# Australia and New Zealand Society for Cell and Developmental Biology

INCORPORATED



### NEWSLETTER - June 2010 Volume 1 of 3

www.anzscdb.org



Edna Hardeman

#### We have grown!

Increasing our membership was a major objective for this year and we are off to an excellent start. A warm welcome to new members and a thank you to those who have renewed! Since January our membership has grown by 30%. Please help us to continue the momentum and encourage staff, students and colleagues to join. Tell them to simply click on http://www.anzscdb. org/ANZSCDB-Membership.html

## Announcement: 2010 Young Investigator Award

The Young Investigator Award recognises an outstanding emerging leader in our disciplines. The applicants for the 2010 award were truly impressive and the selection committee praised the talent that is apparent in the up and coming leaders in our Society. All are to be congratulated on their achievements. The committee is pleased to announce that the recipient of the 2010 YIA is Assoc/Prof Kat Gaus from the Centre for Vascular Research, the University of New South Wales. Kat's research is focused on understanding the organisation of the plasma membrane using a multi-disciplinary approach that includes cell and molecular biology, biphotonics

### IN THIS ISSUE

President's report	1
Editor's column	2
Profile of Prof. Jenny Stow	
Profile ANZSCDB Young Investigator of the Year Award	
A/Prof. Katharina Gaus	. 7
Conference announcements	. 9
ANZSCDB Images for Website Contest:	
Prize winners	. 15
Reports:	
16th NSW Cell and Developmental Biology	
Meeting	
2010 Science Meets Parliament	
60th Meeting of Nobel Laureates	
The Hunter Meeting	19
Letter from the ISDB	
Publications from our members	23
Members in the news	
Executive officers/committee members	30

# President's Report

and surface chemistry. What sets Kat apart is that she is actively engaged in developing new generations of microscopes which allow the analysis of single molecular events. This will transform our understanding of signalling pathways and the organisation of membrane domains and how they regulate signal transmission. One of the major outcomes from Kat's development of these microscopes is the establishment of the Biomedical Imaging Facility at UNSW which is the most advanced fluorescence microscopy centre in Australia and is comparable with the best internationally. Kat will receive her award at OzBio2010 and present her work in the symposium Dynamic subcellular compartments. Kat's scientific journey and accomplishments are profiled in this newsletter.

OzBio2010

Already we've achieved a record attendance at OZBIO2010 (Melbourne, 26 Sept - 1 Oct, 2010) which encompasses the Society's annual meeting COMBIO together with the 12th IUBMB and the 21st FAOBMB. We are indebted to our members on the organising committee Marie Bogoyevitch and David Jans (Cell Architecture & Trafficking Stream) and Helen Abud and Joy Rathjen (Developmental Biology Stream) for organising the symposia as well as securing prominent plenary speakers in our disciplines including Eliza Izaurralde (Scientific Director, Max Planck Inst for Developmental Biology), Tom Rapoport (Harvard) and Lee Niswander (Uni Colorado). This year the Society offered two schemes to engage and support postdoctoral fellow and student attendance at OZBIO. ANZSCDB together with ASBMB offered fellowships that covered

attendance at the Young Scientist Forum (YSF), a conference of 50 international and 10 Australian PhD/early postdoctoral scientists preceding OZBIO, as well as OZ-BIO. The response was high as was the quality of the applications, with approximately 40% of the applicants ANZSCDB members. The selection committee is pleased to announce that National YSF Fellowships were awarded to Stephanie Bannister (CSIRO, VIC) and Cassy Spiller (IMB, QLD). In addition, ANZSCDB offered 20 student bursaries to attend OZBIO (Awardees: Heidi Bildsoe. Teresa Bonello, Yigian Chen, Elizabeth Christie, Baptiste Coxam, Nikki Curthoys, Felix Grusche, Joelle Kartopawiro, Kristie Lee, Jet Phey Lim, Maggie Ma, Sabine Mangold, Rosemary Manhire-Heath, Vicki Metzis, Melissa Pert, Anne Raimondo, Ashesha Sinha, Sophie Wiszniak, Sheena Yao, Hyun Jin Yoo).

#### Sponsored Meetings & Workshops

ANZSCDB's Distinguished Visiting Lectureship in Cell & Developmental Biology Program sponsored Ed Munro (University of Chicago) as the 2010 ANZSCDB HM International Speaker at the Hunter Meeting (http://hcbm.mtci.com.au/) as well as an invited speaker at the 16th NSW Cell and Developmental Biology Meeting, UNSW. Ed was an outstanding contributor at both meetings offering two distinct talks that demonstrated the emerging power of a systems biology approach to fundamental questions in cell and developmental biology. Please do submit suggestions for future ANSCDB speakers at the Hunter Meeting.

The 6th Developmental Biology Workshop (Oct 24-27, Yarra Valley Conference Centre) provides a unique opportunity to engage with international leaders in Developmental Biology and receive in depth tutelage in prominent topics. Organisers Peter Currie, Peter Koopman, Richard Harvey, Rob Saint and Patrick Tam have organised an outstanding program including invited speakers Margaret Buckingham, Stephen Cohen and Austin Smith. Further information is available in this newsletter and on our website http://www.

anzscdb.org/groups/anzscdb/ wiki/0b07d/Events.html

#### State/NZ Chapter Activities

Our program to support local and state Cell and Developmental Biology themed activities is strong with one state-based meeting already held and two organised for later in the year. NSW state representatives Thomas Fath (UNSW) and Bill Phillips (USyd) organised the 16th NSW Cell & Developmental Biology Meeting on March 15th held in the School of Medical Sciences, UNSW and jointly sponsored by ANZSCDB and ASBMB. Over 100 attendees enjoyed an excellent scientific program which included invited speakers Ed Munro (Uni Chicago), Patrick Humbert (Peter Mac) and Miles Davenport (CVR, UNSW). Meetings organised include the 20th Annual Combined Biological Sciences Meeting - Megan Lloyd, August 27th, University of Western Australia and we are delighted to support the 1st Brisbane Cell & Developmental Biology Meeting - Dagmar Wilhelm, Eva Kovacs and Kelly Smith, Oct 22nd, Institute for Molecular Bioscience, University of Queensland. Further information is available in this newsletter.

#### **Affiliated Societies**

As a member society of the National Committee for Biomedical Sciences (NCBMS) of the Australian Academy of Science, we were able to nominate student members to attend the 60th meeting of Nobel Laureates in Lindau, Germany in 2010. We were delighted when 2 of our nominees, Duncan Mortimer (UQ) and Denise Miles (Murdoch Institute) were selected by the international Council for Lindau Nobel Laureate Meetings. They will attend the meeting in Germany from June 27 to July 2 courtesy of the Australian Academy of Science and the Lindau Committee.

Our thanks go to Marie Bogoyevitch and Aleksandra Filipovska who represented ANZSCB at FASTS' Science Meets Parliament 2010. SmP provides valuable insight into how science is perceived at the federal level and provides practical instruction on how to represent science to the politicians.

#### EDITOR'S COLUMN

In this issue of the newsletter we highlight the winners of the photography competition which will feature as the new ANZSCDB banner. Lots and lots of conferences coming up! A detailed outline of the OzBio2010 program is shown

Congratulations to A/Prof. Kat Gaus whom is the winner of the ANZSCDB Young Investigator of the Year Awardee.

Please keep me updated with any comments, news and research advances that you would like to share with other ANZSCDB members.

Megan Chircop (nee Fabbro) mchircop@cmri.org.au

As a member society of the International Society for Developmental Biology we have been asked by the President, Claudio Stern, to comment on the activities of the ISDB (see letter in this newsletter). Please do take this opportunity to respond as to how the ISDB can best meet the needs of our members.

#### A Bit of Fun

To showcase our excellent cell and developmental biology research and the innate artistry of our disciplines, we called upon our members to submit images to use in the heading of the Society's homepage on the web and in printed material advertising our Society. Thank you to all who submitted images to our 2nd ANZSCDB Image Competition and enjoy the winning images shown in this newsletter and of course, on our homepage http://www.anzscdb.org/

Edna Hardeman

# Profile



### Prof. Jenny Stow

Institute of Molecular Biosciences

University of Queensland

## JENNY STOW

### The joys of directing traffic

As far back as Jenny Stow can remember, she and everyone else in her life assumed that she would end up as a vet. Of course, life generally bowls the wrong'un at some stage, and the year that she applied, the vet school in Melbourne took in 17 men...and noone else. So, despite getting the marks, Stow's plans for the future were already set to change. One look at her CV, however, tells you that it turned out OK.

Just a few years later, Professor Stow is an NH-MRC Principal Research Fellow with the Institute for Molecular Bioscience (IMB) at The University of Queensland where she heads a busy and productive research group in cell biology. Her contributions to protein trafficking research over the past two and a half decades and her role as a key driver of Australian cell biology are recognised internationally. She has over 100 peer-reviewed publications to her name and a career-long list of fellowships, national and international grants, appointments, prominent invitations and awards. In her spare time, she has also been Deputy Director (Research) of the IMB since 2008.

#### The lure of research land

So, going back to those summer salad days in the 70s, Stow started a science degree at Monash University with the intention of reapplying for vet science the next year...instead, she discovered basic science, and particularly immunology. She went on to do Honours in the Pathology department at Melbourne's Alfred Hospital, where she spent the year trying to make antibodies in rabbits to different neuronal populations in the brain. Besides gaining several pet rabbits from that year, Stow also came to realise how much she loved doing research...and the dye was set. After Honours, Stow took up a PhD project with Eric Glascow in the Anatomy Department at Monash University and Bob Atkins at Prince Henry's Hospital. "They worked on renal cell populations in glomerulonephritis (GN) and my job was to characterise those cell populations by culturing cells from glomeruli that I isolated from patient biopsies. This work introduced me to clinical-type research and to electron microscopy, both of which I loved."

Stow completed her PhD quite quickly, but had realised towards the end that her research existence was very cloistered, with little exposure to the idea of the next step or how to get there. "By happenstance, not long before I finished my PhD I was in the department next door

I finished my PhD I was in the department next door trying to find out about some extracellular matrix proteins in my samples, and I met Vince Hascall who was on sabbatical from NIH. He was, and still is, one of the world experts on proteoglycan matrix proteins. Vince sort of took me under his wing from then on, and he was the one who suggested that I write to Marilyn Farquhar at Yale University – she was working on both kidney and proteoglycans. So I did…and that is how ended up with a Fogarty Fellowship and on a plane to the USA the day after I handed in my thesis."

#### Finding the Holy Land

The Department of Cell Biology at Yale University School of Medicine was a 'mecca' of cell biology at that time and presented a very steep learning curve for Stow in terms of the research world. "For one thing, I was already months behind on journal articles because Australia got them months after the published date back then. It was a very eye-opening but also very exciting time, and it was when I discovered cell biology. My peers at Yale were fantastic and I learnt heaps about science from Marilyn and from everyone around me. Being a medical school it was also rife with these incredibly dedicated and very smart clinician scientists, and the intensity of that working environment has stood me well ever since."

In Farquhar's lab, Stow set about trying to characterise glomerular basement membrane proteoglycans using biochemistry and electron microscopy. The work yielded several publications, including a couple that turned out to be quite famous papers about proteoglycans making up the glomerular filter for plasma proteins in the kidney. Towards she end of her time at Yale, Stow collaborated with Michael Caplan, then a postgraduate student, to look at polarized secretion in kidney epithelial cells. This successful collaboration yielded a paper in Nature and ignited Stow's growing interest in the mysteries of post-Golgi trafficking. After six highly productive years, Stow left Yale to take up her first Faculty position at Massachusetts General Hospital (MGH)/Harvard Medical School in the Departments of Medicine and Pathology, and moved to Boston. This was to begin an equally productive period and working relationship with renal clinician and department head Dennis Ausiello and fellow group leader, Dennis Brown, working on heterotrimeric G proteins and trafficking. "They were both fantastic mentors who provided loads of support and advice while I was starting my own lab." There, she continued with the kidney trafficking theme, but shifted to more of a Golgi focus.

Again finding herself in an incredibly exciting and stimulating research environment, this next period at MGH was also pivotal for Stow's future career success as a cell biologist. "The Harvard appointment allowed me to establish an independent research group with funding from NIH and various foundations.

I became an investigator on two NIH programs – one in renal cell biology with others in the Renal Unit, and another in digestive diseases, for which I also ran a morphology core facility. Then, with help from the Dennis's, I also wrote and got my own RO1 grant from the NIH. I certainly learnt lots during that time about funding...and it solidified for me the absolute value of basic research for advancing medical science." Stow's work in Boston produced several key findings on trafficking and polarity in kidney cells, revealing new functions for G proteins at the Golgi. These findings transpublications that attracted not during 2005. only international recognition but also a 5-year fellowship

lated into several high-profile Jenny with her family on a holiday in Uluru

ology there all by herself seemed rather daunting. "I knew that David and I would at least talk the same language. NHMRC funding was also state-based in those days and so the competition was a little easier in Qld compared to NSW or Victoria, especially for emerging fields...and this was a big incentive. Of course what I thought at the time was that I would go to Brisbane, get a couple of grants and work there for a while, then go home to Melbourne...and here I still am, interestingly along with most of the other group leaders in the CMCB at the time who had similar plans!"

#### The culture shock that was Brisvegas

So in 1994, Stow arrived in sunny Brisbane with container loads of stuff and a Harvard undergrad student called Brandon to help her set up. Brisbane was a bit of a shock work-wise for Stow (heavily censored summary of what she really said). Not only was she situated on an undergraduate university campus for

the first time in her research career, surrounded by typical undergraduate students and all their trappings (bless them), she missed the support and intensity of having a strong cell biology community around her. "CMCB back then was home to a range of different scientists and research areas, and there was no natural or obvious home for my science. David James and I really were a bit of an outreach station for cell biology."

Stow quickly recruited some great local talent for the lab. some of whom are amazingly still around, and got going. Together, she and James began attracting other like-minded souls to join

from the American Heart Association.

#### Homeward bound...sort of

After 13 or so years in the US, Stow felt it was time to head home, and a Wellcome Trust Fellowship helped her do just that. "Getting the Wellcome fellowship was really a big thrill – it was my ticket back to Australia. I settled on Brisbane after David James contacted me and asked me to join him at the CMCB (now the IMB) at UQ. He had also recently returned from the US to establish a focus in cell biology there, and was looking for company. So, although Brisbane was kind of a strange destination for me (coming originally from Melbourne), it was one I chose rather deliberately." Cell biology was not well recognised as a field in Australia at that time (early 1990s) and research funding was very tied up with a lot of the big interest groups in Melbourne...and starting up a thing called cell bi-

them including Rob Parton from EMBL in 1996 and Alpha Yap from the US a couple of years later, and slowly the cell biology community at UQ started to take shape and to become known. "We built up new staff and equipment, and some renewed enthusiasm, for microscopy, as well as starting new courses to incorporate cell biology into the undergraduate curricula. So, very quickly we established ourselves a cell biology base, which was enormously important for the research to go ahead."

In her spare time, Stow acquired a faithful hound called Wally, untold number of cats, and set about discovering the wonders of camping, scuba diving and snorkelling off the Queensland coast with the University dive club. This led to discovering a guy called Doug, which led to wedding bells, which led to...well, a very little guy called Nick in late 1998. Thus, she also established a family base in Brisbane,

which was enormously important in keeping her sane in the non-work hours.

From then of course the CMCB grew and eventually morphed into the IMB, moving into a shiny new building in 2003. "Through the efforts of people like John Mattick, the institute has grown tremendously from its early days...but at just the right pace to keep most of us here, incorporate new people and grow the science accordingly," says Stow. "The cell biol-

ogy community has also grown enormously in Qld in the time I have been here and there is now a quorum of people and a vibrant, world-class scientific home here now and, Brisbane is still a great place to live."

David James did finally leave the Queensland cell biology family about 10 years later, heading south to a senior role at the Garvan Institute in Sydney, but still remembers those early CMCB challenges very well. "Jenny was a trail blazer. She gave up a fantastic spot at the MGH to move to Queensland, which at the time was a cell biology backwater. She saw the opportunity and knew she had what it needed to get the job done. Today, just 15 vears later, she is the Deputy Head of one of the most outstanding research institutes in the country and cell biology is one of the flagships of the organisation."



Work Xmas parties - a fun tradition in the Stow lab!

chinery that regulate cell polarity and endocytosis in polarized monolayers of epithelial cells, and more recently, in 3D cell culture cyst epithelia models. Stow sees this work on E-cadherin sorting and cell polarity progressing even further with new bioinformatic input available at the IMB and from the increasingly sophisticated capabilities in

role for recycling endosomes in the exocytosis of E-

cadherin using cutting-edge microscopy and image analysis of fixed and live cells. Continuing work in

this area focussed on the trafficking events and ma-

sophisticated capabilities in immunolocalisation and cell imaging.

#### Blast from the past

"Then about 10 years ago, I went even more crazy, and branched out to another cell type," Stow recalls. "Even though I had worked on epithelial-type cells for a very long time, they just didn't traffic enough of anything. So I started almost a side project looking at macrophages, which were actually the cells I had spent many hours of my PhD life staring at down an electron microscope." This work turned out to be a much bigger undertaking than Stow expected, mainly because, as she soon discovered, there was very little known about how macrophages traffic almost anything. It grew into a major focus in the lab, vield-

ing a whole new branch of study and reconnecting Stow with one of her earlier loves – immunology. This work progressed steadily and successfully, attracting national funding from NHMRC and a 5-year grant from the NIH.

Stow's macrophage research concentrated on tumor necrosis factor (TNF) and later, other inflammatory cytokines, as trafficking cargo. "The pathways involved in making sure these molecules get secreted from immune cells are fundamental to immune function and they lie at the heart of pathologies in inflammatory disease." The group's initial work involved characterising previously unknown aspects and machinery of TNF trafficking in macrophages, before going on to reveal new pathways and regulatory mechanisms for TNF secretion via recycling endosomes and phagocytic cups. "We subsequently discovered that subcompartments of recycling endosomes in immune cells regulate the differential sorting and selective secretion of cytokines. Papers coming out of this work were all highly cited and are widely recognised for making major changes in our basic knowledge of cell biology."

Rachael Murray was a crucial part of 'team macro-

#### New home...new directions

Scientifically in Brisbane, Stow continued with the G protein work and was immediately successful in securing NHMRC funding to do so. "It was also a time, however, when I did sit back and really think about what I was working on," she recalls. "I realised it was going to be more difficult to continue the renal stuff here - there were no readily available colleagues doing renal cell biology and clinical ties were hard to maintain with the campus being so far from the major teaching hospitals." So, a mixture of the circumstances and some new cell biologists appearing (eg. Alpha Yap) facilitated her decision to pick up trafficking a bit more broadly. "I kept working with post-Golgi trafficking in epithelial cells, but added E-cadherin as a membrane protein cargo to her soluble proteoglycans.

In 1999, Stow's group in collaboration with Yap made the key finding that epithelial adhesion is maintained by dynamic recycling of cadherin complexes, opening up a whole new project stream for the group in polarized trafficking. Later work also revealed a novel phage' in Stow's lab and remembers the heady days surrounding the key recycling endosome findings (published in Science). "Jenny has real passion and enthusiasm for science. Her brain seems to work on overtime and I am always amazed at the ideas she came up with. She is a fountain of knowledge and never ceased to surprise me with it." Murray now has her own group at The Children's Hospital in Sydney and values the lessons learned at the IMB with Stow. "Jenny is an excellent mentor to both her lab and other people in the department who can often be found in her office asking for career guidance and advice."

#### Stronger together

Collaborations with other cell biologists in Australia and with colleagues overseas have been very important to Stow's research here over the last 15 or

so years. In fact, a recent and particularly exciting advent for her group grew out of a longstanding collaboration with Paul Gleeson at Bio21 in Melbourne. "We have been able to manipulate a Golgi trafficking protein in a mouse and show for the first time that it really does affect TNF trafficking in vivo," Stow explains. "So we know now that when we mess around with trafficking in our cells in culture, it is translatable to something we can do in whole bodies." The potential of these findings for eventual applications in disease is very exciting – to circumvent trafficking that has gone awry or been affected by a genetic defect and to manipulate thing of diseases. "Paul was one of (in Stow's lab) ... the first people that I started collaborating with when I came

back to Australia – I knew of him already because he was one of the few people here with Golgi as a key word associated with their name. It has been a particularly enjoyable and successful collaboration."

The collaborative camaraderie and respect that Stow engenders is clearly mutual. According to Gleeson, "Jenny is a passionate advocate and driving force for the field of cell biology in Australia. She is also a very generous collaborator who is extremely willing to share new ideas and approaches – I always enjoy discussing science with her." Another Melbournian and long-time collaborator Christina Mitchell echoes those sentiments. "Jenny is a fabulous collaborator who is at the same time extremely honest and generous. This is a great combination as it means she will always be completely candid about the data and its interpretation, and then help you with her expertise and reagents to improve the story."

#### Keeping ahead of the pack

Something that Stow has always tried to do in her research is maintain the significance and scientific edge of her group's work, including investing in new technologies and expertise. Her group was one of the first in Australia to take up and promote livecell fluorescence microscopy in trafficking studies, whereby Stow guickly established a strong reputation and niche for her research. "Fluorescence imaging, especially of live cells and organisms has made it possible to really see and understand many cell processes that previously could only be inferred from 'test tube' experiments," Stow explains.

One of Stow's former PhD students, John lock, was responsible for setting up and mastering these new approaches in microscopy using equipment that Stow funded and brought to the IMB. It was Lock's

> cycling endosome as a novel route for post-Golgi exocytic trafficking to the cell surface. He is now a postdoctoral fellow at the Karolinska Institute in Stockholm, being recruited particularly for his microscopy knowledge and technological expertise. He credits Stow's forward thinking attitude in research for guiding his own career path. "I worked with Jenny for 6 years, as an undergraduate and then postgraduate student. I came to realise that what sets her apart from other scientists is her highly attuned strategic sense," Lock says. "Jenny was consistently aware of developing trends, and set about recruiting people and assembling capabilities to preempt these trends and stay in front of the pack. In

work that established the re-



like cytokines in a whole bunch ... And crazy people from around the world

the competitive environment of biological research, the ability to switch from the day to day minutia of research challenges to a strategic overview of important questions and powerful techniques is an enduring and vital talent that Jenny has in spades. It is also a living lesson from which her students can benefit immensely."

Stow takes her mentoring role in science very seriously and in turn gets a lot of reward from it. She consciously tries to impart the same sort of enthusiasm and awe of research to her own students and postdocs that she gained from her earlier training years, especially the importance of basic research that became part of her psyche at Yale and Harvard. "To make the biggest changes in medicine we have to make the biggest discoveries at the most fundamental level...and so while we work on very basic biology, its relevance to cancer, inflammation and other diseases is something that should never be far from what we want out of the work and from what it is really designed to do." She also offers some advice to any scientist just starting out on the road – "Be savvy about science – it's your business and you have to know how it works. Learn everything you can from those around you at every step in your career."

#### Taking up the lead and the challenge

These days Stow has to be particularly savvy about science herself with her dual leadership role as the head of a research group and Deputy Director of the IMB. "Actually, I don't think many people become scientists because they want to be a 'boss'," she says when asked about these roles. "It's the sort of thing that you wake up one day and find has happened to you. However, although it means a lot more red tape and a lot less looking down microscopes, it does give you the opportunity to tackle increasingly big and complex questions in science and to influence research on a larger scale." Stow adds that she certainly did not plan the Deputy Director move, but having been invited to take up the challenge, is attracted by the prospect of have an even broader impact on science - coming from the whole institute - than she could from her own research lab. "Again, I have looked back to my training years and the inspiration I took from the many people I encountered who could do things like this...take a whole institute like the IMB and really make it hum. That is my challenge."

In short, Stow has no regrets about her very early career-plan hiccup. "I love doing science as a career - doing something that is for the good of humanity, being in awe of nature, looking down a microscope at cells, having something to think about ALL the time and, most of all, working amongst a melting pot of interesting, smart, dedicated and crazy people from around the world (and that is just in her group)." What really keeps Stow going though is what she calls her 'main' job...that is, as a "dotty and doting wife and mother" to Doug, their son Nick and to several other, furry 'children', who together, complete her big picture.

Fiona Wylie

# ANZSCDB Young Investigator Award



### A/Prof. Katharina Gaus

Centre for Vascular Research

University of New South Wales

rina Gaus.

Gaus heads the Cellular Membrane Biology Lab in the Centre for Vascular Research (CVR) at UNSW. According to the website, her scientific quest is "to determine how specialised membrane domains organise signalling pathways." The main tool of choice for Gaus in this quest is microscopy, from simple flu-

### **KAT GAUS**

# Pushing the boundaries is all in a day's work

Advances over the past decade have pushed the spatial resolution of light microscopy to the limits of optical diffraction, according to the laws of physics. Now, the so-called "super-resolution" techniques are going around these theoretical limits into the sub-micron range, and in Australia, no comprehensive discussion of super-resolution microscopy is complete without mentioning this year's ANZSCDBI Young Investigator Award winner, Associate Professor Kathaorescence to a very sophisticated level with the new super-resolution techniques, which her group was the first in Australia to adopt. "To understand how the organisation of a membrane affects a certain process we have to actually image it as it is happening, and this makes high-resolution fluorescence microscopy the way to go."

Gaus started her life in science doing undergraduate mathematics and physics at the University of Heidelberg in Germany. However, she was soon heading further afield – to Cambridge University in the UK as an exchange student - and found herself making the switch to a more biological focus. "I discovered that maths and physics were a little bit too dry for me – my real fascination was always with the building blocks of life." Of course, it is hard to shake the past, and Gaus admits that her early academic background carries through to today and is extremely useful as she studies cell biology using biophysical techniques.

Gaus stayed on in Cambridge to do a Masters and PhD at the Institute of Biotechnology. During this time, she designed and developed a biosensor that could detect protein-protein interactions and a ligand library to look for lipoproteins as a diagnostic tool. "This was a fantastic project for me because it introduced me to many different aspects of cellular biology, and also to lipid-protein interactions. From then on, I became interested in the fundamental principles of how cells behave and how lipids play a role in that."

Moving to Australia happened 10 years ago when Gaus won a German postdoctoral fellowship to join the group of Wendy Jessup at the Heart Research Institute in Sydney. There she took on her "first real cell biology project", looking at lipid metabolism in macrophages. Two years later, Gaus successfully applied for an ARC grant and a postdoctoral fellowship, and away she went. It was also around this time that Gaus moved with Jessup's group to UNSW to join the Centre for Vascular Research (CVR).

#### Out on her own

In 2005, Gaus set up her own membrane biology group within the CVR after receiving a Career Development Award from the NHMRC, progressing to Senior Research Fellow in 2009. She is currently a chief investigator on several national and international grants including a 3-year grant from the Human Frontier Science Program to look at the structure and function of olfactory receptor neurons.

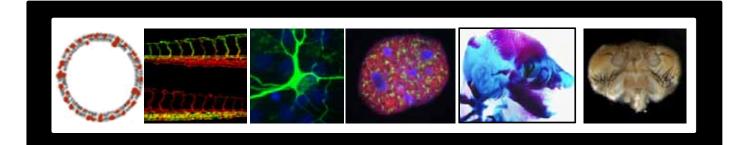
A current focus for Gaus is trying to understand how lipids and particularly dietary lipids affect T cell signalling during an infection, so linking immune dysfunction and obesity. "We think that we have worked out these molecular mechanisms in T cells in the laboratory, and are now taking it to an animal model. So, we feed mice a high-fat diet, take the T cells and basically look at them – and it seems that signalling processes are compromised."

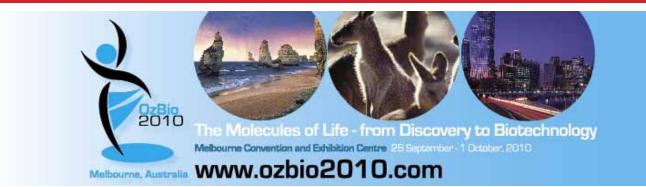
"The work on basic membrane biology and specifically, how lipids modulate the behaviours of signalling proteins continues. We still don't fully understand how membrane organisation affects the efficiency of signal transduction," Gaus says. "We have set up single-molecule imaging techniques to really get a handle on the molecular steps involved." The superresolution optical microscope that Gaus has in her lab is a PALM system (photoactivation localisation microscopy) developed by Zeiss. "By building up an image literally molecule by molecule, we can 'simply' count how many molecules take place in a signalling event and ask a whole new set of questions like is there a fixed ratio of receptors to kinases and how do lipids interfere with this delicate balance."

In terms of future goals, Gaus remains driven by understanding how a signalling cascade that has up to 100 proteins involved can be coordinated by something as simple as a membrane bilayer, and then to come up with some way of describing that process in its entirety rather than one protein at a time. "In a way we are looking for a new language to describe signal transduction. The model that I envisage of 'seeing' the cascade will obviously have to be based on empirical data and then driven by the intermolecular interactions...and our microscopy techniques will help us to quantify these interactions. So, if we understand how molecules behave, we can then zoom out to see it all in time and space."

"In a nutshell, my work really spans anything from the mathematical modelling of conceptual frameworks to single-molecule imaging to teach us about cell biology...and all for the end-of-life type aim of understanding the medical implications of obesity on T cell function"...easy!!

Fiona Wylie

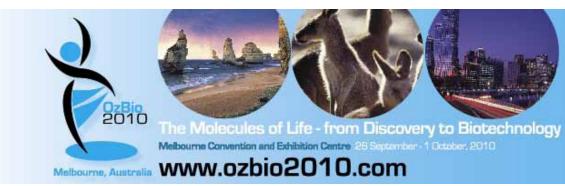


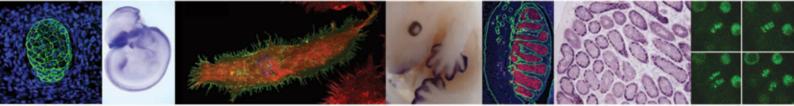


#### OzBio2010 - Program at a glance (subject to change)

Sunday <sup>3pm-7pm</sup> Monday	Registration										
-am	Registration & C	<b>)pening</b> (10.00 a	m)								
Plenary	Peter Doherty (A	lust)									
Plenary Lunch & Posters	Richard Harvey			Schroeder Botany Lecture	(USA)						
-pm Symposia 1	Stem cells & differentiation K. Loveland/A. Perkins Amander Clark (USA) Huck Hui Ng (Hong Kong) Gary Hime (Aust) +Selected from abstracts	Organelles T. Lithgow/K. Tru Heidi McBride (Can Naoko Imamoto (Ja +Selected from abs) +Selected from abs)	ada) pan) tracts	Plant me biology M. Gilliham/S Steve Tyerman Yong-Ling Ruan +Selected from +Selected from	. <i>Tyerman</i> (Aust) (Aust) abstracts	, 1 1	TBA +Selected f		A .Fox, Angus I John M Paul Fo	/P. Lewis Lamond (USA) attick (Aust) ster (Aust) oag (Aust)	Signaling across membranes M.Parker/T. Rapaport Angel Lopez (Aust) Steven Petrou (Aust) TBA +Selected from abstracts
Symposia 2	Abiotic stress: drought & salinity R. Munns /M. Tester John Mullet (USA) Stuart Roy (Aust) Lynne McIntyre (Aust) +Selected from abstracts	Cellular ima L. Tilley/B. Marsh Sam Hess (USA) Brad Marsh (Aust) Cynthia Whitchurch +Selected from abstr	(Aust)	Current adv neural reger J. Vickers/S. Duni James Fawcett (US Lindy Fitzgerald (A Tracey Dickson (Au +Selected from abs	neration <sup>(op</sup> (A) (ust) (st)	n i S E S T	n syste 5. Forrest/ Edward Ber Gean Grimn TBA	ng technolo ems biology (G. Spangenberg tram (Aust) nond (Aust) rom abstracts	gies	Immunology B. Heath/R. O'Heir Chris Goodnow (Aust) Fabienne Mackay (Aust) Lauren Ely (USA) Charles Mackay (Aust)	Signaling N. Harvey/T. Pawson Ralf Adams (Germany) Ben Hogan (Aust) Leigh Coultas (Aust) +Selected from abstracts
<b>Tuesday</b> -am	Welcome Mixer										
Plenary	Elisa Izaurralde	(Germany)	Nobo	ru Mizushin	<b>1a</b> (Japan	1)					
Plenary	Tom Rapaport (	JSA)	Nancy	<b>y Bonini</b> (USA	()						
Symposia 3	Apoptosis R. Kluck/C. Hawkins Richard Youle (USA) Ruth Kluck (Aust) Sharad Kumar (Aust)	Cytoskeleta networks A. Yap/E. Hardenn Ueli Aebi (Switzerlan Jake Baum (Aust) Greg Goodall (Aust)	an d)	Plant genet P. Langridge/P. G Scott Jackson (USA) Tim Sutton (Aust) Brett Ferguson (Aus	t)	NEU J. Got Kevin I Lars Iti Robert	Irodeg z/N. Boni Barnham (J ner (Aust) t Richards (	Aust) (Aust)	M .L Mana Natal TBA	otein-protein eractions awrence/U. Hartl ajit Hayer-Hartl (USA) lie Borg (Aust)	The transcriptome K.K. Khana/T. Gonda Huck Hui Ng (Hong Kong) Frances Shannon (Aust) Merlin Crossley (Aust)
Lunch & Posters -pm	+Selected from abstracts	+Selected from abstr	acts	+Selected from abst	racts I	David	Small (Aus	t)	+Sele	cted from abstracts	+Selected from abstracts
Symposia 4	Bioinformatics J. Whisstock/M. Bellgard Milton Saier (USA) Ashley Buckle (Aust) +Selected from abstracts +Selected from abstracts	Education H.E. Koon/S. Howitt Masatomo Maeda (Jap Peter Cartwright (Aust) TBA +Selected from abstract	SM P. W an) Marc Jean Juliar	genetics & all RNAs ( <i>aterhouse/T. Milla</i> : sots (Belgium) Finnegan (Aust) n Tonti-Filippini (Aust teted from abstracts	Mary-Eller TBA ) TBA	Ge urn/G n Harp	<i>. Cooney</i> per (Canada)	Helen MacNeil (C Kieren Harvey (A TBA	ation wgreen Canada) ust)	SYNAPSE P. Beart/N. Ip Nancy Ip (Hong Kong) Eric Klann (USA)	he Understanding enzymes <i>B. Kemp/A. Vrielink</i> Ivan Rayment (USA) John Whitehead (Aust) Paul Attwood (Aust) Hazel Holden (USA)
Symposia 5	R. Devenish/ N. Mizushima Sharon Tooze (UK) Richard Youle (USA) Mark Prescott (Aust)	Drug discovery & design . Street/H. Blanchard Billy Denny (NZ) Graeme Stevenson (Aust) Ashley Bush (Aust) -Selected from abstracts	inte R. And James Andy P Alex M	t-pathogen ractions dres/H. Drummer Paton (Aust) Poumbourios (Aust) aiter (Aust) ted from abstracts	Secreto pathwa J. Stow/R. J. Frances Brou Len Kritharie Peter Thorn +Selected fre	AY Murro dsky (l des (Au des (Aust)	ay USA) .ust)	Plant energ biology S. Smith/B. Pogso Ian Graham (UK) Bob Furbank (Aust) TBA +Selected from abs	on	Genes, develop ment & disease P. Koopman/C. Smith TBA Craig Smith (Aust) Carol Wicking (Aust) +Selected from abstracts	R. Hannon/J. Wilce Elisa Izaurralde (Germany) Tom Preiss (Aust) Rick Pearson (Aust)
Wednesday -am	Happy Hour										
Plenary	ANZSCDB President Lecture (TBA)			eter Goldacre e (TBA)	Award &	L					
Plenary	Greg Winter (UK) Biochemical Society/ASBMB [		Evan	Eichler (USA)							

Colloquia	Biotechnology Selected from abstracts	Cell bio develo			ne regula			int stems sted from abstracts	& fı	tein structure Inction	Signaling & disease Selected from abstracts
Lunch & Posters -pm	Selected non abstracts	Selected no		Jeice			Jeret		Scietta		
Plenary	Andrew Hill (Aust) ASBMB Merck Medal Lecture										
Symposia 6	Antibody & protein         Molecular bit           engineering         of cancer           D. Christ/G. Coia         A. Strasser/V. Dixit           Lutz Jermutus (UK)         Shigekazu Nagata (Jap           Greg Winter (UK)         Andreas Strasser (Auss)           George Kospidas         John Silke (Aust)		ancer asser/V. Dixit azu Nagata (Japan) as Strasser (Aust)	health D. Topping/S. Jobling an) Matthew Morell (Aust) bavid Topping (Aust) Melissa Fitzgerald (Philipp				Membran dynamics R. Parton/K. Ge Robert Nabi (USA Daniel Abanka (A Matthew Prior (A +Selected from a	aus A) Aust) Aust)	Stress signaling J. Atkin/T. Biden Randal Kaufman (USA) Julie Atkin (Aust) TBA +Selected from abstracts	RNAİ R. Dickins/K. Simpson Rod Bejersbergen (Netherlands) Patrick Humbert (Aust) Darren Saunders (Aust) +Selected from abstracts
Thursday -am	+Selected from abstracts	+Selec			+Selected Iro	in abstracts	•	+selected from a	DSUALIS		+Selected from abstracts
Plenary	Chris Somerville (USA) Brenda Schulman (USA)										
Plenary	Ulrich Hartl (Gerr	nany)	Vishva	a Dixi	t (USA)						
Symposia 7	Dynamic subcellul compartments T. Preiss/E. Izaurralde Archa Fox (Aust) David James (Aust) Kat Gaus (Aust) +Selected from abstracts	dev L. Rick Kozo H Chinfe Ben Er	Iral signaling elopment hards/K. Kaibuch (aibuchi (Japan) ii Chen (USA) mery (Aust) ited from abstracts	i	Patterning morphoge S. Dunwoodle/ Lee Niswander Ruth Arkell (Aus Heather Young +Selected from	ENESIS (L. Niswand (USA) (St) (Aust)	er P. Shi Joh	ant-microbe teractions Dodds/B. Howlet auna Somerville (US nud Bernoux (Aust) in Rathjen (Aust) elected from abstra	t M. GA) Carl Char Brid	Dtein-nucleic aci eractions Wilce/J. Matthews os Barbas (USA) die Bond (Aust) get Mabbutt (Aust) ected from abstracts	d Ubiquitin and Ub- like modifications <i>S. Kumar/B. Schulman</i> David Komander (UK) Catherine Day (NZ) Sandra Nicholson (Aust) +Selected from abstracts
Lunch & Posters - <b>pm</b> Plenary	Johann Deisenho	ofer (USA)	Lee Ni	iswar	nder (USA)						
Symposia 8	Haematopoiesis: Functional systems approaches genomics B. Kile/R. Ramsay F. Shannon/L. O'Conn Harvey Lodish (USA) Tom Gonda (Aust) Doug Hilton (Aust) Barry Pogson (Aust) Carola Vinuesa (Aust) +Selected from abstract		mics non/L. O'Connor nda (Aust) gson (Aust)	David Smyth (Aust) Christine Beveridge (Aust) Anna Koltunow (Aust)			D. Gardner/J B. Mabbutt/M. Hinds TBA Paul Robson ( TBA TBA TBA TBA			onic development	Subcellular targeting D. Jans/N. Imamoto Yoshihiro Yoneda (Japan) Shige Noshimura (Japan) Beric Henderson (Aust) Heeleted from abstracts
Friday	Society Annual G Conference dinn		<b>1</b> eetings								
-am Plenary	Tony Pawson (Ca	nada)	Peter	Wate	erhouse (/	Aust)					
Symposia 9	Biomarkers N. Hoogenraad/B. Cocks Richard Simpson (Aust) Tim Green (Aust) TBA +Selected from abstracts	3. Cocks & polarity S. Russell/N. Waterhouse ust) Minoru Yoshida (Japan) Wolfgang Weninger (Aust) Megan Fabbro (Aust)		Meural function       M         M. Morris/A. Gundlach       M         Alon Chen (Israel)       M         Savin McNally (Aust)       T         Margaret Morris (Aust)       T		h Aust N. Cou Matt V TBA TBA	<i>N. Cowieson</i> Matt Wilce (Aust) TBA TBA		G. / G. / Chri Vine Mik	ONE Wall mposium Fincher/F. Pettolino is Somerville (USA) cent Bulone (Sweden) e Gidley (Aust) lected from abstracts	Yeast biology A. Munn/I. Dawes Charlie Boone (Canada) Marc Wilkins (Aust) Snezhana Olifernko (Singapore) +Selected from abstracts
Plenary	Harvey Lodish (USA) Jian-Kang Zhu (USA)										
-pm	Awards & close of main program (1pm)										
Special Workshops (2-6pm)	Biotechnology Speakers: TBA Careers Speakers: TBA			 s (/	Education: Enquiry-based learning Speakers: Trevor Anderson (Sth Africa); Pauline Ross (Aust); Kristine Elliott, Helen Irving (Aust), Liz Johnson (Aust); Janet Macauley (Aust)				Super-resolution Microscopy Speakers: Sam Hess (USA); Guy Cox (Aust); Cynthia Whitchurch (Aust); Trevor Smith (Aust); Kat Gaus (Aust); Leann Tilley (Aust); Min Gu (Aust)		





# October 22 2010 9 - 5:30pm Cell & Developmental Biology

Institute for Molecular Bioscience The University of Queensland St Lucia, Australia Meeting

**Professor Stephen Cohen**, Temasek Life Sciences Laboratory Exploring growth regulation and metabolism

**Professor Jeff Hardin**, University of Wisconsin Mechanisms of Morphogenesis

# **Dr Ian Smyth**, Monash University Developmental Biology of the Skin

6 postdocs and 6 PhD students will be chosen from abstracts to present a 15 minute talk: PRIZES WILL BE AWARDED TO THE BEST TALKS AND POSTERS

please submit abstracts of up to 200 words by September 10th to either:

Dagmar Wilhelm: d.wilhelm@imb.uq.edu.au Eva Kovacs: e.kovacs@imb.uq.edu.au Kelly Smith: k.smith@imb.uq.edu.au

### Registration is FREE - please do so by emailing Dagmar, Eva, or Kelly

Refreshments and prizes provided by our sponsors: for further information please visit www.imb.uq.edu.au-bcdbm







# 6th Australian Developmental Biology Workshop Yarra Valley, Vic. 24th-27th Oct. 2010.

## **Guest Speakers:**

### Margaret Buckingham (Institute Pasteur, France)

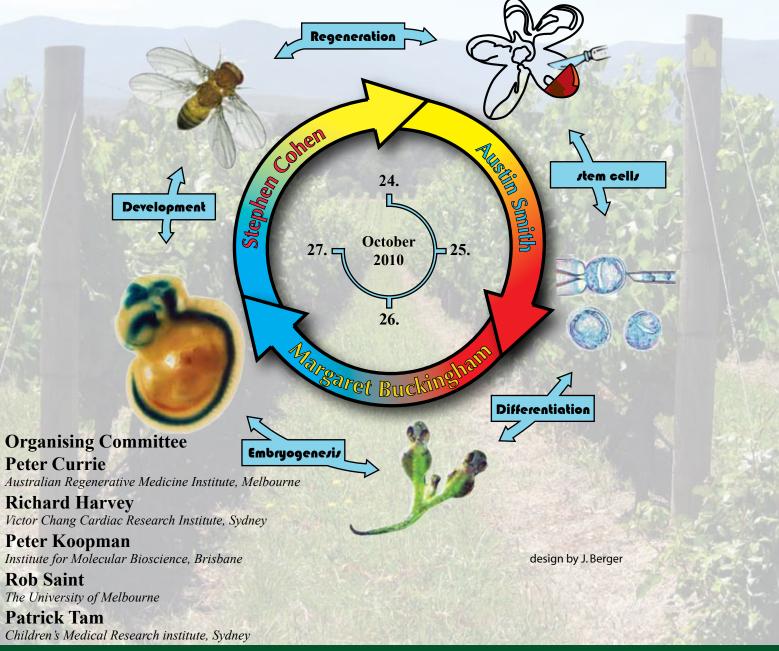
An expert in the application of mouse biology to dissection of genetic control of skeletal and cardiac muscle formation. Her laboratory has identified key regulatory genes and cellular processes that act in muscle development.

#### Stephen Cohen (Institute of Molecular and Cell Biology, A\*Star, Singapore)

A renowned Drosophila developmental geneticist, who undertook ground-breaking studies in the field of microRNAs with universal impact on understanding gene regulation.

#### Austin Smith (Wellcome Centre of Stem Cell Research, University of Cambridge, U.K.)

A world leader in the area of embryonic stem cell research. His laboratory has characterised many of the cellular and molecular mechanisms governing formation, self-renewal and differentiaion of pluripotent and tissue-restricted stem cells.



The aim workshop aims to promote Developmental Biology by providing students and scientists that are activelyworking in this research field with the chance to gain knowledge and practical skills. It is an intimate and highly interactive meeting that allows unparalleled access to world leaders in Developmental Biology for scientists at all levels of their career.

The workshop will consist of in-depth plenary research talks by high profile international speakers, round table discussions on experimental approaches, analysis of model systems, and broad career issues for Developmental Biologists. The number of participants is strictly limited to 24. A mix of students, post docs, and lab leaders will be selected on the basis of their demonstrated interest in Developmental Biology, track record, and the perceived benefit they will obtain from the workshop.

Applications must consist of the applicant's CV (max three pages) and a one page cover letter addressing the above criteria. Please email your application to applications@armi.monash.edu.au. Applications close 31 July 2010. For further information visithttp://www.armi.org.au/About Us/news/Whats On/devbiowkshp.aspx

WHEN DECEMBER 11–15, 2010 WHERE Philadelphia, PA Pennsylvania Convention Center



THE AMERICAN SOCIETY FOR

CELL BIOLOGY

# DON'T MISS IT!

#### WHY • Showcases the most exciting cell biology research

- Offers scientific breadth, depth, and multiple discussion opportunities
- Provides tested **teaching strategies** you can use today
- Builds careers with sessions on funding, job options, strategies, networking, and more
- WHAT Learn about Improving Cancer Chemotherapy
  - Investigate Cytoskeletal Dynamics
  - Consider Cell Biology to Therapeutics
  - Explore In Vivo Imaging
  - HOW Submit your abstract for poster or oral presentation consideration by July 29
    - Suggest a member-organized special interest subgroup by July 29
    - Apply for travel and childcare awards by September 1
    - Take advantage of discounted registration fees by October 7



# All meeting forms and special hotel rate information will be available on the ASCB website, www.ascb.org/meetings. Or email ascbinfo@ascb.org or call 301-347-9300. The ASCB is *your* community, sharing your values. **JOIN US!**

#### Official ASCB Housing Company

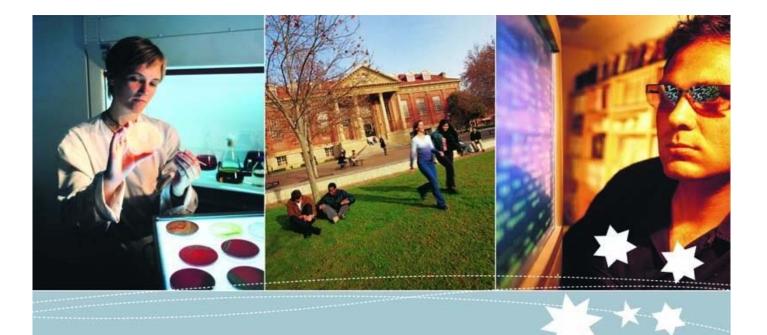




## School of Molecular & Biomedical Science SMBS



# *Considering a PhD in Molecular Biology or Biomedical Science in 2010 or 2011?*



We will fly you to Adelaide to learn about opportunities for PhD study in the SMBS

We will pay airfare and accommodation costs for potential students to visit Adelaide for up to three nights to learn about our research, PhD programs and scholarship opportunities (including relocation assistance), to meet potential supervisors and to sample our fantastic lifestyle. This scheme is open to Australian and New Zealand citizens and Australian permanent residents.

The School of Molecular and Biomedical Science is one of Australia's leading clusters of researchers in the life sciences with a unique collection of advanced facilities and capabilities. We offer PhD programs in Biochemistry, Genetics and Microbiology & Immunology. In the past 5 years all RTS-eligible\* students accepted into a PhD program in the School have been awarded a scholarship that provides full remission of tuition fees and a living allowance currently AUD\$22,500 tax free per annum in 2010.

Scholarships may be available throughout 2010. Applications can be made at any time.

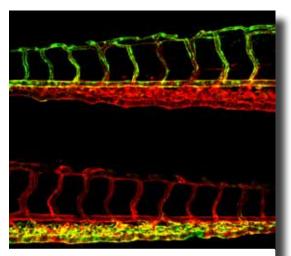
For more information on this scheme, the research areas offered by our School and to download an application form visit the website below.

(Note: You are free to extend your stay in Adelaide at your own expense.)

\* See the website below for more information

### www.adelaide.edu.au/mbs/prospective/pg/visit

## Winners of the "Call for Images for the Website"

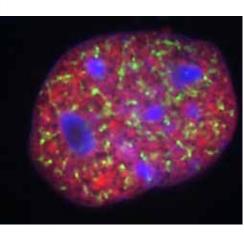


#### 1 Benjamin Hogan

Molecular Genetics of Vascular Development Unit

Institute for Molecular Bioscience, The University of Queensland

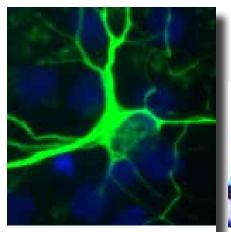
Description: Composite of two live, double transgenic zebrafish tails with complementary fluorescent labeling of embryonic arteries and veins.



#### 3 Jocelyn Widagdo

Neuromuscular & Regenerative Medicine Unit School of Medical Sciences, The University of New South Wales Description: Nuclear localisations of the Williams Syndrome-related protein, GTF2IRD1 (green) and a histone lysine methyltransferase (red) were analysed by co-expression in COS-

7 cells (DAPI, blue). GTF2IRD1, our protein of interest, was shown to interact with the histone lysine methyltransferase through a yeast library screen, implicating GTF2IRD1 in histone/chromatin remodelling and suggesting a possible mechanism for pathologies of Williams Syndrome, a human neurodevelopmental disorder.



**2 Dana Bradford** Neural Migration Laboratory Queensland Brain Institute, The University of Queensland Description: Cultured from the neurogenic region of the adult mouse brain, this differentiated neurosphere is stained to show cell nuclei (DAPI, blue) and immunolabeled with a marker for early neurons (TuJ1, green) showing new neurons sending out processes to explore the environment.



**4 Vicki Metzis** Developmental Genes and Human Disease Unit Institute for Molecular Bioscience, The University of Queensland Description: Skeletal preparation of a prenatal mouse embryo showing the bone (Alizarin Red) and cartilage (alcian blue) of the craniofacial skeleton.



**5 Felix Grusche** Cell Cycle & Development Unit Peter MacCallum Cancer Centre,

Description: The Salvador-Warts-Hippo (SWH) pathway controls tissue growth during development. Partial loss of Expanded in Drosophila, a negative regulator of the SWH pathway, results in tissue overgrowth and the transformation of some eye tissue into excess cuticle and hairs.

### 16th NSW Cell and Developmental Biology Meeting

#### **THOMAS FATH and BILL PHILLIPS**

This year the 16th Annual New South Wales Cell Developmental Biology Meeting was held as a jointly sponsored meeting by ANZSCDB and ASBMB at the University of New South Wales. The meeting focused on Systems Biology and Cell Polarity and was very well received with over 100 attendees from universities and institutes within NSW and the ACT, and we especially welcomed the attendance by Alpha Yap, former President of ANZSCDB and convenor of the 2010 Hunter Cell Biology Meeting. We were delighted that Ed Munro from the University of Chicago could join us as plenary speaker at this year's meeting whose talk entitled "Dynamics of cell polarization in C. elegans: Models and experiments" reflected an impressive comprehensive approach to cell polarity utilizing fundamentals of developmental biology, cytoskeletal dynamics and mathematical modeling. Ed's participation in this meeting was made possible by the ANZSCDB as part of their visiting fellows program that sponsors an international plenary speaker at the Hunter Cell Biology Meeting with associated visits to institutions within Australia. Invited plenary speakers Miles Davenport (Centre for Vascular Research, Lowy Centre, UNSW) gave a challenging introduction to Systems Biology by discussing the dynamics of chronic and acute infectious diseases in his talk entitled "Complex Systems - from the molecule to the disease" and Patrick Humbert (Peter Mac-Callum Cancer Centre, Melbourne) shared his latest research outcome from a large scale RNAi screen on polarity genes with us in his talk entitled "Building tissues and moving them: Cell polarity genes in migration and invasion". Further talks were given by Vladimir Sytnyk (School of Biotechnology and Biomolecular Sciences, UNSW) and PhD students and postdocs from institutions around Sydney including the Victor Chang Cardiac Research Institute, the Children's Medical Research Institute, the Children's Cancer Institute Australia for Medical Research, the University of Sydney, the Centre for Vascular Research and the University of New South Wales. For the first time prizes for the best PhD and postdoc presentations were introduced which were sponsored by ANZSCDB and ASBMB. The ANZSCDB prize for best PhD presentation was awarded to Nikki Curthoys (UNSW/USyd) and the best postdoc prize to Romaric Bouveret (Victor Chang Cardiac Research Institute). We also want to thank Invitrogen and Qiagen for their generous and continued support of the meeting.

#### Thomas Fath and Bill Phillips

Reports

#### 2010 Science meets Parliament

#### **ALEKSANDRA FILIPOVSKA**

On March 9th and 10th the Federation of Australian Scientific and Technological Societies (FASTS) held the 11th Science meets Parliament (SmP) in Canberra. Marie Bogoyevitch and I attended as representatives of ANZSCDB and had the opportunity to participate in a range of workshops and debates on the importance of science representation in parliament and government. This was the first time I had heard of FASTS, which is an organization formed 25 years ago because it was felt that the voices and opinions of scientists were lacking in government. The main consensus over the years was that there was a language barrier between scientists and government officials; the scientists use jargon and too much detail and the government officials are interested in immediate outcomes that would positively affect their imminent need for re-election. One of the ways FASTS facilitates the communication between science and government is by organizing an annual two-day SmP meeting. During the first day the scientists take part in a series of workshops that coach them to communicate their concerns and thoughts to parliamentarians during their meetings which take place on the second day of SmP.



Delegates gather at the Great Hall, Parliament House, for dinner with MPs and science representatives.

Even at the start of the meeting it was obvious to most of us scientists that voicing our concerns about the current state of science to government representatives was going to be challenging. We were quizzed on our understanding of the structure of government and the role of different government departments, the Australian people and the media, which was perceived very differently by most groups. We had media presentations from television, radio and newspaper journalists and gained more insight into how news is delivered to the public. During this presenta- people to enter science professions, but more importion it was highlighted that there is a need for individ- tantly to keep them in their positions by providing job

uals who can bridge communication the gap between scientists and the public, and the media could be a potential facilitator in this process. Although there are many journalists that are interested in taking on such positions, the scientists had concerns about the sensationalism aspect of the media that often misrepresents the scientific discoveries to the public. The debate between the scientists and the journalists was like a

tug of war. The scientists maintained that without the science there would not be any stories; however, the journalists pointed out that without them scientific discoveries would unnoticed and go as a consequence the government will not be aware of the progress and significance of scientific discoveries.

Our main concern as scientists was engaging the parliamentar-



Delegates meet Senator Kim Carr, Minister for Innovation, Industry, Science and Research



**Delegates of SmP2010** 

ians in our research interest with the ultimate goal of sage that came from him was that scientists don't supporting scientific endeavour and basic research. need to sacrifice their integrity to counter the misin-We were advised by several government officials, public servants and political advisors with a wealth of experience in dealing with politicians in three different workshops. The main message that came from all of them was to maintain the interest of the senators and members of parliament by having a succinct story with a practical outcome. We had an hour to practise our pitch for the parliamentarians after which we were taken to Parliament for our dinner, which gave us the opportunity to chat informally to some of the government officials. The following day included meetings with members of Parliament and senators. I had a meeting with an MP from the Liberal party from Western Australia who had an interest in biomedical research and supporting women in science. We discussed the need to maintain a Photo credits: Lorna Sim/FAST vigorous, innovative and competitive research environment, which depends on encouraging talented

formation that floods the media daily.

The SmP experience was very interesting and gave me an insight into how science and research are seen by the government and our public representatives in Parliament. The FASTS is very well organised and run by committed board members and executives. The main goals of FASTS and the SmP meetings are to reinforce the idea that we as scientists have to tailor our messages according to the different audiences if we want our voices to be heard, much like the politicians when they need our votes.

#### Aleksandra Filipovska

Western Australian Institute for Medical Research

security. We discussed the need to support basic research. which has been the most reliable source of knowledge that has led to important discoveries, furthered technology and medicine. which ultimately leads to economic growth, sustainability and leadership.

The lunchtime meeting at the National Press Club was one of the highlights of SmP. The keynote speaker was the author and science writer Chris Mooney who gave a realistic view of the state of science in the United States. He discussed the need for scientists to speak out and put an end to anti-scientific campaigns spurred on by misinformed members of the public. Perhaps the most important mes-

### Student members to attend the 60th Meeting of Nobel Laureates in Lindau, Germany

Congratulations to Duncan Mortimer (Queensland Brain Institute) and Denise Miles (Murdoch Childrens Research Institute) for being chosen to attend the 60th meeting of Nobel Laureates in Lindau, Germany from June 27 to July 2, 2010. As a corresponding society of the National Committee for Biomedical Sciences of the Australian Academy of Science, the ANZSCB Executive was able to nominate postgraduate member applicants to be considered for this honour. The Council for Lindau Nobel Laureate Meetings then selected 13 candidates from all corresponding Academies of Science with an even representation across the disciplines of Medicine or Physiology, Chemistry and Physics. Prior to the Lindau meeting, Duncan and Denise will attend Science at the Shine Dome, the annual celebration of science held at the Academy of Science, Canberra, 5-7 May 2010. Expenses are paid by the Australian Academy of Science and the Lindau Committee. Well done!

(taken from http://www.science.org.au/internat/ lindau.html)

The 2010 meeting of Nobel Laureates in Lindau is a multi-disciplinary meeting dedicated to chemistry, physics and medicine or physiology. With the support of the Academy, thirteen of Australia's brightest early-career researchers will be attending the meeting to be held in Lindau Germany, from 27 June to 2 July. The group will be accompanied by Professor Kurt Lambeck FAA. Not only will this meeting provide the participants with the opportunity to meet and hear from Nobel laureates, it will also provide a fantastic opportunity for our early-career researchers to meet, discuss ideas and to potentially form beneficial relationships with peers from across the globe.

Initiated in 1950 as a European conference of leading scientists in medicine, the guiding principle of the first Lindau Congress of Nobel Laureates was to encourage international scientific dialogue with Nobel Laureates. The first congress of medical specialists has since evolved into open-minded encounters between Laureates, young researchers and students in other fields of the natural sciences: for over half a century now, Nobel Prize winners in medicine/physiology, chemistry and physics convene, alternating, for one week at the end of June each year at Lindau, Germany.

Usually about twenty Nobel Prize winners attend, to share insights into their life's work and to interact with 600 outstanding young researchers and graduate students from around the world (up until now mainly from Western Europe, but increasingly in recent years from the USA, Central America, Latin America, Asia, the Middle East, Russia, Eastern Europe Africa

#### and Australia).

In line with the initiators' concept of building bridges between nations and cultures, the Lindau forum offers the upcoming generation of scientists an opportunity to instigate and foster what can be called a 'Dialogue among Cultures', both on scientific issues and on a more peaceful future. The bridges they build have a lasting impact beyond the Meetings. In the course of their personal encounters, participants jointly develop new ideas, and create networks of cooperation and international friendship, to the benefit of scientific progress.



Left: Duncan Mortimer (Queensland Brain Institute) Right: Denise Miles (Murdoch Childrens Research Institute)

### THANKS TO OUR SUSTAINED SPONSORS

SIGMA Aldrich PTY LTD sponsors of the ANZSCDB President's Medal

Carl Zeiss sponsors of the YIA

Animal Resources Centre Australian Proteome Systems Beckman Coulter Australia BioScientific Millipore Australia Fronine Laboratory Supplies Geneworks Interpath Services Merck Australia Olympus Australia Promega Corporation Australia

#### The Hunter Meeting

#### **ALPHA YAP**

The 10th anniversary edition of the Hunter Meeting was held in the bucolic surrounds of the Sebel-Kirkton Park between March 16 and 19th. As in earlier editions, the aim of the meeting was to provide a forum for cutting edge work in the country (both through presentations and informal discussions) and a dialogue with international experts (a record 11 invited speakers this year). Preceded by an afternoonlong workshop on Imaging, the main meeting began with the now-traditional Keith Stanley Lecture. In her lecture, Sandy Schmid (Scripps Institute) provided a stirring overview of her decades-long efforts to understand the molecular and cellular basis of endocytosis. Illustrating the inter-disciplinary philosophy of the Hunter Meeting, Schmid outlined how her research has moved freely between cellular and detailed structural mechanisms. It provided a scintillating start to the whole meeting, some of the highlights of which are now summarized by the session chairs.

#### Biology/pathology of the vascular system.

Like many others in the meeting, this session featured a series of complementary talks from different disciplines. Martin Schwartz (University of Virginia) spoke about the mechanical forces that are generated within endothelial cells and how such forces are transduced in vascular biology. In a tour de force of state of the art of cell biology his talk showcased a number of novel approaches to assay intracellular forces using protein sensors of force and imaging techniques. Richard Harvey (The Victor Chang Cardiac Research Institute) spoke of his evaluation of stem cells in the mouse heart. He detailed an interesting vascular-associated cell type, which demonstrated clonability and the ability to generate cardiospheres in vivo, that shared many characteristics with pericytes, but potentially acted as a pericyte progenitor population. Ben Hogan (IMB Queensland) talked on the use of zebrafish to study development of the lymphatic system. He detailed an interesting zebrafish mutant that disrupted the ccbe1 gene, which he also showed was mutated in a human lymphatic disease. Madhavi Maddugoda (INSERM, Nice) spoke on the role of membrane dynamics during transendothelial permeability, revealing novel pathways that cells use to regulate membrane dynamics. (Chair: Peter Currie).

#### Protein transport in health and disease

William (Bill) Balch (Scripps Research Institute, La Jolla) talked about the importance of protein homeostasis (which he calls proteostasis) in health and disease. Bill discussed how the proteostasis network is required for proper protein folding, removal of misfolded proteins and for protein transport. Disruption of the proteostasis network can thus potentially result in disease. He then discussed how small molecules that regulate proteostasis can be useful therapeutics in a wide range of folding diseases. Continuing the protein folding theme, John Bateman (Murdoch Children's Research Institute, Melbourne) discussed the role of endoplasmic reticulum (ER) stress in the pathophysiology of skeletal dysplasia. He presented his elegant studies to show how ER stress, caused by mutant misfolded extracellular matrix (ECM) proteins, contributes to the molecular pathology of disease. Peter Thorn (School of Biomedical Sciences, University of Queensland) then presented his elegant work on granule fusion with the cell membrane using live imaging with 2-photon microscopy and confocal microscopy following tissue fixation. Using various different dyes Peter and colleagues have developed methods to enable identification of whether the fusion pores are open or closed. He showed that secretion is not only regulated by the control of the numbers of fusion events but also by the dynamics of granule fusion itself. Antony Cooper (Garvan Institute of Medical Research, Sydney) talked about the role of synuclein in Parkinson's disease (PD). Synuclein associates with synaptic vesicles, its mutations and allele duplication result in early onset PD and it is the main constituent of Lewy bodies. (Chair: Sharad Kumar)

#### **Quantitative Biology**

This session focused on the potential for quantitative and computational approaches to contribute to analysing complex biological phenomena. Ed Munro (University of Chicago) discussed his lab's ongoing efforts to dissect the complex process of endoderm invagination, a key early morphogenetic process in metazoa. Using ascidian embryos, his lab showed that invagination involved complex changes in cell shape, encompassing both apical constriction and shortening of the cell bodies. Computational modelling allowed his group to make testable predictions about the roles of Myosin and Rho signaling in these processes. Geoff Goodhill (University of Queensland) described experimental and computational approaches to studying axonal chemotaxis. Remarkably, axonal growth cones sense very subtle gradients and this is often assumed to involve axonal turning. Instead, he showed that modulation of growth rates may be key when axons respond to shallow gradients. Katerina Gaus (University of NSW) discussed the use of super-resolution imaging and quantitative analysis of signal cluster formation to understand how probabilistic elements contribute to T-cell signaling. (Chair: Alpha Yap)

Ed Munro's participation in the Hunter Meeting was generously supported by the Australia and New Zealand Society for Cell and Developmental Biology. We thank the ANZSCDB for their ongoing support of the Hunter meeting.

#### Cell division and polarity

This session focused on the regulation of cell proliferation, particularly highlighting the importance of cell polarity regulators and the regulation of the mitotic spindle in cell division. Cayetano Gonzalez (Institute for Research in Biomedicine, Barcelona, Spain) spoke about the importance of the microtubule cytoskeleton in defining the plane of cell division in Drosophila neural stem cells (neuroblasts) of the developing brain. Using sophisticated live cell imaging, he showed how the plane of division was mostly randomized if a microtubule-based cue was disrupted, but when reestablished a new plane of division was established at an axis parallel to the newly born daughter cell. This study revealed a potential role for the newly born daughter as an external cue defining the plane of division. Current research in the Gonzalez lab will focus on determining the developmental consequences of altering the plane of division of the neural stem cells. Kum Kum Khanna (Queensland Institute of Medical Research, Brisbane) discussed the DNA damage checkpoint in mammalian cells, focusing on the critical regulator ATM. Kum Kum also described a novel centrosomal protein (Centrobin) that has a role in stabilizing the microtubule structure and in centrosome integrity during mitosis. Patrick Humbert (Peter MacCallum Cancer Center, Melbourne), described the importance of the polarity regulator, Scribble, in cooperative tumourigenesis with Ras signaling in human epithe lial cells and mouse models, particularly focusing on the outcome of a high throughput shRNA screen. Using new generation sequencing and systems biology analysis, this screen revealed novel pathways, including asymmetric cell division regulators, involved in Scribble mediated tumourigenesis in human epithelial cells. (Chair: Helena Richardson)

#### **Signaling and cancer**

The presentations in this session approached the problem of understanding cancer from many perspectives. Margaret Frame (University of Edinburgh) gave a wide-ranging talk that encompassed the roles for integrin signaling through Src and FAK; the impact of drug inhibitors on tumor progression and invasion; and the use of intravital imaging to characterize tumor cell invasion and its responsiveness to therapy. Mathias Ernst (Ludwig Institute for Cancer Research, Melbourne) discussed the relationship between inflammatory cytokines and tumor progression, using a range of mouse models that his lab has developed. Sharad Kumar (Center for Cancer Biology, Adelaide) approached the problem from the perspective of regulated cell death. He described studies in Drosophila that identified a caspase-independent autophagic mechanism for regulated cell death in tissues. Finally, Terry Kwok-Schuelein (Monash University) took us back to integrin signaling, to illustrate how this process is co-opted by pathogenic products of Helicobacter pylori, a known risk factor for gastric cancer. (Chair: Alpha Yap)

Building tissues, preserving tissues The session presented us with three quite different approaches to understanding principles of tissue structure. The underlying theme which emerged was the diversity of roles to which the cytoskeleton has been recruited in the process of morphogenesis. Our international speaker, Mark Peifer (University of North Carolina), presented his elegant work using Drosophila to dissect early processes involved in establishing the body plan with particular reference to the role of the cytoskeleton. Patrick Tam (CMRI, Sydney) described the role of WNT signalling in head morphogenesis in the mouse embryo. He also introduced the role of rhou and its effects on the actin cytoskeleton as an important regulator in early embryogenesis. Helena Richardson (Peter MacCallum Cancer Center, Melbourne) returned us to Drosophila and the integrated roles of polarity regulators in controlling decision making during morphogenesis. The session served to emphasise the power of these two developmental model systems and the increased sophistication of experimental approaches to fundamental questions regarding the building of the body plan. (Chair: Peter Gunning)

#### The EMBO Lecture

A consistent highlight of recent Hunter Meetings has been the plenary lecture sponsored by the EMBO World Programme. 2010 was no exception. This year's EMBO Lecture was presented by Michael Way (London Research Institute, CRUK) who was, indeed, the international speaker at the first Hunter Meeting. Michael gave the meeting a panoramic overview of his lab's work on understanding how cellular biology is co-opted by pathogenic vaccinia viruses. In the process, he illustrated both the ingenious mechanisms that these viruses use to co-opt host machinery, and also how much insight into cellular mechanism comes from studying how pathogens interact with their host cells. We thank the EMBO World Program for their support of the Hunter Meeting. (Chair: Alpha Yap)

#### Cell origin, cell fate

This session addressed key issues of signaling and cell migration in development. Anna-Katerina Hadjantonakis (Memorial Sloan-Kettering Cancer Center, New York) demonstrated that the gastrulating mouse embryo is no longer refractory to live imaging. Using sophisticated genetic manipulation, ex utero embryo culture and microscopy, her group has shown that the visceral endoderm is dispersed by the addition of definitive endoderm directly from the progenitor epiblast. These findings impact on our understanding of how the digestive and respiratory tracts form. John McAvoy (Save Sight Institute, Sydney) described how planar cell polarity (PCP) signaling polarizes fiber cells of the lens using genetically modified mouse models and lens explants. He showed that the asymmetric distribution of cilia and PCP proteins polarize lens fibers allowing their alignment and orientation to lens poles. Martin Lackmann (Monash Universi-

ty, Melbourne) used fluorescent resonance energy transfer (FRET) and Fluorescent lifetime imaging microscopy (FLIM) to determine how Eph receptors switch from promoting cell-cell adhesion to cell signaling. He demonstrated that the A-Disintegrin-And-Metalloprotease (ADAM) releases Eph receptor ligands and thus provides the switch between cellcell adhesion and signaling. Importantly, he showed that the membrane-proximal localisation of the latent kinase domain prevents ephrin ligand shedding in trans. Pamela Stanley (Albert Einstein College of Medicine, New York) described the essential role of O-fucose glycans on Notch signaling. Notch signaling is required to direct cell fate during embryonic development, and receptors lacking O-fucose are unable to signal. She discussed how that glycosyltransferase required for O-fucosylation of the Notch receptor also plays a role in chaperoning the receptor to the cell surface. (Chair: Sally Dunwoodie)

We hope that these vingnettes illustrate the diverse scientific range that distinguished Hunter 2010, as it has earlier meetings. They cannot give any sense of the energy and enthusiasm that pervaded the poster sessions, trade meetings, and informal discussions over coffee, hikes, and just sitting in the sun. We hope that you will experience that directly at Hunter 2011 (which will be held from March 22-25, convened by Phil Robinson and Peter Gunning).



### ANZSCDB and the Internation- ity is to co-sponsor/co-organise 3 regional meetings al Society for Developmental **Biology**

In 2009, the ANZSCDB became a member society of We would also like to continue to provide small grants the International Society for Developmental Biology. The Executive decided that this elevation in membership status was in keeping with the prominence of Developmental Biology in the Asia-Pacific region and the role that our Society has in promoting this discipline. We have received a letter from the President of the ISDB, Claudio Stern, describing the activities of the ISDB and calling on us to provide feedback. Please do take this opportunity to respond either directly to Claudio or you can respond to a member of the ANZSCDB Executive.

#### Dear colleague,

As you may know, I was elected President of the In-ternational Society for Developmental Biology ISDB in Edinburgh last September and started on the job in January this year. I do not intend to make it a habit via your national society. We will consider all sug-to send emails like this but feel that it is important to gestions and I will do my best to acknowledge every re-introduce the ISDB briefly and to give individual de-relation of the children of the second seco velopmental biologists an opportunity to have a say in what we can do for you and the community of developmental biologists over the next 4 years and beyond. Claudio Stern

As you probably know, the ISDB is not strictly a society but rather a federation of national and regional societies (see below for a list); you are receiving this email because you are a member of the ANZCDB. The main mission of the ISDB is to further interactions between developmental biologists on a wider, more international level than is possible through local societies. It has done this mainly through its meeting which occurs once every 4 years. The last meeting was in Edin-burgh in 2009 and the next will be in Cancun, Mexico, in 2013. These meetings are very successful, typically attended by more than 1000 participants and the atmosphere is very lively. In addition to the main meeting the ISDB has been co-sponsoring other occasional meetings on an ad-hoc basis but in my opinion this has been less successful because few people knew The current officers of the ISDB are: about it and because there is not enough funding.

The ISDB is funded from two sources. One is an annual contribution from Elsevier arising from the royalties of the journal "Mechanisms of Development" (MOD), which is the official journal of the ISDB. The other source is an almost nominal capitation fee payable by the member societies. The ISDB is very unusual in that it does not spend any money on administration or on itself - all of the income goes to fund activities that benefit the developmental biology community around the world. Every penny therefore counts. Until recently I was one of the Editors of MOD, but I stepped down to avoid a conflict of interest when I became President of ISDB. The relationship with MOD and the income it brings to the ISDB is crucianic. .... society - the more income, the more the society can up for the community (your support of MOD is therefore tel: (+44-20) 7679 3340 very important so please consider submitting some FAX: (+44-20) 7679 2091 email: < mailto:c.stern@ucl.ac.uk> lab home page: http://www.cdb.ucl.ac.uk/research/

one meeting every 4 years. We would like to increase the frequency of meetings and start other initiatives to serve the international community of developmental biologists. Of course all of this will require improving our funding base considerably, but both need to be developed at the same time for this to work. One possibil-

(covering Europe, pan-American and Asia-Pacific re-gions respectively), each every 4 years, out of cycle so that there is a major regional meeting somewhere every year, supporting a local organizing committee.

to invite one or more major international speakers, and/or to sponsor attendance of particularly promising students or postdocs or even mid-career developmental biologists (who are neither new investigators nor the most famous ones - the forgotten majority, which no other scheme wants to fund), to develop-mental biology meetings. A requirement of this is that these lectures and grants will be advertised widely as "MOD-ISDB lecture", etc.

That's enough. I hope that this email will arouse your interest in the ISDB and that you will contribute to its development in the future, alongside your national and regional societies. You may well have some ideas of your own concerning how the ISDB may be improved to become more visible and to serve you better - if so, we want to hear from you. Please write to me either directly (mailto:c.stern@ucl.ac.uk) or gestions and I will do my best to acknowledge every

Best wishes,

The current member societies of the ISDB are: SDB (USA), BSDB (UK) and the Societies for Develop-mental Biology of Japan, Germany (GfE), France (SFBD), Portugal, Latin-America and Australia/ New Zealand (ANZCDB) and the Asia-Pacific net-work (ADDPN) We are also currently talking to a work (APDBN). We are also currently talking to a few more national societies and encouraging them to join formally: the SDBs of Finland, Belgium, Italy, Israel, Spain, India, Mexico and Hong Kong. We are also currently in discussions with other countries that have a sizeable developmental biology community but not yet a society, including South Africa, Singapore, China, Taiwan and Iran, as well as possibly a European or East European grouping.

Claudio Stern (UK) (President) Marianne Bronner-Fraser (USA) (Secretary) Stefan Schulte-Merker (Netherlands) (Treasurer) Douglas Sipp (Japan) (Business Manager) David Wilkinson (UK) is also being included in this executive board in his capacity as Chief Editor of "Mechanisms of Development".

Claudio D Stern Department of Cell & Developmental Biology, University College London, Gower Street (Anatomy Building), London WC1E 6BT,

# Publications from our Members

Buchert, M, Athineos, D, Abud, H E, Burke, ZD, Faux, MC, Samuel, MS, Jarnicki, AG, Winbanks, CE, Newton, IP, Meniel, VS, Suzuki, H, Stacker, SA, Nathke, IS, Tosh, D, Huelsken, J, Clarke, AR, Heath, JK, Sansom, OJ and Ernst, M. Genetic dissection of differential signaling threshold requirements for the Wnt/beta-catenin pathway in vivo PLoS Genet (2010) 6 (1) e1000816 (\*authors contributed equally)

Chircop (nee Fabbro), M, Malladi, C, Lian, A, Zavortink, M, Gordon, CP, McCluskey, A and Robinson, PJ. Calcineurin activity is required for the completion of cytokinesis. Cell. Mol. Life Sci. Accepted 3/5/2010

Davies, B, Elwood, NJ, Li, S, Cullinane, F, Edwards, GA, Newgreen, DF, Brizard, CP (2010) Human cord blood stem cells enhance neonatal right ventricular function in an ovine model of right ventricular training. Ann Thorac Surg 89, 585-593, 593 e581-584.

Dunwoodie, SL. (2009). The Role of Hypoxia in Development of the Mammalian Embryo Developmental Cell 17, 755-773

Groszmann M, Bylstra J, Lampugnani ER & Smyth DR (2010) Regulation of tissue-specifc expression of SPATULA, a bHLH gene involved in carpel development, seedling germination, and lateral organ growth in Arabidopsis. Journal of Experimental Botany 61, 1495-1508

Grounds MD and Relaix F (2010) Myogenic Precursor Cells, in Section I- Scientific basis of muscle disease in 8th Edition of Disorders of Voluntary Muscles (Eds Hilton-Jones D, Griggs RC, Bushby K and Karpati G.) Cambridge University Press. Chapter 2, pp20-36

Harvey, KF. (2010). Bunched and Madm: a novel growth-regulatory complex? J Biol. 11: 8.

Hotta, R, Anderson, RB., Kobayashi, K, Newgreen, DF and Young, HM, (2010) Effects of tissue age, presence of neurones and endothelin-3 on the ability of enteric neurone precursors to colonize recipient gut: implications for cell-based therapies. Neurogastroenterol Motil.22, 331-e86.

Joshi, S, Perera, S, Gilbert, J, Smith, CM, Gordon, CP, McCluskey, A, Sakoff, JA, Braithwaite, A, Robinson, PJ and Chircop (nee Fabbro), M. The dynamin inhibitors MiTMAB and OcTMAB induce cytokinesis failure and inhibit cell proliferation in human cancer cells. Mol Cancer Ther. Accepted 3/5/2010

Kerr MC, Wang JT, Castro NA, Hamilton NA, Town L, Brown DL, Meunier FA, Brown NF, Stow JL, Teasdale RD. (2010) Inhibition of the PtdIns(5) kinase PIKfyve disrupts intracellular replication of Salmonella. EMBO J. 2010 Accepted Mar 18.

Leong GM, Kee AJ, Millard SM, Martel N, Eriksson N, Turner N, Cooney GJ, Hardeman EC, Muscat GE. The Ski proto-oncogene regulates body composition and suppresses lipogenesis. Int J Obes (Lond). 2010 Mar;34(3):524-36.

McDonald CJ, Jones MK, Wallace DF, Summerville L, Nawaratna S, Subramaniam VN. (2010). Increased iron stores correlate with worse disease outcomes in a mouse model of schistosomiasis infection. PLoS One. 5(3):e9594.

McMahon C, Shavlakadze T, Grounds M. (2010). Role of IGF-1 in age-related loss of keletal muscle mass and function. Book chapter in Sarcopenia, (ed. G. S. Lynch). Springer.

Medic S, and Ziman, M. (2010). PAX3 expression in normal skin melanocytes and melanocytic lesions (naevi and melanomas). PloSONE. In press

Meunier, FA, Nguyen, TH, Colasante, C, Luo, F, Sullivan, RKP, Lavidis, NA, Molgó, J, Meriney, SD and Schiavo, G. (2010). Sustained synaptic vesicle recycling by bulk endocytosis contributes to maintaining high-rate neurotransmitter release stimulated by Glycerotoxin J. Cell Sci. accepted Jan.

Milton, CC, Zhang, X, Albanese, NO, and Harvey, KF (2010). Differential requirement of Salvador-Warts-Hippo pathway members for organ size control in Drosophila melanogaster. Development 137: 735-743.

Monk, Adrian C.; Siddall, Nicole A.; Volk, Talila; Fraser, Barbara; Quinn, Leonie M.; McLaughlin, Eileen A.; Hime, Gary R. (2010). How is required for stem cell maintenance in the Drosophila testis and for the onset of transit amplifying divisions Cell Stem Cell 6: 348-360

Palmer SJ, Santucci N, Widagdo J, Bontempo SJ, Tay ES, Hook J, Lemckert F, Gunning PW, Hardeman EC. (2010). Negative auto-regulation of GTF2IRD1 in Williams-Beuren syndrome via a novel DNA binding mechanism. J Biol Chem. 285(7):4715-24.

Sarraj, MA, Escalona, RM, Umbers, A, Chua, HK, Small, C, Griswold, M, Loveland, K, Findlay, JK and Stenvers, KL. (2010). Fetal testis dysgenesis and compromised Leydig cell function in Tgfbr3 (beta glycan) knockout mice. Biol Reprod. 82(1):153-62.

Shavlakadze T, Chai J, Maley K, Grounds G, Winn N, Rosenthal N, Grounds M. (2010). A growth stimulus is needed for IGF-1 to induce skeletal muscle hypertrophy in vivo. Journal of Cell Science 123:960-71.

Shavlakadze T. and Grounds M. (2010) IGF-1 is a major regulator of muscle mass during growth but not for adult myofibre hypertrophy. Invited commentary for Point: Counterpoint 'IGF is / is not the major physiological regulator of muscle mass.' Journal of Applied Physiology. (ePress).

Shavlakadze T, McGeachie J, Grounds MD.

# **Publications from our Members**

(2010) Delayed but excellent myogenic stem cell response of regenerating skeletal muscles in geriatric mice. Biogerontology. [Epub ahead of print]

atric mice. Biogerontology. [Epub ahead of print] Sparrow, DB, Sillence, D, Wouters, MA, Turnpenny, PD and Dunwoodie, SL. Two novel missense mutations in HAIRY-AND-ENHANCER-OF-SPLIT-7 in a family with spondylocostal dysostosis. European Journal of Human Genetics (2010), 1–6

Stenvers KL, Findlay JK. (2010). Inhibins: from reproductive hormones to tumor suppressors. Trends Endocrinol Metab. 21(3):174-80

Theard, D, F Labarrade, M Partisani, J Milanini, H Sakagami, EA Fon, SA Wood, M Franco and F Luton (2010) Spatio-temporal protection from proteasomal degradation of EFA6 by the deubiquitinase USP9x contributes to tight junction assembly. In press EMBO J.

Wallace DF, Harris JM, Subramaniam VN. (2010). Functional analysis and theoretical modeling of ferroportin reveals clustering of mutations according to phenotype. Am J Physiol Cell Physiol. 298(1):C75-84.

Waters F, Shavlakadze T, McIldowie MJ, Piggott MJ, Grounds MD. (2010) Use of pifithrin to inhibit p53 mediated signalling of TNF in dystrophic muscles of mdx mice. Molecular and Cellular Biochemistry 337:119-131

Xu, Z, B Xia, Q Gong, J Bailey, B Groves, M Radeke, SA Wood, KK Szumlinski, D Ma (2010) Identification of a deubiquitinating enzyme as a novel AGS3-interacting protein. PLoS One 5(3):e9725

Zhang DC, Brinas IM, Binder, B, Landman, KA and Newgreen DF (2010) Neural Crest Regionalisation for Enteric Nervous System Formation: Implications for Hirschsprung's Disease and Stem Cell Therapy. Dev Biol 339: 280-294.

Zhang X, J Zhou, MH Chin, AA Schepmoes, VIA Petyuk, KK Weitz, BO Petritis, ME Monroe, DG Camp, SA Wood, WP Melega, DJ Bigelow, DJ Smith, W-J Qian, and RD Smith (2010) Regionspecific protein abundance changes in the brain of MPTP-induced Parkinson's Disease mouse model. Journal of Proteome Research, in press.

### **MEMBERS IN THE NEWS**

Prof. Roger Reddel has recently been elected as a Fellow of the Australian Academy of Science!

This is the highest accolade that could be bestowed on scientists in Australia. Election to the Academy recognizes Roger's outstanding achievements in biomedical research and seminal contribution to scientific knowledge in cancer biology.

Paul Thomas and Jeff Schwartz have compiled a Special Review Volume of Molecular & Cellular Endocrinology on "The Developmental Biology of Endocrine Organs" due to be published next month. This volume contains a series of informative reviews from leading researchers on a broad range of endocrine tissues including the pancreas, pituitary, gonads and placenta.

Professor Jenny Graves, a founding member of ANZSCDB, was made an Officer of the Order of Australia (AO), for 'service to scientific research in the field of genetics, particularly of Australian marsupials and monotremes, and their relevance to international understanding of human evolution'.

Professor Richard Harvey, Deputy Director of the Victor Chang Cardiac Research Unit and 2009 recipient of the ANZSCDB President's Medal has been awarded the 2010 Lemberg Medal by the Australian Society for Biochemistry and Molecular Biology.

# **Enjoy first-time success**



Gene expression and function analysis sample and assay technologies by QIAGEN

Rely on QIAGEN's manual and automated workflow solutions for:

- Sample collection and disruption, and RNA purification
- Real-time PCR and RT-PCR, and gene expression assays
- RNAi and gene silencing

GEFPortfO

- miRNA purification and assays
- Methylation analysis in epigenetics research
- Protein sample preparation and assays

Making improvements in life possible — <u>www.qiagen.com</u>



Sample & Assay Technologies

# Everyday needs. Everyday

answers.

For exciting promotions and offers for your everyday needs, please visit www.invitrogen.com/everydayneeds.



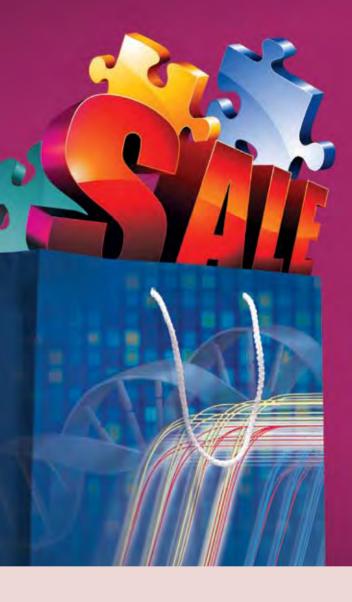
#### Easily find your favorite brands of reagents, kits, and benchtop devices at www.invitrogen.com/everyday.

<b>Ambion<sup>®</sup></b>	<b>Dynal<sup>®</sup></b>	<b>Gateway<sup>®</sup></b>	GIBCO <sup>®</sup>	Lipofectamine <sup>™</sup>
RNAi and RNA sample prep	Magnetic beads for separation	Cloning and protein expression	Cell culture	Transfection reagents
<b>Molecular Probes<sup>®</sup></b>	<b>Novex<sup>®</sup></b>	SuperScript <sup>®</sup>	<b>TaqMan<sup>®</sup></b>	<b>TOPO<sup>®</sup></b>
Fluorescent dyes and probes	Protein separation and blotting	cDNA synthesis reagents	Real-time PCR	PCR cloning

For research use only. Not intended for any animal or human therapeutic or diagnostic use, unless otherwise stated.

© 2010 Life Technologies Corporation or their respective owners. TaqMan is a registered trademark of Roche Molecular Systems, Inc. These products may be covered by one or more Limited Use Label Licenses (see the Invitrogen catalog or our website, www.invitrogen.com). By use of these products you accept the terms and conditions of all applicable Limited Use Label Licenses. CO13093 0410







**Gene Expression Reagents** You Can Count On

Offers run from April 1<sup>st</sup> – June 30<sup>th</sup> 2010

#### Quote PROMO-AJ10 when you order by fax, email or online, to take advantage of these special offers

Technical enquiries: 1800 645 619 Order Enquiries: 1800 802 409 Fax: 1800 066 598 Email: australia.customerservice@roche.com

More offers available on the website. Visit our Special Offer webpage at: www.roche-applied-science.com/specials

#### Offers run from April 1<sup>st</sup> – June 30<sup>th</sup> 2010

- **30% Off Transfection Reagents**
- 30% Off RNA Preparation Kits
- 30% Off cDNA Synthesis Kits
- 30% Off PCR Amplification Kits
- Save On Real Time PCR Kits

#### **TERMS AND CONDITIONS**

- Offers valid for orders received by fax, email or online while stocks last
- Offers are valid only for dates stated in the offer
- Roche may, at its sole discretion, terminate an offer at any time. The special offer discount is taken off the list price.
- Other discount agreements do not apply for these special offers.
  The promotional code must be provided when the order is placed. Promotional codes provided after ord placements are invalid.

#### TRADEMARK

FASTSTART, HIGH PURE, LIGHTCYCLER, MAGNA PURE, REALTIME READY, and TAQMAN are trademarks of Roche. SYBR is a registered trademark of Pholecular Probes, Inc. PAXgene is a registered trademark of PrAnalytiX GmbH. Other brands and product names are trademarks of their respective holders. For life science research only. Not for use in diagnostic procedures.

For full terms and conditions, please visit **www.roche-applied-science.com** 

# G-STORM GSI

# "The C-Storm CSI is everything you need from a thermal cycler"



### PURCHASE THE G-STORM GS1 FOR AN UNBEATABLE PRICE:

\$8450\*

normally \$11761 (with Standard 96 well or 96/48 combi block)

\$8800\*

normally \$12608 (with standard 384 block) \*Price excludes GST

GeneWorks Pty Ltd

- P 08 8234 2644
- FC 1800 882 555
- customerservice@geneworks.com.au
- www.geneworks.com.au

Interchangeable gradient capable blocks

- Colour touch screen display
- Simple yet powerful software
- Ramp rate of up to 3°CS<sup>-1</sup> (standard blocks) up to 6°CS<sup>-1</sup> (new fast block)

Offer expires 30 June, 2010

G-STOR

# GeneWorks

# Where did your bio begin?

Your work impacts new and exciting advancements in research. We want to know what started you on the path to understanding biology.

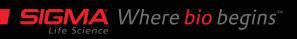


#### Share your story with us and enter to win!

Capture where your bio began using video, text or pictures. Visit *wherebiobegins.com/contest* to upload your story through May 31st, 2010. Your biostory could win you a CompoZr<sup>®</sup> Custom Zinc Finger Nuclease or other biocool prizes such as an Apple<sup>®</sup> iPad or free registration to a Sigma Technical Workshop!

All Australian and New Zealand entries will be eligible to go in the draw to win \$2000 worth of RNAi products. Offer expires July 31st, 2010.

Visit often to learn about the biostories of researchers like you.



SIGMA-ALDRICH®

CompoZr is a registered trademark of Sigma-Aldrich and Sigma-Aldrich Biotechnology, LP. IPad is a registered trademark of Apple Computer Inc. All rights reserved. Apple is not a participant or sponsor of this promotion. For complete contest rules and regulations, please visit wherebiobegins com

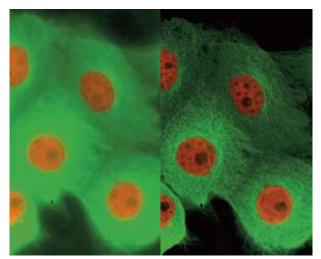
# VivaTome

Discover the Dynamics of Life

#### Fast Optical Sectioning Made Easy

VivaTome makes fast optical sectioning easy and affordable, providing clear optical sections at frame rates high enough for imaging of dynamic events using just a normal white light source.

The Aperture Correlation approach combines the light efficiency of structured illumination with the speed of the Spinning Disk approach. All information necessary to calculate an optical section and a widefield image are captured simultaneously, preventing motion artifacts and giving you access to all the information. The VivaTome fits many different microscope stands and provides the ideal entry-level solution to optically sectioned live cell imaging.



A. Widefield image

B. VivaTome optical section



VivaTome

#### Benefits

- Simple and affordable instrumentation optical sectioning using a normal white light source
- Fast image acquisition observe dynamic events with brilliant contrast
- True optical sectioning efficiently remove out-of-focus blur
- Two different grid patterns guarantee **optimal performance** with the whole range of objectives
- Widefield and optical section simultaneously don't loose any information

#### Applications

- Visualise organelle movement over time
- Follow developmental processes in four dimensions
- Capture **subcellular structures** in three dimensions with an easy-to-use setup and short acquisition times
- Complement your **physiology experiments** with fast fluorescence imaging

#### Contact us now for more information

Carl Zeiss Pty Ltd

Ph: 1300 365 470 micro@zeiss.com.au Ph: 02 9020 1333 www.zeiss.com.au



We make it visible.

# ANZSCDB Executive Officers & Committee Members 2009-2010

#### PRESIDENT: Edna Hardeman

Department of Anatomy School of Medical Sciences Wallace Wurth Building G27 The University of New South Wales Sydney NSW 2052 Tel 02 9385 3760 Fax 02 9385 8016 e.hardeman@unsw.edu.au

#### PRESIDENT ELECT: Peter Currie

Australian Regenerative Medicine Institute Monash University Clayton VIC p.currie@victorchang.unsw.edu.au

#### **TREASURER:** Katharina Gaus

Centre for Vascular Research School of Medical Sciences University of New South Wales Sydney NSW 2052 k.gaus@unsw.edu.au

#### SECRETARY : Geraldine O'Neill

Oncology Research Unit The Children's Hospital at Westmead Westmead NSW 2145 Tel 02 9845 3116 geraldio@chw.edu.au

#### ANZSCDB Secretariat: Ros Barrett-Lennard

Magic Touch Consultancies Tel +612 9524 1799 Fax +612 9524 1744 Mob +61 (0) 419 688 581 anzscdb@mtci.com.au

#### IMMEDIATE PAST PRESIDENT: Alpha Yap

Institute for Molecular Bioscience Queensland Bioscience Precinct The University of Queensland St Lucia QLD 4072 Tel 07 3346 2013 Fax 07 3346 2101 a.yap@imb.uq.edu.au

#### NEWSLETTER EDITOR: Megan Chircop

Chldren's Medical Research Institute Locked Bag 23 Westmead NSW 2145 Tel 02 9687 2800 Fax 02 9687 2120 mchircop@cmri.com.au

#### STATE REPRESENTATIVES:

ACT: Radiya Ali Australian National University radiya.ali@anu.edu.au

**NSW: Bill Phillips** University of Sydney billp@physiol.usyd.edu.au

NSW: Thomas Fath University of NSW CNS.Cytoskeleton@gmail.com

**QLD: Eva Kovacs** University of Queensland e.kovacs@uq.edu.au

**QLD: Dagmar Wilhelm** Institute for Molecular Biosciences University of Queensland d.wilhelm@imb.uq.edu.au

VIC: Ian Smyth Monash University ian.smyth@med.monash.edu.au

VIC: Peter Farlie peter.farlie@mcri.edu.au

**SA: Yessim Khew-Goodall** Yeesim.KHEW-GOODALL@health. sa.gov.au

**SA: Bryan Haines** School of Molecular and Biomedical Sciences University of Adelaide bryan.haines@adelaide.edu.au

WA: Cecilia Prele

Telethon Institute for Child Health Research Tel: +61 8 9489 7884 ceciliap@ichr.uwa.edu.au

WA: Aleksandra Filipovska West Australian Institute for Medical Research afilipov@waimr.uwa.edu.au

NZ: Christine Jasoni Centre for Neuroendocrinology University of Otago Christine.jasoni@stonebow.otago. ac.nz

NZ: Maria Flores m.flores@auckland.ac.nz