CORRIGENDUM: Too much, too soon?: Commercial provision of noninvasive prenatal screening for subchromosomal abnormalities and beyond

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In the published version of this article, there was an error in row one, column four of Table 1. There was a typographical error in the final microdeletion, 11q (Jacobsen syndrome). The corrected table appears below. The authors regret the error.

Table 1 Commercial offering of noninvasive prenatal screening for subchromosomal abnormalities

Company	Test name	Trisomies	Microdeletions	Testing option
Seqeunom	MaterniT21Plus[a,]	21, 18, 13, 16, 22	22q11.2 (DiGeorge or velocardiofacial syndrome), 1p36 deletion, 5p (Cri-du-chat syndrome), 15q11.2 (Angelman and Prader-Willi syndrome) 4p (Wolf-Hirschhorn syndrome), 8q (Langier-Gideon syndrome), 119 (Jacobsen syndrome)	Opt out
Natera	Panorama[b]	21, 18, 13	22q11.2 (Di George or velocardiofacial syndrome), 1p36 deletion, 5p (Cri-du chat syndrome), 15q11.2 (Angelman and Prader-Willi syndrome)	Opt in
BGI	NIFTY Plus[c,d,]*	21, 18, 13	5p-, 1p36, and 2q33.1 deletions	N/A
Igenomix[e]	Nace Plus[f,]*	18, 13, 21, 9, 16	22q11.2 (DiGeorge syndrome), 1p36, 15q11.2 (Angelman, Prader-Willi syndromes) 5p (Cri-du-chat syndrome), and 4p (Wolf-Hirschhorn syndrome)	N/A
Illumina	Verifi[g]	18,13, 21, 9,16	22q11.2 (DiGeorge syndrome), 1p36, 15q11.2 (Angelman, Prader-Willi syndromes) 5p (Cri-du-chat syndrome), and 4p (Wolf-Hirschhorn syndrome)	Opt in

*In these cases the tests are differentiated by name to specifically order the microdeletion testing content.

N/A, not applicable.

[a]http://laboratories.sequenom.com/maternit21plus/prenatal-test-information-for-providers. [b]http://www.panoramatest.com/en/healthcare-provider/#about. [c]http://www.niftytest.com/wp-content/uploads/2014/09/BGIDX_NIFTY_Leaflet_24.06.2014_New_Code.pdf. [d]http://www.bgilearning.com/down/NIFTY_PPT.pptx. [e]Previously Iviomics. [f]http://www.igenomix.com/wp-content/uploads/NACE-gynecologist-brochure-ENG.pdf. [g]Unpublished personal communication. See also http://progenity.com/sites/default/files/resources/GeneticCarrierPrenatal%20Req_082014-FINAL.pdf#view=Fit.

ERRATUM: A genome sequencing program for novel undiagnosed diseases

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In the published version of this article, the PhD degree after Dr. Torkamani's name was repeated twice. The publisher regrets the error.