

Table 3 Important parameters for each peak calling algorithm

Algorithm	Important parameters
CCAT	<p>Minimum score: minimum score of normalized difference</p> <p>Minimum count: minimum number of read counts at the peak</p> <p>Moving Step: step of window sliding</p> <p>SlidingWinSize: size of sliding window</p> <p>Bootstrap pass: number of passes in the bootstrapping process</p>
MACS	<p>NoLambda: if True, MACS will use fixed background lambda as local lambda for every peak region</p> <p>NoModel: whether or not to build the shifting model</p> <p>MFold: regions within MFOLD range of high-confidence enrichment ratio against background to build model</p> <p>PValue: <i>p</i>-value cutoff for peak detection</p>
SICER	<p>WindowSize: size of the windows to scan the genome width</p> <p>GapSize: allowed gap in base pairs between islands</p> <p>FDR: false discovery rate controlling significance</p>
ZINBA	<p>Selectmodel: Specifying select model = FALSE skips the model selection process altogether and may save a significant amount of time</p> <p>extension: average fragment library length (size selected)</p> <p>winSize: Selecting a larger window size increases speed of analysis but decreases resolution and sensitivity to detect enrichment</p> <p>offset: Smaller non-zero offset distances increase sensitivity but also increase computational burden</p> <p>FDR: FDR = TRUE specifies the model to use the FDR threshold rather than posterior probabilities. This typically results in more liberal peak calls. If false, then uses posterior probability to threshold peaks using 1-threshold.</p>
F-seq	<p>FeatureLength: feature length</p> <p>Threshold: standard deviations</p>