ERRATUM

Erratum to: Relationship between enzyme properties and disease progression in Canavan disease

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In the published original article, Table 1, the header "Specific activity" is not given correct. "(U/mg)" should be placed above the left column, "% of native" should be placed above the right column.

The correct table is given here.

The online version of the original article can be found at http://dx.doi.org/ 10.1007/s10545-012-9520-z.

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Table 1 Biochemical properties of aspartoacylase clinical mutants

| Aspartoacylase mutant | Disease phenotype | Specific activity a | | Thermal stability (°C) | Conformational |
|---------------------------|--|---------------------|-------------|------------------------|----------------|
| | | (U/mg) | % of native | | stability (mM) |
| Native | _ | 7.5 | 100 % | 59.9 | 1,150 |
| Mild disease phenotype | | | | | |
| G274R | mild with K213E ^b | 2.6 | 35 % | 58.3 | 650 |
| P181T | mild with E285A b | 2.4 | 32 % | 58.3 | 600 |
| Y231C | mild | 1.8 | 24 % | 49.6 | 1,050 |
| P257R | mild with A305E ^b | 1.6 | 21 % | 50.6 | 1,000 |
| I143T | mild | 1.2 | 16 % | 51.9 | 1,000 |
| K213E | mild with G274R ^b | 1.16 | 15 % | 58 | 750 |
| R71H | mild | 0.83 | 11 % | 57.2 | 650 |
| Y288C | mild | 0.29 | 4 % | 56.7 | 250 |
| | mild with F295S & A305E b | | | | |
| Variable/undetermined dis | sease phenotype | | | | |
| D249V | variable ^c | 2.2 | 29 % | 58.6 | 720 |
| N121I | undetermined ^d | 1.2 | 16 % | 47.1 | 500 |
| A305E | variable ^c mild with P257R & Y288C ^b | 0.78 | 10 % | 58.9 | 900 |
| F295S | variable ^c mild with Y288C ^b | 0.75 | 10 % | 56.6 | 1,050 |
| E285A | variable ^c | 0.02 | 0.3 % | 48.5 | 125 |
| Severe disease phenotype | | | | | |
| I143F | severe | 0.08 | 1 % | 51.2 | 600 |
| C152W | severe | 0.07 | 1 % | 58.5 | 1,100 |
| C152R | severe | 0.04 | 0.5 % | 51.2 | 300 |

 $[^]a$ standard errors on the specific activity measurements are $\pm 5~\%$



^b compound heterozygote patients

^c variable severity and disease progression reported in different patients

^d reported from a single patient of undetermined phenotype