QUINTEST® is a five-test skin test device designed to apply allergenic extract using the puncture technique.

QUINTEST® is designed for use by allergy practitioners who are trained in the application and interpretation of the puncture technique and who are trained in the recognition and treatment of adverse allergic reactions should they occur.

QUINTEST® Multiple Skin Test System consists of three components: The QUINTEST® skin test device (Figure 1), the QUINTIP® reservoirs (Figure 2) that contain the extracts, and the tray (Figure 3) that holds the allergenic extracts. The test devices are packaged two per sterilized blister pack container, 16 units per shelf pack. The QUINTEST® covered tray (Figure 3) is provided as a single unit that holds 60 QUINTIP® reservoirs (Figure 2) and accommodates 12 QUINTEST® skin test devices (Figure 1).
Each tray holds 12 QUINTEST® devices. Once loaded, it can be used immediately or covered and stored in the refrigerator until needed. When the extract-filled tray is not in use, store in the refrigerator with the test devices submerged in the reservoirs. Because of the breakaway design, if less than five test sites are desired, the undesired probe(s) can be removed before use.

SKIN TEST PROCEDURE
Prior to testing, clean the test area with alcohol and allow to dry. (Refer to Hollister-Stier Laboratories’ “Allergenic Extracts for Scratch and Prick/Puncture Testing” or other manufacturer’s package insert for more detailed instructions.) Remove a QUINTEST® skin test device from a prepared tray by lifting it vertically, being careful not to touch the tips on the reservoirs. With the QUINTEST® perpendicular to the skin, press down on the skin with medium pressure and rock the device end to end, without lifting the device from the skin, twice to ensure all tines have contacted the skin. Remove by lifting vertically and discard the QUINTEST®. Faint circles about 4mm in diameter created by the device’s depth control guards should remain at the test site indicating the correct amount of pressure was applied. The visibility of the circles will vary between patients according to the thickness, fragility, or pigmentation of their skin. The amount of pressure needed to produce a satisfactory test site requires that all of the five depth guards come in full contact with the skin. Following the above instructions will produce five uniform skin tests all at the proper depth without inducing bleeding. All test sites are three centimeters apart.

PRODUCT DESCRIPTION
The QUINTEST® device is a molded plastic handle with five testing probes (Figure 1). Inserted into each testing probe is a stainless steel lancet tip that protrudes from the probe enough to give the proper testing depth (Figure 4). The testing probes have a breakaway design for flexibility in use and easy disposal. The QUINTEST® device is to be used once and discarded.

SET-UP INSTRUCTIONS
Remove the QUINTEST® covered tray from the box. The QUINTEST® covered tray is supplied with pressure-sensitive labels. Each label strip has five designated areas corresponding to the reservoirs. Write the name of each antigen in the area provided, remove the label backing and place on the tray to correspond with the antigen. The reservoirs are provided in a sterile pouch of 30. Peel back the paper label as indicated on the pouch of the reservoirs. Firmly place a reservoir into each hole in the QUINTEST® covered tray. Fill the appropriate reservoir with approximately 12 drops (about 0.5mL) using the dropper from the Hollister-Stier 5mL scratch test extract vial.

Once the tray is prepared, remove a QUINTEST® device from the blister pack by peeling off the label. Pick up the device by the handle and place the tips in the reservoirs. With the QUINTEST® device on the tray, the reservoirs are covered to prevent foreign matter from entering the extract. To assure that test extracts are applied to the proper test sites, it may be helpful to always place the test device handle with the QUINTEST® name facing outward from the center of the tray.

READING SKIN TEST REACTIONS
Measure wheal and erythema responses for histamine positive control at 10 minutes and allergens at 15 minutes. To improve accuracy, precision and uniformity of diagnostic testing and eliminate confusion as to the quantitation of the allergic response, we recommend recording wheal and erythema measurements. Reactions from the QUINTEST® device may be graded as follows:

STORAGE AND DISPOSAL
Once you have completed a patient’s test session, place new QUINTEST® devices in the antigen tray, cover and store at 2°-8°C until the next test session. QUINTEST® devices are designed to be used once and discarded (in accordance with all local, state and federal regulations). The breakaway tips may be removed before discarding, and the handle may be disposed of via non-biohazard routes. These devices cannot be cleaned sufficiently to prevent cross contamination from repeated use or to protect subsequent patients from possible transfer of serum hepatitis or other bloodborne pathogens. The reservoirs in the QUINTEST® tray should be replaced periodically. For best results, discard and replace the reservoirs as they are emptied and refill with fresh extract. If a used QUINTEST® device is inadvertently placed in a reservoir(s), the individual reservoir(s) should be discarded and a new reservoir(s) prepared prior to the next test session.