

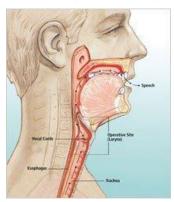
To my healthcare providers: I had a total laryngectomy

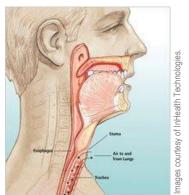
Without a voice prosthesis

My name:
My date of birth:
My personal health number (PHN):
My ear, nose, and throat surgeon:

My airway

I had surgery to remove my larynx. My trachea now leads to a permanent opening (stoma) in my neck.





Before surgery

After surgery

This means my mouth and nose no longer connect to my lungs.

I am a total neck breather.

If I need extra oxygen, ventilating, or intubating, it **must** go through the stoma in my neck. It **must not** go through my mouth and nose.



Tracheostomy tubes

My stoma is permanent. I do not need a tracheostomy tube to keep my airway open after surgery.

Communicating

I speak differently from the way I spoke before surgery because my vocal cords were removed with my larynx.

How I usually communicate:

- ☐ Mouthing words
- ☐ Writing
- ☐ Esophageal speech
- ☐ Text-to-speech
- ☐ Electrolarynx (see image below)

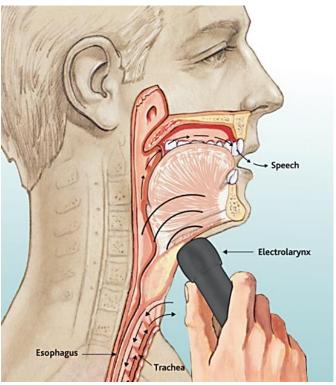


Image courtesy of InHealth Technologies

I had a total laryngectomy without a voice prosthesis - continued

Caring for my stoma and voice prosthesis

To protect my stoma, I usually wear:	☐ LaryTube and heat moisture exchanger
	☐ LaryButton and heat moisture exchanger
	☐ Adhesive baseplate and heat moisture exchanger
	☐ Foam or cloth stoma cover
	☐ Nothing — I leave my stoma uncovered.
To care for my stoma, I usually:	☐ Manage it myself.
	☐ Need some help.
	☐ Need someone to do it for me.

To learn more, see Clinical Skills on Fraser Health Pulse. Find *Total Laryngectomy: Stoma Care and Voice Prosthesis Cleaning.*

For more information

Speak with the hospital Speech-Language Pathologist (SLP). They can help the healthcare team understand my care needs, such as how to use heat moisture exchangers and care for my stoma.

Note: I see the SLPs at Surrey Memorial Hospital.

They have the additional training needed to help patients who had a total laryngectomy.

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