

## Characterization of single-nucleotide polymorphisms in coding regions of human genes

Michele Cargill *et al.*

*Nature Genet.* **22**, 231–238 (1999).

We inadvertently omitted Nila Shaw from our list of authors. The correct author list follows.

Michele Cargill, David Altshuler, James Ireland, Pamela Sklar, Kristin Ardlie, Nila Patil, Nila Shaw, Charles R. Lane, Esther P. Lim, Nilesh Kalyanaraman, James Nemesh, Liuda Ziaugra, Lisa Friedland, Alex Rolfe, Janet Warrington, Robert Lipshutz, George Q. Daley & Eric S. Lander

## Griselli disease maps to chromosome 15q and is associated with mutations in the Myosin-Va gene

Elodie Pastural *et al.*

*Nature Genet.* **16**, 289–292 (1997).

One of the two Myosin Va mutations described, the Arg→Cys alteration at codon 1,246 (C→T transition at nt 3,736), was found to be a relatively common polymorphism. Several healthy individuals are homozygous carriers of the Cys1246 allele. This fact was first pointed out to *Nature Genetics* by Jo Lambert, Jean Marie Naeyart, Anne de Paepe, Rudy van Coster, Alina Ferster, Michele Song and Ludwine Messiaen of the University of Gent, Gent, Belgium. It was subsequently acknowledged by Pastural *et al.*

## Suppression of the novel growth inhibitor p33<sup>ING1</sup> promotes neoplastic transformation

Igor Garkavtsev *et al.*

*Nature Genet.* **14**, 415–420 (1996).

Due to a cloning error, the sequence reported for *ING1* was incorrect. The error appears to have been a result of a compression introducing a frameshift and of the *ING1* gene encoding several differentially spliced isoforms that contain a common 3' exon, one of which is of a size very similar to that reported in the publication above. The original (*ING1a'*, previously called *ING1*) and the corrected (*ING1a*) sequences of the isoform of *ING1* that was first reported are shown at right (a). The sequence that we reported was frameshifted and truncated at the 5' end, and we now know that it encodes a protein with a predicted mass of 46,751 rather than the predicted 33,253 daltons. A truncated *ING1a* message also encodes an expressed protein of 23,656 daltons that results from initiation at the first internal ATG in the conserved exon. An alternatively spliced isoform, *ING1b*, shares a common 3' exon with *ING1a* and encodes a protein with a predicted mass of 31,843 daltons. We now refer to the proteins encoded by this gene as p47<sup>ING1a</sup>, p33<sup>ING1b</sup> and p24<sup>ING1c</sup> due to their relative electrophoretic mobilities. These polypeptides are found in the majority of primary and established tissue culture cells examined, including those of fibroblast, epithelial and glial origin. A schematic diagram of the proteins encoded by the two major transcripts of *ING1* are shown (b). All three proteins contain a region with a high degree of homology to PHD fingers that are implicated in transcriptional regulation.

The complete, corrected nucleotide sequences for the cDNAs encoding *ING1a* and -b are available at accession numbers AF181849 (for *ING1a*, formerly listed as AF001954) and AF181850 at GenBank. We regret any inconvenience that may have resulted from the sequencing error. Expression constructs encoding full-length *ING1a*, *ING1b* and *ING1c* proteins are available from Dongping Ma, Denise Lawless and Karl Riabowol, Department of Biochemistry and Molecular Biology, University of Calgary HSC, 3330 Hospital Dr. NW, Calgary, Alberta, Canada T2N 4N1.

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ING1 a : tgcctcgggg gggggcgggg cagatcgtcg gotttgagag gaotctggca ggtgagagga
ING1 a : cctgtgogtc gttctctgca gaacctggcc cccggggtgt cagagagagg tggcagtttc
ING1 a : gtgtccgocg ggaattgttg gotgttgggg aaactttcct gogaggtcag toaaggotttt
ING1 a : gggggctctg ttttgaatgt ggatcaccac toggagttta ctaagtgtta caaggctggg
ING1 a : cagtagggaa accggaagagt tgggtggggg caaaaaaaa aattgaccgc tgtcccggaa
ING1 a : agtaactagc goctctgocg ggaaggoccc cctgocggtt ctatccagaa cgtagotttg
ING1 a : cagogaattt tataggaact toattgcaat attatggaaq gtcccgcctc agcccocccag
ING1 a : tagttggctg tgaggTCCCTT CGTGAAATGT CCTTATCAAT CCCCCTGGGA ACGATTGGTC
ING1 a : GCTGAGCGCG ATGAAAGCGG GCCTAGCGCA ATRACTGGTA TGGGCTGTGT TTTCCGCTGT
ING1 a : CTTCTTTTTT CTTTTTCGGG GAGGAGCGGG GGGGAGGGTG GACGAGTTGA TTTGAAACGC
ING1 a : TTCGGTCCGC TCGSCCTCCA GCCTTGGATT GGTTCCTCTC GCTGCTGGGG CGGGCCGTGC
ING1 a : TCTTCCGCCC TCGGCTGTGG TTGGTCTCC TCTTCCGCTC CCCCCCTCAA ATCCGGCATT
ING1 a : CCCATAGGCG GCGSCTCTCG GGGTCCGGGG CAGATCTCCC GCTGSCCTCC TCCCATTGG
ING1 a' : ----- -gagtaacc gataataggc CATTGTGCAC GCGACGAGA
ING1 a : CTGAGGCCCT GCGGGGTGTC GCCCCGGCCC CTTCCCCCGC TCAAGCCGGC CACTTTCGGG
ING1 a' : ATTCCAGAT ATAGCAGTAG CAGTGAATCC GSCCTGTGG CTTGGGGCCC GSCCTGCAGT
ING1 a : CCGGAAATTA TAGCAGTAGC AGTGAATCCC GSCCTGTGG CTTGGGGCCC GSCCTGCAGT
ING1 a' : TCGACCGCC TCCCGCACC CCGGGGCCG GCTCGGAGC AGTTTCAGGC CCACTCTTGG
ING1 a : TCGACCGCC TCCCGCACC CCGGGGCCG GCTCGGAGC AGTTTCAGGC CCACTCTTGG
ING1 a' : CTTGACCGAG GGTGGGGCCG CCGTGGCCG TGGAAACGGA TCCTGAA
ING1 a : CTTGACCGAG GGTGGGGCCG CCGTGGCCG TGGAAACGGA TCCTGAA

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5' unique exon ← 3' common exon

