

Evaluation and application of an efficient plant DNA extraction protocol for laboratory and field testing

Qi WANG¹, Xiaoxia SHEN², Tian QIU¹, Wei WU³, Lin LI¹, Zhi'an WANG², Huixia SHOU¹✉

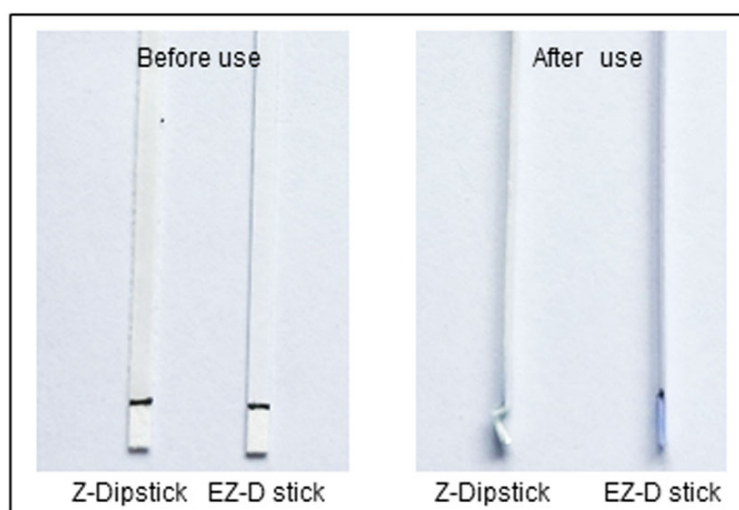


Fig. S1 Comparison of Z-Dipstick and EZ-D stick during DNA extraction.

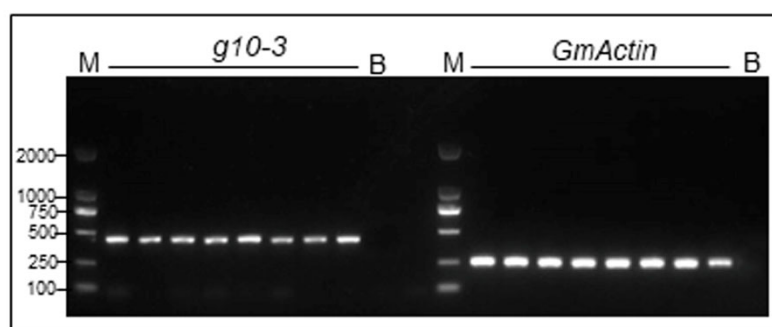


Fig. S2 DNA content in a single EZ-D stick can be used for at least 16 PCR reactions. An EZ-D stick containing the genomic DNA from transgenic soybean ZUTS-33 was dipped sequentially into 16 PCR tubes containing only water, before adding the rest of the PCR components. M, DL2000 marker; B, blank control.

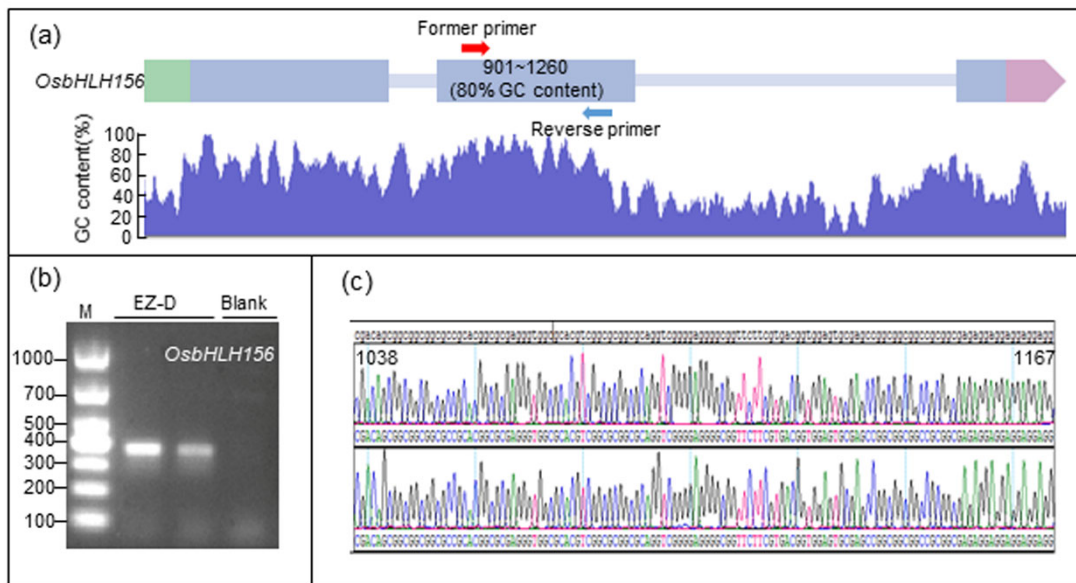


Fig. S3 DNA extracted by EZ-D can be used for PCR amplification of fragment with high GC content. (a) The amplification region of *OsbHLH156* has a high GC content. (b) The high GC content region of *OsbHLH156* was successfully amplified by the EZ-D based PCR method. M, DL2000 marker; B, blank control. (c) Partial sequencing results of *OsbHLH156* amplified products.