Soft-ECC®-S

SFT-2100

Endocervical Curettage Device Designed for the Narrow, Short, or Shallow Cervical Os

056-0010 rev A

Physician and Clinician Primer Ver: Jan 2014



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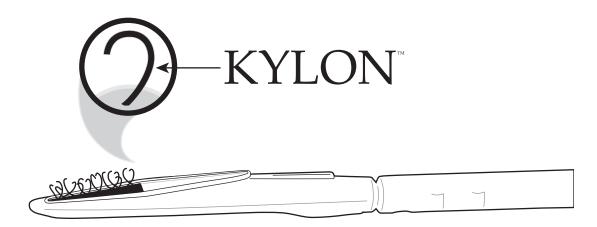
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Q 1: What is Kylon® and how does it work to obtain a biopsy?

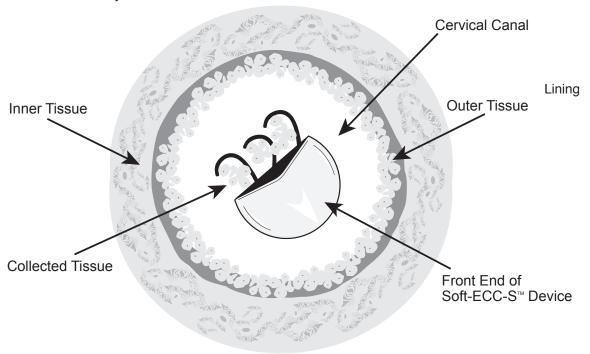
Kylon® is a hooked nylon fabric that can be mounted to the applicator stick, that when pressed on tissue surfaces, allows the hook tip to contact the target tissues, and with pressure and rotation, a trans-epithelial biopsy is obtained (Figure 1). The procedure of applying, pressing, and rotating a shorter pad Kylon® covered tapered tip device into the endocervical os and rotating it to obtain a curettage specimen is called a **Soft-ECC®-S** (Figure 2). The procedure of directing a **Kylon®** tipped applicator directly on an epithelial lesion is called a "**SoftBiopsy®**" and is described in another primer.

Question 1 - Figure 1



Question 1 - Figure 2

The Kylon[®] gently collects tissue samples from the outer layer of the cervical canal.



Q 2: Why were biopsy tools using this patent pending "hooked" curettage fabric called "Kylon®" developed?

Biopsy of mucosal tissues inside a body cavity, such as the cervix or vagina can be traumatic, physically (with evidence of pain and bleeding) and psychologically (fear of cancer, fear of pain or trauma). If trauma is "perceived" by the clinician or the patient, either or both might be less inclined to perform necessary biopsies or seek care to receive them (health seeking behavior may decrease). Neither is good for the quality of the care rendered or received. A minimally invasive biopsy tool was developed with the intent to take a **high quality trans-epithelial tissue biopsy while simultaneously collecting (and storing for transport) most of the specimen inside the fabric**. The device is intended to be minimally invasive, keeping the patient care experience in mind. The goal is also to reduce or eliminate the sampling error associated with biopsy procedures by <u>harvesting and retaining</u> the sample optimally, assuring the fabric tip container is returned to the lab for emptying, processing, and analysis. Thus, both the clinician and the patient are inclined to deliver and receive the necessary care to make the diagnosis and guide therapy.

Clinical trials have been performed, and are ongoing to demonstrate the intended goals of using the Kylon fabric **system** to obtain a biopsy, store the specimen, transport it within the fabric in the preservative vial safely and completely to the laboratory.

Q 3: What is Soft-ECC®-S?

Soft-ECC-S® is a tapered, nylon hooked fabric tipped acrylic tissue sampling system with two componenets; a smaller pad triangular pad biopsy device and a tissue containment detachable tip for storage and transport to the lab. It is intended to be gently inserted into any size endocervical canal and used to obtain an endocervical biopsy / curettage when indicated. The fabric tip is pressed against the endocervical canal, and then rotated 3 complete revolutions clockwise, then 3 complete revolutions counterclockwise, and removed. The tip should be inspected to be sure the specimen fills the fabric base container, then snapped apart from the handle at the fracture line, and placed in a pathological fixative vial. The vial with the sample inside the fabric container tip is processed at the Pathology lab using the SOP to empty the specimen, then the specimen is processed in similar fashion to the conventional endocervical curettage specimen.



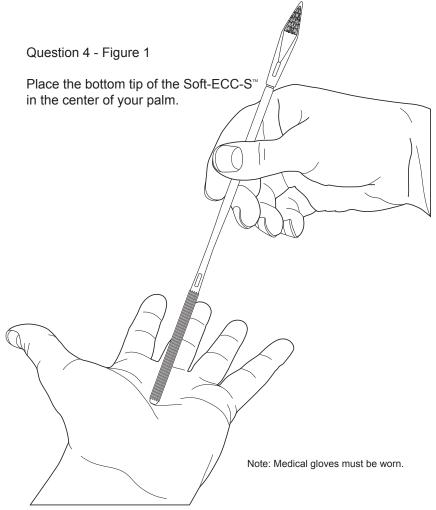
Q 4: How do I use the Soft-ECC®-S device to obtain an endocervical curettage?

Place the handle of the device in the palm and hold the near portion of the shaft in between the thumb and first two fingers (Fig 1). This will permit a twirling motion during the rotational biopsy. (Fig 2) Gently insert the tapered tip into the endocervical canal and press, wiggle, or slightly agitate it so there is a tight fit, or the KYLON pad has completely entered the endocervical canal (Fig 3 and Fig 4). Make note of the placement of the KYLON pad and gently press the pad laterally against the canal surface (Fig 5). Twirl or rotate the device two to three complete 360 degree turns clockwise, then 3 complete revolutions counterclockwise (Fig 6).

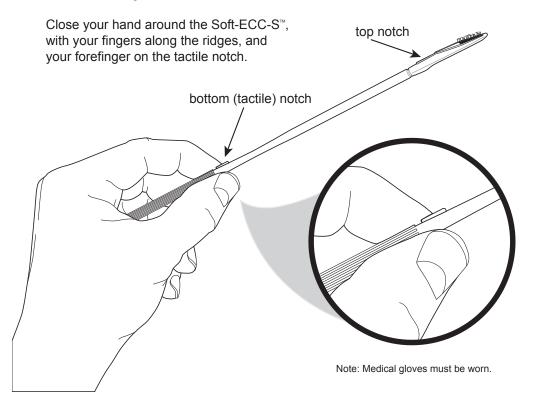
If the fabric pad on the tip does at least enter partially, a sample can still be obtained, but note that the sample is being removed from the outer endocervical canal. The smaller triangular shaped pad in the Soft-ECC®-S should recess completely in shallow cervical canals without exposure of hooks to the outer portio of the cervix.

Alternatively, on could rotationally agitate (like a washing machine effect) 5-8 times while rotating the surface to contact all four quadrants inside the canal.

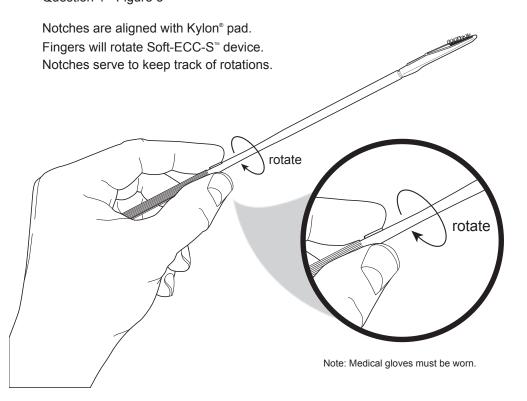
Once an adequate tissue specimen is obtained, simply snap off the device head (Figure 7) from the shaft of the device at the scored area. The tip now contains the specimen completely and can be placed it in the fixative vial provided by the laboratory (Figure 8).



Question 4 - Figure 2

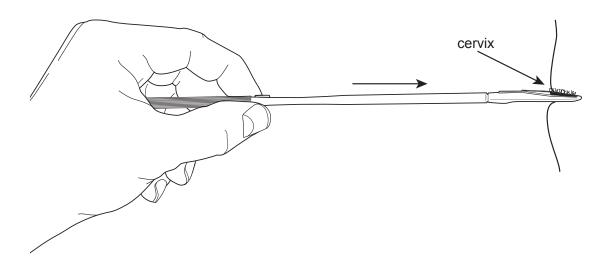


Question 4 - Figure 3



Question 4 - Figure 4

Gently but firmly insert head of device into cervix.

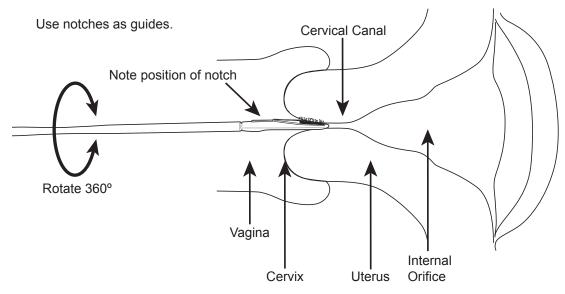


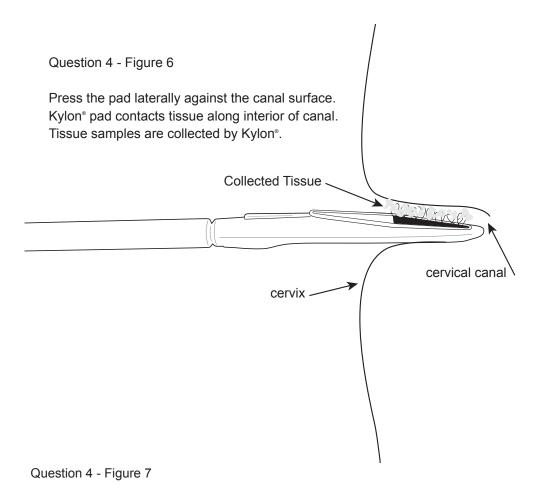
Question 4 - Figure 5

Note: Medical gloves must be worn.

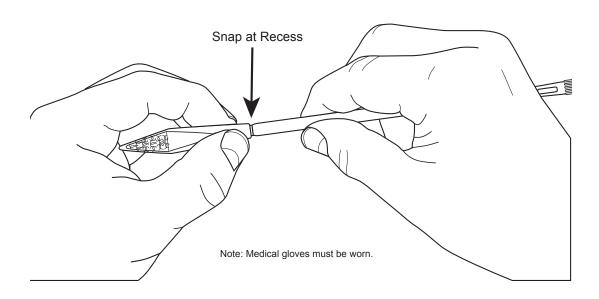
Kylon[®] pad on Soft-ECC-S[™] head is fully positioned inside cervix.

- 1. Rotate clockwise 360° three complete turns.
- 2. Rotate counterclockwise three complete turns.



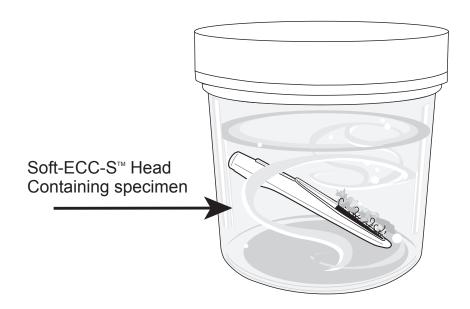


After gently removing Soft-ECC-S™ from cervix, separate head (with Kylon® pad and tissue samples) from body of device by snapping at recessed joint.



Question 4 - Figure 8

Place the head of the Soft-ECC-S™ (with Kylon® pad and tissue samples) into vial of fixative liquid and seal tightly.

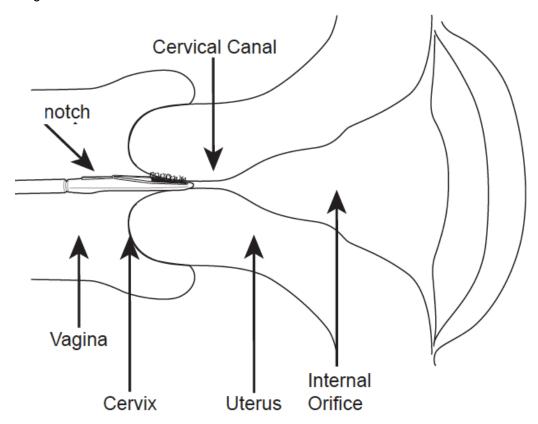


Physician /Clinician Primer : Soft-ECC®-S

Q 5: Why is the tip of the Soft-ECC® curette so pointed and how much of the curette should enter the canal to be considered adequate?

The device is designed to enter almost any endocervical canal. The tip is sufficiently rounded as not to puncture tissue surfaces, but the clinician should be <u>careful</u> to enter the canal in alignment with the direction of the endocervical canal, and <u>not force</u> the tool into the canal. If applied with gentle pressure, a good portion of the **Kylon®** pad will enter into the canal and contact the endocervical epithelial surface. In the majority of cases, at least one third of the **Kylon®** pad should enter the canal to obtain an adequate sample for analysis. However, the likelihood the tapered tip will successfully enter the canal is higher than other rigid conventional endocervical biopsy tools such as the Kevorkian Curette.

Question 5 - Figure 1



DO NOT FORCE the Soft-ECC-S ® curette into the os and avoid advancing it past the internal os.

Q 6: Do the hooks puncture the tissue when pressed? How does the abrasive fabric compare with other biopsy devices?

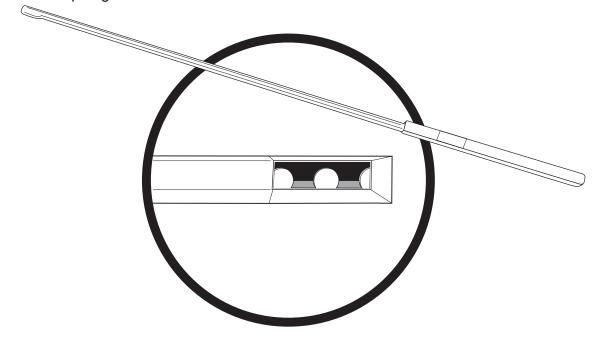
No, the tips of the hooks angle away from the tissue surface before pressure is applied. Once pressed, the hooks open to allow the nylon fibers to contact the tissue surface to perform the curettage. With agitational or rotational forces, the hook tips then frictionally abrade the tissue surface, causing it to buckle and fracture. The hooks then recess under the basement membrane lifting the pieces of trans-epithelial tissue into the fabric. The device edge also abrades the tissue as a curette. The rows hooks function as the basket container to catch and retain tissue fragments. Reearch has shown this cotainment and storage innovation to be responsible for an abundant and highly correlative sample to the underlying pathology.

Other curettes that use a cutting surface, like a forceps, a curette, or a stiff bristle brush are more rigid and are not curved, thus, with pressure, they will puncture or enter the surface with initial placement or pressure. With rotation or lateral motion, the tissue is first sheared, then torn free from the underlying stroma.

With a conventional curettage sharp or bristle brush instrument, the tissue commonly falls free from a curette or straight bristle, and may be incompletely collected in the specimen. If the cutting surface is sharp, the tissue will be shaved or dissected in intact fragments. Often, the clinician has to use a soft nylon cytobrush or forceps to retrieve an endocervical curettage specimen specimen, taking additional time. Not so for the **Soft-ECC®-S** device that simultaneously collects the specimen inside the hooks of the fabric. Unlike the conventional curette, the tip fabric container is snapped free and sent to the lab in a second preservative vial container for tissue removal.

Question 6 - Figure 1

This rectangular metal Kevorkian curette tip scrapes the canal with a sharp edge.



Q 7: How do I apply the device and how much pressure do I apply when pressing the applicator stick and Kylon® pad on to the tissue surface?

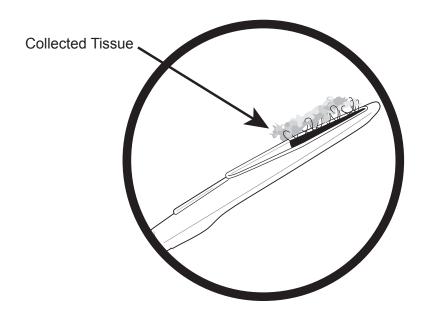
Pressure on a flat, irregular, or rounded tissue surface (like exocervix or vagina) is applied perpendicularly, while pressure inside a canal (radially mounted **Kylon®** pad like the **Soft-ECC®-S** endocervical canal biopsy device) is applied laterally against the canal epithelium. The amount of pressure needed to expose the hook surface to the epithelium would be equivalent of a light massage, depressing the device into the tissue surface so it is firmly applied, but not so hard as to depress the tissue surface more than one or two millimeters.

Q 8: How do I know I have obtained a sufficient sample? What do I do to save the sample for analysis?

After performing the biopsy procedure, gently remove the tip from the endocervix. The device is now non sterile, containing the specimen within the fabric base container. You will note the "excavated" tissue has recessed between the fiber hooks, and will be combined with some blood tinged mucous. The tip is designed to be a dual purpose container, released (snapped off) from the handle with moderate lateral traction, decoupled from the device, and dropped in a vial of fixative. If you do not see any tissue or mucous in the fabric container, repeat the procedure before separating the tip from the device handle.

Question 8 - Figure 1

After performing the biopsy procedure, gently remove the tip from the endocervix. You will note the "excavated" tissue has recessed between the fiber hooks, and will be combined with some blood tinged mucous.

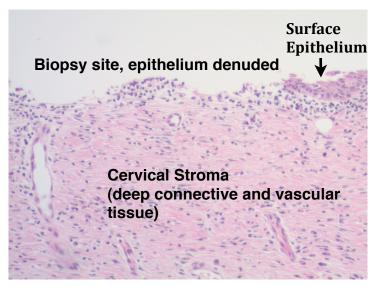


Q-9: Is there Evidence that the biopsy taken with Soft-ECC® or SoftBiopsy® is trans-epithelial (full thickness)?

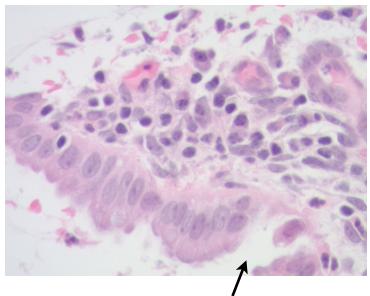
Kylon Based Trans-epithelial Biopsy Sample

Research has shown that the biopsy sample removed with frictional abrasion using the KYLON® hooked fabric is deep enough to remove the full thickness of the epithelium of the cervix into the upper stroma. This includes **both** the glandular endocervical and squamous exocervical tissue and stromal tissue. Both columnar/glandular and squamous epithelium exist together, often near the canal, at the "transformation zone" of the cervix.

Transformation Zone



Endocervical Biopsy

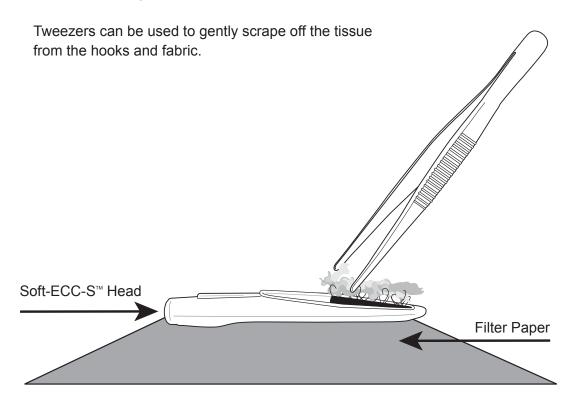


Biopsy: Glands and Stromal Tissue (Intact Architecture)

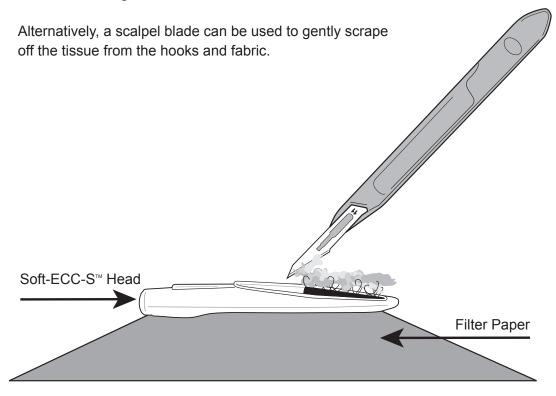
Q 10: How is the specimen taken off the fabric so it can be processed in the Pathology Lab?

A: The pathologist has several methods available to remove the tissue from the fabric tip container. Tweezers or a scalpel blade can be used to gently scrape off the tissue from the hooks and fabric. Alternatively, a small comb has been developed that can be raked across the hooks repeatedly under the surface of the fixative while still in the vial, thus "combing" the tissue from the fabric into the vial. Once the tissue is released, it resembles conventional curettings one would obtain during endocervical curettage. The specimen can then be processed by the lab in an identical manner that is customary for the lab, including cell block if desired.

Question 10 - Figure 1



Question 10 - Figure 2



Question 10 - Figure 3

Another viable technique is to comb the tissue out of the fabric container. Combing is done under the liquid fixative.

