

Supplementary Information

A Sensitive Aptasensor Using Biotin-Streptavidin System for Patulin Detection in Apple Juice

Xiaoqian Tang ^{1,2,3,4,5,6}, Qi Zhang ^{1,2,3,4,5,*}, Maria Isabel Pividori ⁷, Zhaowei Zhang ^{1,2,3}, Jean-Louis Marty ⁸ and Gaëlle Catanante ^{6,*}

¹ Oil Crops Research Institute, Chinese Academy of Agricultural Sciences, 430062 Wuhan, China; wtxqtutu@163.com (X.T.); zhaowei_zhang@126.com (Z.Z.)

² Key Laboratory of Detection for Mycotoxins, Ministry of Agriculture and Rural Affairs, 430062 Wuhan, China

³ Laboratory of Risk Assessment for Oilseeds Products, Ministry of Agriculture and Rural Affairs, 430062 Wuhan, China

⁴ Key Laboratory of Biology and Genetic Improvement of Oil Crops, Ministry of Agriculture and Rural Affairs, 430062 Wuhan, China

⁵ Quality Inspection and Test Center for Oilseeds Products, Ministry of Agriculture and Rural Affairs, 430062 Wuhan, China

⁶ Laboratoire BAE-LBBM USR 3579, Université de Perpignan Via Domitia, 52 Avenue Paul Alduy, CEDEX, 66860 Perpignan, France

⁷ Grup de Sensors i Biosensors, Departament de Química, Universitat Autònoma de Barcelona, 08193 Bellaterra, Spain; Isabel.Pividori@uab.cat

⁸ Sensbiotech, 21 rue de Nogarede, 66400 Ceret, France; jlmarty@univ-perp.fr

* Correspondence: zhangqi@oilcrops.cn(Q.Z.); gaelle.catanante@univ-perp.fr (G.C.)

Buffers and solutions:

The composition of different buffer were prepared as the follows:

Coating buffer: 0.01 mol/L phosphate buffer saline (PBS), pH 7.4; supplemental with 0.5 % w/v BSA

Conjugate diluting buffer: 0.002 mol/L borate pH 7.0; supplemental with 10% w/v sucrose.

Binding buffer: 0.1 mol/L phosphate buffer, pH 7.4, supplemental with 0.005 mol/L MgCl₂.

Running buffer: 0.1 mol/L binding buffer, pH 7.4; supplemental with 5% BSA (w/v), 0.05 % Tween 20 (v/v)

Table S1. Optimize different ratio of BIO-aptamer and DIG-aptamer.

Ratio (mol:mol)	BIO-Aptamer (μmol/L)	DIG-Aptamer (μmol/L)	Inhibition Value (%)
1:1	0.025	0.025	13
1:2	0.025	0.050	12.3
1:4	0.025	0.125	None
1:6	0.025	0.150	None

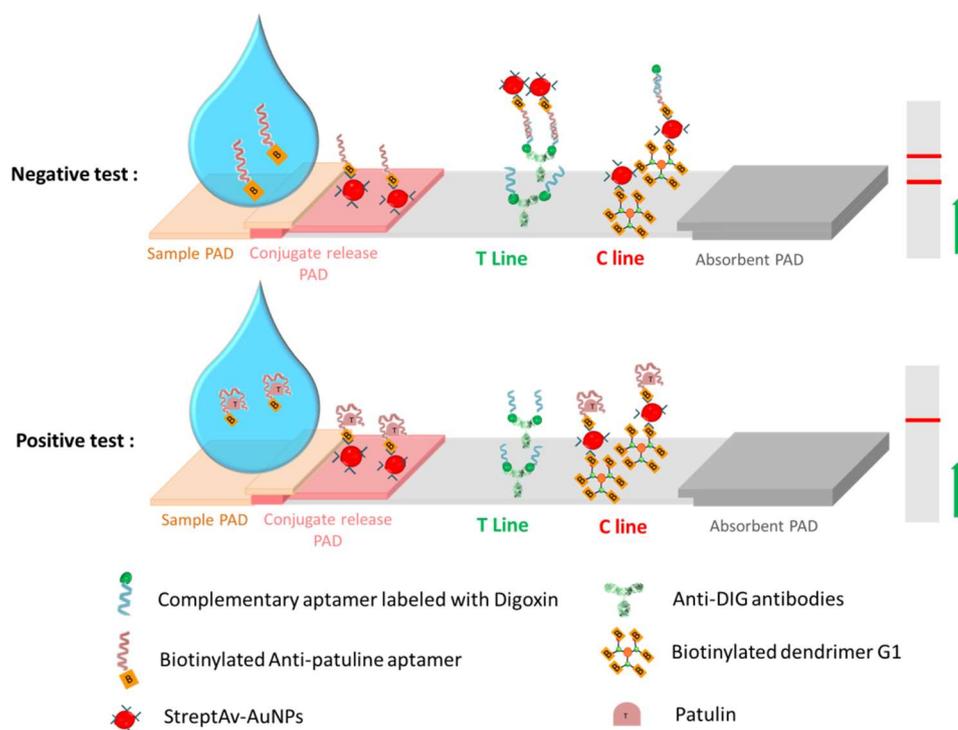


Figure S1. Schematic of the Method II, the different with Method I was the DIG-aptamer was fixed on the T line according the anti-DIG antibody.

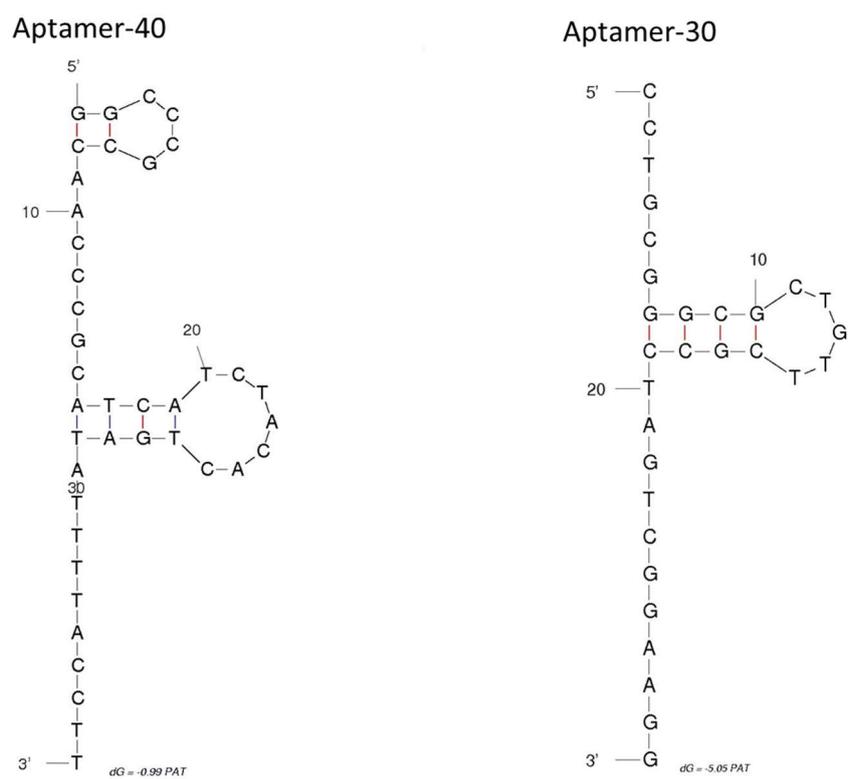


Figure S2. The structure of aptamer-40 and aptamer-30.