



## EDITORIAL

# Neural-mesodermal progenitor interactions in pattern formation: an introduction to the collection [version 1; peer review: not peer reviewed]

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Mesodermal and spinal cord progenitors originate from common founder cells from which they segregate during development. Moreover, neural and mesodermal tissues closely interact during embryogenesis to ensure timely patterning and differentiation of both head and trunk structures. For instance, the fate and morphogenesis of neural progenitors is dependent on signals produced by mesodermal cells and vice-versa. While some of the cellular and molecular signals that mediate these interactions have been described, much more remains to be uncovered. The scope of this collection will cover these interactions between neural (CNS or PNS) and mesodermal progenitors in patterning body plans and specific body systems in vertebrate embryos. This includes, but is not limited to, interactions influencing the formation of body axes, neural tube formation, neural crest migration, gut development, muscle patterning and myogenesis.



This article is included in the [Neural-mesodermal progenitor interactions in pattern formation](#) collection.

**Not Peer Reviewed**

This article is an Editorial and has not been subject to external peer review.

Any comments on the article can be found at the end of the article.

**Corresponding author:** Chaya Kalcheim ([kalcheim@cc.huji.ac.il](mailto:kalcheim@cc.huji.ac.il))**Competing interests:** No competing interests were disclosed.**Grant information:** The author(s) declared that no grants were involved in supporting this work.**Copyright:** © 2014 Kalcheim C and Storey KG. This is an open access article distributed under the terms of the [Creative Commons Attribution Licence](#), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.**How to cite this article:** Kalcheim C and Storey KG. [Neural-mesodermal progenitor interactions in pattern formation: an introduction to the collection \[version 1; peer review: not peer reviewed\]](#) F1000Research 2014, 3:275 (<https://doi.org/10.12688/f1000research.5657.1>)**First published:** 14 Nov 2014, 3:275 (<https://doi.org/10.12688/f1000research.5657.1>)

## Editorial

Clonal analysis indicated that trunk neural and mesodermal progenitors share a common lineage at an early developmental stage (Tzouanacou *et al.*, 2009). These dual-fated neuromesodermal precursors then segregate and further differentiate into either spinal cord or somite precursors. With the aim of elucidating how these fates bifurcate, researchers recently set out to recapitulate their differentiation pathways *in vitro* departing from embryonic stem cells (Gouti *et al.*, 2014). Notably, the above lineages born from common precursors exhibit close interactions during later development that play pivotal roles in patterning body plans and specific body systems. For example, graded mesodermal signals control the timing of neuronal differentiation in the spinal cord (Wilson *et al.*, 2009); somitic signals control the proliferation, delamination, segmental migration and patterning of neural crest-derived peripheral ganglia and, reciprocally, neural crest cells pattern the organization of mesodermal derivatives (i.e; muscles) in both the head and trunk (Kalcheim, 2011).

Thus, the cross-talk between these developing systems is highly dynamic and reiterative. Emerging data highlight additional interactions between mesoderm and neural progenitors in development of gut innervation, craniofacial patterning, etc. and much more is yet to be discovered.

This collection aims to be a platform for original reports, review articles and opinions that expand our understanding on tissue patterning during development including the generation of these tissues and the molecular nature of their interactions.

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No competing interests were disclosed.

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## References

Gouti M, Tsakiridis A, Wymeersch FJ, *et al.*: **In vitro generation of neuromesodermal progenitors reveals distinct roles for wnt signalling in the specification of spinal cord and paraxial mesoderm identity.** *PLoS Biol.* 2014; **12**(8): e1001937.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)

Kalcheim C: **Regulation of trunk myogenesis by the neural crest: a new facet of neural crest-somite interactions.** *Dev Cell.* 2011; **21**(2): 187–188.  
[PubMed Abstract](#) | [Publisher Full Text](#)

Tzouanacou E, Wegener A, Wymeersch FJ, *et al.*: **Redefining the Progression of lineage segregations during mammalian embryogenesis by clonal analysis.** *Dev Cell.* 2009; **17**(3): 365–376.

[PubMed Abstract](#) | [Publisher Full Text](#)

Wilson V, Olivera-Martinez I, Storey KG: **Stem cells, signals and vertebrate body axis extension.** *Development.* 2009; **136**(10): 1591–1604.  
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