

Two weeks ago, I picked up a complimentary copy of Volume 1, Number 1 *Genetics in Medicine* at the American Society of Human Genetics Annual Meeting in Denver. Did the genetic world need another journal, I wondered. Upon my return to Boston, the answer became quickly apparent.

When the medical-pediatric resident doing a genetics elective with our team bemoaned her limited perception of genetics services, we referred her to the similar experience in the Pacific Northwest.¹ To prepare for the clinic patient, an infant with isolated lissencephaly, the fellows and I reviewed the paper on *LISI* mutation rates.² The NICU consult billed as hypotonia and dysmorphic features was an African-American whose karyotype showed the deletion of Prader-Willi syndrome. Needless to say, the brief report about phenotypic differences and unusual features was timely.³ In the future, medical students on genetics elective will be handed a copy of the clinical objectives proposed by the task force.⁴ During a teaching session with the current students, I reminded them about the recommendations for folic acid and referred them to the most up to date published statement.⁵

At the risk of being a cheerleader, I heartily praise the first issue of *Genetics in Medicine* for its immediate applicability to clinical genetic practice. Not bad for a beginner.

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