Supporting Information

For

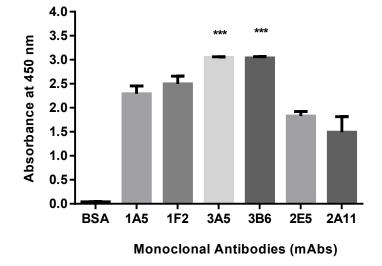
Development and validation of a novel lateral flow immunoassay device for detection of aflatoxins in soy-based foods

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S.1 Specificity of the mAbs for AFB1 by ELISA assay.

High binding EIA/RIA microplates were coated with 0.5 μ g of AFB1-BSA conjugate and kept overnight at 4°C. The plates were blocked with 3% gelatin in PBST. An aliquot of 100 μ L of each mAbs (1:3000) was added for 1h at 37°C. The binding was identified using a goat anti-mouse IgG-HRP (1:5000) and developed using 100 μ L of 4-(4-amino-3,5-dimethylphenyl)-2,6-dimethylaniline (TMB). The reaction was stopped by the addition of 50 μ L of 2M sulfuric acid, and the absorbance was measured at 450 nm. All statistic tests were performed using by two-way ANOVA. ***Denotes p<0.05.