

Two other morphological features of enteropneusts further suggest that the dorsal surfaces of enteropneusts and chordates are homologous. First, enteropneusts have a putative homologue of the chordate notochord on the dorsal side of the animal, the stomochord<sup>10</sup>. Second, enteropneusts have U-shaped brachial skeletons of identical design and orientation with chordates<sup>6-8</sup>.

Before one can conclude that an inversion of the dorsoventral axis occurred at the time of origin of the chordates, minimally it must be shown that: (1) *BMP-2* is expressed ventrally in vertebrates, and member(s) of the *dpp* subfamily<sup>2</sup> are expressed ventrally in 'protochordates'; (2) the dorsal nerve cord of enteropneusts and chordates are not homologues; (3) the stomochord is not the homologue of the notochord; and (4) the pharyngeal skeletons in enteropneusts and chordates arose independently. Until then, Geoffroy's original hypothesis

remains simply a matter of historical significance.

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## Cation selectivity in ion channels

SIR — Valera *et al.*<sup>1</sup> and Brake *et al.*<sup>2</sup> have reported the amino-acid sequences of members of a novel class of ligand-gated ion channel, namely the ATP-gated channel or P<sub>2x</sub> receptor. The channel dis-

in alignment of the highly conserved TTTXGXG motif in the H5 region of potassium channels<sup>5</sup>. Of particular interest is the GXG motif within the H5 inwardly-rectifying and voltage-gated potassium channels tyrosine is conserved in the central position. In the *ether-à-go-go* channel<sup>6</sup>, tyrosine is replaced by phenylalanine, while in the P<sub>2x</sub> receptor<sup>1,2</sup> serine or valine present.

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	*	22	23	24	25	26	27	28	29	30								
IRK1 K <sup>+</sup> channel	T	A	A	F	L	F	S	I	E	I	T	Q	T	T	I	G	Y	G	F	R																		
IRK2 K <sup>+</sup> channel	M	A	A	F	L	F	S	I	E	I	T	Q	T	T	I	G	Y	G	L	R																		
KATP K <sup>+</sup> channel	V	S	A	F	L	F	S	I	E	I	T	Q	T	T	I	G	Y	G	F	R																		
ROMK1 K <sup>+</sup> channel	T	S	A	F	L	F	S	I	E	I	T	Q	T	T	I	G	Y	G	F	R																		
Shaker A K <sup>+</sup> channel	P	D	A	F	W	W	A	V	V	I	T	M	T	V	G	Y	G	D	M																			
DRK1 K <sup>+</sup> channel	P	A	S	F	W	W	A	T	I	T	M	T	V	G	Y	G	D	I																				
<i>ether-à-go-go</i>	V	T	A	L	Y	F	T	M	T	C	M	T	S	V	G	F	G	N	V																			
P <sub>2x</sub> PC12 cells	I	P	T	I	N	L	A	T	A	L	T	S	I	G	V	G	S	F																				
P <sub>2x</sub> vas deferens	K	A	G	K	F	D	I	P	T	M	T	T	I	G	S	G	I	G																				

Alignment of the H5 region of inwardly-rectifying potassium channels (IRK1, IRK2, KATP and ROMK1), voltage-gated potassium channels (DRK1, Shaker A and *ether-à-go-go*) and ATP-gated channels (P<sub>2x</sub> receptors). Boxed region, highly conserved eight-residue potassium-channel signature sequence<sup>5</sup>. Asterisk, denotes the central position of the GXG motif.

plays cation selectivity when expressed in *Xenopus* oocytes, and both groups of authors propose a transmembrane topology similar to that of a recently cloned family of inwardly-rectifying potassium channels<sup>3</sup>, with two potential membrane-spanning helices (M1 and M2) and an intervening region, H5. The sequence alignment proposed by Valera *et al.*<sup>1</sup> for P<sub>2x</sub> cloned from rat vas deferens is very similar to inward-rectifiers and voltage-gated potassium channels within its H5 region.

We would like to propose an alternative assignment for the H5 region of P<sub>2x</sub> cloned from rat PC12 cells to that offered by Brake *et al.*<sup>2</sup>. Our assignment was obtained by alignment of the proposed transmembrane regions of voltage-gated and inwardly-rectifying potassium channels with the P<sub>2x</sub> sequences<sup>1,2</sup>, using the multiple sequence alignment package, AMPS<sup>4</sup>. Our analysis (see figure) results

As proposed by Kumpf and Dougherty<sup>7</sup>, the presence of aromatic amino acids within H5 may result in potassium-selective 'cation-π' interactions in potassium channels. Thus, highly potassium-selective inward-rectifiers and voltage-gated channels exhibit a conserved tyrosine in GXG (or phenylalanine in *ether-à-go-go*), whereas in the non-specific cation channels encoded by P<sub>2x</sub> tyrosine is replaced by a smaller, non-aromatic residue<sup>1,2</sup>.

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## Bet on positional information

SIR — One idea of how patterns of cellular differentiation are specified during development is that first, positional information is set up in a population of cells, and second, that each cell interprets this information according to its genetic constitution and developmental history<sup>1</sup>. Recent studies have suggested that a related set of homeotic genes encode positional identity along the body axis of many animals, and it has even been suggested that this defines a zootype that all animals share<sup>2</sup>. In insects, for example, one can think of the homeotic genes as acting together to give genetic addresses, which establish segment identity<sup>3,4</sup>. Martinez Arias<sup>5</sup> has challenged these ideas on the basis of the work of Warren *et al.*, who have studied the expression of the homeotic gene *Ultrabithorax (Ubx)* in butterflies<sup>6</sup>.

In *Drosophila*, *Ubx* is expressed in the third thoracic segment which forms a haltere. In butterflies the third thoracic segment carries a wing. According to Martinez Arias, "most punters would have put their money on the absence of expression" of *Ubx* in the third thoracic segment of butterflies because he thinks the homeotic genes are associated with specific structures. But I, and other colleagues I have asked, would not have put down even a penny in support of such an idea, for the identity of a segment as specified by homeotic genes says nothing about what structures it will develop. That is the central idea of positional information. Indeed Warren *et al.*'s finding that in butterflies the third thoracic segment expresses *Ubx* strongly supports the idea of positional identity being specified by the homeotic genes. What has changed in evolution of the basic pattern of butterflies and flies is not the positional identity of the thoracic segments nor the deployment of homeotic genes, but their downstream targets.

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