Genetics inMedicine

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CORRECTION

Correction to: Three years of clinical experience with a genome-wide cfDNA screening test for aneuploidies and copy number variants

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Unfortunately, an error occured in Fig. 2. The corrected Fig. 2 is given below.

The original article has been corrected.

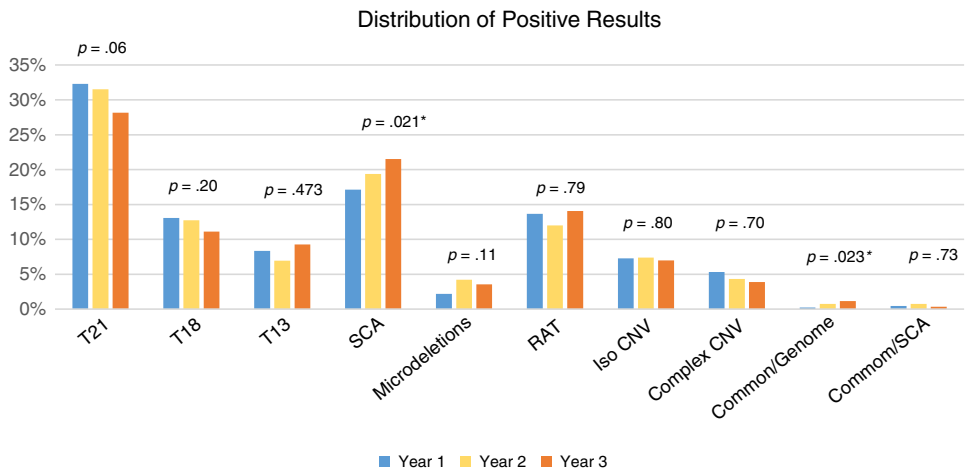

Fig. 2 Graphic depicting the distribution of positives by year. Rare autosomal trisomies (RAT) also include two cases that were monosomies of autosomes. Microdeletions refer to the select list of microdeletions <7 Mb as described in "Materials and Methods". Common/Genome refers to cases positive for a common trisomy and a genome-wide event, while Common/SCA refers to cases positive for a common trisomy and a sex chromosome aneuploidy. Categories with an asterisk (*) show a significant trend, although given the small sample size of the Common/Genome category, significance should be interpreted with caution. Corresponding Z-scores can be found in Table S3. CNV copy-number variant.

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