

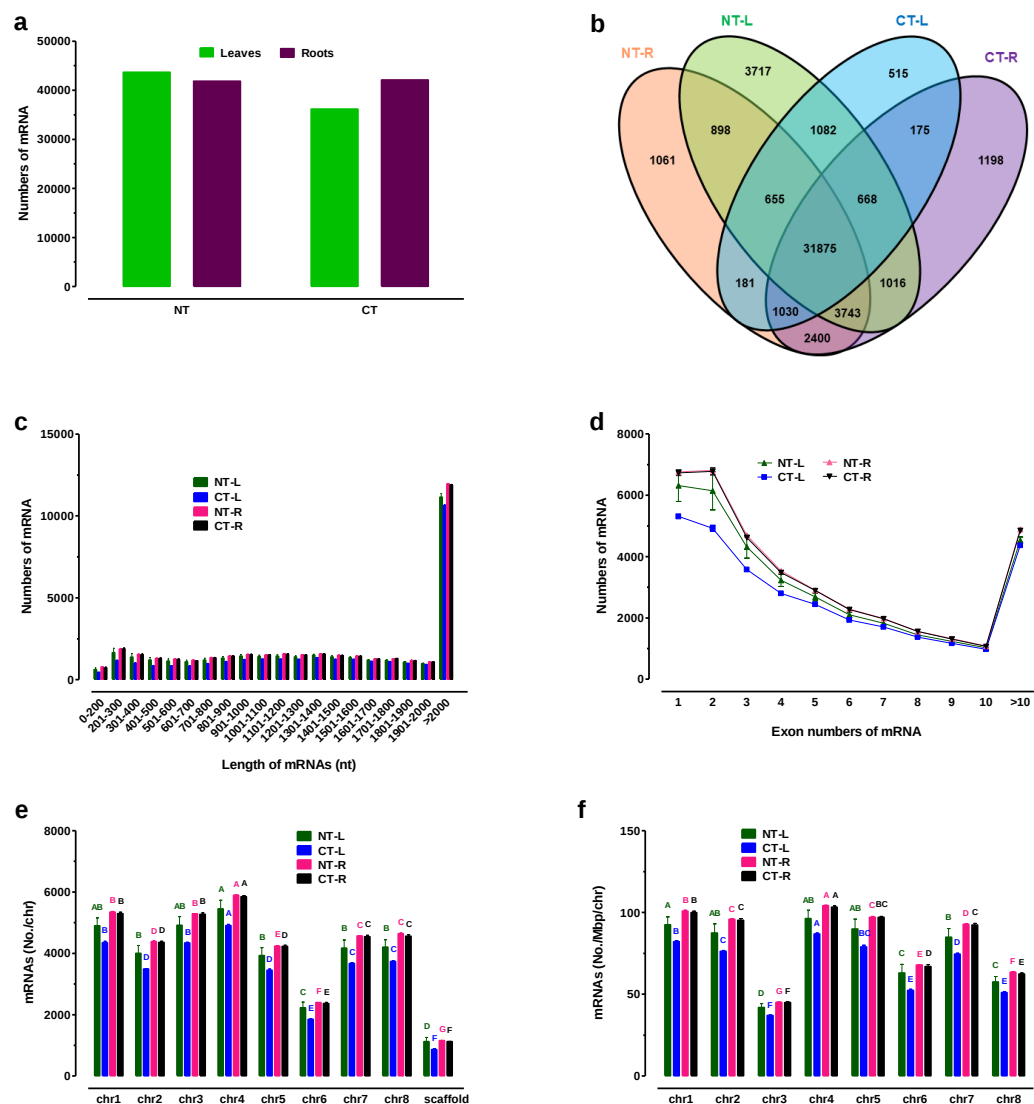


Figure S3. Identification of mRNAs by high-throughput sequencing in *M. truncatula* seedlings. **(a)** Total numbers of mRNA identified in leaves and roots. **(b)** Numbers of common/specific mRNA identified in NT-L, CT-L, NT-R and CT-R. The values shown in **(a)** and **(b)** were obtained from three independent RNAseq experiments. **(c)** Length distribution of mRNAs in leaves and roots. **(d)** Numbers of mRNA containing different numbers of exon in NT-L, CT-L, NT-R and CT-R. **(e)** Number distribution of mRNAs on eight chromosomes of leaves and roots with and without cold treatment. **(f)** Density distribution of mRNAs on eight chromosomes of leaves and roots with and without cold treatment. Capital letters indicate significant difference at $P < 0.05$ according to the t -test between chromosomes in the same treatment samples. Values

shown in (c) (d) (e) and (f) were means $\pm$ SE with three independent RNAseq experiments. NT: non-cold treated; CT: cold treated; L: leaves; R: roots.