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Gold nanoparticle biosensors, a novel application in gene transformation and expression

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Abstract	15
The conventional techniques of PCR, Southern blot, Northern blot, in situ hybridization, and	16
RNase protection assay have long been used to investigate transformation and expression of	17
genes, but most of them are time-consuming and have relatively low sensitivity. In recent years,	18
applying biosensors for molecular identification of biomolecules has been expanding	19
significantly. Hence in this study, Zabol mildew melon was used as a model plant to introduce	20
new DNA and RNA-based biosensors for confirming gene transformation and expression. First,	21
the melon seeds were grown in vivo and Agrobacterium tumefaciens LBA4404 was used to	22
introduce GUS reporter gene to the plant. In order to analyze GUS gene transformation and	23
expression, probes were designed based on DNA, RNA, and cDNA of GUS gene sequence.	24
Then, the analysis was performed using probes attached to gold nanoparticles to observe color	25
change of the solution in presence of the target biomolecules. Hybridization of the probes with	26
target molecules was evaluated at a wavelength of 400 to 700 nm and maximum change was	27
observed in the wavelength range of 550 to 650 nm. In addition, detection limit of the assay was	28
0.25 mg/µL and linear regression showed the relationship between different concentrations of the	29
genomic DNA and absorbance. Consequently, results showed that application of detectors	30
attached to gold nanoparticles for investigation on gene transformation and expression is more	31
rapid, specific and economic compared to the biochemical and molecular techniques. These tests	32
can be carried out with initial optimization at research centers using the least facilities; hence	33
there will be no need for special equipment.	34
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Keywords: gold nanoparticles, unmodified probe, GUS reporter gene, PCR, RT-PCR	36
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