CORRECTION

ENZYME REPLACEMENT AND ENHANCEMENT THERAPIES: LESSONS FROM LYSOSOMAL DISORDERS

Robert J. Desnick and Edward H. Schuchman

Nature Reviews Genetics **3**, 954–966 (2002).

The entry for Niemann–Pick Type C disease in Table 1 was incorrect. It should have read as follows:

Disease/ subtypes	Deficient enzyme	Inheritance	Residual activity	CNS involvement	Primary site of pathology	Principal manifestations	Animal models
Type C	NPC1 (lipid permease), NPC2/HE1 (lysosomal cholesterol binding protein)	AR	–	+	N, RES	Progressive ND, HSM, gait disturbance and ocular abnormality	M

M, naturally occurring mouse.

The entries for Gaucher type 1 and MPSII in Table 2 were incorrect. They should have read as follows:

Disease	Human enzyme replaced/source	Trial	Sponsor	No. of patients	Monthly dose (mg kg^{-1})	Status	References
Gaucher type 1	Acid β-glucosidase/CHO	Phase 1/2, OL	GEN	12	3.2	Approved (1991)	23–25
MPS II	α-*L*-Iduronidate sulphatase/HF	Phase 1/2, R, DB, PC	TKT	12	0.3, 1.0, 3.0	ND	44

In addition, the second sentence in the section 'Enzyme replacement therapy: clinical trials' contained an error and should have read: Preclinical studies in knockout mice or other naturally occurring animal models (such as the dog model of mucopolysaccharidosis I) are under way for several other disorders, including type B Niemann–Pick disease[42], galactosialidosis[86], Wolman disease (cholesteryl ester storage disease)[59] and mucopolysaccharidosis VII[87,88].

CORRECTION

PREIMPLANTATION GENETIC DIAGNOSIS

Peter Braude, Susan Pickering, Frances Flinter and Caroline Mackie Ogilvie

Nature Reviews Genetics **3**, 941–953 (2002).

In this review, reference 91 was cited incorrectly. The correct reference 91 should have been:

Wilton, L., Williamson, R., McBain, J., Edgar, D. & Voullaire, L. Birth of a healthy infant after preimplantation confirmation of euploidy by comparative genomic hybridization. *N. Engl. J. Med.* **345**, 1537–1541 (2001).