

# Edoxaban

**Catalog No: tcsc1331**



## Available Sizes

**Size:** 5mg

**Size:** 10mg

**Size:** 50mg

**Size:** 100mg



## Specifications

**CAS No:**

480449-70-5

**Formula:**

$C_{24}H_{30}ClN_7O_4S$

**Pathway:**

Metabolic Enzyme/Protease

**Target:**

Factor Xa

**Purity / Grade:**

>98%

**Solubility:**

10 mM in DMSO

**Alternative Names:**

DU-176

**Observed Molecular Weight:**

548.06

## Product Description

Edoxaban(DU-176) is an oral factor Xa (FXa) inhibitor in clinical development for stroke prevention

IC50 Value:

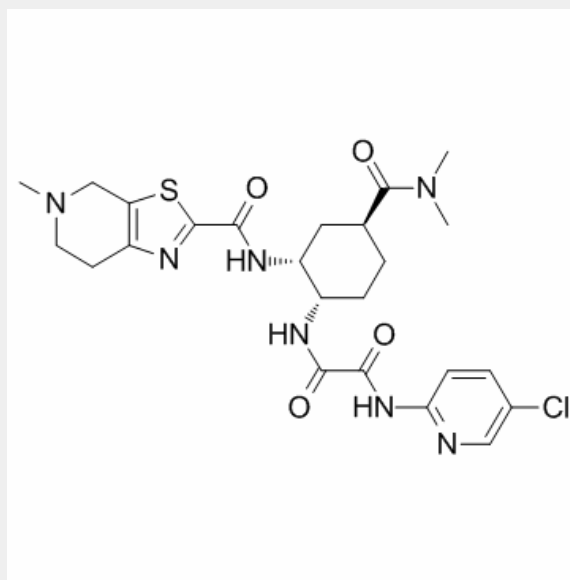
Target: factor Xa

Edoxaban is an oral factor Xa (FXa) inhibitor in clinical development for stroke prevention in patients with atrial fibrillation, an elderly population that frequently receives aspirin (ASA) and/or nonsteroidal anti-inflammatory drugs for concurrent illnesses[1].

in vitro: Edoxaban PK was not affected by concomitant low-dose ASA or naproxen, but high-dose ASA increased systemic exposure of edoxaban by approximately 30%. The effects of edoxaban on prothrombin time, activated partial thromboplastin time, international normalized ratio, anti-FXa, and intrinsic FXa activity were not influenced by administration with ASA or naproxen. Inhibition of platelet aggregation by high-dose ASA, low-dose ASA, or naproxen was not affected by edoxaban[1].

in vivo: Forty-eight subjects, aged 18 to 45 years, received either edoxaban 60 mg once daily × 7 days (n = 24) or digoxin 0.25 mg twice daily × 2 days and once daily × 5 days (n = 24) and then concomitantly for 7 days. Serial blood and urine samples were collected for digoxin and edoxaban concentrations on days 7 and 14. Serial coagulation assays were measured for edoxaban on days 7 and 14. Edoxaban PK parameters demonstrated mild increases in area under the curve and peak concentrations of 9.5% and 15.6%, respectively[2].

Clinical trial: Pharmacokinetics, biotransformation, and mass balance of edoxaban, a selective, direct factor Xa inhibitor, in humans was reported[3].



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