitochondria, which themselves contain between two and 10 copies of the 16.6 kb closed-circle mtDNA genome (91). Thus, even if the mtDNA genome is not the target, it can easily be obtained through indirect methods such as exome capture with an average coverage of 100× (90) and totalling approximately 1–5% of the obtained sequencing reads (86). Therefore, off-target mtDNA represents a precious source of information for many sequencing investigations, as demonstrated by Griffin et al. (92) and Zhang et al. (93), who showed that the mtDNA obtained through WES allows reliable detection of sequence variants with a low error rate, similar to conventional mitochondrial DNA Sanger sequencing. This capacity of exome capture to reliably enrich both targeted exome and off-target mtDNA has been applied for the detection of mutations involved in mitochondrial diseases. For example, through performing this simultaneous dual-genome analysis and identifying mutations in both exome and mtDNA, Dinwiddie et al. (94) successfully diagnosed four patients with distinct mitochondrial disorders for whom conventional molecular testing had previously failed to produce a diagnosis. Such evaluations of mtDNA based on off-target data can be facilitated by dedicated bioinformatics tools such as MitoSeek, which extracts mitochondrial genome information from exome data and analyses associated mutations and structural variants (95).

Based on the potency of this method for expanding knowledge beyond target DNA regions, studies have used...
hybridization capture strategies to study the unknown flanking regions of genes or particular genomic regions. For example, Duncavage et al. (96) employed a capture strategy to identify the sites of Merker cell polyomavirus (MCPyV) integration in the genomes of Merkel carcinoma patients. Targeting the virus genome, these authors achieved up to 37,000-fold coverage of the MCPyV genome, making detection of viral insertion sites possible through the analysis of off-target flanking regions. Similarly, Ma et al. (97) investigated cis-regulatory elements (CREs) surrounding plant microRNA (miRNA) genes via miRNA hybridization capture. They demonstrated the specificity and sensitivity of this enrichment method in *Arabidopsis thaliana* by assembling regions spanning 10–20 kb flanking miRNAs. Finally, Platt II et al. (98) identified retrotransposon insertion sites in the bat genus *Myotis* through this approach to construct a robust transposable element insertion-based phylogeny.

Therefore, off-target data appears to be a rich source of diverse information, making the study of nontargeted reads an important field to explore. Nontargeted reads may contain information that is not covered in the current reference genomes and may also provide information about flanking sequences that are yet to be discovered, leading to new insights into genome exploration. These findings show that hybridization capture can expand our knowledge beyond the currently available information on studied organisms, and this potential has been demonstrated through the application of this strategy to enrich genomic regions of interest from organisms without reference sequences or genomes.

**NONREFERENCE SPECIES CAPTURE**

Despite the intensive use of NGS technologies, there are still no established reference genomes or draft genomes for most nonmodel organisms. Accessing the genetic makeup of these organisms, referred to as nonreference species, can be achieved through *de novo* whole-genome sequencing, but sequencing costs and difficulties in assembly can be prohibitive, without necessarily providing access to complete information (99). Therefore, hybridization capture represents a good alternative for accessing targeted informative subsets of these genomes (Figure 2D). Nevertheless, these methods require *a priori* knowledge of the targeted sequences which is, by definition, unavailable for nonreference species. Different probe design strategies for enriching genomic regions of interest have been developed to overcome this limitation and hybridization capture has been used to reveal new sequence information for a broad range of organisms (99). The first approach for probe design circumventing the need for a reference genome relies on PCR amplification. Due to the large amount of sequence data and numerous primers available in the literature and in databases, obtaining PCR primers for informative loci has become quite easy. These PCR products amplified from a small set of targeted loci can thus be used as capture probes for nonreference species (100). Another strategy leverages partial sequencing data to design probes. Thus, the design can be generated based on low-coverage *de novo* DNA sequencing assemblies (27,101), RNA-Seq transcriptome (94–97,102), expressed sequence tag (EST) data (103–105) or clone libraries (106) which provide core information about the targeted species. Finally, probe design can be based on orthologous genomic regions between divergent reference organisms (107–109). In this case, enrichment relies on the ability of probes designed according to a single reference genome to specifically capture DNA from a broad range of other species. The impact of sequence divergence on the efficiency of hybridization capture has been evaluated in many model and nonmodel species (e.g. 64,73–77).

For example, Hedtke et al. (110) used exome capture to enrich DNA across frog species spanning approximately 250 million years of evolutionary divergence (up to approximately 10% divergence), thus testing the limits of this approach. They demonstrated that the success of enrichment depends on the divergence time between a given frog species and the reference sequence and that even if there is some missing data, the method appears to be effective between organisms that shared a common ancestor up to 200 million years ago.

Hybridization capture for nonreference species has been proven to be an appealing method addressing various biological issues. It was first used to greatly expand the repertoire of available genomic sequences for nonmodel organisms. Indeed, as illustrated by Neves et al. in two different studies (104,111), hybridization capture through exome targeting can efficiently enrich large portions of complex genomes. In these studies, hybridization capture data provided information to guide the assembly of reference lobolly pine genome which was previously largely uncharacterized despite attempts at sequencing, and significantly improved our knowledge of the pine gene space. Similarly, Rosani et al. (112) made use of hybridization capture to increase current knowledge of the unknown Mediterranean mussel genome, George et al. (113) applied this methodology to generate high-quality, full-length sequences for nonhuman primate species and Seoane-Zonjic et al. (114) used it to establish gene structures without reference genomes in the maritime pine mega-genome. Hybridization capture using probes targeting genomic regions from divergent nonreference species has also been broadly employed to define the gene variants at the origin of the process of species divergence. Using this approach, Hebert et al. (103) identified SNPs in genes potentially involved in whitefish divergence and reproductive isolation; Nadeau et al. (115) captured patterns of divergence in radiating geographical species of Heliconius butterflies; and Cosart et al. (116) identified thousands of candidate SNPs in livestock and wild cattle species, revealing the detailed genetic basis of adaptive differentiation and speciation. Similarly, with this approach, Schneider et al. (117) identified mutations at the origin of polymorphism for melanism in three South American feld species. Finally, this hybridization capture approach for nonreference species can also provide information in the field of phylogenetics. Indeed, it enables the collection of data on hundreds of biomarkers across the genomes of many individuals, which is necessary for the establishment of robust phylogenetic relationships. For example, Stephens et al. (118) used this approach to enrich approximately 200 specific nuclear genes from 75 accessions of the North American carnivorous pitcher plant genus *Sarracenia*. Using the multilocus data obtained through hybridization capture, these authors elucidated relationships
within the *Sarracenia* genus, which had not been possible in previous attempts using conventional approaches targeting only a few informative sites. Similarly, based on 259 targeted nuclear loci, Prum *et al.* (119) reconstructed the phylogeny of 198 living bird species, revealing some erroneous relationships in the previously established phylogeny and identifying many new clades. In two analogous studies, Faircloth *et al.* (120,121) designed probes targeting ultraconserved nuclear DNA element (UCEs) that are shared among ray-finned fishes and the insect order Hymenoptera. With these UCEs loci, the authors accurately resolved phylogenetic relationships at both shallow and deep time-scales within the studied organisms. Likewise, in a study conducted by Wang *et al.* (122), hybridization capture was used to sequence mtDNA segments from different accessions of 36 pine species and to gain insights into the mtDNA-based phylogeny of the genus *Pinus*. They revealed that this phylogeny differed from that based on chloroplast and nuclear DNA and, thus, identified a series of mtDNA capture events during pine evolution.

The power of hybridization capture in the field of nonreference species is such that the method has been widely and successfully used to study highly divergent samples from extinct organisms or historical remains.

**ANCIENT DNA CAPTURE**

Ancient DNA extracted from paleontological remains or museum specimens is a rich source of information for understanding the evolutionary history of lineages of both extinct and extant species (123–125). However, because of postmortem preservation conditions, aDNA samples have particular characteristics that make them difficult to analyse using conventional molecular approaches. Indeed, most ancient samples contain short DNA fragments averaging 30–300 bp (126), resulting from the degradation of genomic DNA by cellular endonucleases and exogenous micro-organisms that fragment DNA and depending strongly on the specific geochemistry of the environment (18). Consequently, ancient samples contain a low percentage of endogenous DNA and are highly contaminated by environmental bacteria and fungi (127–129). For example, one study revealed that 55,000-year-old bone fragments from a Neanderthal individual were composed of a complex mixture of approximately 1% degraded human DNA and 99% contaminant environmental DNA (130). Only ideally preserved permafrost remains exhibit little damage and can yield fragments of nearly 2 kb, with less contamination (131). Another significant source of contamination of aDNA samples is modern human DNA introduced during excavation and museum or laboratory handling (124,132). Finally, postmortem nucleotide changes occur and introduce errors in sequencing reads, thus requiring particular attention during data analysis. The most common changes are cytosine deamination to uracil, which is then converted to thymine by DNA polymerase, and the transition of guanine to adenine, which can be used to reliably differentiate between ancient and contaminating modern DNA sequences (18,133,134).

Until recently, aDNA studies were restricted to PCR amplification of short overlapping DNA fragments due to the length of aDNA fragments. This approach enabled the reconstruction of long DNA sequences and mitochondrial genomes, but hindered complete genome reconstruction (135). The introduction of NGS, based on the sequencing of short sequencing reads that are problematic for PCR amplification, has consequently greatly facilitated aDNA studies (94–96). Nevertheless, because of the very low proportion of endogenous DNA in ancient samples and the considerable sequencing effort required to access it, NGS approaches are usually not sufficiently efficient or financially unfeasible. Therefore, to overcome these methodological limitations, hybridization capture approaches have often been employed (Figure 2E). Due to the ability of hybridization capture to specifically enrich targeted DNA from massive amounts of contaminating environmental DNA with probes designed based on very distant species, this strategy has quickly become a gold standard for ancient aDNA research. Its utility was first demonstrated by Noonan *et al.* (136) in a pioneering study showing that it was possible to reconstruct up to 65 kb of nuclear DNA targets from a Neanderthal library using hybridization capture, compared with only 5 kb when using conventional approaches. The efficiency of hybridization capture has since been widely demonstrated, and was especially well illustrated by Carpenter *et al.* (129) who showed that this method enabled 6- to 159-fold enrichment of ancient human DNA using modern human DNA probes and that targeted DNA represented up to 60% of the obtained reads, versus the average of 1.2% observed before hybridization capture. This revolutionary approach in the field of paleogenomics has been widely employed to effectively target full mitochondrial genomes (e.g. 128,131) as well as nuclear DNA (e.g. 129,132) from human, animal, vegetal or microbial species.

This technological advance in the area of aDNA research has allowed better characterization of ancient extinct organisms and led to more precise definition of phylogenetic links between ancient and modern species. For example, Castellano *et al.* (137) used hybridization capture to enrich the exomes of two Neanderthals from Spain and Croatia. Comparisons with other genomes revealed that the diversity among Neanderthals was remarkably low compared with that found in modern humans and that amino acid substitutions underlie the phenotypic differences between Neanderthals and present-day humans. Similarly, Burbano *et al.* (138) generated data from megabase-size genomic regions from Neanderthal DNA samples. Comparison of these data with sequences of present-day humans at the same positions allowed them to identify 88 amino acid substitutions that have occurred since the population split of modern humans and Neanderthals. In nonhuman species, this approach enabled the resolution of many evolutionary histories through the capture of mitochondrial genomes from aDNA. Llamas *et al.* (139) provided a reliable phylogeny of the Australian megafauna and clarified the phylogenetic position of the extinct giant short-faced kangaroo and giant wallaby, while Vilstrup *et al.* (140) elucidated the taxonomic radiation that gave rise to modern horses, zebras and donkeys by applying a capture strategy to three extinct equid lineages. In addition, hybridization capture provides additional information on the adaptation of species to their environment over time. For instance, Immel *et al.* (141) targeted and recon-
conceived these limitations through the shotgun sequencing of metagenomic samples, which enabled the association of micro-organismal community structure with realized metabolic functions (156,157). However, the establishment of such link based on large genomic sequences or genome reconstruction requires sufficient sequencing coverage to be available for dominant micro-organisms, or a considerable sequencing effort to access less abundant and rare organisms (158).

In contrast, Single Cell Genomics (SCG) recovers genomes from targeted individual cells present in complex environments and precisely identifies their metabolic functions; however, this technique is not always practical and is not sufficiently scalable to provide an exhaustive overview of microbial communities (159,160). More innovative, but still anecdotic, targeting enrichment approaches in the field of metagenomics, such as hybridization capture, have been developed to fill this gap in the exploration of microbial communities (Figure 2F). Indeed, although these communities are more complex considering that they are composed of a multitude of organisms, metagenomic samples are similar to nonreference species and aDNA samples in the sense that the genetic information they contain is only suspected and is not well known. The demonstrated potential of hybridization capture to enrich divergent sequence targets in such cases has therefore been applied to metagenomic samples to enrich biomarkers of interest that are known to be present but, whose variability and diversity are not known (158).

Denonfoux et al. (161) described the first adaptation of SHS to a complex metagenomic sample. The methodology was used to explore the methanogenic communities present in a lacustrine environment by targeting the methyl coenzyme M reductase subunit A (mcrA) gene with a set of nonoverlapping probes, which targeted both known sequences and potential undescribed variants of the mcrA gene. The mcrA sequences represented more than 40% of the obtained sequences after two cycles of capture, revealing enrichment compared with shotgun sequencing, in which only 0.003% of the sequences corresponded to the target gene. This approach efficiently enriched complete targets but also recovered off-target sequences adjacent to the mcrA gene such as parts of the mcr operon. The extended mcrA flanking regions showed evidence of the association of the mcr operon with Fe metabolism genes that are usually located tens or hundreds of kilobases downstream. This finding suggested that this genomic organization, which has never been described previously in methanogens, is a consequence of the adaptation of the methanogenic population to the studied environment. In addition, because mcrA and 16S gene phylogenies are congruent, this approach allowed the methanogen community to be described and revealed higher diversity than previously observed with other methods. Indeed, hybridization capture recovered sequences from the Methanobacteriales order, belonging to the rare biosphere, which were not detected through direct sample sequencing due to the sequencing depth, or through PCR amplification, due to possible primer bias. Using the same approach, Biderre-Petit et al. (162) identified new Dehalococcoidia reductive dehalogenase homologous sequences from a metagenomic sample targeting rdhA genes or insertion sequence (IS) elements located nearby.
Finally, Manoharan et al. (163) applied hybridization capture to metagenomic soil samples. These authors highly enriched functional genes encoding carbohydrate-active enzymes and secretory proteases, while preserving the diversity of targeted biomarkers and the taxonomic distribution of micro-organisms. When applied to other genomic loci, such as 16S or 18S rRNA genes, in other metagenomic samples, the performance of hybridization capture enrichment has been shown to be greater than 90% (unpublished data from our lab). This strategy provides a better taxonomic description of ecosystems because of the more reliable diversity observed compared with conventional approaches, such as barcoding, through access to complete biomarkers and their subsequent accurate affiliation. These promising initial studies offer new strategies for the exploration of ecosystems and allow access to complete phylogenetic or functional biomarker diversity as well as providing a first glimpse into the identification of metabolic capabilities through the association of biomarkers.

To further explore metagenomic samples, we have recently adapted SHS to very large DNA fragments containing targeted biomarkers to reconstruct complete genomic regions and initiate targeted genome reconstruction (unpublished data from our lab). Sequencing data analysis revealed efficient enrichment of sequences from targeted genomes, leading to the assembly of nearly 100 kb DNA fragments including complete plasmids containing the targeted biomarker, with a very high sequence coverage. This allowed the reconstruction of metabolic pathways via biomarker association and linkage between identities and realized metabolic functions, representing a huge step forward in understanding the functions of microbial communities.

Environmental DNA is recognized as a rich source of natural products such as biocatalysts and bioactives, which are of great interest for industrial applications. Commonly, evaluation of the diversity of natural products synthesized by complex microbial communities is achieved more generally through the generation of clone libraries from metagenomic DNA and their subsequent expression in heterologous hosts (164,165). After functional screening for the desired activity, clones of interest are sequenced to determine the gene sequences encoding the metabolites of interest. Although effective, this strategy can be very tedious depending on the complexity of the metagenomic sample because of the screening step in which clones of interest must be identified among thousands of other possible clones (166). To ease this screening step, hybridization capture has been applied to environmental samples to enrich genes involved in specific environmental processes prior to clone library construction and analysis. Thus, Bragulini et al. (167) employed SHS to isolate glycoside hydrolase cDNAs from forest soil samples using explorative nonoverlapping capture probes targeting referenced and nonreferenced genes variants. The captured fragments were cloned into an expression vector and transformed into yeast for heterologous cDNA expression. Sequencing of randomly selected clones revealed that, after enrichment, targeted genes represented more than 90% and up to 100% of the obtained sequences, among which approximately 70% were of full-length, allowing their expression in the heterologous system. Hence, up to 25% of the screened recombinant yeasts exhibited secretion of endoxylanases, greatly reducing the depth required for library screening. Finally, as previously observed, hybridization capture also succeeded in selecting phylogenetically diverse representatives of the targeted gene family because none of the captured sequences were identical to the sequences described in databases and used for probe design.

As illustrated here, the success of hybridization capture experiments for the exploration of metagenomic samples strongly depends on probe design, including variant-specific and explorative probes to provide comprehensive diversity. Dedicated software has been developed to meet this specific need, such as PhylArray (168), MetabolicDesign (169), HiSpOD (170) and KApoD (171). Therefore, hybridization capture for the exploration of metagenomic samples appears to be a particularly promising strategy with diverse applications.

CONCLUSION

In the light of the studies performed to date, hybridization capture appears to be a particularly innovative, useful and effective approach for studying key genomic subsets in numerous biological domains. In resequencing approaches for medical research, hybridization capture through exome targeting has proven to be an extremely useful tool for the identification of mutations underlying Mendelian diseases as well as complex diseases and cancers, which has shed light on the ability of this approach to diagnose pathologies. This partitioning approach has also led to applications in the agronomical field to associate genetic variants with phenotypic traits of agricultural interest in crops and livestock. Moreover, because of the ability of hybridization capture to enrich numerous defined targets as well as their flanking regions, this strategy has broadened our knowledge of relevant underexploited fractions of the genomes of various models. Its capacity to extend what is already known from the genomes of organisms has been demonstrated through its application to nonreference species and paleogenomic samples on the basis of restricted genomic information. Finally, even if the use of hybridization capture to explore metagenomic samples remains anecdotic, its application allows us to better understand these complex ecosystems beyond the limits of conventional molecular approaches.

Hybridization capture strongly depends on NGS technologies and greatly benefits from the massive amounts of enriched sequence data provided. However, as sequencing technologies rapidly improve and the cost per sequenced nucleotide plummet, the price of whole-genome sequencing could quickly become comparable to the cost of exome sequencing, and deep insights into complex metagenomic samples could be achieved with few sequencing runs. Nevertheless, a niche may always remain for this technology because many biological questions do not require exhaustive sequences, but only restricted and targeted information. In addition, decreasing sequencing costs will be associated with an increase in the amount of data requiring processing and storage, with additional costs. Finally, the third-generation sequencing platforms with the possibility of sequencing longer DNA fragments provides real benefits to hybridization capture to span repetitive sequences.
It is therefore suitable for studying complex chromosomal structural variations, especially those involving repeats. But other genomic research applications, such as small insertion and deletion validation, haplotype phasing and microbial genome reconstruction from metagenomics samples could also benefit from this technology. In the latter case, these data will help us to improve our knowledge of the functioning of ecosystems, which remains a less prominent field, but likely one of the richest.

Hybridization capture is a particularly powerful method with a firmly established role in genome research. It has allowed the discovery of many unsuspected findings that would have been difficult to obtain using other approaches. New technical improvements can be made, but the application of this methodology to other biological fields is still underexplored and will provide influential results.

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