In the article: D'Alessandro LC, Mital S. Pediatric transplantation: opportunities for pharmacogenomics and genomics. *Personalized Medicine* 10(4), 397–404 (2013) (www.futuremedicine.com/doi/full/10.2217/PME.13.26), on page 400 the following sentence incorrectly appeared as:

Adult studies have identified two SNPs in the *SLCO1B1* gene to be significantly associated with statin myopathy: a noncoding SNP (**-11187G>A** rs4363657) in intron 11 and a nonsynonymous SNP (521T>C Val174Ala rs4149056) in exon 6, with highest risk (18%) in those with the 521CC genotype [39].

The correct sentence is shown below:

Adult studies have identified two SNPs in the *SLCO1B1* gene to be significantly associated with statin myopathy: a noncoding SNP (**1498-1331T>C** rs4363657) in intron 11 and a nonsynonymous SNP (521T>C Val174Ala rs4149056) in exon 6, with highest risk (18%) in those with the 521CC genotype [39].

The authors would like to sincerely apologize for any confusion or inconvenience caused.