

## **Systematic myostatin expression screening platform for identification and evaluation of myogenesis-related phytogetic in economic animal**

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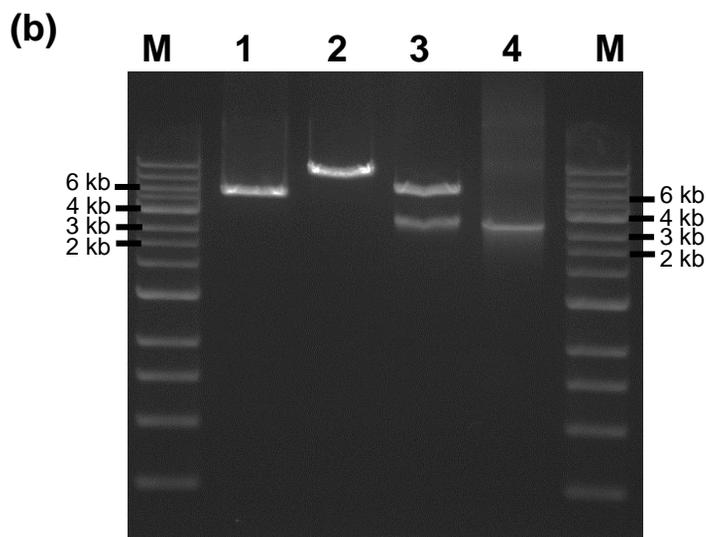
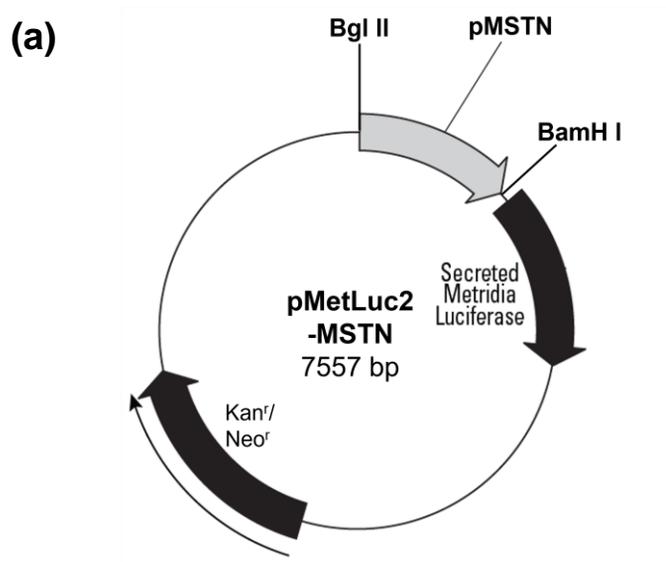
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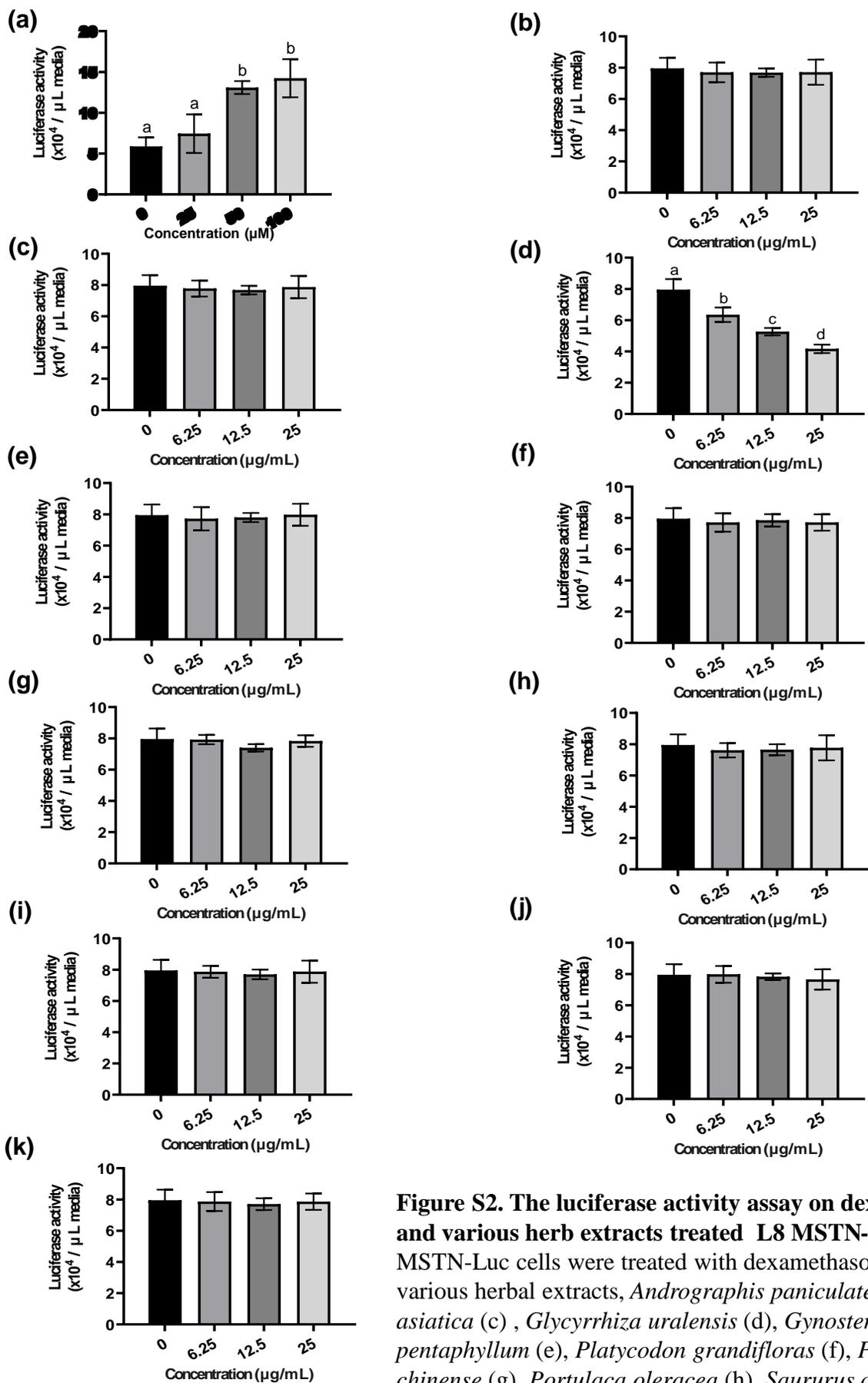
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**Figure S1. Myostatin promoter-reporter plasmid.** The 5'-flanking region (2.46 kb) upstream of the translation start site of the MSTN gene (GenBank accession no. AY204900) was inserted into pMetLuc2 plasmid (a), and confirmed by restriction enzyme digestion (b). Lane 1: pMetLuc2 plasmid after *Bgl*III digestion. Lane 2: pMetLuc2-MSTN plasmid after *Bgl*III digestion. Lane 3: pMetLuc2-MSTN plasmid after *Bgl*III and *Bam*H1 digestions. Lane 4: PCR amplification product of the 5'-flanking region (2.46 kb) upstream of the translation start site of MSTN gene. Lane M: 1-kb DNA ladder.



**Figure S2. The luciferase activity assay on dexamethasone and various herb extracts treated L8 MSTN-Luc cells.** L8 MSTN-Luc cells were treated with dexamethasone (a) or various herbal extracts, *Andrographis paniculate* (b), *Centella asiatica* (c), *Glycyrrhiza uralensis* (d), *Gynostemma pentaphyllum* (e), *Platycodon grandifloras* (f), *Polygonum chinense* (g), *Portulaca oleracea* (h), *Saururus chinensis* (i), *Smilax china* (j) and *Taraxacum campyloides* (k). L8 MSTN-Luc cells luciferase activity was measured after 24h treatment. Each data point represents the mean  $\pm$  SEM. Different letters (a-d) denote significant differences ( $p < .05$ ) amount treatments.