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Development



Cover: Fz5/8 is required for archenteron invagination in the sea urchin embryo. DIC images of a larva in which the Fz5/8 pathway has been inhibited (right) and of a larva displaying a normal phenotype (left) after being rescued by co-injection of the wild-type form of the receptor.

See article by Croce et al. on p. 547.

REVIEW

385 The interpretation of morphogen gradients Ashe, H. L. and Briscoe, J.

RESEARCH ARTICLES

395 lessen encodes a zebrafish trap100 required for enteric nervous system development
Pietsch, J., Delalande, J.-M., Jakaitis, B., Stensby, J. D., Dohle, S., Talbot, W. S., Raible, D. W. and Shepherd, I. T.

407 Cholesterol modification is necessary for controlled planar long-range activity of Hedgehog in *Drosophila* epithelia Gallet, A., Ruel, L., Staccini-Lavenant, L. and Thérond, P. P.

The *Cdx4* mutation affects axial development and reveals an essential role of Cdx genes in the ontogenesis of the placental labyrinth in mice van Nes, J., de Graaff, W., Lebrin, F., Gerhard, M., Beck, F. and Deschamps, J.

429 Timing of identity: spatiotemporal regulation of hunchback in neuroblast lineages of Drosophila by Seven-up and Prospero Mettler, U., Vogler, G. and Urban, J.

439 The Dictyostelium bZIP transcription factor DimB regulates prestalk-specific gene expression
Zhukovskaya, N. V., Fukuzawa, M., Yamada, Y., Araki, T. and Williams, J. G.

bZIP transcription factor interactions regulate DIF responses in *Dictyostelium* Huang, E., Blagg, S. L., Keller, T., Katoh, M., Shaulsky, G. and Thompson, C. R. L.

459 Drosophila Cornichon acts as cargo receptor for ER export of the TGF α -like growth factor Gurken Bökel, C., Dass, S., Wilsch-Bräuninger, M. and Roth, S.

471 Hedgehog lipid modifications are required for Hedgehog stabilization in the extracellular matrix
 Callejo, A., Torroja, C., Quijada, L. and Guerrero, I.

Ras-dva, a member of novel family of small GTPases, is required for the anterior ectoderm patterning in the *Xenopus laevis* embryo

Tereshina, M. B., Zaraisky, A. G. and Novoselov, V. V.

Neurogenin 2 is required for the development of ventral midbrain dopaminergic neurons
 Kele, J., Simplicio, N., Ferri, A. L. M., Mira, H., Guillemot, F., Arenas, E. and Ang, S.-L.

Development of the mesencephalic dopaminergic neuron system is compromised in the absence of neurogenin 2
 Andersson, E., Jensen, J. B., Parmar, M., Guillemot, F. and Björklund, A.

The Sonic hedgehog pathway independently controls the patterning, proliferation and survival of neuroepithelial cells by regulating Gli activity
Cayuso, J., Ulloa, F., Cox, B., Briscoe, J. and Martí, E.

EVELOPMENT

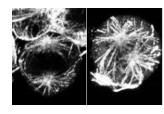
529 Extrinsic cues orient the cell division axis in *Drosophila* embryonic neuroblasts Siegrist, S. E. and Doe, C. Q.

537 Massive loss of Cajal-Retzius cells does not disrupt neocortical layer order Yoshida, M., Assimacopoulos, S., Jones, K. R. and Grove, E. A.



How do responding cells perceive and interpret the concentration-dependent information provided by a morphogen? Ashe and Briscoe review recent work on the molecular features of morphogen signalling and identify some emerging common principles. See review article on p. 385.

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In the absence of extrinsic cues, neuroblast cortical polarity does not form until metaphase. Left, a neuroblast cultured in contact with other cell types at prophase; right, a neuroblast cultured in isolation at prophase. Both are stained for α -tubulin. See article by Siegrist and Doe on p. 529.

Frizzled5/8 is required in secondary mesenchyme cells to initiate archenteron invagination during sea urchin development
 Croce, J., Duloquin, L., Lhomond, G., McClay, D. R. and Gache, C.

DEVELOPMENT AND DISEASE

- A dominant-negative form of the E3 ubiquitin ligase *Cullin-1* disrupts the correct allocation of cell fate in the neural crest lineage Voigt, J. and Papalopulu, N.
- GLI3-dependent transcriptional repression of Gli1, Gli2 and kidney patterning genes disrupts renal morphogenesis
 Hu, M. C., Mo, R., Bhella, S., Wilson, C. W., Chuang, P.-T., Hui, C.-c. and Rosenblum, N. D.