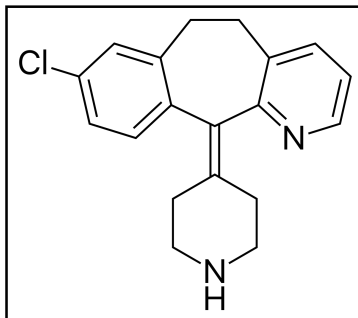


**Desloratadine**  
**2.50 mg in Co-formulation Tablet with 120 mg of Pseudoephedrine**

**Structure:**



**Molecular Formula and Mass:** C<sub>19</sub>H<sub>19</sub>ClN<sub>2</sub> – 310.82

**Category:** Antihistamine

**Sample:**

Grind one tablet and dissolve in 7.50 mL methanol. Shake at least 10 min and filter. Final concentration of sample solution = 2.50 mg/7.50 mL = 0.333 mg/mL, which is the required concentration representing 100%.

**Standards:**

High Standard:

The high limit is 115%; therefore the concentration of the high standard = (0.333 mg/mL X 1.15 = 0.383 mg/mL. Weigh approximately 11.5 mg of standard. If you weighed 11.6 mg of standard, dissolve it in: (11.6 mg)/(0.383 mg/mL) = 30.3 mL of methanol. This makes the high standard solution concentration equal to 0.383 mg/mL.

Low Standard:

The low limit is 85%; therefore the concentration of the low standard = (0.333 mg/mL X 0.85 = 0.283 mg/mL. Dilute 1.00 mL of high standard to 1.35 mL by adding 0.35 mL of methanol (1.15/0.85 = 1.35).

**Spotting:**

Spot on the 5 X 10 cm silica gel TLC aluminium plate with 3.00 µL aliquots as follows:

Left spot	low standard (85%) = 0.850 µg
Center Spot	100% sample = 1.00 µg
Right Spot	high standard (115%) = 1.15 µg

**Development:**

Mix 24.0 mL of ethyl acetate, 3.00 mL methanol and 1.00 mL concentrated ammonia. Develop the plate in a small glass chamber with approximately 20.0 mL of this solution until the solvent front reaches within 1 cm of the top of the TLC plate.

(R<sub>f</sub> = 0.050)

**Detection:**

UV:

Dry the plate and observe under ultraviolet light at 254 nm. Observe the intensities and the sizes of the spots.

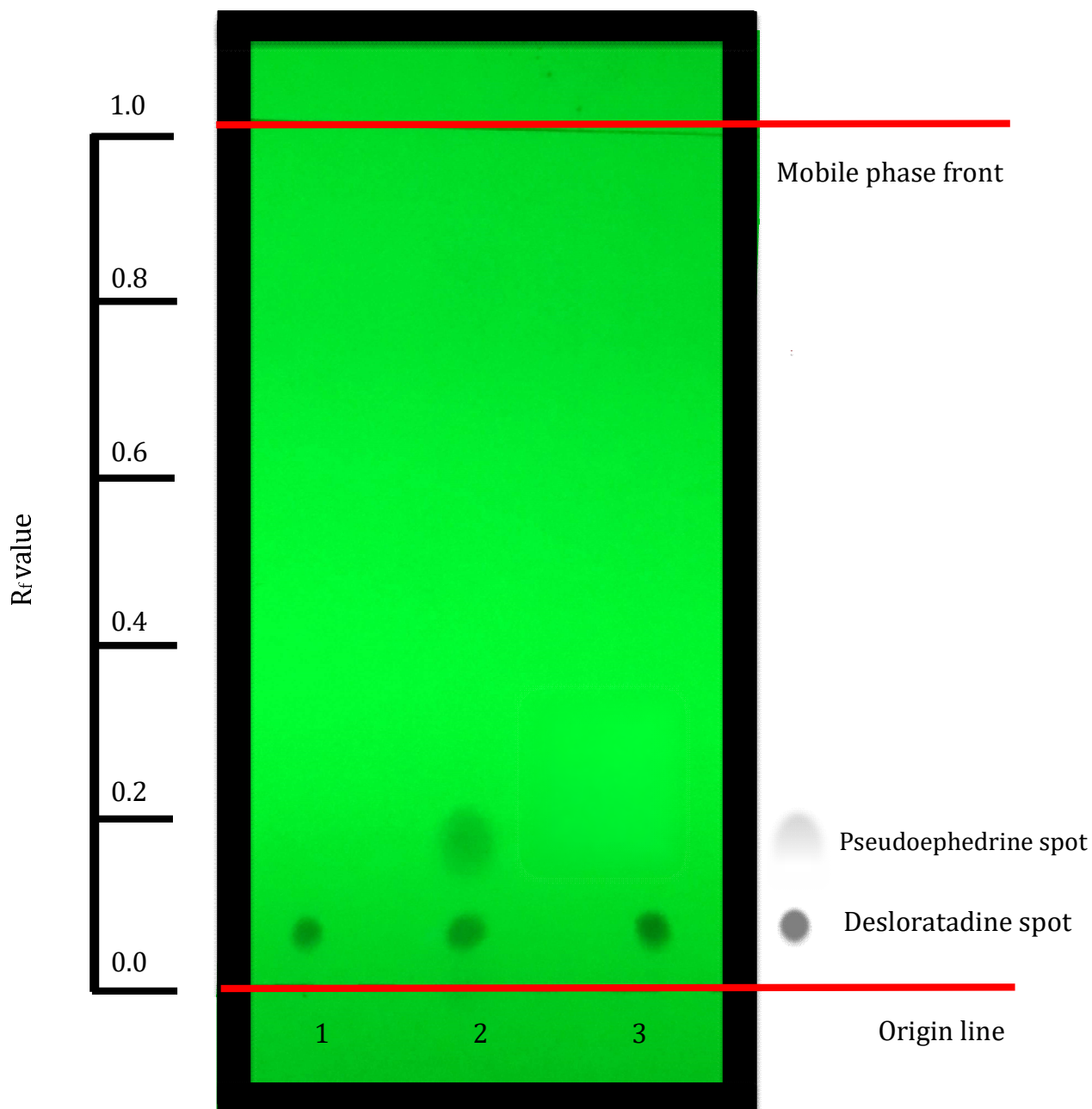


Plate observed under ultraviolet light at 254 nm

Lane 1: Low standard (85%) = 0.850  $\mu\text{g}$

Lane 2: 100% sample = 1.00  $\mu\text{g}$

Lane 3: High standard (115%) = 1.15  $\mu\text{g}$