

Regulation of cyclin D1 expression in ES cells: the role of the Ras/MAPK pathway

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Pluripotent Stem Cells: Biology and Applications

Organizers
Calvin B. Harley, John Gearhart, Rudolf Jaenisch,
Janet Rossant and James Thomson

Sponsored by Amgen Inc.

Sheraton Tamarron Resort • Durango, Colorado February 6 - 11, 2001

Celebrating 30 Years of Connecting the Scientific Community

A Keystone Symposium

Pluripotent Stem Cells: Biology and Applications

Organizers: Calvin B. Harley, John Gearhart, Rudolf Jaenisch, Janet Rossant and James Thomson

Sheraton Tamarron Resort, Durango, Colorado • February 6 - 11, 2001

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Tuesday, February 6		
2 - 7pm 6:15 - 7:15pm	Registration Welcome	Alcove Windom Peak
7:15 - 7:30pm 7:30 - 8:30pm	Orientation Keynote Address Robert A. Weinberg, Whitehead Institute for Biomedical Research (001) Rules Governing the Creation of Human Tumor Cells	Silverton Silverton
	Wednesday, February 7	
6:30 - 8am	Breakfast	Windom Peak/Antlers/Kiva
8 – 11am	Embryonic Cells: Origin, Derivation, Properties I James A. Thomson, University of Wisconsin (002) Derivation and Properties of Human Embryonic Stem Cells Austin G. Smith, University of Edinburgh (003) Determinants of Embryonic Stem Cell Pluripotency Coffee Break Hans R. Schöler, University of Pennsylvania (004) Oct-4 Gene Function and Regulation in Pluripotent Stem Cells Alan Trounson, Monash Institute for Reproduction and Development (00) Human Embryo, Embryonic Stem Cells and Differentiation	Silverton Windom Peak 05)
11am - 1pm	Poster Setup	Innisbrook/Animas
4 – 6pm	Poster Session 1: ES, EG and Adult Stem Cells	Innisbrook/Animas
5 – 6pm	Social Hour	Windom Peak
8 – 10pm	Embryonic Cells: Origin, Derivation, Properties II Coffee Available ~ Windom Peak John Gearhart, Johns Hopkins University Hospital Derivation and Properties of Human Embryonic Germ Cells Peter J. Donovan, Thomas Jefferson University (007)	Silverton

Roger A. Pedersen, University of California-San Francisco

Pluripotent Embryonic Germ Cell Lines

Embryonic Approaches to Stem Cell Differentiation

Regulation of cyclin D1 expression in ES cells : the role of the RAS \rightarrow MAPK pathway.

Marielle Afanassieff, Ludmila Jirmanova, Ludovic Vallier, Suzy Markossian, Jacques Samarut, and Pierre Savatier. Ecole Normale Supérieure de Lyon, CNRS UMR5665 – INRA LA913, Lyon, France.

The cellular machinery that is organized to collect extracellular signals and transduce them via tyrosine kinase receptors and the SOS-RAS-MEKK-MAPK pathway seems to be dedicated largely to driving RB phosphorylation. Cyclin D1 is likely to play a key role in this process as (i) cyclin D1 expression is regulated by MAPK-dependent transcription factors (jun, fos, ets) and (ii) cyclin D1/CDK4 complexes are involved in RB phosphorylation during the G1 phase of the cell cycle. This circuitry appears to be operative in virtually all cell types and its relevance is illustrated by its disruption in many types of human tumors. Using ES cells which express the neo' gene driven off the oct4 promoter (IOUD2 cells) so as to kill all spontaneously differentiating cells, we show that ES cells express low, although constant, level of cyclin D1. This results from low activity of the cyl1 promoter. Promoter deletion analysis shows that this background activity originates from a proximal region of the promoter, and that distal regions known to be involved in cyclin D1 expression in response to serum stimulation are not required. Treatment of IOUD2 cells with wortmannin (an inhibitor of RAS activation) or PD98059 (an inhibitor of MEK) does not inhibit cyclin D1 expression. None of these inhibitors induce growth retardation. Induction of differentiation with retinoic acid (RA) up-regulates cyclin D1 expression, and distal - serum-responsive - elements of the cyll promoter are involved. Differentiating cells become sensitive to inhibitors of the Ras → ERK cascade. Both wortmannin and PD98059 inhibit cell growth and down-regulate expression of cyclin D1. Hence, cyclin D1 expression seems not to be regulated by the RAS -> MEK -> ERK phosphorylation in ES cells. This regulation is likely to be restored upon differentiation.

- (1) Marielle AFANASSIEFF
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- (3) meeting C1
- (4) poster session I



KEYSTONE SYMPOSIA

Connecting the Scientific Community

Thursday, November 16, 2000

Marielle Afanassieff Laboratoire de Biologie Ecole Normale Superieure 46 Allee d'Italie, Cedex 07 Lyon, 69364 France

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Your abstract has been accepted as

Poster Number 102 in

POSTER SESSION 1: ES, EG and Adult Stem Cells

Wednesday, February 7, 2001

Pluripotent Stem Cells: Biology and Applications (C1)

February 6 - February 11, 2001

Sheraton Tamarron Resort, Durango, Colorado Early Registration Deadline: December 6, 2000

- Please remember that Abstract Submission / Poster Acceptance does not guarantee registration.
- If you have not already registered for this meeting, please do so as soon as possible.
- You may register at www.keystonesymposia.org
- Poster sessions will be held from 4:00 PM 6:00 PM
- You may set up your poster after 11:00 AM on Wednesday, February 7.
- The poster must be removed by the end of the poster session.
- Each poster will have 4 feet by 4 feet of space available for display.
- Thumbtacks will be available for mounting your materials.
- Your poster will be viewed from a distance of several feet; use bold type and large figures.
- Keystone Symposia staff will put poster numbers on the display boards before you set up your poster.
- Your abstract will be published in a book which will include all poster and speaker abstracts received for this meeting.
- This book will be distributed to all registered conferees upon check-in.

Sincerely,

Keystone Symposia on behalf of Calvin B. Harley, John Gearhart, Rudolf Jaenisch, Janet Rossant and James Thomson, Organizers Pluripotent Stem Cells: Biology and Applications