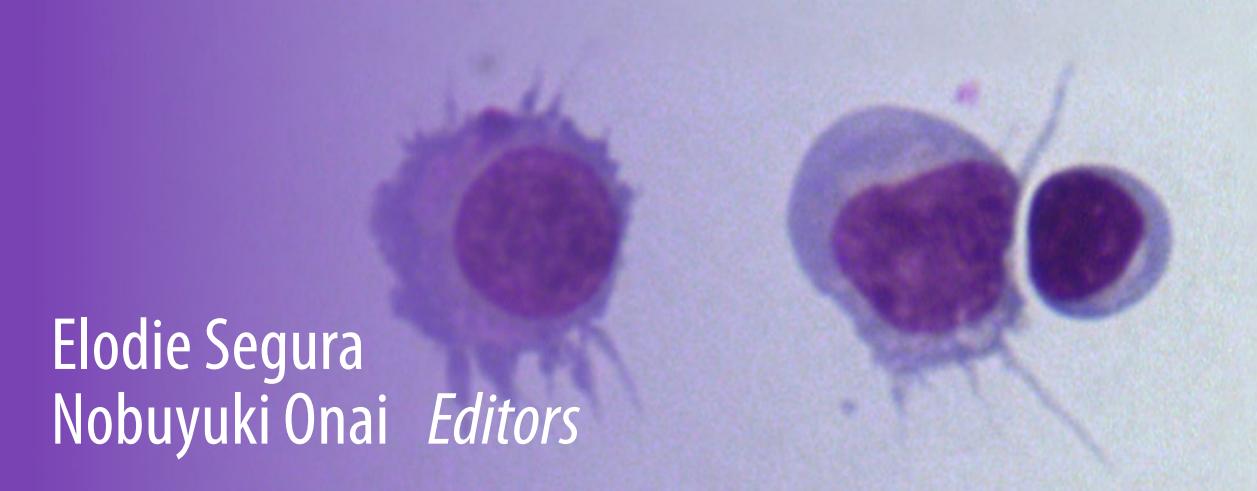


Methods in  
Molecular Biology 1423

Springer Protocols



A light micrograph showing three dendritic cells stained with purple. They have large, dark purple nuclei and prominent, thin, hair-like processes called dendrites extending from their peripheries.

Elodie Segura  
Nobuyuki Onai *Editors*

# Dendritic Cell Protocols

*Third Edition*

 Humana Press

# METHODS IN MOLECULAR BIOLOGY

*Series Editor*

John M. Walker

School of Life and Medical Sciences

University of Hertfordshire

Hatfield, Hertfordshire, AL10 9AB, UK

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# **Dendritic Cell Protocols**

**Third Edition**

Edited by

**Elodie Segura**

*Pavillon Pasteur, Institut Curie INSERM U932, Paris, France*

**Nobuyuki Onai**

*Department of Biodefense, Tokyo Medical and Dental University, Tokyo, Japan*

 **Humana Press**

*Editors*

Elodie Segura  
Pavillon Pasteur  
Institut Curie INSERM U932  
Paris, France

Nobuyuki Onai  
Department of Biodefense  
Tokyo Medical and Dental University  
Tokyo, Japan

ISSN 1064-3745                    ISSN 1940-6029 (electronic)  
Methods in Molecular Biology  
ISBN 978-1-4939-3604-5            ISBN 978-1-4939-3606-9 (eBook)  
DOI 10.1007/978-1-4939-3606-9

Library of Congress Control Number: 2016939414

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Printed on acid-free paper

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## Preface

Dendritic cells are fascinating cells but can be challenging to work with. In this new edition of *Dendritic Cell Protocols*, we aimed at complementing the previous edition in order to provide both beginners and more experienced researchers a choice of methods to isolate and analyze dendritic cells. An introductory review provides an overview of recent advances in the characterization of DC subsets in mouse and human.

Dendritic cells are rare, fragile, and their isolation is often a tedious procedure. For those who wish to generate dendritic cells *in vitro*, two chapters provide methods to culture human and mouse dendritic cells. Several chapters are devoted to protocols for the isolation of dendritic cells from various organs and tissues (lymphoid organs, intestine, skin, lung, liver), both in mouse and human. In addition, a chapter describes the isolation of dendritic cell progenitors from mouse, and another chapter the purification of dendritic cells from human blood.

Dendritic cells are often described as conductors of the immune response and, as such, perform a wide range of functions. We have compiled cutting-edge methods for the analysis of dendritic cell properties *ex vivo*. Some of these methods can be applied to dendritic cells from any species. We also included a “bioinformatics tutorial” chapter for the analysis of dendritic cell transcriptome by nonexperts.

In vivo mouse studies have significantly contributed to our knowledge of dendritic cells homeostasis, interactions with other immune cells, and division of labor between dendritic cell subsets. The last section contains several protocols for the *in vivo* analysis of dendritic cells through cell ablation, adoptive transfer, infection models, or *in vivo* imaging. Finally, we provide a protocol for the generation of humanized mice for analyzing human dendritic cells in a physiological setting.

The editors wish to thank all authors for their excellent contributions. We hope that this Protocols book will be a valuable tool for a better understanding of dendritic cell biology.

*Paris, France*  
*Tokyo, Japan*

*Elodie Segura*  
*Nobuyuki Onai*



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## Contributors

- HANS ACHA-ORBEA • *Department of Biochemistry CIIL, University of Lausanne, Epalinges, Switzerland*
- SOLANA ALCULUMBRE • *INSERM U932 Immunity and Cancer, Paris, France; Institut Curie, Centre de recherche, Paris, France*
- SEBASTIAN AMIGORENA • *INSERM U932, Institut Curie, Centre de recherche, Paris, France*
- KEIICHI ARIMURA • *Division of Immunology, Department of Infectious Diseases, Faculty of Medicine, University of Miyazaki, Miyazaki, Japan*
- DEVIKA ASHOK • *Department of Biochemistry CIIL, University of Lausanne, Epalinges, Switzerland*
- TEGEST AYCHEK • *Department of Immunology, The Weizmann Institute of Science, Rehovot, Israel*
- SREEKUMAR BALAN • *Centre d'Immunologie de Marseille-Luminy (CIML), Marseille, France; INSERM U1104, Marseille, France; CNRS UMR7280, Marseille, France*
- BIANA BERNSTEIN • *Department of Immunology, The Weizmann Institute of Science, Rehovot, Israel*
- CATERINA CURATO • *Department of Immunology, The Weizmann Institute of Science, Rehovot, Israel*
- MARC DALOD • *Centre d'Immunologie de Marseille-Luminy (CIML) UNIV UM2, Aix Marseille Université, Marseille, France; INSERM U1104, Marseille, France; CNRS UMR7280, Marseille, France*
- CLAIRE DUMONT • *Department of Biochemistry and Molecular Biology, Bio21 Molecular Science and Biotechnology Institute, The University of Melbourne, Parkville, VIC, Australia*
- MÉLANIE DURAND • *INSERM U932, Paris, France; Institut Curie, Centre de recherche, Paris, France*
- GYOHEI EGAWA • *Department of Dermatology, Kyoto University Graduate School of Medicine, Kyoto, Japan*
- JANA M. ELLEGAST • *Division of Hematology, University Hospital Zurich, Zurich, Switzerland*
- TOMOHIRO FUKAYA • *Division of Immunology, Department of Infectious Diseases, Faculty of Medicine, University of Miyazaki, Miyazaki, Japan*
- FLORENT GINHOUX • *Singapore Immunology Network (SIgN), Agency for Science, Technology and Research (A\*STAR), Singapore, Singapore*
- MARTIN GUILLIAMS • *Laboratory of Immunoregulation, VIB Inflammation Research Center, Ghent University, Ghent, Belgium; Department of Biomedical Molecular Biology, Ghent University, Ghent, Belgium*
- MERRY GUNAWAN • *Human DC Lab, Institute of Cellular Medicine, Newcastle University, Newcastle upon Tyne, UK*
- MUZLIFAH HANIFFA • *Human DC Lab, Institute of Cellular Medicine, Newcastle University, Newcastle upon Tyne, UK*

HIROAKI HEMMI • *Laboratory for Immune Regulation, WPI Immunology Frontier Research Center, Osaka University, Osaka, Japan; Laboratory of Inflammatory Regulation, RIKEN Center for Integrative Medical Sciences, Kanagawa, Japan; Department of Immunology, Institute of Advanced Medicine Wakayama Medical University, Wakayama, Japan*

SANDRINE HENRI • *Centre d'Immunologie de Marseille-Luminy (CIML), UM2 Aix-Marseille Université, Marseille Cedex 9, France; INSERM U1104, Marseille Cedex 9, France; CNRS UMR7280, Marseille Cedex 9, France*

KATSUAKI HOSHINO • *Laboratory for Immune Regulation, WPI Immunology Frontier Research Center, Osaka University, Osaka, Japan; Laboratory of Inflammatory Regulation, RIKEN Center for Integrative Medical Sciences, Kanagawa, Japan; Department of Immunology, Faculty of Medicine, Kagawa University, Kagawa, Japan*

BAPTISTE JANELA • *Singapore Immunology Network (SIgN), Agency for Science, Technology and Research (A\*STAR), Singapore, Singapore*

LAURA JARDINE • *Human DC Lab, Institute of Cellular Medicine, Newcastle University, Newcastle upon Tyne, UK*

ANGUS P.R. JOHNSTON • *Drug Delivery, Disposition and Dynamics, Monash Institute of Pharmaceutical Sciences, Monash University, Parkville, VIC, Australia; ARC Centre of Excellence in Convergent Bio-Nano Science and Technology, Monash University, Parkville, VIC, Australia*

STEFFEN JUNG • *Department of Immunology, The Weizmann Institute of Science, Rehovot, Israel*

KENJI KABASHIMA • *Department of Dermatology, Kyoto University Graduate School of Medicine, Kyoto, Japan; Precursory Research for Embryonic Science and Technology, Japan science and Technology Agency, Saitama, Japan; Singapore Immunology Network (SIgN) and Institute of Medical Biology, Agency for Science, Technology and Research (A\*STAR), Biopolis, Singapore*

TSUNEYASU KAISHO • *Laboratory for Immune Regulation, WPI Immunology Frontier Research Center, Osaka University, Osaka, Japan; Laboratory of Inflammatory Regulation, RIKEN Center for Integrative Medical Sciences, Kanagawa, Japan; Department of Immunology, Institute of Advanced Medicine, Wakayama Medical University, Wakayama, Japan*

HAIYIN LIU • *Department of Biochemistry and Molecular Biology, Bio21 Molecular Science and Biotechnology Institute, The University of Melbourne, Parkville, VIC, Australia; Drug Delivery, Disposition and Dynamics, Monash Institute of Pharmaceutical Sciences, Monash University, Parkville, VIC, Australia*

JOAO G. MAGALHAES • *INSERM U932, Institut Curie, Centre de recherche, Paris, France*

CAMILLE MALOSSE • *Centre d'Immunologie de Marseille-Luminy (CIML), UM2 Aix-Marseille Université, Marseille Cedex 9, France; INSERM U1104, Marseille Cedex 9, France; CNRS UMR7280, Marseille Cedex 9, France*

MARKUS G. MANZ • *Division of Hematology, University Hospital Zurich, Zurich, Switzerland*

NAOMI McGOVERN • *Singapore Immunology Network (SIgN), Agency for Science, Technology and Research (A\*STAR), Singapore, Singapore*

SIMON W.F. MILLING • *Centre for Immunobiology, Institute of Infection, Immunity and Inflammation, College of Veterinary, Medical and Life Sciences, University of Glasgow, Scotland, UK*

JUSTINE D. MINTERN • *Department of Biochemistry and Molecular Biology, Bio21 Molecular Science and Biotechnology Institute, The University of Melbourne, Parkville, VIC, Australia*

- ALLAN McI MOWAT • *Centre for Immunobiology, Institute of Infection, Immunity and Inflammation, College of Veterinary, Medical and Life Sciences, University of Glasgow, Scotland, UK*
- TOSHIAKI OHTEKI • *Department of Biodefense, Medical Research Institute, Tokyo Medical and Dental University, Tokyo, Japan*
- NOBUYUKI ONAI • *Department of Biodefense, Medical Research Institute, Tokyo Medical and Dental University, Tokyo, Japan*
- LUCIA PATTARINI • *INSERM U932 Immunity and Cancer, Paris, France; Institut Curie, Centre de recherche, Paris, France*
- MATTEO PIGNI • *Department of Biochemistry CIIL, University of Lausanne, Epalinges, Switzerland*
- CHRISTIANE RUEDL • *School of Biological Sciences, Nanyang Technological University, Singapore, Singapore*
- YASUYUKI SAITO • *Division of Hematology, University Hospital Zurich, Zurich, Switzerland; Division of Molecular and Cellular Signaling, Department of Biochemistry and Molecular Biology, Kobe University Graduate School of Medicine, Kobe, Japan*
- KATSUAKI SATO • *Division of Immunology, Department of Infectious Diseases, Faculty of Medicine, University of Miyazaki, Miyazaki, Japan*
- ANDREAS SCHLITZER • *Singapore Immunology Network (SIgN), Agency for Science, Technology and Research (A\*STAR), Singapore, Singapore*
- CHARLOTTE L. SCOTT • *Centre for Immunobiology, Institute of Infection, Immunity and Inflammation, College of Veterinary, Medical and Life Sciences, University of Glasgow, Scotland, UK; Laboratory of Immunoregulation, Inflammation Research Centre (IRC), VIB Ghent University, Ghent, Belgium; Department of Respiratory Medicine, Ghent University Hospital, Ghent, Belgium*
- ELODIE SEGURA • *INSERM U932, Paris, France; Institut Curie, Centre de recherche, Paris, France*
- HIDEAKI TAKAGI • *Division of Immunology, Department of Infectious Diseases, Faculty of Medicine, University of Miyazaki, Miyazaki, Japan*
- SIMON TAVERNIER • *Laboratory of Immunoregulation, VIB Inflammation Research Center, Ghent University, Ghent, Belgium; Department of Respiratory Medicine, Ghent University, Ghent, Belgium*
- PIOTR TETLAK • *School of Biological Sciences, Nanyang Technological University, Singapore, Singapore*
- TOMOFUMI UTO • *Division of Immunology, Department of Infectious Diseases, Faculty of Medicine, University of Miyazaki, Miyazaki, Japan*
- LIANNE VAN DE LAAR • *Laboratory of Immunoregulation, VIB Inflammation Research Center, Ghent University, Ghent, Belgium; Department of Internal Medicine, Ghent University, Ghent, Belgium*
- OMAR I. VIVAR • *INSERM U932, Institut Curie, Centre de recherche, Paris, France*
- DAVID VREMEC • *The Walter and Eliza Hall Institute of Medical Research, Parkville, VIC, Australia*
- THIEN-PHONG VU MANH • *Centre d'Immunologie de Marseille-Luminy (CIML) UNIV UM2, Aix Marseille Université, Marseille, France; INSERM U1104, Marseille, France; CNRS UMR7280, Marseille, France*
- PAMELA B. WRIGHT • *Centre for Immunobiology, Institute of Infection, Immunity and Inflammation, College of Veterinary, Medical and Life Sciences, University of Glasgow, Scotland, UK*



# **Part I**

## **Introduction**

