On the Cover: Secondary cell wall thickenings from cultured *Zinnia elegans* tracheary elements are rich in lignocellulose and autofluorescent due to the presence of lignin (top image). The topography of the secondary cell wall can be visualized after generating cell fragments, which include rings of these secondary wall thickenings (bottom left; atomic force microscopy image). Closer examination of a region (box) within this ring reveals cellulose fibrils that are embedded in a granular matrix and organized in a parallel, concentric orientation (bottom right; atomic force microscopy image). To investigate the composition and structure of the cell wall of individual tracheary elements, Lacayo et al. (pp. 121–133) employed three imaging platforms in conjunction with chemical extraction and a fluorescent cellulose binding protein. Cultured tracheary elements from *Z. elegans* can provide important insights into cell wall development, organization, and dynamics, which are especially important in research on lignocellulosic biofuel production. Fluorescence microscopy image and cover design by Dr. Catherine Lacayo; atomic force microscopy images by Dr. Alexander Malkin.

ON THE INSIDE

Peter V. Minorsky

GENOME ANALYSIS


The Association of Multiple Interacting Genes with Specific Phenotypes in Rice Using Gene Coexpression Networks. Stephen P. Ficklin, Feng Luo, and F. Alex Feltus

BIOINFORMATICS

iTILLING: A Personalized Approach to the Identification of Induced Mutations in Arabidopsis. Susan M. Bush and Patrick J. Krysan

Combining Machine Learning and Homology-Based Approaches to Accurately Predict Subcellular Localization in Arabidopsis. Rakesh Kaundal, Reena Saini, and Patrick X. Zhao

RESEARCH ARTICLES

BIOCHEMICAL PROCESSES AND MACROMOLECULAR STRUCTURES

Virus-Induced Gene Silencing of Plastidial Soluble Inorganic Pyrophosphatase Impairs Essential Leaf Anabolic Pathways and Reduces Drought Stress Tolerance in *Nicotiana benthamiana*. Gavin M. George, Margaretha J. van der Merve, Adriano Nunes-Nesi, Rolene Bauer, Alisdair R. Fernie, Jens Kossmann, and James R. Lloyd

Enzymatic Functions of Wild Tomato Methylketone Synthases 1 and 2. Geng Yu, Thuong T.H. Nguyen, Yongxia Guo, Ines Schauvinhold, Michele E. Auldridge, Nazmul Bhuiyan, Imri Ben-Israel, Yoko Iijima, Eyal Fridman, Joseph P. Noel, and Eran Pichersky

Continued on next page
A Glucurono(arabino)xylan Synthase Complex from Wheat Contains Members of the GT43, GT47, and GT75 Families and Functions Cooperatively. Wei Zeng, Nan Jiang, Ramya Nadella, Tara L. Killen, Vijayanand Nadella, and Ahmed Faik

Identification and Characterization of Proteins Involved in Rice Urea and Arginine Catabolism. Feng-Qiu Cao, Andrea K. Werner, Kathleen Dahncke, Tina Romeis, Lai-Hua Liu, and Claus-Peter Witte

A Copal-8-ol Diphosphate Synthase from the Angiosperm Cistus creticus subsp. creticus Is a Putative Key Enzyme for the Formation of Pharmacologically Active, Oxygen-Containing Labdane-Type Diterpenes. Vasiliki Falara, Eran Pichersky, and Angelos K. Kanellis

A Genome-Scale Metabolic Model Accurately Predicts Fluxes in Central Carbon Metabolism under Stress Conditions. Thomas C.R. Williams, Mark G. Poolman, Andrew J.M. Howden, Markus Schwarzlander, David A. Fell, R. George Ratcliffe, and Lee J. Sweetlove

CYP93G2 Is a Flavanone 2-Hydroxylase Required for C-Glycosylflavone Biosynthesis in Rice. Yegang Du, Hung Chu, Ivan K. Chu, and Clive Lo

Cooperation of LPA3 and LPA2 Is Essential for Photosystem II Assembly in Arabidopsis. Wenhe Cai, Jinfang Ma, Wei Chi, Meijuan Zou, Jinkui Guo, Congming Lu, and Lixin Zhang

Combined Effects of CO2 and Light on the N2-Fixing Cyanobacterium Trichodesmium IMS101: Physiological Responses. Sven A. Kranz, Orly Levitan, Klaus-Uwe Richter, Ondřej Prašil, Ilana Berman-Frank, and Björn Rost

Homomeric Interaction of AtVSR1 Is Essential for Its Function as a Vacuolar Sorting Receptor. Hyeran Kim, Hyangju Kang, Mihue Jang, Jeong Ho Chang, Yansong Miao, Liwen Jiang, and Inhwan Hwang

OsC6, Encoding a Lipid Transfer Protein, Is Required for Postmeiotic Anther Development In Rice. Dasheng Zhang, Wanqi Liang, Changsong Yin, Jie Zong, Fangwei Gu, and Dabing Zhang


Successful Reproduction Requires the Function of Arabidopsis YELLOW STRIPE-LIKE1 and YELLOW STRIPE-LIKE3 Metal-Nicotianamine Transporters in Both Vegetative and Reproductive Structures. Heng-Hsuan Chu, Jeff Chiecko, Tracy Punshon, Antonio Lanzirotti, Brett Lahner, David E. Salt, and Elisabeth L. Walker

TaCHP: A Wheat Zinc Finger Protein Gene Down-Regulated by Abscisic Acid and Salinity Stress Plays a Positive Role in Stress Tolerance. Cuiling Li, Jian Lv, Xin Zhao, Xinghui Ai, Xinlei Zhu, Mengcheng Wang, Shuangyi Zhao, and Guangmin Xia

Arabidopsis Plants Acclimate to Water Deficit at Low Cost through Changes of Carbon Usage: An Integrated Perspective Using Growth, Metabolite, Enzyme, and Gene Expression Analysis. Irène Hummel, Florent Pantin, Ronan Sulpice, Maria Piques, Gaëlle Rolland, Myriam Daouzat, Angelique Christophe, Marjorie Perent, Marie Boutellé, Mark Stitt, Yves Gibon, and Bertrand Muller

Functional Analysis of the Group 4 Late Embryogenesis Abundant Proteins Reveals Their Relevance in the Adaptive Response during Water Deficit in Arabidopsis. Yadira Olvera-Carrillo, Francisco Campos, José Luis Reyes, Alejandro Garcia-Rubio, and Alejandra A. Covarrubias

RTM3, Which Controls Long-Distance Movement of Potyviruses, Is a Member of a New Plant Gene Family Encoding a Meprin and TRAF Homology Domain-Containing Protein. Patrick Cosson, Luc Sofer, Quang Hien Le, Valérie Léger, Valérie Schurdi-Leeraud, Steven A. Whitham, Miki L. Yamamoto, Suresh Gopalan, Olivier Le Gall, Thierry Candresse, James C. Carrington, and Frédéric Revers

Plant Immunity Directly or Indirectly Restricts the Injection of Type III Effectors by the Pseudomonas syringae Type III Secretion System. Emerson Crabill, Anna Joe, Anna Block, Jennifer M. van Rooyen, and James R. Alfano


Ethylene Signaling Regulates Accumulation of the FLS2 Receptor and Is Required for the Oxidative Burst Contributing to Plant Immunity. Sophia Mersmann, Gildas Bourdais, Steffen Rietz, and Silke Robatzek

Cryptochrome as a Sensor of the Blue/Green Ratio of Natural Radiation in Arabidopsis. Romina Sellaro, María Crepy, Santiago Ariel Trupkin, Elizabeth Karayekov, Ana Sabrina Buchovský, Constanza Rossi, and Jorge José Casal

Overexpression of the Epidermis-Specific Homeodomain-Leucine Zipper IV Transcription Factor OUTER CELL LAYER1 in Maize Identiﬁes Target Genes Involved in Lipid Metabolism and Cuticle Biosynthesis. Marie Javelle, Vanessa Vernoud, Nathalie Depège-Fargeix, Christine Arnould, Delphine Oursel, Frédéric Domergue, Xavier Sarda, and Peter M. Rogowsky

An Atlas of Type I MADS Box Gene Expression during Female Gametophyte and Seed Development in Arabidopsis. Marian Bemer, Klaas Heijmans, Chiara Airoldi, Brendan Davies, and Gerco C. Angenent

Multiple Regulatory Elements in the Arabidopsis NIA1 Promoter Act Synergistically to Form a Nitrate Enhancer. Rongchen Wang, Peizhu Guan, Mingsheng Chen, Xiujuan Xing, Yali Zhang, and Nigel M. Crawford

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