

Figures

Figure 1

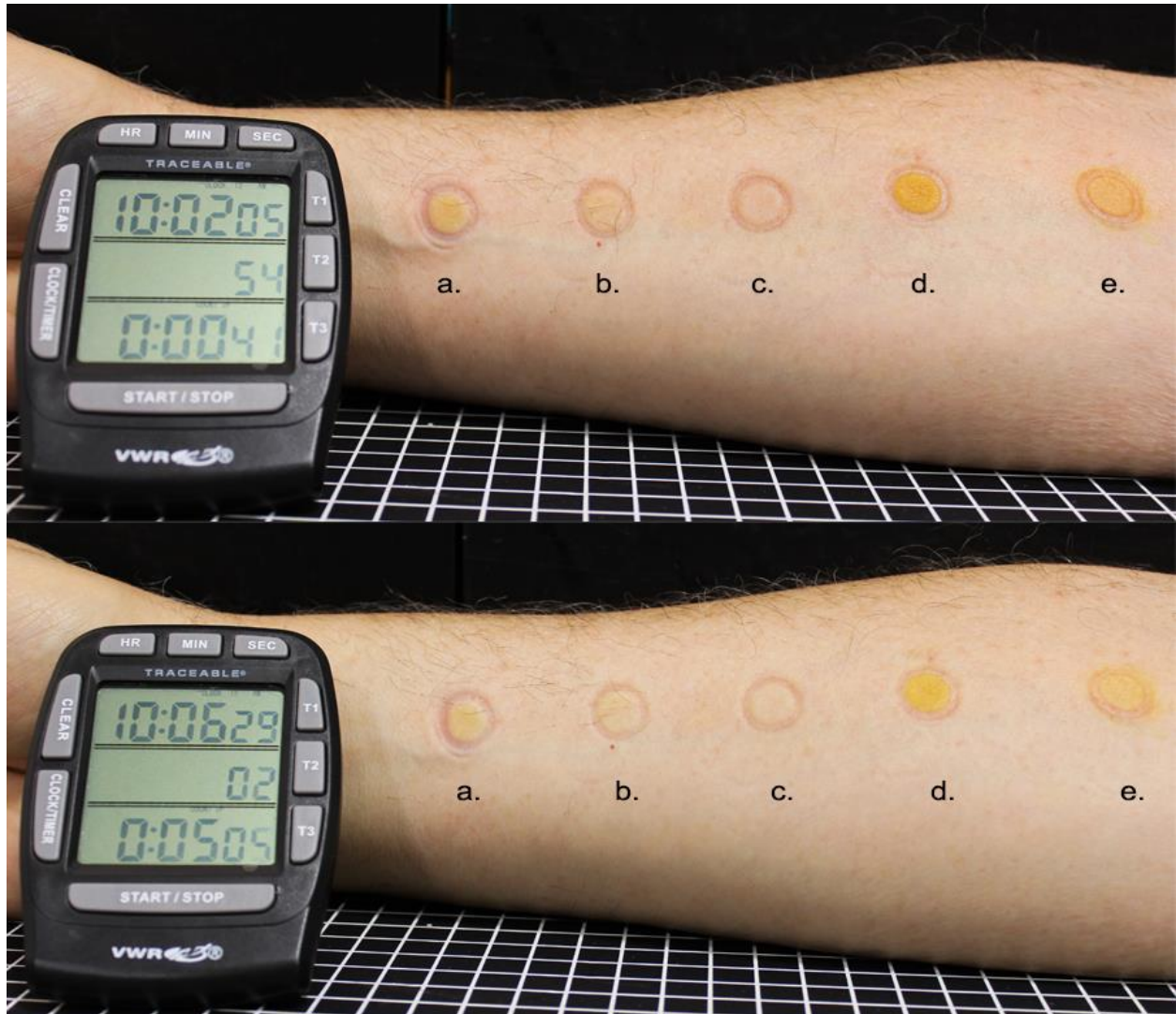


Figure 2

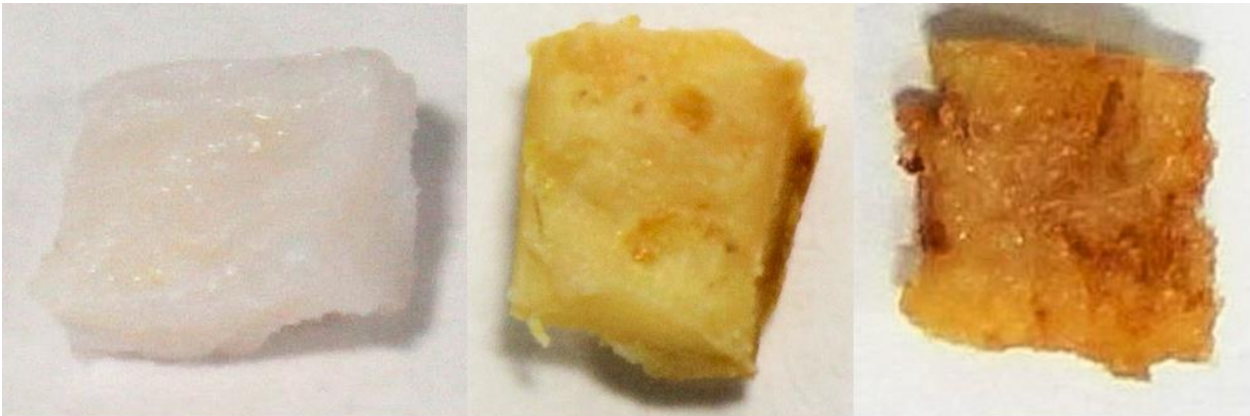


a.



b.

Figure 3



a.

b.

c.

Figure 4

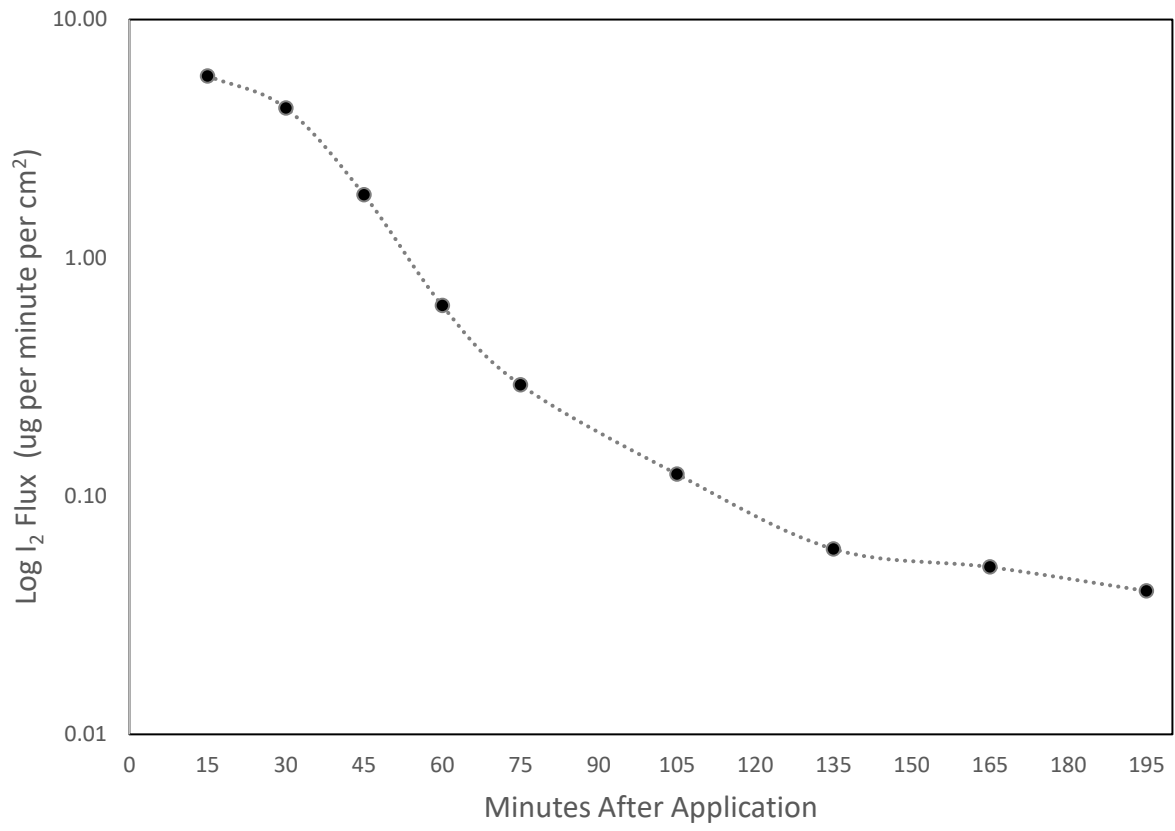


Figure Legends

Figure 1.

One mL of (a) 300 ppm I₂ in water; (b) 10% PVP; (c) 7800 ppm I₂-glycerin, (d) Lugol's solution and (e) were simultaneously contacted to the forearm of a volunteer for 3 minutes and residual was removed with a Dacron alcohol wiped. Images were taken immediately after application (top) and after 5 minutes (bottom).

Figure 2.

Figure 2. (a) Iodine staining of pig skin epidermis after application of 30 uL of 66,000 ppm I₂-glycerin to a 1.77 cm² circular piece of pig skin. (b) I₂ detected (aqua color) in purplish hypodermis tissue using a SenSafe Iodine Check test after removal of the epidermal and dermal skin layers.

Figure 3.

Cubes of hypodermis tissue 5 x 5 mm were sectioned, hydrated for 30 minutes and then patted dry. Cubes of hypodermis tissue were submerged in 1 ml of the following 3 compositions: (a) 10% PVP-I, (b) Lugol's solution, and (c) glycerin with 15,200 ppm I₂. Each piece of hypodermis tissue was removed after 15 minutes and rinsed thoroughly with water until a color change was not detected. The degree of staining was proportional to I₂ exposure.

Figure 4.

I₂ flux (ug/min-cm²) at 34°C from a hydrated 1.77 cm² piece of pig skin treated with 10 ul of a 66,000 ppm I₂-glycerin composition.

