

The Structure, Expression, and Function Prediction of *DAZAP2*, A Down-Regulated Gene in Multiple Myeloma

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In our previous studies, *DAZAP2* gene expression was down-regulated in untreated patients of multiple myeloma (MM). For better studying the structure and function of *DAZAP2*, a full-length cDNA was isolated from mononuclear cells of a normal human bone marrow, sequenced and deposited to Genbank (AY430097). This sequence has an identical ORF (open reading frame) as the NM.014764 from human testis and the D31767 from human cell line KG-1. Phylogenetic analysis and structure prediction reveal that *DAZAP2* homologues are highly conserved throughout evolution and share a polyproline region and several potential SH2/SH3 binding sites. *DAZAP2* occurs as a single-copy gene with a four-exon organization. We further noticed that the functional *DAZAP2* gene is located on Chromosome 12 and its pseudogene gene is on Chromosome 2 with electronic location of human chromosome in Genbank, though no genetic abnormalities of MM have been reported on Chromosome 12. The ORF of human *DAZAP2* encodes a 17-kDa protein, which is highly similar to mouse *Prtb*. The *DAZAP2* protein is mainly localized in cytoplasm with a discrete pattern of punctuated distribution. *DAZAP2* may associate with carcinogenesis of MM and participate in yet-to-be identified signaling pathways to regulate proliferation and differentiation of plasma cells.

Key words: multiple myeloma, *DAZAP2*, SH2/SH3, polyproline

Introduction

Given that there is a large amount of data on the sequence of new genes whose functions are yet unknown in public databases, and hence a great challenge to researchers in the functional study, bioinformatics has become a powerful tool for predicting gene's function. Gene expression patterns in diseases (including cancer) in combination with special analytical software do give scientists directions for further research in acquiring functional information of new genes. Multiple myeloma (MM) is a disorder characterized by the uncontrolled proliferation and accumulation of malignant plasma cells in the bone marrow (1). To identify the MM-associated genes and delineate their pathogenic correlation, we used cDNA profiling to assess the expression of 1,999 genes in bone marrow

mononuclear cells from MM patients and normal subjects. The levels of the most significantly down-regulated gene *DAZAP2* (deleted in azoospermia associated protein 2; ref. 2) were verified by Western blotting analysis of MM samples (7/11 or 63.6%), but all normal subjects (control) expressed *DAZAP2* (3). *DAZAP2* expression at both the mRNA and protein levels was negatively correlated with the pathogenesis of MM. The *DAZAP2* protein had originally been identified as an interacting protein of germ-cell-specific RNA-binding proteins DAZ (deleted in azoospermia; ref. 4). The published *DAZAP2* cDNA comes from a human testis cDNA library, located on Chromosome 2 with unknown function (4). To explore the functional significance of the down-regulated *DAZAP2* gene in MM, we isolated this gene, characterized its product from human bone marrow mononuclear cells of a normal subject, and searched the available databases. By suitable software, we identified its homologues and characterized its molecular phyloge-

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nies. Since the positions of functional and/or structural importance tend to be more conserved in evolution, we may find important functional region of *DAZAP2* by amino acid sequence alignment of 15 vertebrates' *DAZAP2* proteins. Furthermore, the expression pattern of *DAZAP2* may also provide hints of its function. All these analyses tried to shed new light on the function of the *DAZAP2* gene.

Results

Sequence analysis, molecular phylogeny and genomic organization of *DAZAP2*

A full-length *DAZAP2* cDNA was isolated from mononuclear cells of normal human bone marrow, se-

quenced and deposited to Genbank (AY430097). Figure 1 shows its nucleotide and predicted amino acid sequence. The *DAZAP2* gene contains a short 5' UTR, an ORF (open reading frame) from Nucleotides 84-590, and a relatively long 3' UTR plus poly(A) tail. Its ORF is predicted to encode a 17-kDa protein with 168 amino acids and is the same as those of the *DAZAP2* gene (NM_014764) from human testis (4) and D31767 (KIAA0058) from human cell line KG-1 (5). Human *DAZAP2* is highly similar to mouse *Prtb* (proline codon-rich transcript, brain expressed; ref. 6). Further sequence analysis reveals that *DAZAP2* contains a proline-rich region and several potential SH2 (YxxΨ) and SH3 (PxΨP) domain-binding motifs.

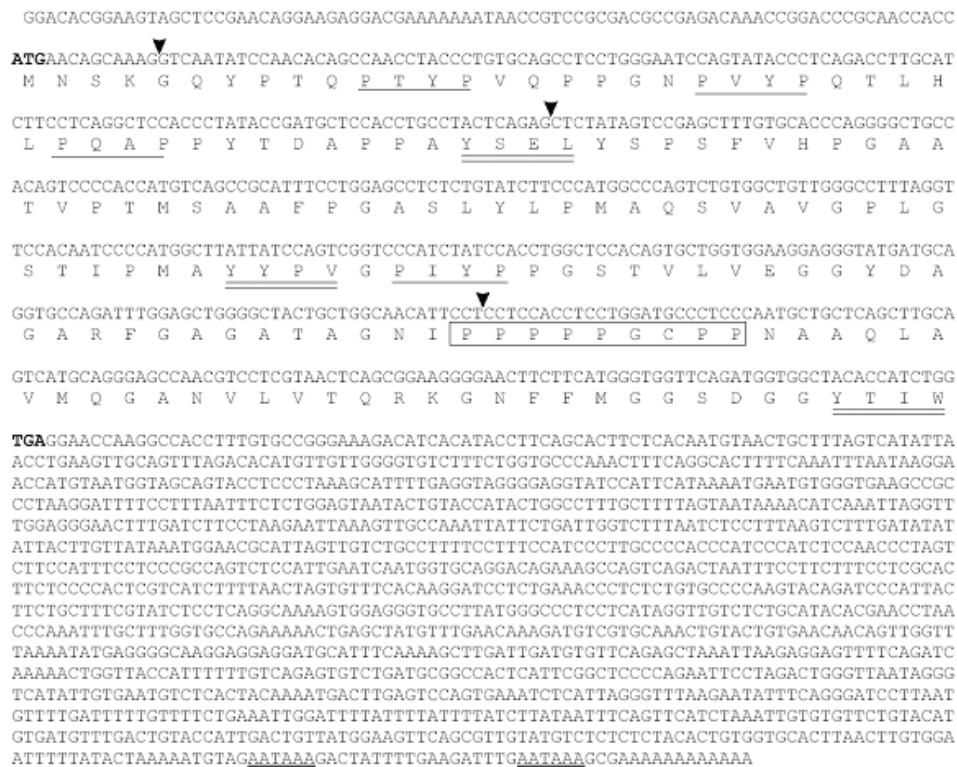


Fig. 1 The nucleotide and predicted amino acid sequence of *DAZAP2*. The cDNA was cloned from total RNA of mononuclear cells of normal bone marrow. The polyproline region is shown by a rectangle. Potential SH2 (YxxΨ) and SH3 (PxΨP) domain-binding motifs are underlined. The initiation and stop codons are in bold. Triangles denote exon boundaries. The 3'-polyadenylation signals are underlined.

As shown in Figure 2A, human *DAZAP2* has a highly related gene in two distant species, *Apis mellifera* (B1515614) and *Ciona intestinalis* (AY490098). This gene is highly conserved in vertebrates from zebrafish to human being whose homologues show a slow evolution. Comparison of 15 vertebrates'

DAZAP2 (Genbank IDs: AU297060, CB553719, XM.217038, NM.011873, BM967886, CD051977, CD467315, CD728885, AY490096, BC042215, BX078169, CB516510, AY490097, and BC059628) reveals that the polyproline region is a conserved structural motif (Figure 2B). Notably, its C-terminus

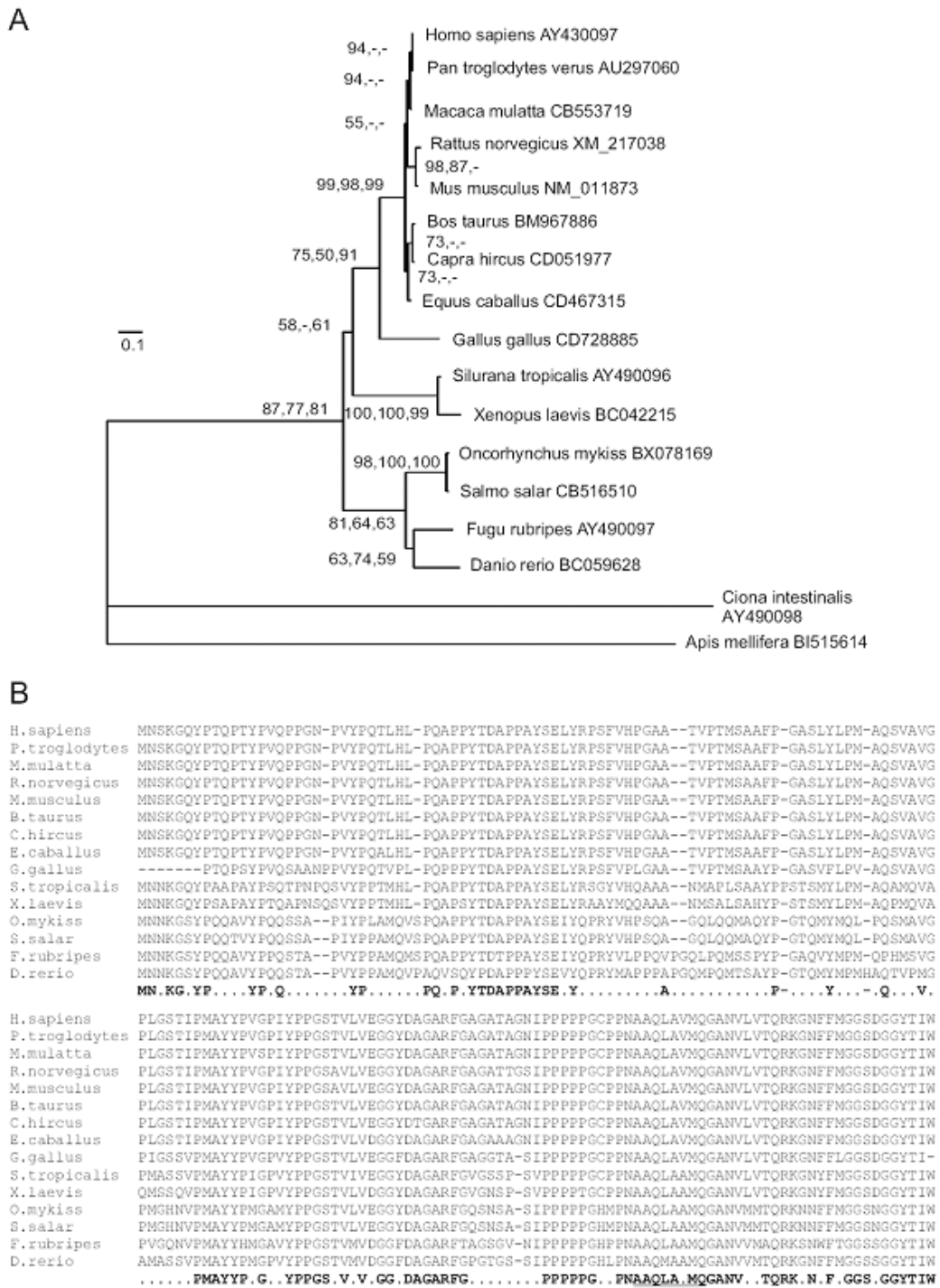


Fig. 2 Phylogenetic analysis and sequence alignment of *DAZAP2*. **A**. The tree topologies. The branch lengths, estimated by ML methods, denote nucleotide substitutions per nucleotide sites. The numbers on nodes are bootstrap test values. *Left*: ML at 1,000 puzzling steps for protein sequences under HKY85 model with five categories of Gamma substitution rate plus invariable sites (5G+I); *Middle*: ML at 1,000 puzzling steps for nucleotide sequences under JTT model with 5G+I; *Right*: MP at 200 replicates for protein sequences under default parameters. GenBank accession numbers are given along the tree. **B**. Sequence alignment of vertebrate *DAZAP2* proteins. Dashes denote gaps or missing residues. The bold amino acids at the bottom represent 90% consensus in the alignment. The 17 prolines are largely conserved and the N-terminus is less conserved than the C-terminus. The single α -helix (underlined) present in the C-terminal end is flanked by potential extended β -sheets (not shown).

DAZAP2 expression in human tissues

To define the pattern of *DAZAP2* expression in various tissues, a human multiple tissue Northern blot was performed. As shown in Figure 4, *DAZAP2* is expressed as a 1.9-Kb transcript in seven normal human tissues, including stomach, thyroid, spinal cord, lymph node, trachea, adrenal gland, and bone marrow. The results indicate a ubiquitous expression of the *DAZAP2* gene *in vivo* and suggest an important functional role of the *DAZAP2* gene in multiple tissues.

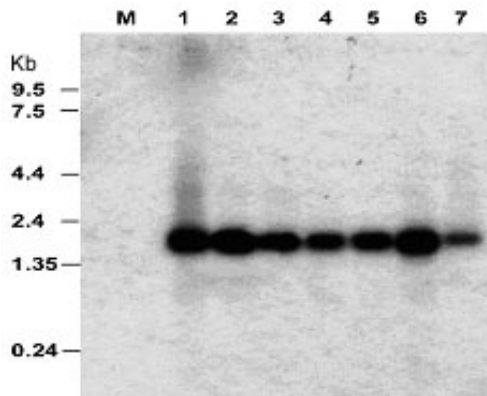


Fig. 4 Expression of *DAZAP2* in human tissues. A human multiple tissue Northern blot was hybridized with a *DAZAP2* cDNA probe. Lane M: size markers. Lanes 1-7: stomach, thyroid, spinal cord, lymph node, trachea, adrenal gland, and bone marrow. A single and abundant transcript of about 1.9 Kb is seen in all tissues. The intensity of β -actin mRNA bands was relatively constant (not shown).

Expression and intracellular localization of DAZAP2 protein

To verify the expression of DAZAP2 protein in eukaryotic cells, pEGFP-N1-DAZAP2 was transfected into COS7 cells. Western-blotting analysis detected a 44-kDa protein in cells transfected with pEGFP-N1-DAZAP2 and a 27-kDa protein in cells transfected with pEGFP-N1 alone by using an anti-GFP antibody. Thus, the DAZAP2 protein expressed in COS7 cells was about 17-kDa (Figure 5), a result that is consistent with ORF prediction and Western blot of native DAZAP2 protein expressed in the bone marrow (data not shown).

The intracellular localization of DAZAP2 protein was next examined by immunofluorescent microscopy. In control cells expressing GFP alone, the green fluorescence was distributed evenly in the cytosol and

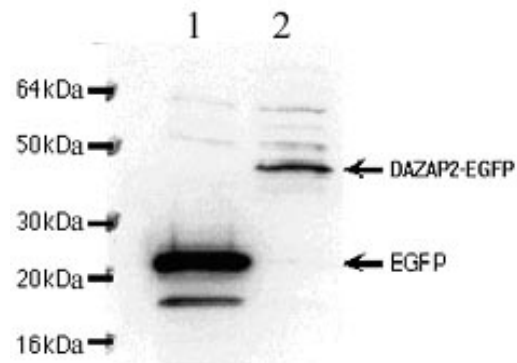


Fig. 5 Immunoblotting analysis of the fusion protein DAZAP2-EGFP expression. Lane 1: EGFP protein; the MW (molecular weight) of EGFP protein is about 27 kDa. Lane 2: DAZAP2-EGFP fusion protein; the MW of DAZAP2-EGFP fusion protein is about 44 kDa.

in the nucleus (Figure 6A). By contrast, COS7 cells expressing DAZAP2-GFP showed a largely cytosolic location with small distinctive punctated staining (Figure 6B, green color). The green fluorescence of DAZAP2-GFP fusion protein showed no apparent overlap with the staining of Golgi (Figure 6B, red color), indicating that DAZAP2 protein mainly resides in the cytoplasm with a discrete pattern of punctated structures.

Discussion

Multiple myeloma is a malignant disorder of plasma cells for which the molecular etiology is still unknown (1). The fact that the down-regulation of *DAZAP2* occurred at both the mRNA (2) and protein levels (3) provides a strong indication for its specific association with MM. A full-length 1,913 bp *DAZAP2* cDNA (AY430097) isolated from human bone marrow mononuclear cells has an identical ORF as the published *DAZAP2* from human testis library and KG-1 cell line. To explore the sequence features and functional significance of *DAZAP2*, we searched available databases and used three program packages and methods to identify its homologues and characterize its molecular phylogenies. A more detailed picture of *DAZAP2* molecular evolution was obtained when we isolated three closely related cDNA fragments of *DAZAP2* from three distant species (AY490096, AY490097, AY490098). The three program packages and methods we employed are TREE-PUZZLE, MEGA2 and T-coffee. TREE-PUZZLE is a quartet-based maximum-likelihood phylogenetic analysis tool, including a broad variety of evolutionary models (7),

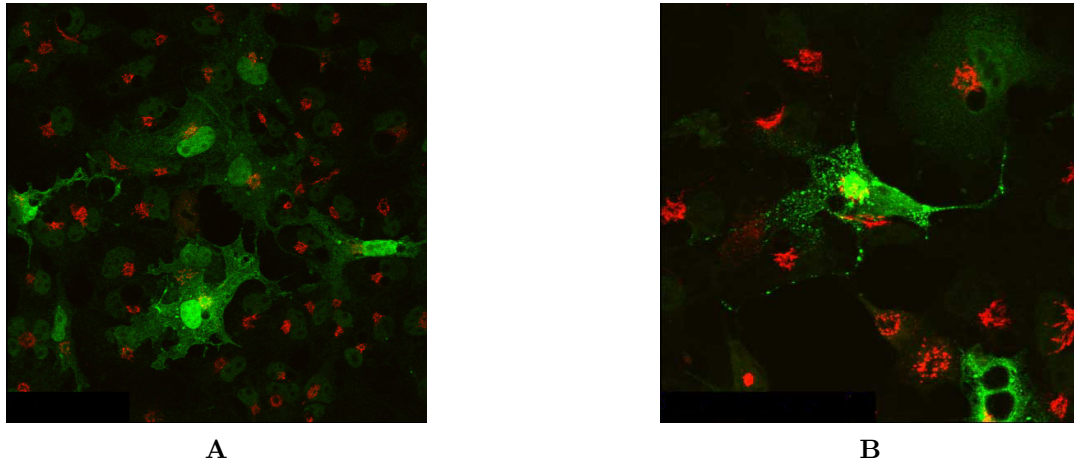


Fig. 6 Intracellular localization of DAZAP2 protein. COS7 cells were transfected with either pEGFP vector or DAZAP2-GFP plasmid and visualized under confocal microscopy. **A.** Expression of GFP in control cells. Note the green fluorescence of GFP is evenly distributed in the cytosol and the nucleus. **B.** The DAZAP2-GFP fusion protein (green) shows largely a cytosolic distribution with small discrete punctuated staining; its green fluorescence appears not to overlap with the staining of Golgi (red).

for example, models for DNA sequences (TN, ref. 8), models for protein sequences (JTT and VT, ref. 9, 10). MEGA2 is a set of software that can compute evolutionary distances based on the observed differences in amino acid and nucleotide sequences (11). The bootstrap test and the interior branch length tests can examine the reliability of the inferred phylogenies (12, 13). Most of the bootstrap test values on nodes of the tree topologies have a high score (Figure 2A), which elucidate the accuracy of the phylogenetic tree (14). From the result of the phylogenetic tree, in which human *DAZAP2* has a highly related gene in two distant species, *Apis mellifera* and *Ciona intestinalis*, we concluded that *DAZAP2* has an early origin. This suggests that *DAZAP2* might have vital cellular functions. T-coffee is a method for multiple alignment that provides a dramatic improvement in accuracy, and its resulting alignments are significantly more reliable as compared to the most commonly used alternatives (15). By this reliable method, we showed that *DAZAP2* was conserved throughout evolution, particularly in the C-terminus of the protein we defined a potentially important function domain for *DAZAP2*. Its vertebrate homologues virtually carried the same genomic organization and structural features (four exons and three introns; the others are 5' and 3' untranslated regions). Moreover, by nucleotide sequence alignment, we found that human has a functional gene on Chromosome 12 and its pseudogene on Chromosome 2, yet Chromosome 12 has not been implicated to harbor MM-associated gene abnormalities (1, 16).

The pseudogene protein expression was inactivated by a nonsense mutation (substitution TAG for TAC). Structure prediction revealed multiple sequence motifs in *DAZAP2* including several potential SH2 and SH3 binding sites, which also conserved throughout evolution. Protein-protein interactions mediated by SH2 and SH3 domains have been shown to play a central role in multiple signaling pathways that regulate cell proliferation and differentiation (17). For instance, the SH3-SH2-SH3 Grb2 adapter links the receptors to the Ras pathway (18); Nck/Dock, composed of three SH3 domains and one SH2 domain, links cell surface receptors to the actin cytoskeleton (19). Hence, it is of great interest to uncover the role of *DAZAP2* in such signaling events in regulating the proliferation of tumor cells in MM, should the disruption of these interactions result in loss-of-function phenotypes.

As shown here and elsewhere (4, 5), *DAZAP2* was broadly expressed in normal tissues and cell lines. Given this ubiquitous occurrence of *DAZAP2* and testis specificity of *DAZ*, their interaction is unlikely to play a critical role in the pathogenesis of MM, although the expression of *DAZ* was not assessed. It is of further interest to note the mouse homologue *Prtb*, which was first cloned from mouse brain (6) and located in the cytoplasm (20). Its expression is increased in osteoclast cell line (MC3P3-E1) upon adhesion to polystyrene with or without surface-adsorbed serum proteins, suggesting that *Prtb* is involved in cell adhesion-mediated function (20). The human 17-kDa

DAZAP2 protein was also mainly localized in cytoplasm as mouse Prtb protein. Nevertheless, whether human *DAZAP2* and mouse or rat *Prtb* genes are functionally equivalent in the setting of MM pathogenesis and progression remains to be elucidated.

Materials and Methods

Bone marrow sample

Normal control bone marrow samples were obtained from the Department of Hematology of Xiangya Hospital, Central South University, China.

Total RNA preparation

Mononuclear cells were isolated from bone marrow samples by the lymphocyte isolation solution. Total RNA was prepared from the bone marrow mononuclear cells using Trizol kit (Invitrogen, Carlsberg, USA). RNA concentration was determined on a UV spectrophotometer. The quality of RNA was assessed by electrophoresis on 1.2% agarose gel.

Isolation and sequencing of *DAZAP2* cDNA from bone marrow

Based on the sequence of *DAZAP2*, gene-specific primers (GSPs) in either sense (S) or antisense (A) were designed to isolate the *DAZAP2* gene from mononuclear cells of bone marrow of normal control by the method of rapid amplification of cDNA ends (RACE). For the 5' RACE, 2.5 μ g of total RNA was reverse-transcribed with the 5'-end phosphorylated GSP (RT1): 5'-ACAACATGTGTCT-3'. The RNA strand of DNA-RNA duplex was digested and the single cDNA strand was circularized as described in the 5'-Full RACE kit (TaKaRa Bio Inc., Dalian, China) (21). The cDNA was then amplified sequentially with two different pairs of GSPs as follows: S1, 5'-GGTATGATGCAGGTGCCAG-3'/A1, 5'-GACCGACTGGATAAT-3'; and S2, 5'-CGTCCTCGTAACTCAGCGG-3'/A2, 5'-GATACA GAGAACTACAGG-3'. For the 3' RACE, 1.0 μ g of total RNA was reverse-transcribed with supplied oligo dT-3 sites adapter primer, and then amplified once with adapter primer 5'-CTGATC TAGAGGTACCGGATCC-3' and GSP (S3), 5'-GGTGGCTACCCATCTGGTG-3', as described in the 3'-Full RACE kit (TaKaRa Bio Inc.). PCR products were analyzed on 1.5% agarose gel. The 5'

and 3' RACE products were purified, subcloned into the pEGM-Teasy vector (Promega, San Luis Obispo, USA), and then sequenced on an automated ABI377 sequencer.

Sequence analysis, structure prediction and phylogenetic analysis

Sequence analysis and structural prediction were performed by searching public databases on the web-site servers of NCBI (<http://www.ncbi.nlm.nih.gov/BLAST/>), NetStar (<http://www.cbs.dtu.dk/services>) and Scan Prosite (<http://us.expasy.org/cgi-bin/scanprosite>). The tree topologies were estimated using maximum likelihood (ML; ref. 7) and maximum parsimony (MP; ref. 11). The sequence alignment was carried out using T-coffee (15) followed by manual inspection.

Analysis of *DAZAP2* expression in human tissues

The human multiple tissue Northern blot with poly(A)⁺ RNA prepared from various tissues (Clontech, Palo Alto, USA) was hybridized with the ³²P-labeled *DAZAP2* cDNA probe. The probe spans the entire ORF of *DAZAP2*. Human β -actin cDNA was used as a control probe. The blot was hybridized and washed under highly stringent conditions.

Transfection, immunoblotting and immunolocalization

The recombinant pEGFP-*DAZAP2* expression vector was constructed as follows. The primers for *DAZAP2* amplification were 5'-CGGAATTCGCCGCCACCATGAACAGCAAAGGTC-3' (sense) and 5'-GCGGATCCCGCCAGATGGTGTAGCCACCATC-3' (antisense), which contain *EcoR* I and *Bam*H I sites (in bold), respectively. The ORF of *DAZAP2* was amplified by PCR using the cDNA template derived from mononuclear cell RNA of normal bone marrow. PCR products were digested with *EcoR* I and *Bam*H I followed by ligation to the pEFGP-N1 vector (Clontech) and transformation into *E. coli* DH5 α cells. The ORF and fusion with the *GFP* gene in the pEGFP-N1-*DAZAP2* vector were verified by *EcoR* I and *Bam*H I digestion followed by DNA sequencing.

COS7 cells (ATCC) were used as a host for

DAZAP2-GFP fusion protein expression. Transfection was performed with LipofectAMINE-2000 using the manufacturer's protocol (Invitrogen, Carlsbad, USA). For immunoblotting analysis, cells were seeded in a 3-cm dish, transfected with DAZAP2-GFP plasmid and cultured for 24 h. The cells were lysed and the total proteins (10 μ g) were separated on a 4%-20% precast Tris-glycine gel, and were then electro-transferred to PVDF (polyvinylidene difluoride) membranes. The protein bands were visualized by using an ECL-plus kit (Amersham, Piscataway, USA). Immunostaining was done as above with the primary anti-GFP antibody (Clontech; ref. 22). Briefly, cells grown on coverslips were fixed with 3.7% paraformaldehyde for 30 min, permeated with 0.1% Triton X-100 for 10 min, and immunostained with the respective antibodies for 1 h after blocking with 5% BSA (bovine serum albumin) for 30 min. Cells were then stained with cy2- or cy3-labeled secondary antibodies (The Jackson Laboratory, Bar Harbor, USA). After several times of washing in PBS (phosphate-buffered saline), coverslips were mounted on slides with anti-fade buffer and observed under confocal microscopy.

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