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Does size matter? : bridging and dose selection in paediatric trials

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Some say that we are what we eat. I'd rather say we are who we meet.

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Curriculum vitæ

Massimo Cella was born on the 13th of November 1978 in Lodi, Italy. He attended the Liceo Scientifico (High School) “G.Gandini” in Lodi, where he obtained his scientific diploma in 1997. Subsequently he started his undergraduated training in Pharmaceutical Chemistry and Technology at the Università di Pavia, where he obtained his MSc in January 2004, followed by a 2nd level University Master degree in Pre-formulation and Drug Development in 2005 and another 2nd level University Master degree in Regulatory Affairs in 2007. In June 2004 he started a research fellowship at the Consorzio per le Valutazioni Biologiche e Farmacologiche (CVBF) in Pavia, under the supervision of Prof. Adriana Ceci. Here he worked on designations and registration dossiers for orphan and paediatric drugs. At the same time he was actively involved in the research activities of the Task Force in Europe for Drug Development for the Young (TEDDY), a Network of Excellence funded under the Sixth EU Framework Programme for Research and Technological Development.

In November 2005 Massimo started his PhD research programme at the division of Pharmacology of the Leiden-Amsterdam Center for Drug Research in Leiden, with Dr. Oscar Della Pasqua as co-promoter and Prof. Meindert Danhof as promoter, which led to this thesis. From September 2009 to May 2010 he was invited to GlaxoSmithKline in Greenford, UK, to work on PK modelling of drug combinations in children.

Since May 2010 Massimo Cella is a post-doc fellow at the Clinical Pharmacology - Modelling and Simulation group of GlaxoSmithKline, Stockley Park, UK, where he is part of the Innovative Medicine Initiative (IMI) funded project *PharmaCog*. Among other tasks, he has taken responsibility for the development of translational and disease progression models in Alzheimer's Disease.

List of publications

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