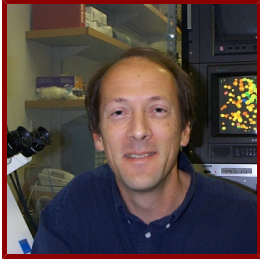


Stanford



Richard Lewis

Professor of Molecular and Cellular Physiology
Molecular & Cellular Physiology

CONTACT INFORMATION

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Bio

ACADEMIC APPOINTMENTS

- Professor, Molecular & Cellular Physiology
- Member, Bio-X
- Member, Stanford Neurosciences Institute

ADMINISTRATIVE APPOINTMENTS

- Chair, Stanford University School of Medicine - Molecular & Cellular Physiology, (2004-2009)

HONORS AND AWARDS

- Junior Faculty Scholar, HHMI (1996-1998)
- MERIT Award, National Institute of General Medical Sciences (July 2012 - present)

PROFESSIONAL EDUCATION

- B.S., Yale University, Molec Biophysics & Biochem (1978)
- Ph.D., California Inst. of Technology, Neurobiology (1985)

LINKS

- Lewis Lab site: <http://med.stanford.edu/lewislab>

Research & Scholarship

CURRENT RESEARCH AND SCHOLARLY INTERESTS

Research in our laboratory is focused on the molecular mechanisms of calcium signaling through store-operated Ca²⁺ channels (SOCs). This class of ion channels is regulated in a unique way, by the depletion of Ca²⁺ from the lumen of the endoplasmic reticulum (ER) which normally occurs following stimulation of cell surface receptors that generate IP₃. They are expressed in practically all cells, where they contribute to diverse functions including secretion, gene expression, and cell differentiation. A SOC called the Ca²⁺ release-activated Ca²⁺ channel, or CRAC channel, is particularly important in T cells, where it generates sustained Ca²⁺ signals

that are essential for triggering T cells to proliferate and carry out immune functions. Loss of function of the CRAC channel by a single mutation in its structural gene leads to a devastating severe combined immunodeficiency (SCID) syndrome in humans.

A major effort in the lab is to understand how the depletion of Ca²⁺ from the ER triggers the opening of CRAC channels in the plasma membrane. Since the recent discovery of genes encoding the ER Ca²⁺ sensor (STIM1) and the CRAC channel (Orai1), we have made rapid progress in revealing the choreographic nature of this process. Following store depletion, STIM1 moves from locations throughout the ER to accumulate in ER subregions positioned within 10-25 nm of the plasma membrane (PM). Simultaneously, the CRAC channel gathers at corresponding sites in the PM directly opposite STIM1, where it is opened through an interaction with STIM1. This is an unprecedented mechanism for channel activation, in which a stimulus acts to bring a channel and its activator/sensor together for interaction across apposed membrane compartments. We are currently studying how changes in ER Ca²⁺ drive the redistribution of STIM1 and the mechanism by which STIM1 and Orai1 interact across the ER-PM gap. For these studies we combine protein engineering with patch-clamp electrophysiology and live-cell imaging using total internal reflection fluorescence (TIRF), confocal, and single-molecule microscopy.

A second major area of interest is how specific information is encoded in Ca²⁺ signals. Specificity is an acute problem for pluripotent messengers like Ca²⁺ that are involved in multiple signaling pathways. We have shown that transcriptional specificity in T cells can be achieved through the amplitude and dynamics of Ca²⁺ signals generated by CRAC channels. We are now studying how these features contribute to cell fate decisions during T cell development. In the thymus, self-reactive thymocytes are deleted through negative selection, while cells with the appropriate avidity for “self” are allowed to mature into T cells and populate the periphery (positive selection). To study the role of Ca²⁺ in this choice, we have developed a novel thymic slice preparation in which we use two-photon microscopy to track and record Ca²⁺ signals in single thymocytes as they migrate through tissue engineered to provide defined selection signals. We have found that positive selection is associated with Ca²⁺ oscillations, which immobilize the cells at locations of self-antigen recognition to promote gene activation. We are currently comparing signaling during positive and negative selection to determine how the Ca²⁺ signal “signature” helps a T cell decide whether to live and prosper or die.

Teaching

COURSES

2016-17

- How Cells Work: Energetics, Compartments, and Coupling in Cell Biology: MCP 156, MCP 256 (Spr)

2015-16

- Imaging: Biological Light Microscopy: BIO 152 (Spr)

2014-15

- How Cells Work: Energetics, Compartments, and Coupling in Cell Biology: MCP 156, MCP 256 (Win)
- Imaging: Biological Light Microscopy: BIO 152, MCP 222 (Win)

STANFORD ADVISEES

Postdoctoral Faculty Sponsor

Ruoyi Qiu, Stijn Van Dorp

GRADUATE AND FELLOWSHIP PROGRAM AFFILIATIONS

- Biophysics (Phd Program)
- Chemical and Systems Biology (Phd Program)
- Immunology (Phd Program)
- Molecular and Cellular Physiology (Phd Program)

- Neurosciences (Phd Program)

Publications

PUBLICATIONS

- **Functional Analysis of Orai1 Concatemers Supports a Hexameric Stoichiometry for the CRAC Channel.** *Biophysical journal*
Yen, M., Lokteva, L. A., Lewis, R. S.
2016; 111 (9): 1897-1907
- **Orai1 pore residues control CRAC channel inactivation independently of calmodulin.** *journal of general physiology*
Mullins, F. M., Yen, M., Lewis, R. S.
2016; 147 (2): 137-152
- **The inactivation domain of STIM1 is functionally coupled with the Orai1 pore to enable Ca²⁺-dependent inactivation.** *journal of general physiology*
Mullins, F. M., Lewis, R. S.
2016; 147 (2): 153-164
- **Calcium influx through CRAC channels controls actin organization and dynamics at the immune synapse.** *eLife*
Hartzell, C. A., Jankowska, K. I., Burkhardt, J. K., Lewis, R. S.
2016; 5
- **STORE-OPERATED CALCIUM CHANNELS** *PHYSIOLOGICAL REVIEWS*
Prakriya, M., Lewis, R. S.
2015; 95 (4): 1383-1436
- **Alternative splicing converts STIM2 from an activator to an inhibitor of store-operated calcium channels.** *journal of cell biology*
Rana, A., Yen, M., Sadaghiani, A. M., Malmersjö, S., Park, C. Y., Dolmetsch, R. E., Lewis, R. S.
2015; 209 (5): 653-670
- **Single-molecule analysis of diffusion and trapping of STIM1 and Orai1 at endoplasmic reticulum-plasma membrane junctions** *MOLECULAR BIOLOGY OF THE CELL*
Wu, M. M., Covington, E. D., Lewis, R. S.
2014; 25 (22): 3672-3685
- **Stoichiometric requirements for trapping and gating of Ca²⁺ release-activated Ca²⁺ (CRAC) channels by stromal interaction molecule 1 (STIM1)** *PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA*
Hoover, P. J., Lewis, R. S.
2011; 108 (32): 13299-13304
- **Essential Role for the CRAC Activation Domain in Store-dependent Oligomerization of STIM1** *MOLECULAR BIOLOGY OF THE CELL*
Covington, E. D., Wu, M. M., Lewis, R. S.
2010; 21 (11): 1897-1907
- **STIM1 Clusters and Activates CRAC Channels via Direct Binding of a Cytosolic Domain to Orai1** *CELL*
Park, C. Y., Hoover, P. J., Mullins, F. M., Bachhawat, P., Covington, E. D., Raunser, S., Walz, T., Garcia, K. C., Dolmetsch, R. E., Lewis, R. S.
2009; 136 (5): 876-890
- **Oligomerization of STIM1 couples ER calcium depletion to CRAC channel activation** *NATURE*
Luik, R. M., Wang, B., Prakriya, M., Wu, M. M., Lewis, R. S.
2008; 454 (7203): 538-U11
- **The molecular choreography of a store-operated calcium channel** *NATURE*
Lewis, R. S.
2007; 446 (7133): 284-287
- **Ca²⁺ store depletion causes STIM1 to accumulate in ER regions closely associated with the plasma membrane** *JOURNAL OF CELL BIOLOGY*
Wu, M. M., Buchanan, J., Luik, R. M., Lewis, R. S.
2006; 174 (6): 803-813

- **The elementary unit of store-operated Ca²⁺ entry: local activation of CRAC channels by STIM1 at ER-plasma membrane junctions** *JOURNAL OF CELL BIOLOGY*
Luik, R. M., Wu, M. M., Buchanan, J., Lewis, R. S.
2006; 174 (6): 815-825
- **Regulation of CRAC channel activity by recruitment of silent channels to a high open-probability gating mode** *JOURNAL OF GENERAL PHYSIOLOGY*
Prakriya, M., Lewis, R. S.
2006; 128 (3): 373-386
- **Calcium oscillations regulate thymocyte motility during positive selection in the three-dimensional thymic environment** *NATURE IMMUNOLOGY*
Bhakta, N. R., Oh, D. Y., Lewis, R. S.
2005; 6 (2): 143-151
- **Calcium oscillations increase the efficiency and specificity of gene expression** *NATURE*
Dolmetsch, R. E., XU, K. L., Lewis, R. S.
1998; 392 (6679): 933-936
- **Abnormal Calcium Handling Properties Underlie Familial Hypertrophic Cardiomyopathy Pathology in Patient-Specific Induced Pluripotent Stem Cells** *CELL STEM CELL*
Lan, F., Lee, A. S., Liang, P., Sanchez-Freire, V., Nguyen, P. K., Wang, L., Han, L., Yen, M., Wang, Y., Sun, N., Abilez, O. J., Hu, S., Ebert, et al
2013; 12 (1): 101-113
- **Store-Operated Calcium Channels: New Perspectives on Mechanism and Function** *COLD SPRING HARBOR PERSPECTIVES IN BIOLOGY*
Lewis, R. S.
2011; 3 (12)
- **Molecular Basis of Calcium Signaling in Lymphocytes: STIM and ORAI** *ANNUAL REVIEW OF IMMUNOLOGY, VOL 28*
Hogan, P. G., Lewis, R. S., Rao, A.
2010; 28: 491-533
- **Differential Contribution of Chemotaxis and Substrate Restriction to Segregation of Immature and Mature Thymocytes** *IMMUNITY*
Ehrlich, L. I., Oh, D. Y., Weissman, I. L., Lewis, R. S.
2009; 31 (6): 986-998
- **STIM1 and calmodulin interact with Orail to induce Ca²⁺-dependent inactivation of CRAC channels** *PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA*
Mullins, F. M., Park, C. Y., Dolmetsch, R. E., Lewis, R. S.
2009; 106 (36): 15495-15500
- **Some assembly required: Constructing the elementary units of store-operated Ca²⁺ entry** *CELL CALCIUM*
Wu, M. M., Luik, R. M., Lewis, R. S.
2007; 42 (2): 163-172
- **New insights into the molecular mechanisms of store-operated Ca²⁺ signaling in T cells** *TRENDS IN MOLECULAR MEDICINE*
Luik, R. M., Lewis, R. S.
2007; 13 (3): 103-107
- **Real-time measurement of signaling and motility during T cell development in the thymus** *SEMINARS IN IMMUNOLOGY*
Bhakta, N. R., Lewis, R. S.
2005; 17 (6): 411-420
- **A severe defect in CRAC Ca²⁺ channel activation and altered K⁺ channel gating in T cells from immunodeficient patients** *JOURNAL OF EXPERIMENTAL MEDICINE*
Feske, S., Prakriya, M., Rao, A., Lewis, R. S.
2005; 202 (5): 651-662
- **Modulation of plasma membrane calcium-ATPase activity by local calcium microdomains near CRAC channels in human T cells** *JOURNAL OF PHYSIOLOGY-LONDON*
Bautista, D. M., Lewis, R. S.
2004; 556 (3): 805-817

- **CRAC channels: activation, permeation, and the search for a molecular identity** *CELL CALCIUM*
Prakriya, M., Lewis, R. S.
2003; 33 (5-6): 311-321
- **Enhancement of calcium signalling dynamics and stability by delayed modulation of the plasma-membrane calcium-ATPase in human T cells** *JOURNAL OF PHYSIOLOGY-LONDON*
Bautista, D. M., Hoth, M., Lewis, R. S.
2002; 541 (3): 877-894
- **Separation and characterization of currents through store-operated CRAC channels and Mg²⁺-inhibited cation (MIC) channels** *JOURNAL OF GENERAL PHYSIOLOGY*
Prakriya, M., Lewis, R. S.
2002; 119 (5): 487-507
- **Potentiation and inhibition of Ca²⁺ release-activated Ca²⁺ channels by 2-aminoethyl-diphenyl borate (2-APB) occurs independently of IP₃ receptors** *JOURNAL OF PHYSIOLOGY-LONDON*
Prakriya, M., Lewis, R. S.
2001; 536 (1): 3-19
- **Calcium signaling mechanisms in T lymphocytes** *ANNUAL REVIEW OF IMMUNOLOGY*
Lewis, R. S.
2001; 19: 497-521
- **Mitochondrial control of calcium-channel gating: A mechanism for sustained signaling and transcriptional activation in T lymphocytes** *PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA*
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2000; 97 (19): 10607-10612
- **Store-operated calcium channels** *ION CHANNEL REGULATION*
Lewis, R. S.
1999; 33: 279-307
- **Mitochondrial regulation of store-operated calcium signaling in T lymphocytes** *JOURNAL OF CELL BIOLOGY*
Hoth, M., Fanger, C. M., Lewis, R. S.
1997; 137 (3): 633-648
- **Differential activation of transcription factors induced by Ca²⁺ response amplitude and duration** *NATURE*
Dolmetsch, R. E., Lewis, R. S., Goodnow, C. C., Healy, J. I.
1997; 386 (6627): 855-858
- **Function follows form: The role of store-operated calcium channels in T-cell activation** *CELLULAR PHYSIOLOGY AND BIOCHEMISTRY*
Fanger, C. M., Zweifach, A., Dolmetsch, R. E., Hoth, M., Lewis, R. S.
1997; 7 (3-4): 203-218
- **Calcium-dependent potentiation of store-operated calcium channels in T lymphocytes** *JOURNAL OF GENERAL PHYSIOLOGY*
Zweifach, A., Lewis, R. S.
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- **CHARACTERIZATION OF T-CELL MUTANTS WITH DEFECTS IN CAPACITATIVE CALCIUM-ENTRY - GENETIC-EVIDENCE FOR THE PHYSIOLOGICAL ROLES OF CRAC CHANNELS** *JOURNAL OF CELL BIOLOGY*
Fanger, C. M., Hoth, M., Crabtree, G. R., Lewis, R. S.
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- **SLOW CALCIUM-DEPENDENT INACTIVATION OF DEPLETION-ACTIVATED CALCIUM CURRENT - STORE-DEPENDENT AND STORE-INDEPENDENT MECHANISMS** *JOURNAL OF BIOLOGICAL CHEMISTRY*
Zweifach, A., Lewis, R. S.
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- **RAPID INACTIVATION OF DEPLETION-ACTIVATED CALCIUM CURRENT (I-CRAC) DUE TO LOCAL CALCIUM FEEDBACK** *JOURNAL OF GENERAL PHYSIOLOGY*
Zweifach, A., Lewis, R. S.

1995; 105 (2): 209-226

- **SIGNALING BETWEEN INTRACELLULAR CA₂⁺ STORES AND DEPLETION-ACTIVATED CA₂⁺ CHANNELS GENERATES [CA₂⁺]_i OSCILLATIONS IN T-LYMPHOCYTES** *JOURNAL OF GENERAL PHYSIOLOGY*

Dolmetsch, R. E., Lewis, R. S.

1994; 103 (3): 365-388

- **MITOGEN-REGULATED CA₂⁺ CURRENT OF T-LYMPHOCYTES IS ACTIVATED BY DEPLETION OF INTRACELLULAR CA₂⁺ STORES** *PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA*

Zweifach, A., Lewis, R. S.

1993; 90 (13): 6295-6299