

Windsor Group Newsletter

Turner Syndrome Society of Canada

January, 2018

Here's to 2018!

One of our members got to meet some Calgary members over the holidays!

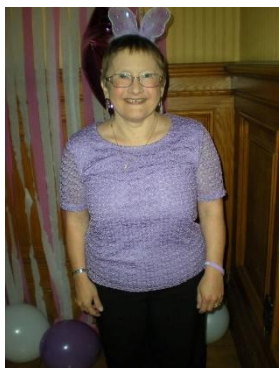
Other news on the Windsor front is three of our members helping out with revamping Strategic Policy Manual and Policy and Procedures.

As well we are very grateful to have an information table at St. Clair College Thursday, February 8 during the College Wellness Week.

We are hoping to arrange for local get together March, June and December.

Sjann Johnson, Rhiannon Cates, Vicki Ming, Dalton William





STORIES

ENDING

My name is Sjann. I'm 56 years old and was diagnosed with Turner Syndrome roughly in 1970. My final height is 4'6". I have been married 30 years and we chose not to adopt. I was on growth hormone, Anavar, before the Premarin, progesterone, ortho novum hormones began.

I do have to say that my friends, family, teachers and employers did not cause me to feel different about myself due to TS. I would say media and advertising were the biggest culprit there. I see what a girl, a woman, an intimate relationship "should" be and to this

day, somewhat believe the lies of Hollywood and advertisers.

I contacted TSSC about 20 years ago on reading in a popular Canadian woman's magazine that fertility specialists were advocating aborting babies with TS. That infuriated me. I dipped in again when I was dealing with more personal issues, and while reading found out that indeed there is information

on us "Golden Butterflies"!

My hope is to build a chapter in Windsor and connect with more "Golden Butterflies".



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