Assay of anthocyanin biosynthetic enzyme activity

Preparation of buffer

1. 25 mM HEPES buffer (pH 7.4) is prepared with EDTA (0.2 mM), AsA (2 mM), and PVP (2%), which was kept in -4°C refrigerator.

Leaf tissue sample extraction

2. 0.2 g of leaf samples were ground in 2 ml of ice-cold 25 mM HEPES buffer (pH 7.4). The homogenates were centrifuged at 4°C for 20 min at the speed of 12,000 r.p.m, and removed supernatants to assay the enzymatic activities of CHS, CHI, F3H, DFR, ANS and ANR, using an ELISA Kit (U.S.A TSZ biological Trade Co., Ltd.) according to the manufacturer’s instructions.

Add standard

3. Set standard wells, testing sample wells. Add standard 50 µl to standard well.

Add sample

4. Set blank wells separately (blank comparison wells don’t add sample and HRP-Conjugate reagent, other each step operation is same). Testing sample well. add sample dilution 40 µl to testing sample wells, then add testing sample 10 µl (sample final dilution is 5-fold), add sample to wells, don’t touch the well wall as far as possible, and mix gently.

Incubate

5. After closing plate with closure plate membrane, incubate for 30 min at 37°C.

Configure liquid

6. 30-fold wash solution diluted 30-fold with distilled water and reserve.
7 Uncover closure plant membrane, discard liquid, dry by swing, add washing buffer to every well, still for 30s then drain, repeat 5 times, dry by pat.

8 Add HRP-Conjugate reagent 50 µl to each well, except blank well.

9 Incubate is operated with step 5, and washing is operated with step 7.

10 Add chromogen solution A 50 µl and chromogen solution B to each well, evade the light preservation for 15 min at 37°C.

11 Add stop solution 50 µl to each well, stop the reaction (the blue color change to yellow color).

12 Take adding stopbland well as zero, read absorbance at 450nm after adding stop solution and within 15 mim

Citation: Assay of anthocyanin biosynthetic enzyme activity. https://dx.doi.org/10.17504/protocols.io.h2mb8c6

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