

Keith

The Populus trichocarpa PtHSP17.8 involved in heat and salt stress tolerances

Plant Cell Rep. 2016 Mar 28. [Epub ahead of print]

Li J1, Zhang J2,3, Jia H1,4, Li Y1, Xu X1, Wang L1, Lu M5,6.

1State Key Laboratory of Tree Genetics and Breeding, Key Laboratory of Tree Breeding and Cultivation of the State Forestry Administration, Research Institute of Forestry, Chinese Academy of Forestry, Beijing, 100091, China.

2State Key Laboratory of Tree Genetics and Breeding, Key Laboratory of Tree Breeding and Cultivation of the State Forestry Administration, Research Institute of Forestry, Chinese Academy of Forestry, Beijing, 100091, China.

3Collaborative Innovation Center of Sustainable Forestry in Southern China, Nanjing Forestry University, Nanjing, 210037, China.

4Collaborative Innovation Center of Sustainable Forestry in Southern China, Nanjing Forestry University, Nanjing, 210037, China.

5State Key Laboratory of Tree Genetics and Breeding, Key Laboratory of Tree Breeding and Cultivation of the State Forestry Administration, Research Institute of Forestry, Chinese Academy of Forestry, Beijing, 100091, China.

6Collaborative Innovation Center of Sustainable Forestry in Southern China, Nanjing Forestry University, Nanjing, 210037, China.

PtHSP17.8 was regulated by various abiotic stresses. Overexpression of PtHSP17.8 enhanced the tolerance to heat and salt stresses through maintain ROS homeostasis and cooperate with stress-related genes in Arabidopsis. Small heat shock proteins (sHSPs) play important roles in response to diverse biotic and abiotic stresses, especially in heat tolerance. However, limited information is available on the stress tolerance roles of sHSPs in woody species. To explore the function of sHSPs in poplar, we isolated and characterized PtHSP17.8 from *Populus trichocarpa*. Phylogenetic analysis and subcellular localization revealed that PtHSP17.8 was a cytosolic class I sHSP. The gene expression profile of PtHSP17.8 in various tissues showed that it was significantly accumulated in stem and root, which was consistent with the GUS expression pattern driven by promoter of PtHSP17.8. The expression of PtHSP17.8 could be induced by various abiotic stresses and significantly activated by heat stress. Overexpression of PtHSP17.8 enhanced the tolerance to heat and salt stresses in Arabidopsis. The seedling survival rate, root length, relative water content, antioxidative enzyme activities, proline, and soluble sugar content were increased in transgenic Arabidopsis under heat and salt stresses, but not in normal condition. The co-expression network of PtHSP17.8 were constructed and demonstrated many stress responsive genes included. The stress-related genes in the co-expression network were up-regulated in the PtHSP17.8 overexpression seedlings. These results suggest that PtHSP17.8 confers heat and salt tolerances in plants.

Gene Expression Profile in the Long-Living Lotus: Insights into the Heat Stress Response Mechanism

PLoS One. 2016 Mar 28;11(3):e0152540.

Liu X1, Du F1, Li N1, Chang Y1, Yao D1.

1Institute of Botany, Jiangsu Province and Chinese Academy of Sciences, Nanjing 210014, China.

Lotus (*Nelumbo Adans*) is an aquatic perennial plant that flourished during the middle Albian stage. In this study, we characterized the digital gene expression signatures for China Antique lotus under conditions of heat shock stress. Using RNA-seq technology, we sequenced four libraries, specifically, two biological replicates for control plant samples and two for heat stress samples. As a result, 6,528,866 to 8,771,183 clean reads were mapped to the reference genome, accounting for 92-96% total clean reads. A total of 396 significantly altered genes were detected across the genome, among which 315 were upregulated and 81 were downregulated by heat shock stress. Gene ontology (GO) enrichment of differentially expressed genes revealed protein folding, cell morphogenesis and cellular component morphogenesis as the top three functional terms under heat shock stress. Kyoto Encyclopedia of Genes and Genomes (KEGG) analysis led to the identification of protein processing in endoplasmic reticulum, plant-pathogen interactions, spliceosome, endocytosis, and protein export as significantly enriched pathways. Among the upregulated genes, small heat shock proteins (sHsps) and genes related to cell morphogenesis were particularly abundant under heat stress. Data from the current study provide valuable clues that may help elucidate the molecular events underlying heat stress response in China Antique lotus.

Arabidopsis AtDJA3 Null Mutant Shows Increased Sensitivity to Abscisic Acid, Salt, and Osmotic Stress in Germination and Post-germination Stages.

Front Plant Sci. 2016 Feb 25;7:220.

Salas-Muñoz S1, Rodríguez-Hernández AA1, Ortega-Amaro MA1, Salazar-Badillo FB1, Jiménez-Bremont JF1.

1Laboratorio de Biotecnología Molecular de Plantas, División de Biología Molecular, Instituto Potosino de Investigación Científica y Tecnológica San Luis Potosí, México.

DnaJ proteins are essential co-chaperones involved in abiotic and biotic stress responses. Arabidopsis AtDJA3 gene encodes a molecular co-chaperone of 420

amino acids, which belongs to the J-protein family. In this study, we report the functional characterization of the AtDjA3 gene using the Arabidopsis knockout line designated j3 and the 35S::AtDjA3 overexpression lines. Loss of AtDjA3 function was associated with small seed production. In fact, j3 mutant seeds showed a reduction of 24% in seed weight compared to Col-0 seeds. Expression analysis showed that the AtDjA3 gene was modulated in response to NaCl, glucose, and abscisic acid (ABA). The j3 line had increased sensitivity to NaCl and glucose treatments in the germination and cotyledon development in comparison to parental Col-0. Furthermore, the j3 mutant line exhibited higher ABA sensitivity in comparison to parental Col-0 and 35S::AtDjA3 overexpression lines. In addition, we examined the expression of ABI3 gene, which is a central regulator in ABA signaling, in j3 mutant and 35S::AtDjA3 overexpression lines. Under 5 μ M ABA treatment at 24 h, j3 mutant seedlings displayed higher ABI3 expression, whereas in 35S::AtDjA3 overexpression lines, ABI3 gene expression was repressed. Taken together, these results demonstrate that the AtDjA3 gene is involved in seed development and abiotic stress tolerance.

Transcriptional induction of the heat shock protein B8 mediates the clearance of misfolded proteins responsible for motor neuron diseases.

Sci Rep. 2016 Mar 10;6:22827.

Crippa V1, D'Agostino VG2, Cristofani R3, Rusmini P3, Cicardi ME3, Messi E3, Loffredo R2, Pancher M2, Piccolella M3, Galbiati M3, Meroni M3, Cereda C1, Carra S4, Provenzani A2, Poletti A3,5.

1Laboratory of Experimental Neurobiology, C. Mondino National Neurological Institute, Pavia, Italy.

2Centre for Integrative Biology (CIBIO), Università degli Studi di Trento, Trento, Italy.

3Dipartimento di Scienze Farmacologiche e Biomolecolari (DiSFeB), Centro di Eccellenza sulle Malattie Neurodegenerative, Università degli Studi di Milano, Milano, Italy.

4Dipartimento di Scienze Biomediche, Metaboliche e Neuroscienze, Università di Modena e Reggio Emilia, Modena, Italy.

5Centro InterUniversitario sulle Malattie Neurodegenerative, Università degli Studi di Firenze, Genova, Roma Tor Vergata and Milano, Italy.

Neurodegenerative diseases (NDs) are often associated with the presence of misfolded protein inclusions. The chaperone HSPB8 is upregulated in mice, the human brain and muscle structures affected during NDs progression. HSPB8 exerts a potent pro-degradative activity on several misfolded proteins responsible for familial NDs forms. Here, we demonstrated that HSPB8 also counteracts accumulation of aberrantly localized misfolded forms of TDP-43 and its 25 KDa fragment involved in most sporadic cases of Amyotrophic Lateral Sclerosis (sALS) and of Fronto Lateral Temporal Dementia (FLTD). HSPB8 acts with BAG3 and the

HSP70/HSC70-CHIP complex enhancing the autophagic removal of misfolded proteins. We performed a high-throughput screening (HTS) to find small molecules capable of inducing HSPB8 in neurons for therapeutic purposes. We identified two compounds, colchicine and doxorubicin, that robustly up-regulated HSPB8 expression. Both colchicine and doxorubicin increased the expression of the master regulator of autophagy TFEB, the autophagy linker p62/SQSTM1 and the autophagosome component LC3. In line, both drugs counteracted the accumulation of TDP-43 and TDP-25 misfolded species responsible for motoneuronal death in sALS. Thus, analogs of colchicine and doxorubicin able to induce HSPB8 and with better safety and tolerability may result beneficial in NDs models.

Oligomers of Heat-Shock Proteins: Structures That Don't Imply Function

PLoS Comput Biol. 2016 Feb 29;12(2):e1004756.

Jacobs WM1, Knowles TP1, Frenkel D1.

1Department of Chemistry, University of Cambridge, Cambridge, United Kingdom.

Most proteins must remain soluble in the cytosol in order to perform their biological functions. To protect against undesired protein aggregation, living cells maintain a population of molecular chaperones that ensure the solubility of the proteome. Here we report simulations of a lattice model of interacting proteins to understand how low concentrations of passive molecular chaperones, such as small heat-shock proteins, suppress thermodynamic instabilities in protein solutions. Given fixed concentrations of chaperones and client proteins, the solubility of the proteome can be increased by tuning the chaperone-client binding strength. Surprisingly, we find that the binding strength that optimizes solubility while preventing irreversible chaperone binding also promotes the formation of weakly bound chaperone oligomers, although the presence of these oligomers does not significantly affect the thermodynamic stability of the solution. Such oligomers are commonly observed in experiments on small heat-shock proteins, but their connection to the biological function of these chaperones has remained unclear. Our simulations suggest that this clustering may not have any essential biological function, but rather emerges as a natural side-effect of optimizing the thermodynamic stability of the proteome.

An atomistic view of Hsp70 allosteric crosstalk: from the nucleotide to the substrate binding domain and back.

Sci Rep. 2016 Mar 30;6:23474.

Chiappori F1, Merelli I1, Milanese L1, Colombo G2, Morra G2.

1Istituto di Tecnologie Biomediche - CNR, Segrate (Mi), 20090 Italy.
2Istituto di Chimica del Riconoscimento Molecolare - CNR, Milano, 20131 Italy.

The Hsp70 is an allosterically regulated family of molecular chaperones. They consist of two structural domains, NBD and SBD, connected by a flexible linker. ATP hydrolysis at the NBD modulates substrate recognition at the SBD, while peptide binding at the SBD enhances ATP hydrolysis. In this study we apply Molecular Dynamics (MD) to elucidate the molecular determinants underlying the allosteric communication from the NBD to the SBD and back. We observe that local structural and dynamical modulation can be coupled to large-scale rearrangements, and that different combinations of ligands at NBD and SBD differently affect the SBD domain mobility. Substituting ADP with ATP in the NBD induces specific structural changes involving the linker and the two NBD lobes. Also, a SBD-bound peptide drives the linker docking by increasing the local dynamical coordination of its C-terminal end: a partially docked DnaK structure is achieved by combining ATP in the NBD and peptide in the SBD. We propose that the MD-based analysis of the inter domain dynamics and structure modulation could be used as a tool to computationally predict the allosteric behaviour and functional response of Hsp70 upon introducing mutations or binding small molecules, with potential applications for drug discovery.

Lens regeneration using endogenous stem cells with gain of visual function

Nature **531**, 323–328 (17 March 2016)

Haotian Lin, Hong Ouyang, Jie Zhu, Shan Huang, Zhenzhen Liu, Shuyi Chen, Guiqun Cao, Gen Li, Robert A. J. Signer, Yanxin Xu, Christopher Chung, Ying Zhang, Danni Lin, Sherrina Patel, Frances Wu, Huimin Cai, Jiayi Hou, Cindy Wen, Maryam Jafari, Xialin Liu, Lixia Luo, Jin Zhu, Austin Qiu, Rui Hou, Baoxin Chen *et al.*

State Key Laboratory of Ophthalmology, Zhongshan Ophthalmic Center, Sun Yat-sen University, Guangzhou 510060, China

Shiley Eye Institute, Institute for Engineering in Medicine, Institute for Genomic Medicine, University of California, San Diego, La Jolla, California 92093, USA

Molecular Medicine Research Center, State Key Laboratory of Biotherapy, West China Hospital, Sichuan University, Sichuan 610041, China

Guangzhou KangRui Biological Pharmaceutical Technology Company, Guangzhou 510005, China

Howard Hughes Medical Institute, Children's Research Institute, Department of Pediatrics, University of Texas Southwestern Medical Center, Dallas, Texas 75390, USA

Department of Ophthalmology, West China Hospital, Sichuan University, Sichuan 610041, China

Division of Genetics, Department of Medicine, Brigham and Women's Hospital and Harvard Medical School, Boston, Massachusetts 02115, USA

Clinical and Translational Research Institute, University of California, San Diego, La Jolla, California 92093, USA

Veterans Administration Healthcare System, San Diego, California 92093, USA

The repair and regeneration of tissues using endogenous stem cells represents an ultimate goal in regenerative medicine. To our knowledge, human lens regeneration has not yet been demonstrated. Currently, the only treatment for cataracts, the leading cause of blindness worldwide, is to extract the cataractous lens and implant an artificial intraocular lens. However, this procedure poses notable risks of complications. Here we isolate lens epithelial stem/progenitor cells (LECs) in mammals and show that *Pax6* and *Bmi1* are required for LEC renewal. We design a surgical method of cataract removal that preserves endogenous LECs and achieves functional lens regeneration in rabbits and macaques, as well as in human infants with cataracts. Our method differs conceptually from current practice, as it preserves endogenous LECs and their natural environment maximally, and regenerates lenses with visual function. Our approach demonstrates a novel treatment strategy for cataracts and provides a new paradigm for tissue regeneration using endogenous stem cells.

Damian

Eroglu, E., Gottschalk, B., Charoensin, S., Blass, S., Bischof, H., Rost, R., Madreiter-Sokolowski, C. T., Pelzmann, B., Bernhart, E., Sattler, W., Hallstrom, S., Malinski, T., Waldeck-Weiermair, M., Graier, W. F., and Malli, R. (2016) Development of novel FP-based probes for live-cell imaging of nitric oxide dynamics, *Nat. Commun.* 7.

Trabjerg, E., Jakobsen, R. U., Mysling, S., Christensen, S., Jørgensen, T. J. D., and Rand, K. D. (2015) Conformational Analysis of Large and Highly Disulfide-Stabilized Proteins by Integrating Online Electrochemical Reduction into an Optimized H/D Exchange Mass Spectrometry Workflow, *Anal. Chem.* 87, 8880-8888.

Jarrett

Elimination of HIV-1 Genomes from Human T-lymphoid Cells by CRISPR/Cas9 Gene Editing

Rafal Kaminski, Yilan Chen, Tracy Fischer, Ellen Tedaldi, Alessandro Napoli, Yonggang Zhang, Jonathan Karn, Wenhui Hu & Kamel Khalili
NATURE Scientific Reports 6, Article number: 22555 (2016)

<http://www.nature.com/articles/srep22555>

Boreal and temperate trees show strong acclimation of respiration to warming

Peter B. Reich, Kerrie M. Sendall, Artur Stefanski, Xiaorong Wei, Roy L. Rich & Rebecca A. Montgomery

Nature 531, 633–636 (31 March 2016)

<http://www.nature.com/nature/journal/v531/n7596/pdf/nature17142.pdf>

Structure of promoter-bound TFIID and model of human pre-initiation complex assembly

Robert K. Louder, Yuan He, José Ramón López-Blanco, Jie Fang, Pablo Chacón & Eva Nogales

Nature **531**, 604–609 (31 March 2016)

<http://www.nature.com/nature/journal/v531/n7596/full/nature17394.html>

Minsoo

1. *Mol Cell*. 2016 Mar 17;61(6):914-24. doi: 10.1016/j.molcel.2016.02.030.

Digital Quantification of Proteins and mRNA in Single Mammalian Cells.

Albayrak C(1), Jordi CA(1), Zechner C(1), Lin J(1), Bichsel CA(1), Khammash M(1), Tay S(2).

Author information:

(1)Department of Biosystems Science and Engineering, ETH Zürich, 4058 Basel, Switzerland. (2)Department of Biosystems Science and Engineering, ETH Zürich, 4058 Basel, Switzerland; Institute for Molecular Engineering, University of Chicago, Chicago, IL 60637, USA. Electronic address: tays@uchicago.edu.

Absolute quantification of macromolecules in single cells is critical for understanding and modeling biological systems that feature cellular heterogeneity. Here we show extremely sensitive and absolute quantification of both proteins and mRNA in single mammalian cells by a very practical workflow that combines proximity ligation assay (PLA) and digital PCR. This digital PLA method has femtomolar sensitivity, which enables the quantification of very small protein concentration changes over its entire 3-log dynamic range, a quality necessary for accounting for single-cell heterogeneity. We counted both endogenous (CD147) and exogenously expressed (GFP-p65) proteins from hundreds of single cells and determined the correlation between CD147 mRNA and the protein it encodes. Using our data, a stochastic two-state model of the central dogma was constructed and verified using joint mRNA/protein distributions, allowing us to estimate transcription burst sizes and extrinsic noise strength and calculate the transcription and translation rate constants in single mammalian cells.

2. *Nat Methods*. 2016 Mar 28. doi: 10.1038/nmeth.3810. [Epub ahead of print]

Robust transcriptome-wide discovery of RNA-binding protein binding sites with enhanced CLIP (eCLIP).

Van Nostrand EL(1),(2),(3), Pratt GA(1),(2),(3),(4), Shishkin AA(5), Gelboin-Burkhart C(1),(2),(3), Fang MY(1),(2),(3), Sundararaman B(1),(2),(3), Blue SM(1),(2),(3), Nguyen TB(1),(2),(3), Surka C(5), Elkins K(1),(2),(3), Stanton R(1),(2),(3), Rigo F(6), Guttman M(5), Yeo GW(1),(2),(3),(4),(7),(8).

Author information:

(1)Department of Cellular and Molecular Medicine, University of California at San Diego, La Jolla, California, USA. (2)Stem Cell Program, University of California at San Diego, La Jolla, California, USA. (3)Institute for Genomic Medicine, University of California at San Diego, La Jolla, California, USA. (4)Bioinformatics and Systems Biology Graduate Program, University of California at San Diego, La Jolla, California, USA. (5)Division of Biology and Biological Engineering, California Institute of Technology, Pasadena, California, USA. (6)Ionis Pharmaceuticals, Carlsbad, California, USA. (7)Department of Physiology, Yong Loo Lin School of Medicine, National University of Singapore, Singapore. (8)Molecular Engineering Laboratory, A*STAR, Singapore.

As RNA-binding proteins (RBPs) play essential roles in cellular physiology by interacting with target RNA molecules, binding site identification by UV crosslinking and immunoprecipitation (CLIP) of ribonucleoprotein complexes is critical to understanding RBP function. However, current CLIP protocols are technically demanding and yield low-complexity libraries with high experimental failure rates. We have developed an enhanced CLIP (eCLIP) protocol that decreases requisite amplification by ~1,000-fold, decreasing discarded PCR duplicate reads by ~60% while maintaining single-nucleotide binding resolution. By simplifying the generation of paired IgG and size-matched input controls, eCLIP improves specificity in the discovery of authentic binding sites. We generated 102 eCLIP experiments for 73 diverse RBPs in HepG2 and K562 cells (available at <https://www.encodeproject.org>), demonstrating that eCLIP enables large-scale and robust profiling, with amplification and sample requirements similar to those of ChIP-seq. eCLIP enables integrative analysis of diverse RBPs to reveal factor-specific profiles, common artifacts for CLIP and RNA-centric perspectives on RBP activity.

March 28 2016

A proposal regarding best practices for validating the identity of genetic stocks and the effects of genetic variants


Detlef Weigel, Joy Bergelson, Edward S. Buckler, Joseph R. Ecker, and Magnus Nordborg

Plant Cell 2016 tpc.15.00502; Advance Publication March 8, 2016;

doi:10.1105/tpc.15.00502 **OPEN**

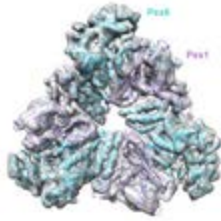
<http://www.plantcell.org/content/early/2016/03/08/tpc.15.00502>

[FEBS Journal Content Alert: 283, 6 \(March 2016\)](#)

 [Structures of the double-ring AAA ATPase Pex1–Pex6 involved in peroxisome biogenesis \(pages 986–992\)](#)

Dongyan Tan, Neil B. Blok, Tom A. Rapoport and Thomas Walz

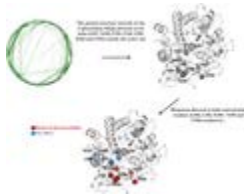
Article first published online: 12 NOV 2015 | DOI: 10.1111/febs.13569



The Pex1/Pex6 complex is a member of the AAA family of ATPases that is involved in peroxisome biogenesis. Recently, cryo-electron microscopy structures have been determined of the Pex1/Pex6 complex in different nucleotide states. This Structural Snapshot describes the structural features of the complex, their implications for its function, as well as questions that still await answers.

[Protein thermal denaturation is modulated by central residues in the protein structure network \(pages 1124–1138\)](#)

Valquiria P. Souza, Cecília M. Ikegami, Guilherme M. Arantes and Sandro R. Marana
 Article first published online: 2 FEB 2016 | DOI: 10.1111/febs.13659



Mutations directed to seven hubs of the protein structure network of the β -glucosidase S β gly reduced its thermostability. Moreover, mutations directed to the vicinity of a hub residue also caused significant decreases in the S β gly thermostability. On the contrary mutations directed to non-hub residues had no effect. These results show that protein structure networks are robust, whereas attacks on central nodes cause network failure.

The Plant Journal Content Alert (New Articles)

[Bacterial Microcompartments as Metabolic Modules for Plant Synthetic Biology](#)

C. Raul Gonzalez-Esquer, Sarah E. Newnham and Cheryl A. Kerfeld

Accepted manuscript online: 16 MAR 2016 08:51AM EST | DOI: 10.1111/tpj.13166

[Transcriptome dynamics of Arabidopsis during sequential biotic and abiotic stresses](#)

Silvia Coolen, Silvia Proietti, Richard Hickman, Nelson H. Davila Olivas, Ping-Ping Huang, Marcel C. Van Verk, Johan A. Van Pelt, Alexander H.J. Wittenberg, Martin De Vos, Marcel Prins, Joop J.A. Van Loon, Mark G.M. Aarts, Marcel Dicke, Corné M.J. Pieterse and Saskia C.M. Van Wees

Accepted manuscript online: 15 MAR 2016 10:40PM EST | DOI: 10.1111/tpj.13167

Journal of Agronomy and Crop Sci... Content Alert (New Articles)

[Susceptibility of Faba Bean \(*Vicia faba* L.\) to Heat Stress During Floral Development and Anthesis](#)

J. Bishop, S. G. Potts and H. E. Jones

Article first published online: 21 MAR 2016 | DOI: 10.1111/jac.12172

**Science, March 25 p. [10.1126/science.aad6253](https://doi.org/10.1126/science.aad6253)
Designing and building a minimal genome**

A goal in biology is to understand the molecular and biological function of every gene in a cell. One way to approach this is to build a minimal genome that includes only the genes essential for life. In 2010, a 1079-kb genome based on the genome of *Mycoplasma mycoides* (JCV-syn1.0) was chemically synthesized and supported cell growth when transplanted into cytoplasm. Hutchison III *et al.* used a design, build, and test cycle to reduce this genome to 531 kb (473 genes). The resulting JCV-syn3.0 retains genes involved in key processes such as transcription and translation, but also contains 149 genes of unknown function.

The Plant Journal Content Alert (New Articles)

Large-scale atlas of microarray data reveals the distinct expression landscape of different tissues in Arabidopsis

Fei He, Shinjae Yoo, Daifeng Wang, Sunita Kumari, Mark Gerstein, Doreen Ware and Sergei Maslov

Accepted manuscript online: 25 MAR 2016 09:45AM EST | DOI: 10.1111/tpj.13175

Current Opinion in Chemical Biology: Alert 25 March-31 March

[Dynamics of co-translational protein targeting](#) Review Article

Pages 79-86

Margaret M Elvekrog, Peter Walter

Development of novel FP-based probes for live-cell imaging of nitric oxide dynamics

[Emrah Eroglu](#), [Benjamin Gottschalk](#), [Suphachai Charoensin](#), et al

Nature Communications 7, Article number: 10623 doi:10.1038/ncomms10623

Nitric oxide (NO[•]) is a free radical with a wide range of biological effects, but practically impossible to visualize in single cells. Here we report the development of novel multicoloured fluorescent quenching-based NO[•] probes by fusing a bacteria-derived NO[•]-binding domain close to distinct fluorescent protein variants. These genetically encoded NO[•] probes, referred to as geNOps, provide a selective, specific and real-time read-out of cellular NO[•] dynamics and, hence, open a new era of NO[•] bioimaging. The combination of geNOps with a Ca²⁺ sensor allowed us to visualize NO[•] and Ca²⁺ signals simultaneously in single endothelial cells. Moreover, targeting of the NO[•] probes was used to detect NO[•] signals within mitochondria. The geNOps are useful new tools to further investigate and understand the complex patterns of NO[•] signalling on the single (sub)cellular level.

Starting too soon: upstream reading frames repress downstream translation

Anna M McGeachy and Nicholas T Ingolia

<http://EMBOJ.embopress.org/content/35/7/699?etoc>

The analysis of ribosome profiling data across three vertebrate species demonstrates the global effect of uORF-dependent translational control of gene expression.

Upstream ORFs are prevalent translational repressors in vertebrates

Timothy G Johnstone, Ariel A Bazzini, and Antonio J Giraldez

Published online 19.02.2016

<http://EMBOJ.embopress.org/content/35/7/706?etoc>

Upstream open reading frames (uORFs) are found in many genes, but their functional impact is unclear. This study explores their abundant translation, conservation, and potent role in shaping the vertebrate translational landscape.

The chloroplast NADPH thioredoxin reductase C, NTRC, controls non-photochemical quenching of light energy and photosynthetic electron transport in *Arabidopsis* (pages 804–822)

Belén Naranjo, Clara Mignée, Anja Krieger-Liszkay, Dámaso Hornero-Méndez, Lourdes Gallardo-Guerrero, Francisco Javier Cejudo and Marika Lindahl
Article first published online: 18 JAN 2016 | DOI: 10.1111/pce.12652

Excess excitation and the means by which plants respond to defend themselves against light-induced damage have been extensively studied, and the importance of qE is well documented. Nevertheless, the success of a plant also depends on the capacity to restrict the induction of acclimation mechanisms, such as qE, in order to avoid wasting the energy absorbed. Given the possible redox control and implication of thioredoxins in protection mechanisms against excess light, we have investigated the role of the chloroplast NADPH-dependent thioredoxin reductase NTRC. This enzyme has been previously implied in peroxide detoxification and abiotic stress tolerance. Surprisingly, NTRC proved to be essential, not for the induction of protection mechanisms under strong irradiance, but for the maintenance of optimal photosynthetic efficiency under normal-light conditions and down-regulation of qE through control of the trans-thylakoid pH gradient.



Analysis of the sodium chloride-dependent respiratory kinetics of wheat mitochondria reveals differential effects on phosphorylating and non-phosphorylating electron transport pathways (pages 823–833)

R. P. Jacoby, M. H. Che-Othman, A. H. Millar and N. L. Taylor
Article first published online: 11 DEC 2015 | DOI: 10.1111/pce.12653

A number of previous studies have documented the gross response of mitochondrial respiration to salinity treatment, but it is unclear how NaCl directly affects the kinetics of plant phosphorylating and non-phosphorylating electron transport pathways. This study investigates the direct effects of NaCl upon different respiratory pathways in wheat, by measuring rates of isolated mitochondrial oxygen consumption across different substrate oxidation pathways in saline media. These data deepen our understanding of how plant respiration responds to NaCl, offering new mechanistic explanations for the divergent salinity responses of whole-plant respiratory rate in the literature.

Nature Genetics Contents: April 2016 pp 343 - 473

***OsSPL13* controls grain size in cultivated rice pp447 - 456**

Lizhen Si, Jiaying Chen, Xuehui Huang, Hao Gong, Jianghong Luo *et al.*

doi:10.1038/ng.3518

Bin Han and colleagues present a genome-wide association analysis of grain size and shape in cultivated rice and identify a major locus for grain size encoding the transcription factor OsSPL13. They find that the large-grain allele in tropical *japonica* cultivars was introgressed from *indica* varieties during selection for improved grain yield.

FEBS Journal Content Alert (New Articles)

Sequestration of cellular interacting partners by protein aggregates: implication in a loss-of-function pathology

Hui Yang and Hong-Yu Hu

Accepted manuscript online: 26 MAR 2016 06:40AM EST | DOI: 10.1111/febs.13722

Protein misfolding and aggregation are a hallmark of several neurodegenerative diseases (NDs). However, how protein aggregation leads to cytotoxicity and neurodegeneration is still controversial. Emerging evidence demonstrates that sequestration of cellular interacting partners by protein aggregates contributes to the pathogenesis of these diseases. Here, we review current research on sequestration of cellular proteins by protein aggregates and its relation to proteinopathies. Based on different interaction modes, we classify these protein sequestrations into four types: protein co-aggregation, domain/motif-mediated sequestration, RNA-assisted sequestration, and sequestration of molecular chaperones. Thus, the cellular essential proteins and/or RNAs hijacked by protein aggregates may lose their biological functions, consequently resulting in cytotoxicity and neurodegeneration. We have proposed a hijacking model recapitulating the sequestration process and the loss-of-function pathology of ND.

Mitochondria and senescence: new actors for an old play

Nicolás Herranz and Jesús Gil Published online 23.02.2016

<http://EMBOJ.embopress.org/content/35/7/701?etoc>

A new study reveals a causal role of mitochondria for cellular senescence phenotypes and ageing.

Mack KL, Shorter J.

Engineering and Evolution of Molecular Chaperones and Protein Disaggregases with Enhanced Activity.

Front Mol Biosci. 2016;3:8. PMID: 27014702 [PubMed]

Molière N, Höymann J, Schäfer H, Turgay K.

Role of Hsp100/Clp Protease Complexes in Controlling the Regulation of Motility in *Bacillus subtilis*.

Front Microbiol. 2016;7:315. PMID: 27014237 [PubMed]

Arosio P, Michaels TC, Linse S, Månsson C, Emanuelsson C, Presto J, Johansson J, Vendruscolo M, Dobson CM, Knowles TP.

Kinetic analysis reveals the diversity of microscopic mechanisms through which molecular chaperones suppress amyloid formation.

Nat Commun. 2016 Mar 24;7:10948. PMID: 27009901 [PubMed - in process]

Cuevas-Velazquez CL, Saab-Rincón G, Reyes JL, Covarrubias AA.

The unstructured N-terminal region of Arabidopsis group 4 Late Embryogenesis Abundant Proteins (LEA) is required for folding and for chaperone-like activity under water deficit.
J Biol Chem. 2016 Mar 22;. [Epub ahead of print]
PMID: 27006402 [PubMed - as supplied by publisher]

Kang CH, Lee YM, Park JH, Nawkar GM, Oh HT, Kim MG, Lee SI, Kim WY, Yun DJ, Lee SY.
Ribosomal P3 protein AtP3B of Arabidopsis acts as both protein and RNA chaperone to increase tolerance of heat and cold stresses.
Plant Cell Environ. 2016 Mar 23;. [Epub ahead of print]
PMID: 27004478 [PubMed - as supplied by publisher]

Paila YD, Richardson LG, Inoue H, Parks ES, McMahon J, Inoue K, Schnell DJ.
Multi-functional roles for the polypeptide transport associated domains of Toc75 in chloroplast protein import.
Elife. 2016 Mar 21;5.
PMID: 26999824 [PubMed - in process]

Romero-Puertas MC, Sandalio LM.
Nitric Oxide Level Is Self-Regulating and Also Regulates Its ROS Partners.
Front Plant Sci. 2016;7:316.
PMID: 27014332 [PubMed]

Silveira NM, Frungillo L, Marcos FC, Pelegriano MT, Miranda MT, Seabra AB, Salgado I, Machado EC, Ribeiro RV.
Exogenous nitric oxide improves sugarcane growth and photosynthesis under water deficit.
Planta. 2016 Mar 22;. [Epub ahead of print]
PMID: 27002974 [PubMed - as supplied by publisher]

Ingolia NT.
Ribosome Footprint Profiling of Translation throughout the Genome.
Cell. 2016 Mar 24;165(1):22-33.
PMID: 27015305 [PubMed - in process]

Nahomi RB, Pantcheva M, Nagaraj RH.
Î±B-Crystallin is Essential for the TGF-Î²-mediated Epithelial to Mesenchymal Transition of Lens Epithelial Cells.
Biochem J. 2016 Mar 17;. [Epub ahead of print]
PMID: 26987815 [PubMed - as supplied by publisher]

Bogamuwa S, Jang JC.
Plant Tandem CCCH Zinc Finger Proteins Interact with ABA, Drought, and Stress Response Regulators in Processing-Bodies and Stress Granules.
PLoS One. 2016;11(3):e0151574. PMID: 26978070 [PubMed - in process]

Lee C, Wigren E, LÃ¼nsdorf H, RÃ¶mling U.
Protein homeostasis-more than resisting a hot bath.
Curr Opin Microbiol. 2016 Mar 11;30:147-154 PMID: 26974352 [PubMed - as supplied by publisher]

Chiappori F, Mattiazzi L, Milanesi L, Merelli I.

A novel molecular dynamics approach to evaluate the effect of phosphorylation on multimeric protein interface: the β -Crystallin case study.
BMC Bioinformatics. 2016 Mar 2;17 Suppl 4:57.
PMID: 26961246 [PubMed - in process]

Arasimowicz-Jelonek M, Floryszak-Wieczorek J.
Nitric Oxide in the Offensive Strategy of Fungal and Oomycete Plant Pathogens.
Front Plant Sci. 2016;7:252. PMID: 26973690 [PubMed]

Niu L, Liao W.
Hydrogen Peroxide Signaling in Plant Development and Abiotic Responses: Crosstalk with Nitric Oxide and Calcium.
Front Plant Sci. 2016;7:230. PMID: 26973673 [PubMed]

Kan Q, Wu W, Yu W, Zhang J, Xu J, Rengel Z, Chen L, Cui X, Chen Q.
Nitrate reductase-mediated NO production enhances Cd accumulation in *Panax notoginseng* roots by affecting root cell wall properties.
J Plant Physiol. 2016 Mar 3;193:64-70. PMID: 26956919 [PubMed - as supplied by publisher]

Yang L, Ji J, Harris-Shultz KR, Wang H, Wang H, Abd-Allah EF, Luo Y, Hu X.
The Dynamic Changes of the Plasma Membrane Proteins and the Protective Roles of Nitric Oxide in Rice Subjected to Heavy Metal Cadmium Stress.
Front Plant Sci. 2016;7:190. PMID: 26955374 [PubMed]

Weis F, Giudice E, Churcher M, Jin L, Hilcenko C, Wong CC, Traynor D, Kay RR, Warren AJ.
Mechanism of eIF6 release from the nascent 60S ribosomal subunit.
Nat Struct Mol Biol. 2015 Nov;22(11):914-9. PMID: 26479198 [PubMed - indexed for MEDLINE]

The Plant Journal Content Alert (New Articles)

[Extending the biosynthetic repertoires of cyanobacteria and chloroplasts](#)

Agnieszka Zygadlo Nielsen, Silas Busck Mellor, Konstantinos Vavitsas, Artur Jacek Włodarczyk, Thiagarajan Gnanasekaran, Maria Perestrello Ramos H de Jesus, Brian Christopher King, Kamil Bakowski and Poul Erik Jensen

Accepted manuscript online: 23 MAR 2016 01:11AM EST | DOI: 10.1111/tpj.13173

Advances in Botanical Research: Alert 19 March-25 March

[Coevolution of Organelle RNA Editing and Nuclear Specificity Factors in Early Land Plants](#) Original Research Article

Pages 37-93

M. Schallenberg-Rüdinger, V. Knoop

Cell: Alert 19 March-25 March

[Types of Ubiquitin Ligases](#)

Pages 248-248.e1

Francesca Ester Morreale, Helen Walden

Current Opinion in Cell Biology: Alert 18 March-24 March

[Shaping the multi-scale architecture of mitochondria](#) Review Article

Pages 45-51

Vaishali Jayashankar, Irina A Mueller, Susanne M Rafelski

Plant, Cell & Environment Content Alert (New Articles)

[Ribosomal P3 protein AtP3B of Arabidopsis acts as both protein and RNA chaperone to increase tolerance of heat and cold stresses](#)

Chang Ho Kang, Young Mee Lee, Joung Hun Park, Ganesh M. Nawkar, Hun Taek Oh, Min Gap Kim, Soo In Lee, Woe Yeon Kim, Dae-Jin Yun and Sang Yeol Lee

Accepted manuscript online: 23 MAR 2016 05:15AM EST | DOI: 10.1111/pce.12742

[A dual system formed by the ARC and NR molybdoenzymes mediates nitrite-dependent NO production in *Chlamydomonas*](#)

Alejandro Chamizo-Ampudia, Emanuel Sanz-Luque, Ángel Llamas, Francisco Ocaña-Calahorro, Vicente Mariscal, Alfonso Carreras, Juan B. Barroso, Aurora Galván and Emilio Fernández

Accepted manuscript online: 17 MAR 2016 05:05AM EST | DOI: 10.1111/pce.12739

FEBS Journal Content Alert (New Articles)

[To deliver or to degrade: an interplay of the ubiquitin–proteasome system, autophagy and vesicular transport in plants](#)

Katarzyna Zientara-Rytter and Agnieszka Sirko

Accepted manuscript online: 17 MAR 2016 01:00AM EST | DOI: 10.1111/febs.13712

Nature Reviews Molecular Cell Biology contents January 2016 Volume 17 Number 4 pp 201-262

Mitonuclear communication in homeostasis and stress

Pedro M. Quirós, Adrienne Mottis & Johan Auwerx

p213 | doi:10.1038/nrm.2016.23

As most mitochondrial proteins are encoded in the nucleus, mitochondrial activity requires efficient communication between the nuclear and mitochondrial genomes. This is mediated by nucleus-to-mitochondria (anterograde), mitochondria-to-nucleus (retrograde) and mitonuclear feedback signalling, as well as the integrated stress response and extracellular communication, which regulate homeostasis and, consequently, healthspan and lifespan.

[Abstract](#) | [Full Text](#) | [PDF](#)

Molecular Cell: Alert 12 March-18 March

[Resources for the Comprehensive Discovery of Functional RNA Elements](#) Original Research Article

Pages 903-913

Balaji Sundararaman, Lijun Zhan, Steven M. Blue, Rebecca Stanton, Keri Elkins, Sara Olson, Xintao Wei, Eric L. Van Nostrand, Gabriel A. Pratt, Stephanie C. Huelga, Brendan M. Smalec, Xiaofeng Wang, Eurie L. Hong, Jean M. Davidson, Eric Lécuyer, Brenton R. Graveley, Gene W. Yeo