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Kaja Kantorska studied Biotechnology in the years 2002-2007 at the International Faculty of Engineering. After graduation she completed an internship in the European Commission, where she works until today. For the first 7 years she was dealing with genetically modified organisms. Since 2016 she has been responsible for the development of the pharmaceutical legislation and policies concerning authorisation and manufacturing and use of medicinal products in the Union, in particular with respect to orphan medicinal products.