

CORRIGENDUM

## Protein synthesis levels are increased in a subset of individuals with fragile X syndrome

Sébastien Jacquemont<sup>1,2,†</sup>, Laura Pacini<sup>3,†</sup>, Aia E. Jønch<sup>4,5,‡</sup>, Giulia Cencelli<sup>3,‡</sup>, Izabela Rozenberg<sup>6</sup>, Yunsheng He<sup>7</sup>, Laura D'Andrea<sup>3</sup>, Giorgia Pedini<sup>3</sup>, Marwa Eldeeb<sup>8</sup>, Rob Willemsen, Fabrizio Gasparini<sup>10</sup>, Flora Tassone<sup>11</sup>, Randi Hagerman<sup>12</sup>, Baltazar Gomez-Mancilla<sup>6,13</sup>, Claudia Bagni<sup>3,14,\*</sup>

<sup>1</sup>Sainte-Justine University Hospital Research Centre, Montreal, QC H3T 1C5, <sup>2</sup>University of Montreal, Montreal, QC H3T 1J4, Canada, <sup>3</sup>Department of Biomedicine and Prevention, University of Rome Tor Vergata, 00133 Rome, Italy, <sup>4</sup>Department of Clinical Genetics, Odense University Hospital, <sup>5</sup>Human Genetics, Department of Clinical Research, University of Southern Denmark, 5000 Odense, Denmark, <sup>6</sup>Neuroscience Translational Medicine, Novartis Institutes for Biomedical Research, Novartis Pharma AG, 4056 Basel, Switzerland, <sup>7</sup>Biomarker Development, Novartis Institutes for Biomedical Research, Cambridge, MA 02139, USA, <sup>8</sup>Medical Investigation of Neurodevelopmental Disorders (MIND) Institute, University of California, Davis Medical Center, Sacramento, CA 95817, USA, <sup>9</sup>Department of Clinical Genetics, Erasmus Medical Center, 1738, 3000 DR Rotterdam, The Netherlands, <sup>10</sup>Neuroscience Discovery, Novartis Institutes for BioMedical Research, 4002 Basel, Switzerland, <sup>11</sup>Department of Biochemistry and Molecular Medicine and Medical Investigation of Neurodevelopmental Disorders (MIND) Institute, <sup>12</sup>Department of Pediatric and Medical Investigation of Neurodevelopmental Disorders (MIND) Institute, University of California Davis, School of Medicine, Sacramento, CA 95817, USA, <sup>13</sup>Department of Neurology and Neurosurgery, McGill University, Montreal, QC H3A 0G4, Canada and <sup>14</sup>Department of Fundamental Neuroscience, University of Lausanne, 1005 Lausanne, Switzerland

\*To whom correspondence should be addressed at: Department of Biomedicine and Prevention, University of Rome, Tor Vergata, Via Montpellier 1, 00133 Rome, Italy. Tel: +390672596063; Fax: +390672596053; Email: claudia.bagni@uniroma2.it; Department of Fundamental Neuroscience, University of Lausanne, Rue du Bugnon 9, 1005 Lausanne, Switzerland. Tel: +41216925120; Email: claudia.bagni@unil.ch

*Human Molecular Genetics*, 2018, 27(12), 2039–2051.  
doi: 10.1093/hmg/ddy099.

This article initially published with incomplete supplementary material. This error has now been corrected, and the correct supplementary material is published.

The authors regret the error.