Corrigendum

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.