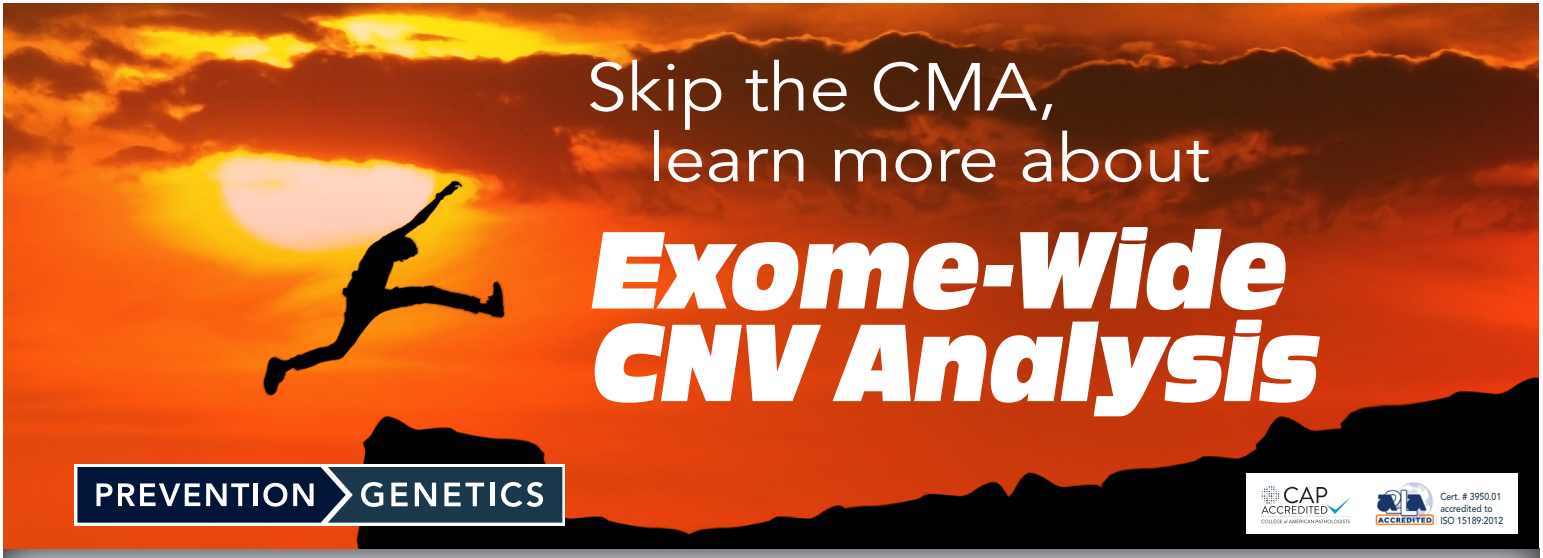

Exome-wide CNV analysis

now available at PreventionGenetics

We are pleased to announce an expanded option for PGxome-based defined and custom panels: Exome-Wide CNV Analysis. PreventionGenetics' Exome-Wide CNV Analysis uses exome sequencing data to identify chromosomal imbalances similar to those detected by CMA.

- Genes from the selected panel receive the same CNV analysis as previously offered (CNVs of four or more exons are detected with sensitivity greater than 95%).
- Exome-Wide CNV Analysis method will identify aneuploidy, triploidy, unbalanced rearrangements, and known microdeletion and microduplication syndromes, as well as unique CNV events.
- Exome-wide CNV targets deletions larger than 250 kb and duplications larger than 500 kb.
- The cost for Exome-Wide CNV Analysis is $250, which is added to the price of the panel.

To confirm if this option is available, visit the panel-specific test descriptions and look for the exome-wide CNV badge:

Exome-Wide CNV
AVAILABLE FOR THIS PANEL

PUT US TO THE TEST **To learn more, visit PreventionGenetics.com**

3800 South Business Park Avenue • Marshfield, Wisconsin 54449 • Phone: (715) 387-0484 • **www.PreventionGenetics.com**