Supporting Information to:

Prevention of Early Liver Injury by Breviscapine in Streptozotocin-Induced Diabetic Rats

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Fig. 1S Histological observation of breviscapine-treated diabetic rat induced by STZ. The liver sections were stained with Oil red O. Representative liver sections are shown (magnification ×100). Breviscapine was orally administrated at 48 h after STZ injection in rats. STZ (65 mg/kg) was administered to rats by intraperitoneal injection. The liver was obtained 8 weeks after STZ injection. A: Control; B: STZ; C: STZ + Breviscapine (20 mg/kg).

Fig. 2S Histological observation of breviscapine-treated diabetic rat induced by STZ. The liver sections were stained with Masson’s trichrome. Representative liver sections were shown (magnification ×100). Breviscapine was orally administrated at 48 h after STZ injection in rats. STZ (65 mg/kg) was administered to rats by intraperitoneal injection. The liver was obtained 8 weeks after STZ injection. A: Control; B: STZ; C: STZ + Breviscapine (20 mg/kg).
Fig. 3S ED-1-immunoreactivity in breviscapine-treated diabetic rat liver induced by STZ. Representative liver sections are shown (magnification ×100).

Breviscapine was orally administrated at 48 h after STZ injection in rats. STZ (65 mg/kg) was administered to rats by intraperitoneal injection. The liver was obtained 8 weeks after STZ injection. **A:** Control; **B:** STZ; **C:** STZ + Breviscapine (20 mg/kg).