



Erratum

Erratum to “Interference free and simplified liquid chromatography-based determination of thiopurine S-methyltransferase activity in erythrocytes”
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The Publisher regrets that there was an error in [Fig. 3](#) and legend to [Fig. 2](#).
Please find the amendments on the following pages.
The Publisher apologises for the error.

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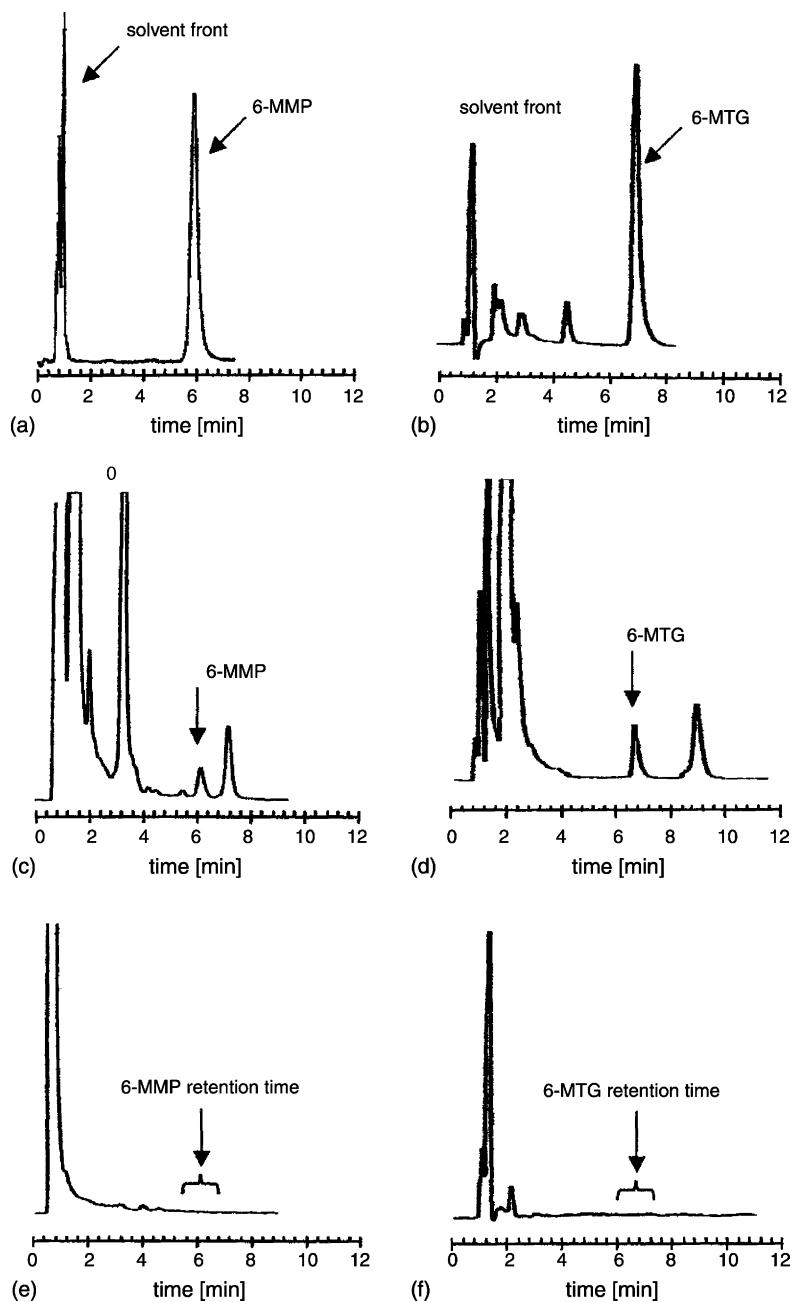


Fig. 2. Chromatographic separation of 6-MMP (a, c) and 6-MTG (b, d). (c and d): perchloric extract of two different representative erythrocyte lysate samples, incubated with 6-MP (c) or 6-TG (d) and SAM for 1 h at 37 °C; substrate- and co-substrate are eluted within 2 min and cannot be determined under these conditions; (e, f): corresponding blank; perchloric extract of a blank lysate (without substrate-/co-substrate) after incubation for 1 h at 37 °C. Chromatographic conditions: Nucleosil 120-3, C18, 70 mm × 3 mm with guard cartridge. The injection volume was 25 μ l and the flow rate adjusted to 0.5 ml/min. The mobile phase consisted of 0.1 M phosphate buffer pH 2.7 acetonitrile (separation of 6-MMP) and 3.5% (separation of 6-MTG), respectively. 6-MMP was measured at wavelength of 290 nm and 6-MTG at 315, respectively. (a and b): Calibration standards of 6-MMP (a: 3.0 μ M) and 6-MTG (b: 5.5 μ M) in assay buffer supplemented by HClO₄ (0.5 μ M). The retention time was 6.0 min, the total run time 8.0 min for 6-MMP, and 7.3 and 11.0 min for 6-MTG, respectively.

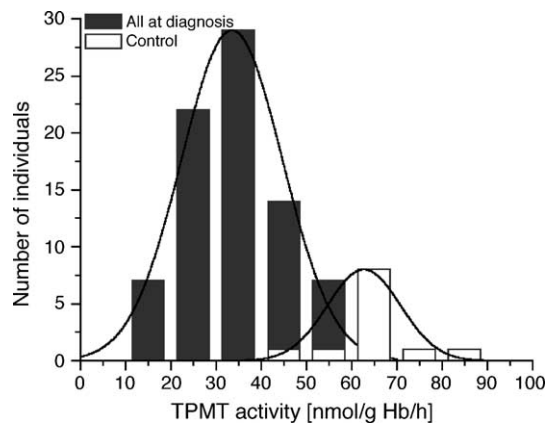


Fig. 3. TPMT activity in erythrocytes. Distribution of TPMT activity in 74 individuals, 64 children with ALL at diagnosis before onset of therapy and 12 healthy adult volunteers.