

Gene-view selection-random effects over sites and branches [BUSTED]



Sites



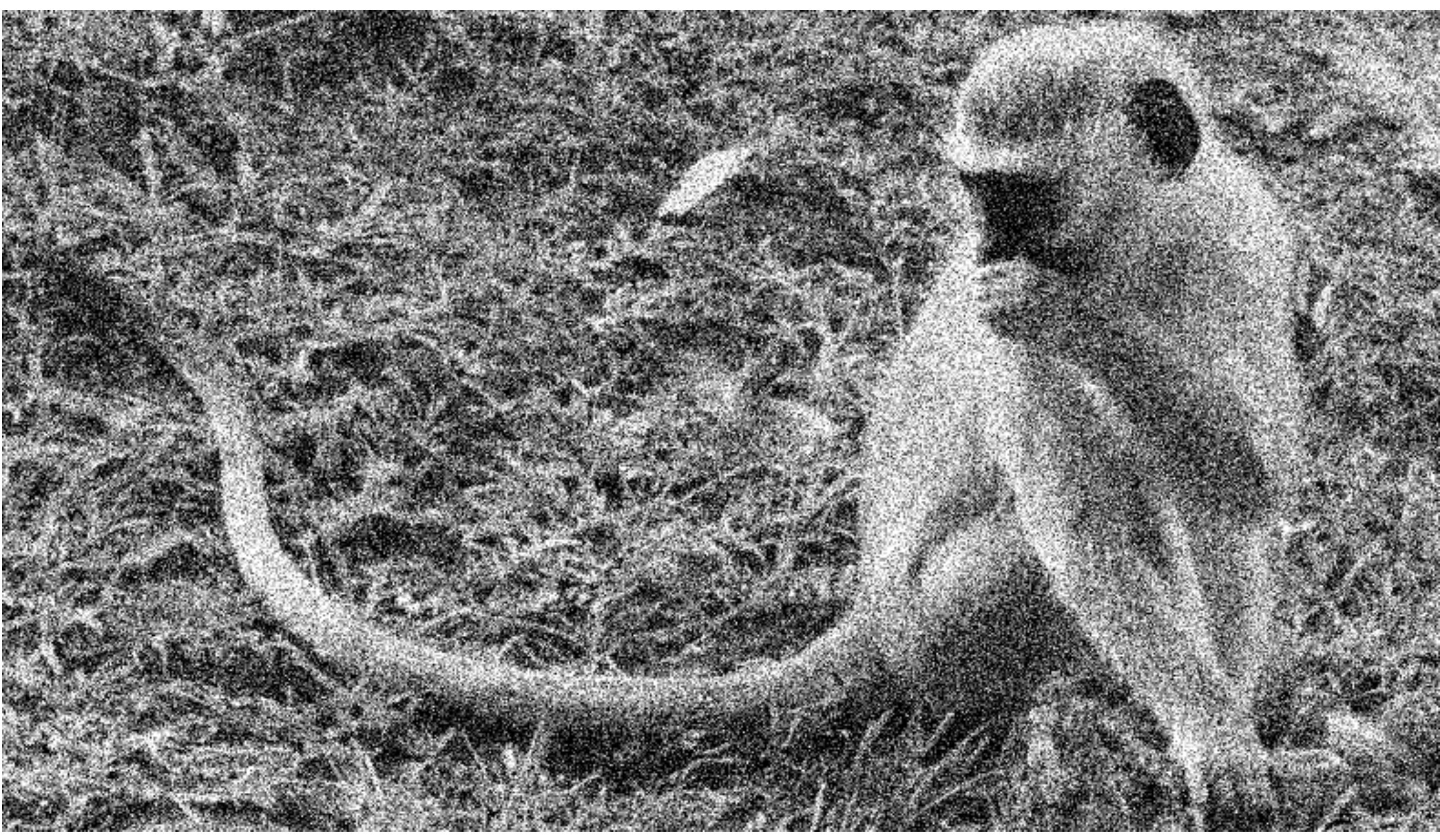


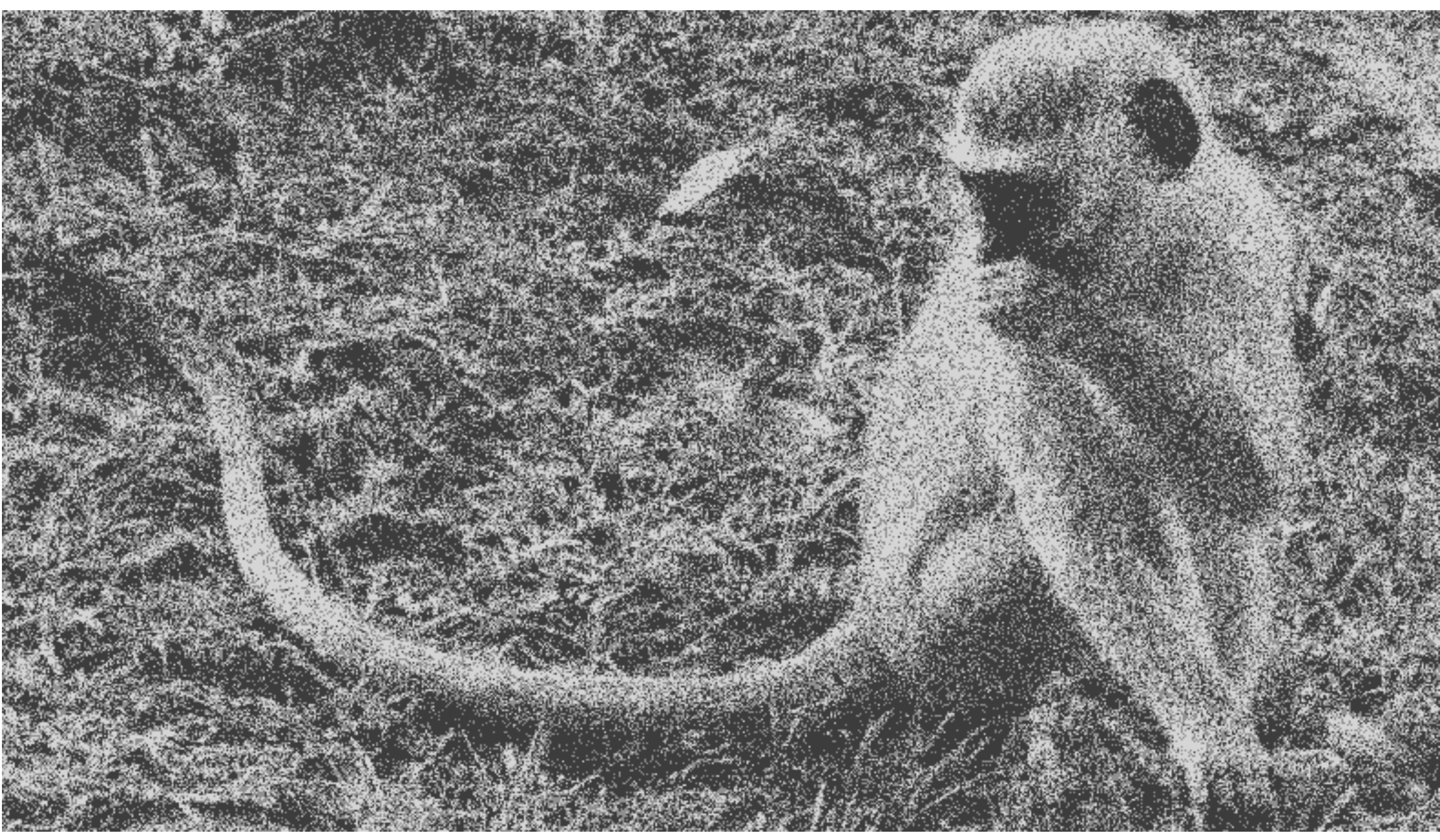


Is there enough **image area** that is sufficiently bright; allow each pixel to be one of K ($=3$) colors, chosen adaptively, e.g. to minimize perceptual differences



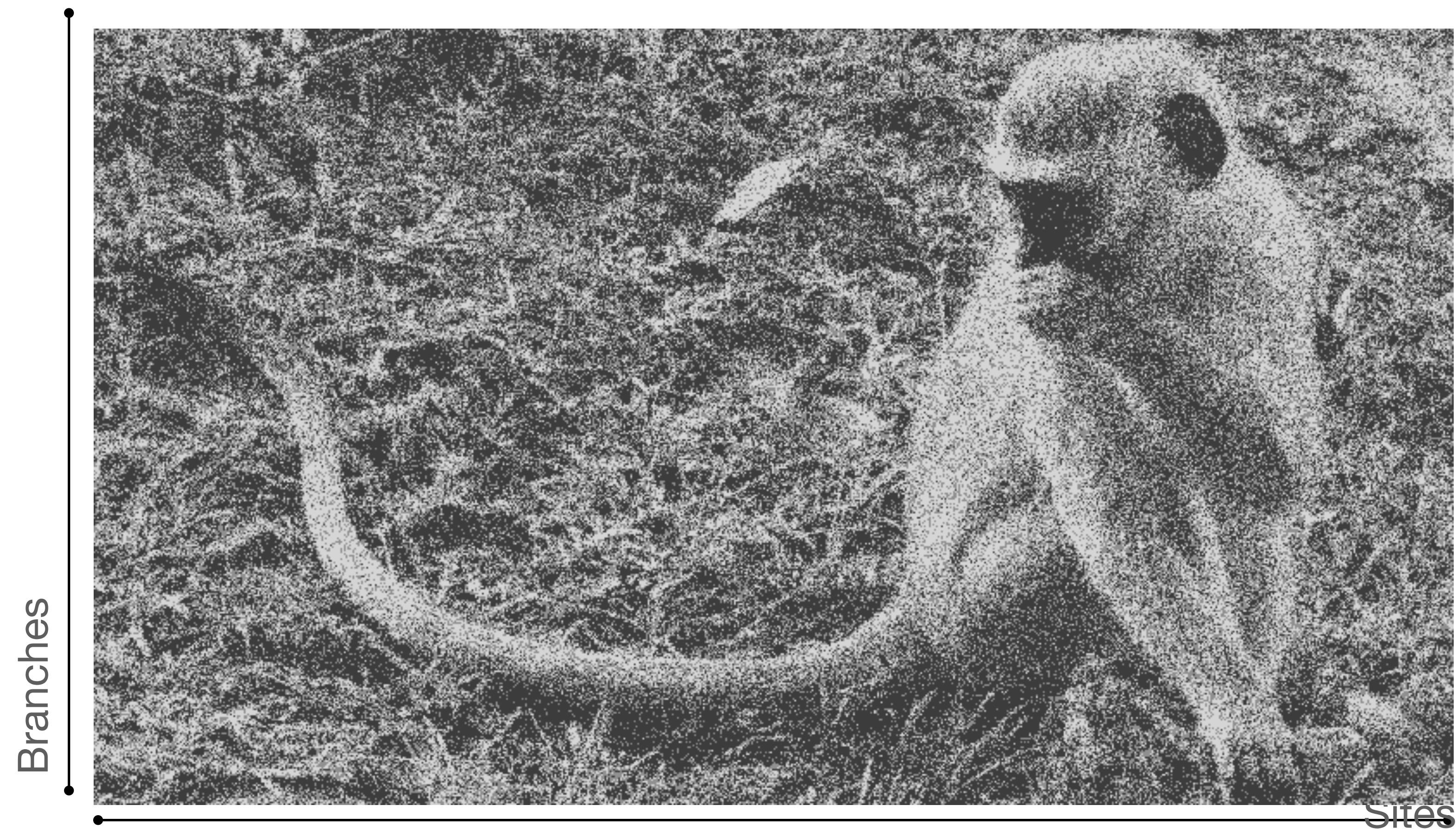
[BUSTED]: each branch-site combination is drawn from a K -bin (dS, dN) distribution. The distribution is estimated from the entire alignment. Tests if $dN/dS > 1$ for some branch/site pairs in the alignment







Gene-wide selection - random effects over sites and branches [BUSTED]



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BUSTED inference

- Because BUSTED is a random-effects method, it pools information across multiple sites and branches to gain power
- The cost to this pooling is lack of site-level **resolution**, i.e., it is not immediately obvious which sites and/or branches drive the signal
- Standard ways to extract individual site contributions to the overall signal is to perform a post-hoc analysis, such as empirical Bayes, or “category loading”
- For BUSTED, “category loading” is faster and experimentally better
- Can also compute exploratory evidence for selection support along individual branches at specific sites