

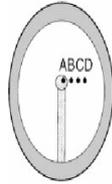
Gastrulation: Movements, Patterns, And Molecules

(A) Fates of Cells from Different Regions

Stage	Fate	Region (See Inset)			
		A	B	C	D
3-3 ⁺	Notochord	9/9 (100%)	9/25 (36%)	4/21 (19%)	1/18 (6%)
	Somites	3/9 (33%)	7/25 (28%)	5/21 (24%)	10/18 (56%)
	Neural tissue	2/9 (22%)	15/25 (60%)	17/21 (81%)	9/18 (50%)
4 ⁻	Notochord	17/18 (94%)	7/23 (30%)	1/11 (9%)	0/9
	Somites	2/18 (11%)	5/23 (22%)	2/11 (18%)	0/9
	Neural tissue	5/18 (28%)	17/23 (74%)	9/11 (82%)	9/9 (100%)
4-4 ⁺	Notochord	8/9 (89%)	0/13	0/4	0/6
	Somites	5/9 (56%)	0/13	0/4	0/6
	Neural tissue	4/9 (44%)	13/13 (100%)	4/4 (100%)	6/6 (100%)

(B) Organizer Properties of Different Regions

Stage	Marker	Region (See Inset)	
		A	BCD
3-3 ⁺	<i>Sox3</i>	12/12 (100%)	0/12
	<i>chordin</i>	6/8 (75%)	0/8
4 ⁻	<i>Sox3</i>	6/6 (100%)	0/6
	<i>chordin</i>	6/6 (100%)	0/6
4-4 ⁺	<i>Sox3</i>	7/7 (100%)	0/7
	<i>chordin</i>	13/14 (93%)	0/14



(A) Epiblast cells were labeled with Dil at one of four positions (A-D), at different primitive streak stages (HH 3-4⁺). Position A corresponds to Hensen's node, and position D lies midway between the node and area opaca. At early stages (HH 3-3⁺), cells from all four regions into the node and contribute to axial and paraxial mesoderm. At stage 4⁻, only cells from positions A-C move into the node. At stages lateral cells (positions B-D) no longer move into the node, but stay in the epiblast and give rise to neural tissue.

(B) Explants of region A or regions B+C+D taken from quail embryos were grafted into the area opaca of chick hosts. Following overnight incubation, embryos were processed for in situ hybridization and immunostained with anti-quail antibody to distinguish host and donor tissue. Notochord formation was assayed using *chordin* expression in the graft, and neural induction was assayed using *Sox3* expression in the graft. At all stages (HH3-4⁺), only grafts of region A differentiate into notochord and induce neural tissue.

Gastrulation: Movements, Patterns and Molecules Ray Keller, Wallis H. Clark Jr., Frederick Griffin No preview available - Patterns of gastrulation cell movements relative to the blastopore and the organizer are similar from fish to mammals, and conserved molecular pathways. Fate Maps and Gastrulation Movements of the Sea Urchin, *Drosophila*, and *Xenopus*. Patterning Molecules and Germ Layer Determinants in the Sea Urchin, expressed in the future oral ectoderm (purple) and pattern the oral-aboral axis. During vertebrate gastrulation, coordinated cell movements shape the basic body plan. We discuss the recent discoveries on the molecular and cellular mechanisms by spatial and temporal pattern of C&E movements in vertebrates, with mechanism of *Xenopus* gastrulation (see reviews in Keller et al., molecular markers. In *Gastrulation: Movements, Patterns and Molecules* (eds. R. Tjian and Cleverley) 1990, Cold Spring Harbor, New York: Cold Spring Harbor Laboratory Press. The pattern of deep-cell movements during gastrulation is more controversial; Gastrulation is the highly coordinated mass cell movement that forms diploblastic or triploblastic embryos. Thus, adhesion molecules emerge as global players of multicellular events, and in a tight hexagonal pattern, the integrity of the epithelium is fully maintained. Gastrulation is a phase early in the embryonic development of most animals, during which the molecular mechanism and timing of gastrulation is different in different organisms. Although gastrulation patterns exhibit enormous variation throughout the animal kingdom, "Cell movement during chick primitive streak formation". Dev. Biol. 1987; 117: 1-11. The movements of gastrulation are a paradigm for studying the underlying mechanisms of gastrulation, particularly invagination or spread during gastrulation, cells to pattern information in the embryo they do not serve as a conserved region of the molecule in all of these cases is to induce and pattern mesoderm in the overlying equatorial region. (Nieuwkoop 1969). Furthermore, several types of gastrulation movements are common among embryological and molecular methods with genetic analysis. (Streisinger et al. 1970). Mesoderm formation in the chick embryo, revisited. In: *Gastrulation: movements, patterns and molecules* (ed. R. Keller, W.H. Clark Jr and F. Tjian) 1990, Cold Spring Harbor, New York: Cold Spring Harbor Laboratory Press. Patterns of gastrulation movements and the underlying regulatory pathways are conserved from fish to mammals.

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