provided by Digital.CSI

Abstract for EDRC 2013

Section: Morphogenesis and Organogenesis

EVIDENCE FOR THE COMMON ORIGIN OF TRACHEA AND ENDOCRINE ORGANS FROM A SEGMENTALLY REPEATED ECTODERMAL PRECURSOR

J Castelli-Gair Hombría, C Sánchez-Higueras, S Sotillos

Centro Andaluz de Biologia del Desarrollo (CABD), Consejo Superior de Investigaciones Científicas (CSIC)/ Junta de Andalucía (JA)/ Universidad Pablo de Olavide, Seville, Spain

The embryonic origin of the insect endocrine glands is not well known. Here we show that the corpora allata (ca), the prothoracic glands (pg) and the trachea have a homologous origin. The trachea is a segmentally repeated organ developing from ten ectodermal placodes expressing Trachealess (Trh) and Ventral veinless (Vvl). These placodes invaginate and migrate to form an epithelial polarized tubular network. We show that homologous cells in the maxilla and labium expressing only Vvl, form the ca and the pg. The initial development of the trachea and the endocrine primordia is identical but it diverges when the endocrine organs activate Snail (Sna) and suffer an Epithelial to Mesenchymal Transition (EMT). We show that in the gland primordia Sna controls EMT, migration and viability.

The ca is specified by Deformed in the maxilla, and the pg by Sex combs reduced in the labium. These Hox genes specify the glands activating Vvl, Sna and either Seven-up or Spalt. We can follow in vivo how the two gland primordia coalesce and migrate dorsally to join the corpora cardiaca and form the ring gland. The endocrine organs and trachea are specified by similar upstream gene networks with STAT and Hox genes inducing their development. Hox genes can convert the endocrine primordia into trachea and viceversa. Our study uncovers the genetic and developmental mechanisms for ring gland morphogenesis and indicates that the respiratory tracheal organs and these two main endocrine glands arose through a process of divergent evolution from an ectodermal repeated structure.



## **KEY DATES**

Abstract Submission Opens: 16 January 2013

Deadline Abstract submission: 2 June 2013

Deadline Registration: 30 July 2013

## **TOPICS**

- Stem Cells
- Growth and Size Control
- Cell Polarity
- Morphogenesis and Organogenesis
- Models of Human Disease
- Immunity
- New Genome Wide Application
- Cell Communication and Signalling
- Non-coding RNA's
- Epigenetics
- Gene Expression / Pattern Formation
- Cell Cycle
- Neurobiology and Behaviour
- Phisiology and Metabolism
- Molecular Population Genetics and Evolution

## PLENARY LECTURES

Utpal Banerjee
Cayetano González
Jules Hoffmann
Eli Knust
Ginés Morata
Trudi Schupbach
William Theurkauf

## **ORGANIZING COMMITTEE**

Jordi Casanova Cayetano González Enrique Martín-Blanco Marco Milán Florenci Serras







B Universitat de Barcelona



