

Regulation of DNA damage and immune response pathways by posttranslational protein modification

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Curriculum vitae

Madelon Dijk was born on June 3, 1988 in Rotterdam, The Netherlands. In 2006 she successfully completed grammar school at Johan de Witt Gymnasium in Dordrecht and started the study of Life Science and Technology at Leiden University (UL) and Delft University of Technology (TU Delft). After studying the behavior of S. cerevisiae at near-zero growth rates during an internship at the department of Industrial Microbiology at TU Delft, she obtained her bachelor's degree cum laude (with honors) in 2009. During her master's education, she investigated catalytic intermediates of cytochrome bd-I oxidases at the former department of Enzymology at TU Delft. Furthermore, she intended to optimize the immobilization of lipases for industrial purposes during an internship at ChiralVision in Leiden. In 2011, the master's degree was obtained cum laude. Subsequently, she joined the former department of Toxicogenetics at Leiden University Medical Center (LUMC) as a PhD student to investigate the regulation of the nucleotide excision DNA repair pathway under the supervision of Prof. dr. Leon H. Mullenders. In addition, she worked with Dr. Navraj S. Pannu at the department of Biophysical Structural Chemistry at UL on a project on C1-inhibitor. From 2014 on, the research on protein regulation during DNA damage repair was continued at the department of Human Genetics at the LUMC under the supervision of Prof. dr. Haico van Attikum. After finishing the practical part of the PhD project, she started working in the field of data analysis at the National Office for Identity Data (Rijksdienst voor Identiteitsgegevens) in Den Haag.

Publications

2018	TRiC controls transcription resumption after UV damage by regulating Cockayne syndrome protein A
	Alex Pines*, Madelon Dijk*, Matthew Makowski, Elisabeth M. Meulenbroek, Mischa G. Vrouwe, Yana van der Weegen, Marijke Baltissen, Pim J. French, Martin E. van Royen, Martijn S. Luijsterburg, Leon H. Mullenders, Michiel Vermeulen, Wim Vermeulen ⁺ , Navraj S. Pannu ⁺ and Haico van Attikum ⁺ *co-first authors, *co-corresponding authors <i>Nature Communications; 9(1): 1040</i>
2016	How dextran sulfate affects C1-inhibitor activity:
	a model for polysaccharide potentiation
	Madelon Dijk, Jolande Holkers, Patrick Voskamp, Bruno M. Giannetti, Willem-Jan Waterreus, Harrie A. van Veen and Navraj S. Pannu
	Structure; 24(12): 2182-2189
2014	Insight in the multilevel regulation of NER
	Madelon Dijk, Dimitris Typas, Leon H. Mullenders and Alex Pines
	Experimental Cell Research; 329(1): 116-123
2012	Oxoferryl-porphyrin radical catalytic intermediate in cytochrome bd
	oxidases protects cells from formation of reactive oxygen species
	Angela Paulus, Sebastiaan G.H. Rossius, Madelon Dijk and Simon de Vries Journal of Biological Chemistry; 287(12): 8830-8838
2011	Extreme calorie restriction and energy source starvation in Saccharomyces
	cerevisiae represents distinct physiological states
	Léonie G.M. Boender, Marinka J.H. Almering, Madelon Dijk, Antonius J.A. van Maris, Johannes H. de Winde, Jack T. Pronk and Pascale Daran-Lapujade
	Biochimica et Biophysica Acta - Molecular Cell Research; 1813(12): 2133-2144
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Acknowledgements

In this thesis I have addressed the importance of regulatory proteins. Without them, pathways would never be properly activated, correctly executed and timely completed. In fact, they may stall halfway or totally derail. I found that the same holds true for a PhD track. Although to the PhD student it may sometimes feel like a solitary expedition, there are some indispensable cofactors that safeguard both the trajectory and the student. On the road to this thesis, I experienced the significance of such factors, embodied by colleagues, friends and family.

Leon, thank you for initiating the pathway and always believing in its successful completion. I appreciate both your scientific enthusiasm and your interest in my personal well-being. Raj, coactivator, I also value your guidance along the way.

As illustrated by many aspects of the DNA damage response, alternative pathways are crucial. Haico, thank you for welcoming me in your lab, providing such routes and being willing to coordinate them as my promotor. Silvère, the adjustment to the friendly, inspiring working environment of your department was easy.

One should also not underestimate the significance of crosstalk with other proteins, even when pathways separate. Alex, I enjoyed collaborating with you and I think we did great at complementing and strengthening each other's work. Romy, Juliette, Leonie and Jenny, thanks for sharing frustrations and breakthroughs while being challenged by comparable internal and external hazards. A warm thanks also goes to my two paranymphs. Hanneke, I was lucky to have you as a constant and close factor in my ever-changing environment. Chantal, although you joined later on, it feels like you have always been there for scientific and emotional support.

Finally, I can't stress enough the importance of the chaperones that are fundamental to a protein's stability. Papa, thank you for being the proud, concerned, encouraging dad I can always depend on. Mama, your love and support will always be remembered. Lisanne, Justin, I feel so fortunate to form this stable, inseparable complex with the two of you. Chris, thanks for knowing me so well, recognizing exactly when I need protection, distraction, motivation or, sometimes, inhibition. I couldn't have done this without you.

While this thesis only describes a fine selection of regulatory mechanisms, it is the combination of all modulating factors that directs successful protein and pathway functioning. Likewise, I owe a big thanks to all colleagues, lab mates, students, staff members, roommates, friends and family that either crossed my path occasionally or supported me for longer. Together, you contributed significantly to this accomplishment.