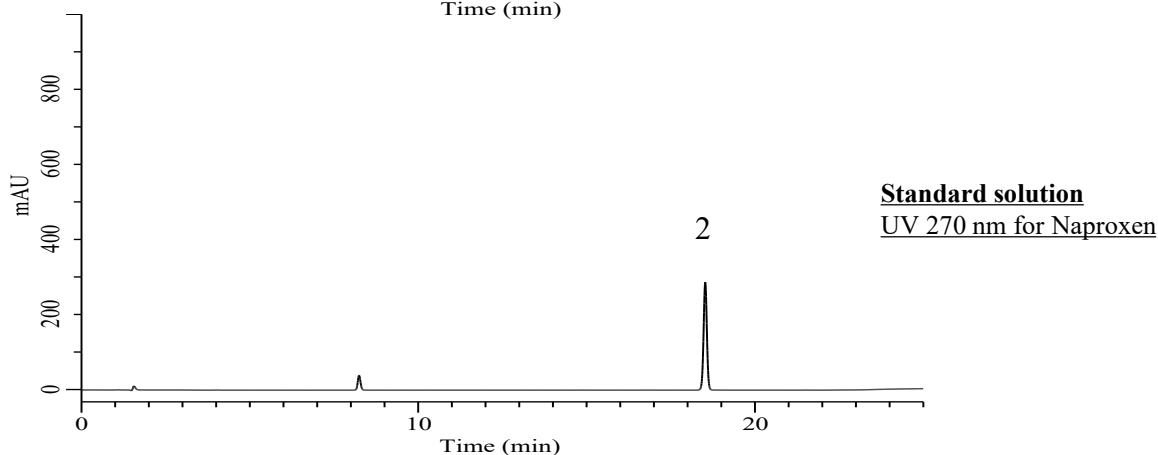
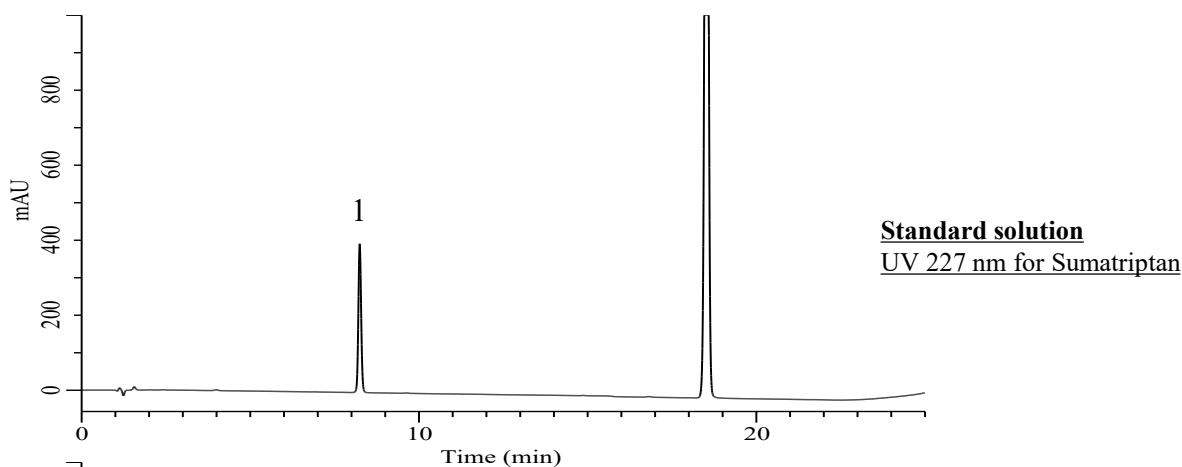


Analysis of Sumatriptan succinate and Naproxen sodium

(Under the Condition of the draft for USP, Sumatriptan and Naproxen Sodium Tablets, Assay)



Conditions

System : Primaide HPLC system (HITACHI)

Column : Inertsil Ph-3 (GL Sciences Inc.)
(3 μ m, 100 x 4.6 mm I.D.)

Column Cat. No. : 5020-05044

Eluent : A) CH₃CN
B) 0.1 % TFA in H₂O

Time (min)	A (vol %)	B (vol %)
0	5	95
20	50	50
22	5	95
25	5	95

Flow Rate : 1.2 mL/min

Col. Temp. : 35 °C

Detection : UV 227 nm for Sumatriptan (1430 DAD)
UV 270 nm for Naproxen (1430 DAD)

Injection Vol. : 10 μ L

Sample : Standard

Analyte:

1. Sumatriptan succinate 0.048 mg/mL

2. Naproxen sodium 0.2 mg/mL

Relative retention time

Sumatriptan succinate : 1.0 (8.2)

Naproxen sodium : 2.25 (18.5)

Tailing factor

peak area of 1 : 1.09 (\leq 1.8)

peak area of 2 : 1.02 (\leq 1.8)

RSD of the

peak area of 1 (%) (n=6) : 0.82 (\leq 1.0)

peak area of 2 (%) (n=6) : 0.77 (\leq 1.0)