1 Introduction

Distinct combinations of histone modifications are associated with different classes of functional genomic elements such as promoters, enhancers, and genes (Consortium et al., 2015). Chromatin immunoprecipitation sequencing (ChIP-seq) experiments are commonly used to obtain genome-wide profiles of histone modifications associated with different types of functional genomic elements. However, the quality of histone ChIP-seq data is affected by a myriad of experimental parameters such as the amount of input DNA, antibody specificity, ChIP enrichment, and sequencing depth. Making accurate inferences from chromatin profiling experiments that involve diverse experimental parameters is challenging.

Results: We introduce a convolutional denoising algorithm, Coda, that uses convolutional neural networks to learn a mapping from suboptimal to high-quality histone ChIP-seq data. This overcomes various sources of noise and variability, substantially enhancing and recovering signal when applied to low-quality chromatin profiling datasets across individuals, cell types, and species. Our method has the potential to improve data quality at reduced costs. More broadly, this approach – using a high-dimensional discriminative model to encode a generative noise process – is generally applicable to other biological domains where it is easy to generate noisy data but difficult to analytically characterize the noise or underlying data distribution.

Abstract

Motivation: Chromatin immunoprecipitation sequencing (ChIP-seq) experiments are commonly used to obtain genome-wide profiles of histone modifications associated with different types of functional genomic elements. However, the quality of histone ChIP-seq data is affected by a myriad of experimental parameters such as the amount of input DNA, antibody specificity, ChIP enrichment, and sequencing depth. Making accurate inferences from chromatin profiling experiments that involve diverse experimental parameters is challenging.

Results: We introduce a convolutional denoising algorithm, Coda, that uses convolutional neural networks to learn a mapping from suboptimal to high-quality histone ChIP-seq data. This overcomes various sources of noise and variability, substantially enhancing and recovering signal when applied to low-quality chromatin profiling datasets across individuals, cell types, and species. Our method has the potential to improve data quality at reduced costs. More broadly, this approach – using a high-dimensional discriminative model to encode a generative noise process – is generally applicable to other biological domains where it is easy to generate noisy data but difficult to analytically characterize the noise or underlying data distribution.

Availability: https://github.com/kundajelab/coda

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2015), and methylation status (Angermueller et al. 2016b): for example, they have been used to predict regulatory sequence determinants of DNA and RNA binding proteins (Alipanahi et al. 2015; Koh et al., 2015; Zhou and Troyanskaya, 2015), chromatin accessibility (Kelley et al., 2015), and methylation status (Angermueller et al., 2016a).

### 2 Methods

#### Model

Coda takes in a pair of matching ChIP-seq datasets of the same histone modifications in the same cell-type – one high-quality and the other noisy – and uses convolutional neural networks (CNNs) to learn a mapping from the noisy to the high-quality ChIP-seq data. The noisy dataset used in training can be derived computationally (e.g., by subsampling the high-quality data) or experimentally (e.g., by conducting the same ChIP-seq experiment with fewer input cells). Once this mapping has been learned, the same mapping can then be applied to new, noisy data in any other cellular context with the same underlying noise structure.

For each type of noise (e.g., due to low cell numbers, sequencing depth, or enrichment) and each target histone mark, we train two separate CNNs to accomplish two tasks: a regression task to predict histone ChIP signal (i.e., the fold enrichment of ChIP reads over input DNA control) and a binary classification task to predict the presence or absence of a significant histone mark peak (Fig. 2). In total, if a given experiment has $M$ marks, then we train 2$M$ models separately (one regression and one classification model for each mark). Each individual model optionally makes use of the noisy ChIP-seq data from all available marks but outputs only one target histone mark. This allows us to learn separate features for each mark and task while still leveraging information from multiple input histone tracks; we find empirically that this improves performance.

For computational efficiency, we first bin the genome into 25bp bins, averaging the signal in each bin. Let $L$ be the number of bins in the genome (i.e., the length of the genome divided by 25). Each individual model takes in an $M \times L$ input matrix $X$ and returns a $1 \times L$ output vector $Y$ representing the predicted high-quality signal (in the regression setting) or peak calls (in the classification setting). It does this by feeding the noisy data through a first convolutional layer, a rectified linear unit (ReLU) layer, a second convolutional layer, and then a final ReLU or sigmoid layer (for regression or classification, respectively). For the first convolutional layer, we use 6 convolutional filters, each 51 bins in length; for the second convolutional layer, we use a single filter of length 1001. Effectively, this means that a prediction at the i-th bin is a function of the noisy data at a 25.025bp window centered on the i-th bin.

The convolutional nature of our models (and the lack of max-pooling layers commonly seen in neural network architectures for computer vision) enables us to do efficient genome-wide prediction, as 98% of the computation required for predicting signal at the i-th bin is shared with the computation required for predicting the $(i+1)$-th bin. In particular, to compute the prediction at the i-th bin, the network needs to perform $6 \times 1001 \times 51$ operations at the first convolutional layer and $6 \times 1001$ operations at the second convolutional layer. To compute the prediction at the $(i+1)$-th bin, the network needs to perform only $6 \times 51$ more operations at the first convolutional layer and $6 \times 1001$ operations at the second convolutional layer, saving $6 \times 1001 \times 50$ operations. Other models, especially non-linear models such as random forests, would require a
completely separate set of computations for each bin and are therefore significantly more computationally expensive when it comes to making predictions across the entire genome.

Training and evaluation
We applied Coda to three distinct sources of noise: low sequencing depth, low cell input, and low ChIP enrichment. In all cases, the inputs to our model were noisy signal measurements of multiple histone marks (see Data Availability and Processing for more details), and we trained separate models to predict the high-quality signal and peak calls for each target mark.

For the regression tasks (predicting signal), we evaluated performance by computing the Pearson correlation and mean squared error (MSE) between the predicted and measured high-quality fold-enrichment signal profiles after an inverse hyperbolic sine transformation, which reduced the dominance of outliers. We compared this to the baseline performance obtained by directly comparing the noisy and high-quality signal profiles of the target mark (after the same inverse hyperbolic sine transformation).

For the classification tasks (predicting presence or absence of a peak), we compared our model's output to peaks called by the MACS2 peak caller (Feng et al., 2012) on the high-quality signal for the target mark. As our dataset is unbalanced – peaks only make up a small proportion of the genome – we evaluated performance by computing the area under the precision-recall curve (AUPRC), a standard measure of classification performance for unbalanced datasets (Davis and Goadrich, 2006). We compared the AUPRC of our model to a baseline obtained by comparing MACS2 peaks on the noisy data for the target mark to those obtained from the high-quality data for the target mark (see Data Availability and Processing for further details on dataset preparation).

We trained our models on 50,000 positions randomly sampled from peak regions of the genome and 50,000 positions sampled from non-peak regions, sampling from each autosome with equal likelihood. We defined peak regions using the output mark of interest and with the high-quality data. Further increasing dataset size did not increase performance; as each sample covered 25.025bp, 100,000 samples had good coverage of the entire genome. We selected the training dataset to be balanced because a uniformly drawn dataset would have had very few peaks; making it difficult for the model to learn to predict at peak regions; however, the test results reported in this paper are on the entire (unbalanced) genome. We used the Keras package (François Chollet, 2015) for training and AdaGrad (Duchi et al., 2011) as the optimizer, stopping training if validation loss did not improve for three consecutive epochs. We did not observe overfitting with our models (train and test error were comparable), and therefore opted not to use common regularization techniques such as dropout (Srivastava et al., 2014).

We chose model hyperparameters and architecture through hold-out validation on the low-sequencing-depth denoising task with GM12787 as the training cell line (Kasowski et al., 2013), holding out a random 20% subset of the training data for validation; this task will be discussed in more detail in the next section. The model architecture described above (6 convolutional filters each 51 bins in length in the first layer, and 1 convolutional filter of length 1001 in the second layer) yielded optimal validation performance out of the configurations we tried (varying the number of convolutional filters and the lengths of the filters by up to an order of magnitude). Adding an additional layer to the neural network brought a modest increase in performance at the cost of more computation time and complexity. To be sure that our model architecture generalized, we used the same architecture and hyperparameters for all denoising tasks without any further tuning.

3 Results
Removing noise from low sequencing depth data
A minimum of 40-50M reads is recommended for optimal sensitivity for histone ChIP-seq experiments in human samples targeting most canonical histone marks (Jung et al., 2014). As adhering to this standard can often be infeasible due to cost and other limitations, a substantial proportion of publicly available datasets do not meet these standards. Motivated by these constraints, we tested whether our model could recover high-read depth signal from low-read depth experiments.

Training and testing on the same cell type across different individuals
We evaluated Coda on lymphoblastoid cell lines (LCLs) derived from six individuals of diverse ancestry (European (CEU), Yoruba (YRI), Japanese, Han Chinese, San) (Kasowski et al., 2013). We used the CEU-derived cell line (GM12787) to train our model to reconstruct the high-depth signal (100M reads per mark; exact numbers in Data Availability and Processing) from a simulated noisy signal derived by subsampling 1M reads per mark. On the other five cell lines, Coda significantly improved Pearson correlation between the full and noisy signal (Fig. 3A, left) and the accuracy of peak calling (Fig. 3A, right). Using just 1M reads per mark, the predicted output of our model was equivalent in quality to signal derived from 15M+ reads (H3K27ac) and 25M+ reads (H3K36me3) (Fig. 3B). Fig. 4 shows how Coda can accurately reconstruct histone modification levels at the promoter of the PAX5 gene, a master transcription factor required for differentiation into the B-lymphoid lineage (Nutt et al., 1999).

We confirmed Coda was not simply memorizing the profile of the training cell line (GM12787) and copying it to the test cell lines by examining differential regions, called by DESeq (Anders and Huber, 2010), between GM12787 and the other cell lines (Kasowski et al., 2013). Coda improved correlation and peak-calling even in those regions (Table 1). Similarly, it also improved correlation on the regions of the genome with enriched signal, i.e., called as statistically significant peaks (Table 2).

Table 1. Denoising differential regions (diff. reg.) between test cell line GM18526 and training cell line GM12787 and training cell line GM12787. Performance reported is improvement of the denoised model over baseline (subsampled reads) on the test cell line. In parentheses we report the baseline results followed by the denoised results. Peak-calling results on H3K27me3 are omitted due to the lack of peak calls in differential regions; all results on H3K36me3 are omitted due to low number of differential regions.

<table>
<thead>
<tr>
<th></th>
<th>MISE (diff. reg.)</th>
<th>Pearson R (diff. reg.)</th>
<th>AUPRC (diff. reg.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>H3K4me1</td>
<td>-85% (4.01, 0.57)</td>
<td>+59% (0.37, 0.59)</td>
<td>+63% (0.93, 0.57)</td>
</tr>
<tr>
<td>H3K4me3</td>
<td>-75% (2.88, 0.70)</td>
<td>+14% (0.63, 0.72)</td>
<td>+11% (0.78, 0.87)</td>
</tr>
<tr>
<td>H3K27ac</td>
<td>-66% (4.43, 0.48)</td>
<td>+39% (0.55, 0.77)</td>
<td>+66% (0.90, 0.95)</td>
</tr>
<tr>
<td>H3K27me3</td>
<td>-30% (0.78, 0.15)</td>
<td>+106% (0.14, 0.30)</td>
<td>-</td>
</tr>
</tbody>
</table>

Training and testing on different cell types across different individuals
We next assessed if Coda could be trained on one cell type in one individual and used to denoise low-sequencing-depth data from a different cell type in a different individual. As above, the model was trained to output high-depth data (30M reads) from low-depth data (1M reads). We used histone ChIP-seq data spanning T-cells (E037), monocytes (E029), mesenchymal stem cells (MSCs, E026), and fibroblasts (E056) from the Roadmap Epigenomics Consortium (Consortium et al., 2015). Coda substantially improved the quality of the low-depth signal on the test cell type for all pairs of cell types (Table 3), illustrating its ability to denoise low-depth data on a cell type even if high-depth training data for that cell type is not available.
we report the baseline results followed by the denoised results. Over baseline (original, subsampled reads) on the test cell line. In parentheses cell line GM12878. Performance reported is improvement of the denoised model on peak regions between test cell line GM18526 and training regions, with similar (very slightly better) MSE and correlation across the whole genome. This implies that Coda is better able to learn to match the exact values of the signal tracks on “difficult” regions (i.e., where there is the greatest deviation from the training signal), even though the linear model matches the rough shape. These regions are important to predict well because they can give insight into the differences between individuals and cell types. We note that many forms of smoothing can be represented via linear regression. For example, a standard Gaussian filter can be interpreted as taking a linear combination of surrounding points with fixed coefficients. The comparison against a linear regression baseline therefore sets an upper limit to the complexity of models that would be practically useful in genome-wide prediction.

**Fig. 3.** Coda removes noise from low-sequencing-depth experiments on lymphoblastoid cell lines derived from different individuals. (A) Compared to the signal from subsampled reads (blue), the denoised signal (green) shows greater correlation with the full signal (left) and more accurate peak-calling (right) across all cell lines. The model was trained on GM12878 and tested on different cell lines, within each column in the plot, each point is a single test cell line. (B) With 1M reads per mark, the denoised H3K27ac data is equivalent in quality to a dataset with 15M+ reads per mark, and the H3K36me3 data is equivalent to a dataset with 21M+ reads per mark. Similar results hold for other marks. These results are from training on GM12878 and testing on GM18526.

**Table 3.** Cross cell-type experiments. Rows are train cell type, while columns are test cell type. In parentheses we report the baseline results followed by the denoised results, averaged across all histone marks used.

<table>
<thead>
<tr>
<th></th>
<th>Monocytes</th>
<th>MSCs</th>
<th>Fibroblasts</th>
</tr>
</thead>
<tbody>
<tr>
<td>PearsonR</td>
<td>+33% (+0.51, 0.67)</td>
<td>+58% (+0.44, 0.70)</td>
<td>+78% (+0.36, 0.65)</td>
</tr>
<tr>
<td>T-cells</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Monocytes</td>
<td>+59% (+0.44, 0.70)</td>
<td>+79% (+0.36, 0.65)</td>
<td>+81% (+0.36, 0.66)</td>
</tr>
<tr>
<td>MSCs</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>AUPRC</td>
<td>+116% (+0.31, 0.66)</td>
<td>+139% (+0.31, 0.73)</td>
<td>+94% (+0.35, 0.69)</td>
</tr>
<tr>
<td>T-cells</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Monocytes</td>
<td>+139% (+0.31, 0.73)</td>
<td>+94% (+0.35, 0.69)</td>
<td>+100% (+0.35, 0.71)</td>
</tr>
<tr>
<td>MSCs</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

Coda outperforms linear baselines

We compared Coda to a linear and logistic regression baseline for signal denoising and peak calling, respectively. In both cases, we used an input region of the same size as Coda (i.e., 25,025bp centered on the location to be predicted, binned into 25bp bins). As noted above, the desire for computational efficiency in making genome-wide predictions across multiple marks limits the complexity of models that would be practically useful in genome-wide prediction.

When evaluated in the same cell type, different individual setting, Coda achieved 3x lower MSE on peak regions and 2x lower MSE on differential regions, with similar (very slightly better) MSE and correlation across the whole genome.
Table 4. Low-cell-input experiments. We report improvement of the denoised model output over baseline (original low-input experiments), as compared to high-input experiments. In parentheses we report the baseline results followed by the denoised results.

<table>
<thead>
<tr>
<th>Mark</th>
<th>MSE</th>
<th>Pearson R</th>
<th>AUPRC</th>
</tr>
</thead>
<tbody>
<tr>
<td>ULI-NChIP</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>H3K4me3-61%</td>
<td>0.19</td>
<td>0.54</td>
<td>+208%</td>
</tr>
<tr>
<td>H3K9me3-46%</td>
<td>0.51</td>
<td>0.27</td>
<td>+28%</td>
</tr>
<tr>
<td>H3K27me3-41%</td>
<td>0.68</td>
<td>0.40</td>
<td>+57%</td>
</tr>
<tr>
<td>MOWChIP</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>H3K4me3-42%</td>
<td>0.18</td>
<td>0.68</td>
<td>+122%</td>
</tr>
<tr>
<td>H3K27ac-21%</td>
<td>1.44</td>
<td>1.14</td>
<td>+159%</td>
</tr>
</tbody>
</table>

Fig. 5. Genome browser tracks for low-cell-input experiments. We compare noisy signal obtained from 100 cells (top) with Coda’s output (middle) and the target, high-quality signal obtained from 10,000 cells (bottom) at the Runx1 gene, a key regulator of the B-lymphoid lineage. (Nechanitzky et al., 2015)
is depleted at DHSs. (Shu et al., 2011) For each of those marks, we compared the average uncorrupted signal, the average denoised signal, and the average low-enrichment signal within 5000 bp of the summits of DNase I hypersensitive peaks in GM12878 from ENCODE data (Bernstein et al., 2012). As expected, the corrupted, low-enrichment signal was biased by the reads from the control experiment and had significantly lower fold enrichment of H3K4me1 and H3K27ac at DHSs, compared to the uncorrupted signal. In contrast, the denoised signal was significantly more enriched at DHSs than the corrupted signal, more closely resembling the uncorrupted signal. Conversely, the corrupted signal had higher levels of H3K27me3 at DHSs, whereas the denoised signal had low levels of

Table 5. Low-enrichment experiments. We report improvement of the denoised model output over baseline (low-enrichment experiments), as compared to high-enrichment experiments. In parentheses we report the baseline results followed by the denoised results.

<table>
<thead>
<tr>
<th>Material</th>
<th>MSE</th>
<th>Pearson R</th>
<th>AUPRC</th>
</tr>
</thead>
<tbody>
<tr>
<td>H3K4me1</td>
<td>-75%</td>
<td>+42%</td>
<td>+215%</td>
</tr>
<tr>
<td>H3K4me3</td>
<td>-86%</td>
<td>+54%</td>
<td>+94%</td>
</tr>
<tr>
<td>H3K27ac</td>
<td>-78%</td>
<td>+37%</td>
<td>+121%</td>
</tr>
<tr>
<td>H3K27me3</td>
<td>-61%</td>
<td>+88%</td>
<td>+225%</td>
</tr>
<tr>
<td>H3K27ac</td>
<td>-44%</td>
<td>+47%</td>
<td>+168%</td>
</tr>
</tbody>
</table>

4 Conclusion
We describe a convolutional denoising algorithm, Coda, that uses paired noisy and high-quality samples to substantially improve the quality of new, noisy ChIP-seq data. Our approach transfers information from generative noise processes (e.g., mixing in control reads to simulate low-enrichment, or performing low-input experiments) to a flexible discriminative model that can be used to denoise new data. We believe that a similar approach can be used in other biological assays, e.g., ATAC-seq and DNase-seq (Buenrostro et al., 2013; Crawford et al., 2006), where it is near impossible to analytically characterize all types of technical noise or the overall data distribution but possible to generate noisy versions of high-quality samples through experimental or computational perturbation. This can significantly reduce cost while maintaining or even improving quality, especially in high-throughput settings or when dealing with limited amounts of input material (e.g., in clinical studies).

An important caveat to our work is that the performance of Coda depends strongly on the similarity of the noise distributions and underlying data distributions in the test and training sets. For example, Coda expects that the relationships between different histone marks should be conserved between the test and training set. Thus, applying a set of trained Coda
models to data that is very different from what it was trained on is unlikely to work. We also assume that the noise parameters in test data are known in advance, e.g., the sequencing depth, the number of input cells, or the level of ChIP enrichment. In some cases (e.g., the low-sequencing-depth and low-cell-input settings) this is true, but in others (e.g., the low-enrichment setting) it is not always possible. An important direction for future work is to make Coda more robust, for example, training a single model over various settings of the noise parameters and various cell types could improve the generalizability of the models. To further improve performance, more complex neural network architectures could also be explored: for example, using recurrent neural networks (Sutskever et al., 2014) to explicitly model long-range spatial correlations in the genome; multi-tasking across output marks instead of training separate models for each mark; or using deeper networks.

Another avenue for future work is exploring using more than just the noisy histone ChIP-seq data at test time. In this work, we use only the noisy data at test time, training our models to transform it into high-quality data. In reality, at test time we might have access to other data: for example, we might also have the DNA sequence of the test sample or access to high-quality ChIP-seq data on a closely related cell type. Other work has used DNA sequence to predict transcription factor binding (Alipanahi et al., 2015; Zhou and Troyanskaya, 2015), chromatin accessibility (Kelley et al., 2015), and methylation status (Angermueller et al., 2016a). A natural next step would be to combine the ideas from these methods with ours, e.g., by having a separate convolutional module in our neural network that incorporates sequence information and joins with the ChIP-seq module at an intermediate layer. Others have also used high-quality ChIP-seq data from closely related cell types for imputation (Ernst and Kellis, 2015); combining this with our denoising approach could help to avoid a potential pitfall of imputation approaches, namely the loss of cell-type-specific signal, while improving the accuracy of our denoised output.

Below, we provide a link to a script that trains a model for low-sequencing-depth noise using the LCL data described above. Since the type of noise can vary from context to context, we also provide the code for the general Coda framework to allow for developers of new protocols (e.g., new low-cell-count techniques) or core facilities that have high throughput to train Coda with data specific to their context.

### Data Availability and Processing

**Datasets**

We used the following publicly-available GEO datasets in this work:

1. GSE50893 for ChIP-seq data on LCLs (Kasowski et al., 2013)
2. GSE63523 for ULI-NChIP-seq data (Brind’Amour et al., 2015)
3. GSE65516 for MOWChIP-seq data (Cao et al., 2015)
4. GSM376620 for DNase hypersensitive peaks (Bernstein et al., 2012)

For the low-sequencing-depth experiments, the full depth for GM12878 (training set) was 171M (million reads) for H3K4me1, 168M for H3K4me3, 328M for H3K27ac, 265M for H3K27me3, and 123M for H3K36me3. The full depth for GM18526 (test set) was 120M for H3K4me1, 136M for H3K4me3, 125M for H3K27ac, 138M for H3K27me3, and 223M for H3K36me3.

For the cross-cell-type experiments, we used the consolidated Roadmap Epigenomics datasets (Consortium et al., 2015), which is publicly available from http://egg2.wustl.edu/roadmap/data/byFileType/alignments/. Each mark is downsampled to a maximum of 30M reads to maximize consistency across marks; we used this as the full depth data, and downsampled to 1M reads for the noisy data. A detailed description of this dataset is available in (Roa, 2015).

**Dataset preparation**

**Fold change signal profiles and peak calling.** For each experiment, we used align2rawsignal (Kundaje, 2013) to generate signal tracks and MACS2 (Fong et al., 2012) to call peaks, as implemented in the AQUAS package (Lee and Kundaje, 2016). For the signal track, we used fold change relative to the expected uniform distribution of reads after an inverse hyperbolic sine transformation (Hoffman et al., 2012). We used the gappedPeaks output from MACS2 as the peak calls. For computational efficiency, we binned the genome into 250bp segments, averaging the signal in each segment.

We evaluated our peak calling on a bin-by-bin basis, i.e., our model output one number for each bin representing the probability that that bin was a true peak, and we treated each bin as a separate example for the purposes of computing AUROC, our metric for peak calling performance. To get ground truth data for our peak calling tasks, we labeled each bin as “peak” or “non-peak” based on whether that bin was part of a peak called by MACS2 on the high-quality data.

Computing AUROC requires predictions to be ranked in order of confidence. For our model, we used the output probabilities for each bin to calculate the ranking. MACS2 outputs both a peak p-value track, assigning a p-value to each genomic coordinate, and a set of binary peak calls. To measure baseline performance on the noisy data, we ranked each bin by the maximum peak p-value assigned by MACS2 to a genomic coordinate in that bin, unless that bin did not intersect with any of the binary peak calls, in which case it was assigned a p-value of −inf (i.e., ranked last). We did this to ensure that the high-quality peak track had an AUROC of 1; empirically, this also improved performance of the noisy MACS2 baseline.

**Histone marks used**

We used different sets of input and output histone marks for different experiments depending on which marks each dataset provided. For the same cell type, different individual experiments (using lymphoblastoid cell lines), we trained and tested on H3K4me1, H3K4me3, H3K27ac, H3K27me3, and H3K36me3; we used the same data for the low-ChIP-enrichment experiments. For the different cell type, different individual experiments (using the uniformly-processed Roadmap Epigenomics Consortium datasets (Consortium et al., 2015)), we trained and tested on H3K4me1, H3K4me3, H3K9me3, H3K27ac, H3K27me3, and H3K36me3. For all of the above experiments, we also used data from the control experiments (no antibody) as input. Lastly, for the low-cell-input experiments, we used H3K4me3, H3K9me3, and H3K27me3 from the ULI-NChIP-seq dataset and H3K4me3 and H3K27ac from the MOWChIP-seq dataset.

**Low-cell-input datasets**

The ULI-NChIP-seq (Brind’Amour et al., 2015) and MOWChIP-seq (Cao et al., 2015) papers provided several datasets corresponding to different numbers of input cells used. For each protocol, we used the datasets with the lowest number of input cells as the noisy input data (ULI-NChIP-seq: 102 cells for H3K9me3 and H3K27me3, 5x103 cells for H3K4me3; MOWChIP-seq: 102 cells) and the datasets with the highest number of input cells as the gold-standard, high-quality data (ULI-NChIP-seq: 106 cells for H3K9me3, 105 cells for H3K4me3 and H3K27me3; MOWChIP-seq: 104 cells). The ULI-NChIP-seq data had matching low- and high-input experiments only for a single cell type, so we divided it into chr5-19 for training, chr3-4 for validation, and chr1-2 for testing.

**Code, data, and browser track availability**

Our code is available on Github at https://github.com/kundajelab/coda, including a script that downloads pre-processed data to train Coda with data specific to their context.
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references
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2369896
allgnan, b., delong, a., weinach, m. t. and frey, b. j. (2015) predicting the sequence specificities of dna-and rna-binding proteins by deep learning. nature biotechnology, 33, 833–838. url http://dx.doi.org/10.1038/nbt.3302
anand, r., anshu, r., rittenberg, e. et al. (2016) deep speech 2: end-to-end speech recognition in english and mandarin. in international conference on machine learning, pp. 173–182
angermueller, c., parnassa, t., paris, l. and steph, o. (2014b) deep learning for computational biology. molecular systems biology, 10, 978. url http://dx.doi.org/10.1038/msb.2015651
bernstein, b. e., bryne, e., dunham, i. et al. (2012) an integrated encyclopedia of dna elements in the human genome. nature, 490, 55–74. url http://dx.doi.org/10.1038/nature11247
bernstein, b. e., mikkelsen, t. s., xie, x. e. et al. (2006) a bivalent chromatin structure marks key developmental genes in embryonic stem cells. cell, 125, 315–326
bri$aal$amu$er, j., liu, s., hudson, m. et al. (2015) an ultra-low-input native chip-seq protocol for genome-wide profiling of rare cell populations. nature communications, 6, 6033. url http://www.nature.com/articles/ncomms6033
burewosta, j.d., gieru, p.g., zuba, l.c., chang, h.y. and greenleaf, w.j. (2013) transcription of native chromatin for fast and sensitive epigenomic profiling of open chromatin, dna-binding proteins and nucleosome position. nature methods, 10, 1213–1218. url http://www.nature.com/dois/10.1038/nmeth.2688
consortium, r. e., kundaje, a., medvednykh, w. et al. (2015) integrative analysis of 111 reference human epigenomes. nature, 518, 317–330. url http://dx.doi.org/10.1038/nature14488
François Chollet (2015) keras. url https://github.com/fchollet/keras
Gromek, E., pfeiffer, a., matths, h. et al. (2015) conserved epigenomic signals in mice and humans reveal immune basis of alzheimer’s disease. nature, 518, 365–369. url http://dx.doi.org/10.1038/nature14252
mekchin, R., alibas, D., scherer, s. et al. (2013) transcription factor EBFP is essential for the maintenance of B cell identity and prevents progression of alternative fates in committed cells. nature immunology, 14, 867–875. url http://www.nature.com/doifinder/10.1038/ni.2641
Denoising Genome-wide Histone ChIP-seq with Convolutional Neural Networks

