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Real time quantitative imaging of transcriptional activity at the single cell level (bioRxiv)

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How plants transcriptionally operate developmental programs and responses to stress in time and space has been an important question in plant biology. Fluorescent protein reporters are commonly used to address this question, but their performance is limited at short timescales (<30 min) before the proteins get matured. Also, expression of fluorescent proteins can be influenced by various processes before and after maturation (e.g., RNA transport and protein degradation). To overcome these limitations, Alamos et al. implemented an mRNA fluorescent-tagging approach in Nicotiana benthamiana and Arabidopsis thaliana. In this system, the promoter of a gene is tagged with the PP7 (or MS2) viral DNA sequence that when transcribed accumulates fluorescent proteins already expressed in the nucleus. The signal can be immediately detected as a fluorescent spot, and the signal intensity reports the absolute count of RNA polymerase molecules actively transcribing the target gene. The authors generated stably transformed lines of Arabidopsis with construct targeting the heat shock protein HSP101 to monitor plant responses to heat stress.



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