

Supplemental Figures

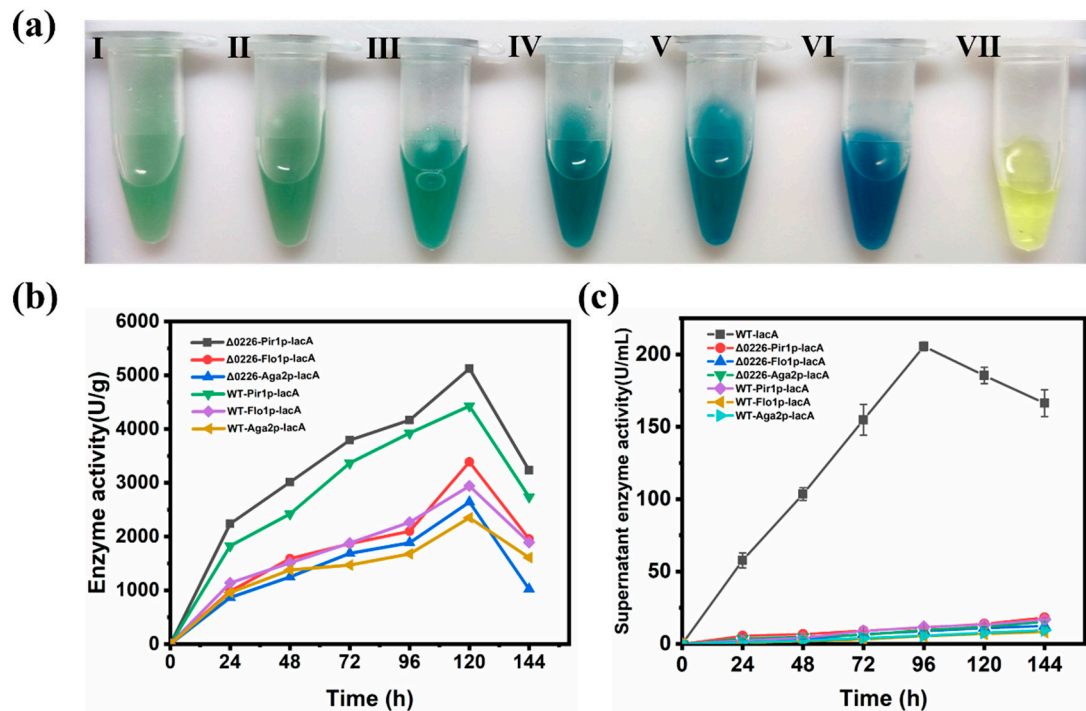


Figure S1 Expression of enzymes from six surface-displayed engineered strains.

(a) I: WT - Aga2p-lacA, II: $\Delta 0226$ -Aga2p-lacA, III: WT-Flo1p-lacA, IV: $\Delta 0226$ -Flo1p-lacA, V: WT-Pir1p-lacA, VI: $\Delta 0226$ -Pir1p-lacA, VII: control. the darker the color is, the better the expression of β -galactosidase. The enzyme activity of $\Delta 0226$ -Pir1p-lacA, $\Delta 0226$ -Flo1p-lacA, $\Delta 0226$ -Aga2p-lacA, WT-Pir1p-lacA, WT-Flo1p-lacA, and WT-Aga2p-lacA in (b) cell precipitates and (c) supernatant, respectively. The enzymatic activity of $\Delta 0226$ -Pir1p/ Flo1p / Aga2p -lacA was higher than that of WT-Pir1p/ Flo1p / Aga2p -lacA, indicating that knockout of $\Delta 0226$ strain could increase the efficiency of surface display and allow more anchoring proteins to be anchored to the cell wall of *P. pastoris*.