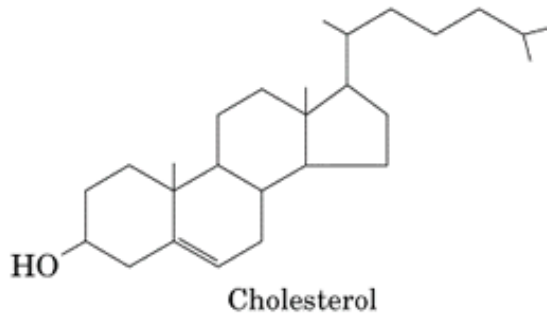


1. 膽固醇如何影響細胞膜流動？



膽固醇會降低細胞膜的流動性，但卻是細胞膜的朔化劑，有膽固醇從中固定，可以使細胞膜不易受外力的影響而破損，導致細胞崩潰。另外，還可增加細胞膜的穩定性，減少在高溫時膜的液化或冷溫時膜固化。

2. DNA 序列上的相同區域，可否因為讀法不同而產生不同蛋白質？

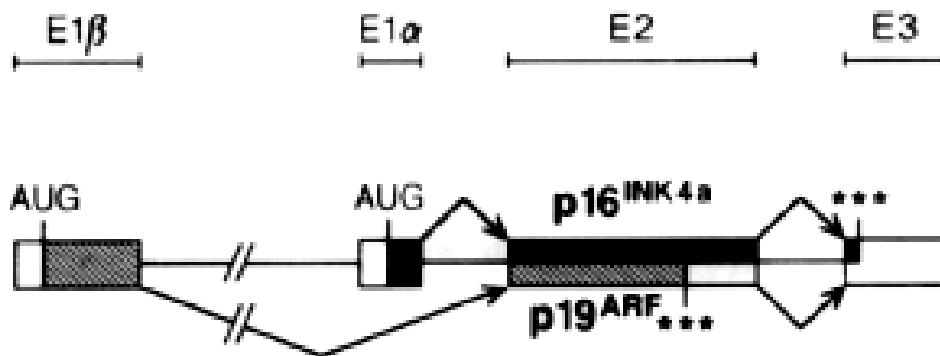


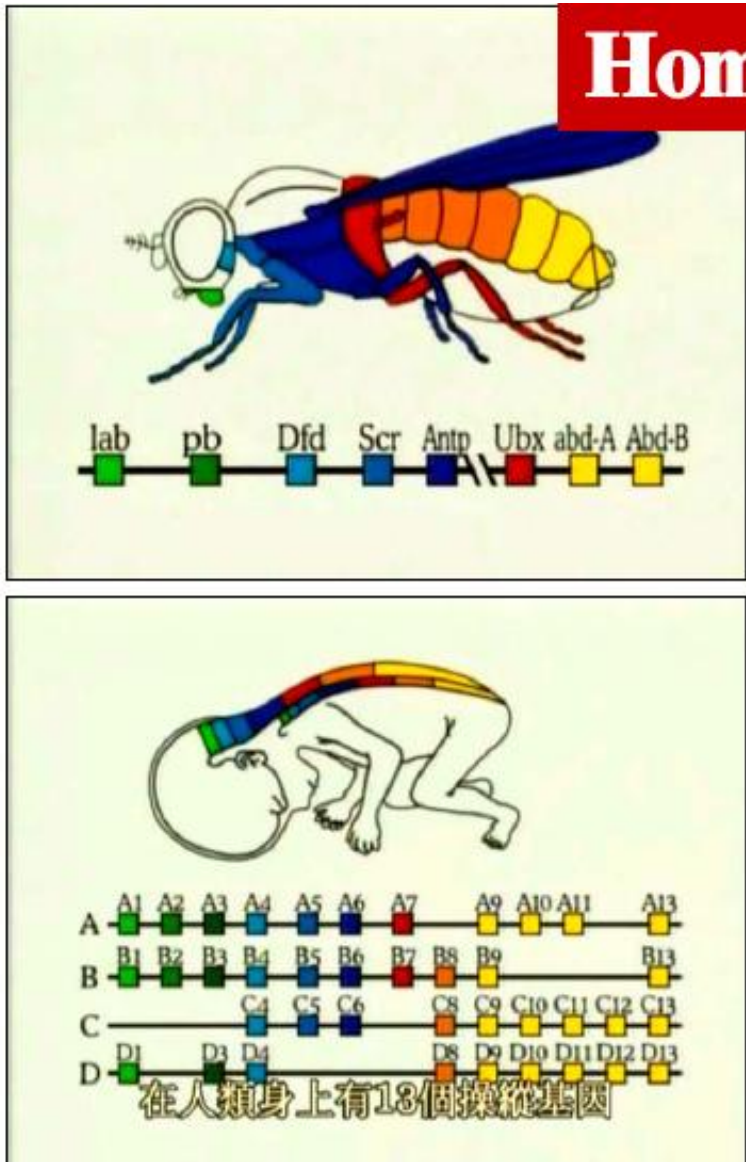
Figure 1. Architecture of the *INK4a* Gene

Transcription of α and β mRNAs encoding p16^{INK4a} and p19^{ARF}, respectively, is initiated at two promoters (Stone et al., 1995; Mao et al., 1995). Exon 1 α and 1 β products are spliced to the same exon 2 acceptor to generate α and β mRNAs of similar lengths but using alternative reading frames for translation. Mouse p19^{ARF} and p16^{INK4a} are each longer than their human cognates, and, like human p16^{INK4a}, the mouse p16^{INK4a} distal C-terminal amino acids are encoded by a third exon (Quelle et al., 1995). Protein coding sequences are depicted by closed (p16^{INK4a}) or hatched (p19^{ARF}) bars, and noncoding sequences by open bars.

Quelle et al. (1995) Alternative Reading Frames of the *INK4 α* Tumor Suppressor Gene Encode Two Unrelated Proteins Capable of Inducing Cell Cycle Arrest. *Cell* (1995) 83: 993-1000

3. 控制器官發育的主鑰匙基因之最新研究？

其中 HOXD13 的突變會造成多指症 (synpolydactyly)；而 HOXA13 的突變則會導致手腳生殖器症 (hand-foot-genital syndrome, HGFS)。另外 HOXB 群集中的 8 個成員會影響紅血球的發育，其中 HOXB4 與 HOXB7 又會影響 T 細胞與 B 細胞。<http://zh.wikipedia.org/zh/Hox%E5%9F%BA%E5%9B%A0>



4. 六根手指頭的基因遺傳？

Polydactyly is the most common congenital digital anomaly of the hand and foot. It may appear in isolation or in association with other birth defects. Isolated polydactyly is often **autosomal dominant** or occasionally random, while syndromic polydactyly is commonly autosomal recessive. According to Muragaki et al, mutations in the **HOXD13** gene are associated with synpolydactyly.

<http://en.wikipedia.org/wiki/Polydactyly>

<http://emedicine.medscape.com/article/1113584-overview>

Muragaki et al. Altered growth and branching patterns in synpolydactyly caused by mutations in HOXD13. *Science*. 1996; 272(5261):548-51.