Infinium Monkeys: Infinium 450K Array for the Cynomolgus macaque (Macaca fascicularis)

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ABSTRACT The Infinium Human Methylation450 BeadChip Array (Infinium 450K) is a robust and cost-efficient survey of genome-wide DNA methylation patterns. Macaca fascicularis (Cynomolgus macaque) is an important disease model; however, its genome sequence is only recently published, and few tools exist to interrogate the molecular state of Cynomolgus macaque tissues. Although the Infinium 450K is a hybridization array designed to the human genome, the relative conservation between the macaque and human genomes makes its use in macaques feasible. Here, we used the Infinium 450K array to assay DNA methylation in 11 macaque muscle biopsies. We showed that probe hybridization efficiency was related to the degree of sequence identity between the human probes and the macaque genome sequence. Approximately 61% of the Human Infinium 450K probes could be reliably mapped to the Cynomolgus macaque genome and contain a CpG site of interest. We also compared the Infinium 450K data to reduced representation bisulfite sequencing data generated on the same samples and found a high level of concordance between the two independent methodologies, which can be further improved by filtering for probe sequence identity and mismatch location. We conclude that the Infinium 450K array can be used to measure the DNA methylene of Cynomolgus macaque tissues using the provided filters. We also provide a pipeline for validation of the array in other species using a simple BLAST-based sequence identity filter.

Biomedical research relies on nonhuman primate models, including the Cynomolgus macaque. Cynomolgus models are used to investigate diseases as diverse as Parkinson (Grabli et al. 2013; Potts et al. 2014; Riverol et al. 2013), influenza (Kobayashi et al. 2013; Pham et al. 2013), asthma (Cheng et al. 2013), diabetes (Kaplan and Wagner 2006), and atherosclerosis (Clarkson et al. 2013; Kaplan et al. 1996). However, the genome of this important model only was elucidated recently (Higashino et al. 2012; Yan et al. 2011), and tools for genomics analysis have consequently been lacking.

The rapidly expanding field of epigenomics has the potential to disentangle the influences of inheritance and environment on complex diseases and greatly increase our understanding, as well as provide biomarkers and therapeutics (Michels et al. 2013; Mill and Heijmans 2013). Therefore, it is desirable to study the epigenomics of Cynomolgus macaque disease models. Although candidate gene approaches have been reported (Rager et al. 2013), genome-wide approaches have been hampered by the lack of suitable tools. Likewise, although high-coverage MeDIP-seq approaches have been reported in the closely related rhesus macaque (Bell et al. 2012; Provencal et al. 2012), this technique is expensive, generally less quantitative, and has lower resolution.

The Infinium Human Methylation450 BeadChip Array (Infinium 450K) is a widely used tool in epigenetics that provides a robust and cost-efficient way to measure genome-wide DNA methylation patterns. It is a hybridization array designed to probe human genomic sequences

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probes were related to humans (chimpanzee, bonobo, gorilla, and orangutans), and probes were filtered for conservation between the human probe sequence and the ape genomic sequence (Hernando-Herraez et al. 2013).

We set out to determine the accuracy of the Infinium 450K in macaque tissues by comparing Infinium 450K and reduced representation bisulftite sequenced (RRBS) DNA methylation profiles generated from the same Cynomolgus muscle biopsies. Here we show that with filtering of the probes for good alignment between the human and Cynomolgus macaque genomic sequence, 51% of the array probes achieve a Pearson correlation R of 0.88 and 86% and 69% of the probes agree within 20% and 10% methylation difference respectively. As we can compare sequence identity with actual measurement variation between Infinium and RRBS, we also provide the sequence identity thresholds to achieve accurate Infinium measures and an in silico pipeline for converting the Infinium 450K to additional species.

MATERIALS AND METHODS

Samples

Muscle biopsies were performed on eleven adult male cynomolgus macaques under anesthesia with ketamine (7 mg/kg intramuscularly), and muscle samples were taken from the quadriceps region using a 3-mm biopsy punch. A single cruciate stitch using 6/0 PDS was placed and muscle samples were taken from the quadriceps region using a 3-mm biopsy punch. A single cruciate stitch using 6/0 PDS was placed and muscle samples were taken from the quadriceps region using a 3-mm biopsy punch. Muscle biopsies were performed on eleven adult male cynomolgus macaques under anesthesia with ketamine (7 mg/kg intramuscularly), and meloxicam (0.2 mg/kg subcutaneously) were given. All experimental procedures approved by SingHealth Institutional Animal Care and Use Committee.

Infinium 450K

After extraction of genomic DNA according to standard procedures, 1 μg genomic DNA was bisulftite converted using EZ DNA Methylation™ Gold Kit (Catalogue No.: D5002, Zymo Research). Successful conversion was confirmed via methylation-specific polymerase chain reaction before proceeding with subsequent steps of the Infinium assay protocol. The bisulftite-converted genomic DNA was isothermally amplified at 37°C for 22 hr, enzymatically fragmented, purified, and hybridized on an Infinium Human Methylation 450 BeadChip (cat. no. WD-314-1002; Illumina Inc.) at 48°C for 18 hr, after which the BeadChip was then washed to remove any unhybridized or non-specific hybridized DNA. Labeled single-base extension was performed on primers hybridized with DNA, and the hybridized DNA was removed. The extended primers were stained with multiple layers of fluorescence, the BeadChip was then coated using a proprietary solution and scanned using the Illumina iScan system. The image data were processed with the Genome Studio Methylation Module software. Further data processing was performed as described previously (Pan et al. 2012).

Reduced representation bisulftite signaling

One microgram of genomic DNA was restriction digested with 60U of MspI (cat. no. R0106S; New England Biolabs) overnight at 37°C. The digested product was purified by conventional ethanol precipitation method. End-repair, addition of ‘A’ tail and adapter ligation were performed using the TrueSeq DNA LT Sample Prep Kit V2 (cat. no. FC-121-2001; Illumina), according to manufacturer’s instructions with slight modifications. During the adapter-ligation step, 2 μL of Early Access Methylation Adapter oligo (cat. no. ME-100-0010; Illumina) was used instead. The adapter-ligated DNA fragments were subsequently purified with AMPure XP beads (cat. no. A63881; Agencourt), as per manufacturer’s protocol. Purified fragments were then bisulftite treated using the EpiTect Bisulftite Kit (cat. no. 59104; QIAGEN GmbH), according to manufacturer’s instructions. The converted DNA was amplified in a final concentration of 1 U of PhiTurbo Cx Hotstart DNA Polymerase and 250 μM of dNTP mix (cat. no. STR_600412; Stratagene), as well as 500 nM each of polymerase chain reaction primers PE 1.0 and 2.0 (Gu et al. 2011). The thermocycling condition used was 2 min at 95°C for heat activation, and 15 cycles of 30 sec at 95°C, 30 sec at 68°C, and 1 min at 72°C, followed by a 10 min final extension at 72°C. The enriched fragments were purified by gel electrophoresis and quantified by Agilent 2100 Bioanalyzer (Agilent Technologies). Sequencing was performed on the Illumina Genome Analyzer Ix platform, as per manufacturer’s instructions. Raw reads were filtered and trimmed (i.e., base quality < 20 were removed, read length < 20 bp were removed and entire read pair is removed if one of the trimmed reads become shorter than 20 bp), and then aligned against the Cynomolgus macaque genome (Yan et al. 2011) using RRBSMAP (Xu et al. 2012) with default settings (-p 8 -v 0.1 -w 100 -s 12 -D C-CGG). We filtered for CpGs

![Figure 1](image-url)
with at least 10x read coverage and estimated their methylation ratios using the python script methratio in RRBSMAP.

Alignment of Infinium probes to Cynomolgus macaque genome

We used blastn (Altschul et al. 1997) to map the 485,512 50 bp Infinium probes onto the Cynomolgus macaque genome using an e-value threshold of e\(^{-10}\). We used the Burrows-Wheeler Alignment tool (Li et al. 2009) to obtain the mismatched base positions and their location with respect to the CpG site being assayed. For each CpG, we checked for corresponding gene name or partial gene name matches between human and Cynomolgus macaque by searching for the closest gene to the CpG site. We further checked for ortholog gene name changes between human and macaque using the gene mapping from ENSEMBL (Hubbard et al. 2002).

RESULTS

Approximately 61% of human Infinium 450K probes could be reliably mapped to the Cynomolgus macaque genome and contain the CpG site of interest

We searched the Cynomolgus macaque genome sequence with the 485,512 50-bp Infinium probe sequences using blastn (Altschul et al., 1997) with an e-value threshold of e\(^{-10}\). This means that e\(^{-10}\) matches would be expected purely by chance and that the candidate match would likely be a bona fide match to the probe sequence. We also ran the Burrows-Wheeler Alignment tool (Li et al. 2009) to determine the mismatch base positions with respect to the location of the CpG site being assayed. Out of all the human probes, 328,091 probes (68%) could be mapped to the Cynomolgus macaque genome with up to four mismatches and up to two gaps of up to 4 bp. In addition, 30,021 (9.2%) of the mapped probes contained a mismatch in the CpG site itself (including those that do not have the CpG site) and are thus omitted from downstream analyses. We will thereafter refer to this set of mapped probes with the CpG site present as “CM probes” \(n = 298,070\). A total of 94% of the CM-probes had unique matches, whereas the rest had at least one alignment with e-value < e\(^{-10}\) (Figure 1). Thus, in total, approximately 61% of the Human Infinium 450K probes could be reliably mapped to the Cynomolgus macaque genome and contain a CpG site of interest.

Approximately 85% of CE probes were the Cynomolgus ortholog of the human gene

We then asked whether the gene annotation for the human probes corresponded to the Cynomolgus macaque gene ortholog for the matching macaque sequence. We searched for the exact gene within which the sequence lies or the closest gene to the matching sequence. Of the CM-probes, 229,505 had an associated human gene name from the Illumina Infinium 450K array annotation. Taking into account repeat matches, partial gene name matches and orthologous gene name changes between human and macaque (Hubbard et al. 2002), we found that 195,072 (85%) of the CM-probes had the same gene name or the equivalent macaque ortholog of the annotated human gene (Figure 1).

Cynomolgus tissues run on Infinium 450K arrays passed quality control, and approximately 80% of all probes yielded robust and detectable signal

We ran the Infinium 450K array on eleven Cynomolgus muscle tissues with one technical replicate. The raw data were analyzed using the Genome Studio Methylation Module software. The Illumina standard controls, including staining, extension, hybridization, target removal, bisulfit conversion, specificity I and II, nonpolymorphic controls all passed quality checks. Similarly, the negative controls, which target bisulfite-converted sequences that do not contain human CpG dinucleotides, and thus measure background intensity, had on average a very low intensity of around 200 AU and 250 AU for the green and red channels respectively (recommended background intensity from Illumina is <300 AU) (Supporting information, Figure S1), suggesting that the negative control probes work well for Cynomolgus macaques. The signal to noise ratio for target probes against

![Figure 2](image-url)  
**Figure 2** Hierarchical clustering of 12 Infinium 450K data of Cynomolgus macaque muscle tissues with a technical replicate.

![Figure 3](image-url)  
**Figure 3** Histograms of alignment bitscore for Infinium probes with detection \(P < 0.05\) (red), and \(> 0.05\) (blue).
negative green and red control probes were, on average, 10.6- and 8.3-fold. The two technical replicates clustered together on the dendogram (Figure 2). The global methylation correlation between the technical replicates was \( R = 0.97 \), whereas that between two different muscle samples was \( R = 0.94 \) (Figure S2). Notably, on average, 79.1 \( \pm \) 1.3\% of probes across all 12 sample profiles, including the two replicates, had the desired degree of hybridization with a \( P \)-value detection of less than 0.05 and approximately 73.4 \( \pm \) 1.6\% of probes had a \( P \)-value detection of less than 0.01. Interestingly, this is for the whole array including probes that do not align well to the Cynomolgus macaque genome, suggesting that some of these non-conserved probes still hybridize efficiently. When the analysis is restricted to CM probes only, on average, 86.8\% and 82.4\% had a \( P \)-value of \(<0.05\) and \(<0.01\), respectively. We then ran the Infinium 450K processing pipeline as described by (Pan et al. 2012) to obtain the beta values for each CpG on the array.

The degree of hybridization is positively related to the sequence identity between the human probe and Cynomolgus macaque genome

To assess whether the degree of hybridization of the probes is affected by sequence identity between human and Cynomolgus macaque, we took the CM-probes and computed the correlation between their detection \( p \)-values and alignment quality. Detection \( p \)-values were positively correlated with alignment \( e \)-value scores (Spearman rho = 0.39) and inversely correlated with the percent identity scores and bitscores (Spearman rho = -0.32 and rho = -0.39, respectively). When we excluded probes (\( n = 21,569 \)) with a mismatch at +/- 1 bp from the CpG site, the correlation between the degree of hybridization and alignment quality improved (Spearman rho = 0.42, -0.36, and -0.42 for \( e \)-value, percent identity, and bitscore, respectively).

When we grouped probes according to their detection \( p \)-values, those with \( P > 0.05 \) tended to have lower alignment bitscores compared to probes with \( P < 0.05 \) (Figure 3).

Concordance of Infinium 450K data and RRBS measurements within Cynomolgus macaque tissues and in relation to probe sequence divergence

We ran the same 11 Cynomolgus muscle samples using RRBS to determine the degree to which the Infinium 450K data correspond...
with methylation measurements from an independent sequencing-based technology. We searched each unique sample for CpGs present in both Infinium 450K and RRBS data and found that, on average, there were 4730 such CpGs for each sample. The Bland-Altman plot for one sample shows that the majority of the CpGs lie on x-axis (Figure 4). And, consistent with what has been reported previously by Pan et al. (2012), CpGs with intermediate methylation levels tend to show the biggest discrepancies between the two technologies.

To determine the relationship between the degree of agreement of RRBS and Infinium 450K data with sequence divergence, we stratified these shared CM-probe CpGs according to the quality of alignment of the Infinium probes to the Cynomolgus macaque genome. We found that across the 12 paired samples, probes with relatively poorer alignment scores consistently had lower RRBS-Infinium correlations compared with probes with greater alignment scores (Figure 5A). On average, the percent of shared probes with RRBS and Infinium 450K DNA methylation values within 20%, 10%, and 5% of each other were relatively greater for those with greater alignment scores (Figure 5B). The range across individuals is small except at very low alignment bitscores (Table S1). We further stratified CpGs by their average methylation levels and found that the concordance of the majority of unmethylated or methylated CpGs were much better as compared to CpGs with intermediate average methylation levels (Figure 5C).

We also examined whether the concordance between the two technologies is dependent on whether the sequence match is unique. We observe that probes with unique matches tended to have better performance (greater correlation and smaller % error) compared to probes that could be reliably aligned to multiple regions of the Cynomolgus macaque genome (Figure 6). We also examined whether the agreement in methylation values is affected by the position of the mismatch from the CpG site of interest. Probes with no mismatches at both the CpG site and +/- 1 bp had much better correlation and agreement compared with the probes that had either a mismatch at the CpG site itself or close to it. Probes with mismatches at either +/- 1 bp from the CpG tended to perform as badly as probes with mismatches exactly at the CpG site (Figure 7).

We examined whether the concordance between the RRBS and Infinium data differed depending on whether data were from a type I or type II Infinium probe. Type II probes have been shown to have a smaller dynamic range than type I probes (Dedeurwaerder et al. 2011), although this effect is corrected in our data processing. We could not detect any difference in the sequence alignment or agreement between RRBS and Infinium data from type I and type II probes (Table S2 and Figure S3). Studies have reported extensive probes that target human polymorphic CpGs in the Infinium 450K array (Chen et al. 2013; Price et al. 2013). We observed no difference in the RRBS and Infinium 450K agreement between polymorphic and nonpolymorphic probes (Figure S4). In addition to absolute DNA methylation levels, we also analyzed the difference in methylation levels across different individuals measured using the same technology, and compared the concordance of this difference across the two platforms. The differences between inter-individual differences measured on Infinium 450K assays compared to inter-individual differences measured using RRBS were within 20% for 98% of probes, within 10% for 87% of probes and within 5% for 74% of probes (see Figure S5 for an example).
Probe selection criteria for best concordance and coverage

We observed that the agreement between RRBS and Infinium data can be increased by taking into account sequence divergence of the CM probe, uniqueness of the alignment as well as the position of the mismatch relative to the CpG site being measured. We then assessed the coverage of probes on the array as a function of these three levels of alignment filters and their performance (Table 1). Filtering for uniquely matched probes with no mismatch at 1 bp away from the CpG site and with bitscore >70 (Figure S6), we observed a substantial improvement in the agreement between Infinium and RRBS data, whereas there was a substantial decrease in the number of probes retained at >75 bitscore (Figure S7). When we adjusted the threshold from 60 to 70, the percent of usable probes changed only slightly from 54% to 51% after we applied the three-step filter. As we further increased the bitscore threshold to 75, the performance increased only marginally, but the coverage decreased dramatically to only 41%. Thus, to maximize coverage while maintaining accuracy, we recommend an alignment bitscore threshold of 70.

In summary, to obtain reasonably accurate Cynomolgus macaque methylation results from the Human Infinium 450K array, we suggest a three-step probe filter: bitscore >70, uniqueness of sequence match, and no mismatches near to the CpG site. We reason that the use of these procedures should allow for the effective and informative utilization of the Infinium 450K array in species with sequenced genomes. Furthermore, it would be ideal to follow up on the significant results from the array using another method such as pyrosequencing as an additional technical validation. We have provided a comprehensive Cynomolgus macaque annotation file for the Infinium 450K array incorporating all the parameters discussed previously (File S1).

DISCUSSION

We demonstrated here for the first time that the use of the human Infinium 450K array is feasible in Cynomolgus macaque tissues. Using in silico analysis with careful probe filtering, we found that approximately 61% of all human probes could be mapped to the macaque genome and contained an informative CpG, with the vast majority of these probes reporting on orthologous genes. Applying the Infinium 450K array to measure methylation patterns of DNA derived from macaque muscle tissues, we found that a large percentage of all probes passed quality controls and yielded robust DNA methylation signal, which positively correlated with the degree of sequence conservation. Comparing the array-based DNA methylation profiles with those obtained by RRBS provided overall high concordance, suggesting that these technologies can be benchmarked vs. one another.

Not surprisingly, we showed that the agreement between RRBS and Infinium 450K data was most dependent on the sequence divergence of the human probes compared with Cynomolgus macaque, the uniqueness of the probe sequence to the Cynomolgus genome, and the position of the mismatch relative to the target CpG. Approximately 250K probes on the array reported less than 20% methylation difference between Infinium and RRBS data. We also observed that the concordance between the two technologies is much better for fully methylated and unmethylated CpGs than the partially methylated CpGs. Finally, we provided a bioinformatics blueprint for the adaptation of Infinium 450K array technology to other species with sequenced genomes.

Our study is a logical extension of previous work using the Infinium 27K array to study DNA methylation in Cynomolgus macaque. The advantages of using the Infinium 450K for such investigations mirror those in human tissues and include the dramatically increased number of alignment levels, the ability to filter based on sequence identity, and the inclusion of single-nucleotide polymorphisms. These features allow for a more comprehensive assessment of methylation patterns across the entire genome, which is particularly important in the study of species with complex and diverse genetic backgrounds. Furthermore, the use of Infinium data in conjunction with RRBS provides a robust method for validating results and identifying novel methylation sites that may be specific to the Cynomolgus macaque species.
and broader genomic representation of CpG loci. Others have similarly applied the Infinium 450K array to measure the cytosine methylation profiles of great apes, which are evolutionarily closer to human compared to macaques. From their analysis, approximately 200K (44%) probes were retained for orangutans, which is comparable with the 51% we obtained for Cynomolgus macaque. Great apes are evolutionarily closer to humans than macaques, so the greater percentage of probes retained for macaques might be due to the less-stringent choice of alignment filters in our study. Regardless, these studies collectively open new areas for evolutionary research with broad biological implications. Specifically, using the Infinium 450K array on non-human model systems such as Cynomolgus macaque and great apes offers the opportunity to directly compare at single nucleotide resolution the DNA methylation profiles across evolutionary trees, including within-species comparisons of geographically diverse populations. Likewise, the potential for cross-species comparisons is most exciting, given the thousands of human DNA methylation profiles that are being created using this technology.

To better benchmark the performance of the Infinium 450K array we compared resulting data directly to those obtained using RRBSseq performed on the same samples. Even with aggressive filtering we do not achieve the agreement between Infinium and RRBS that has been seen for human tissues i.e., correlation of 0.97 and 84% of probes within 10% methylation error (Pan et al. 2012). This discrepancy could be due to experimental variation or sequencing errors in the unfinished Cynomolgus genome. Invalid sequence (or indeed polymorphism) would affect the accuracy of our alignment calls and could have led to the inclusion of mismatched probes. Nevertheless, this cross-technology comparison strongly supports the technical and biologically validity of the Cynomolgus macaque DNA methylation profiles obtained from the Infinium 450K array.

Various sequencing methods for profiling DNA methylation at the genome-wide or whole-genome scale are available, however, these are expensive and require high quality genome sequence as a reference. The popular platforms include RRBS and MeDIP-seq. MeDIP-seq or MBD-seq uses an antibody or methyl-binding protein to create a genomic library of methylated regions that are then sequenced. It tends to bias toward regions that are moderately or highly methylated and does not cover the hypomethylated regions that are thought to be greatly methylated CpGs (Ziller et al. 2013).

Array-based methods offer researchers familiar protocols, simpler data analysis and relatively low costs compared with sequencing. The Infinium array also generally requires less starting material than standard sequence based methods. Each Infinium 450K run costs about half that of RRBS and MeDIP-seq. This cost-saving would be most pertinent to large epigenome-wide association studies involving Cynomolgus macaques. Finally, researchers who wish to profile the DNA methylome in Cynomolgus macaque tissues should use the Infinium 450K array with some caution, using the filters described here and confirming locus-specific results with an independent technique. We suggest that these considerations permit the use of the Human Infinium 450K array with Cynomolgus macaque tissues, an approach that should, among other possibilities, enable affordable and powerful studies of tissue specificity in a complex, non-human primate model with considerably larger sample sizes than would be afforded by sequencing protocols.

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