The Human *CD53* Gene, Coding for a Four Transmembrane Domain Protein, Maps to Chromosomal Region 1p13

M. Eugenia Gonzalez, Fernando Pardo-Manuel de Villena, Elena Fernandez-Ruiz, Santiago Rodriguez de Cordoba, and Pedro A. Lazo

Unidad de Genética Molecular (CSIC), Centro Nacional de Biología Celular y Retrovirus, Instituto de Salud Carlos III, 28220 Majadahonda, Madrid, Spain,

Centro de Investigaciones Biológicas, CSIC, Velazquez 144, and Servicio de Inmunología, Hospital de la Princesa, Universidad Autónoma de Madrid, 28006 Madrid, Spain

The rat OX44/CD53 protein is the prototypic member of a "novel" family of proteins. These proteins are characterized by four highly hydrophobic transmembrane domains, two small extracellular domains, one of which is extensively N-glycosylated, and both the amino and the carboxy terminus intracytoplasmic. The function of these proteins remains elusive and several possible functions have been suggested depending on the experimental system used, but all of them are somehow implicated in the control of cell proliferation by binding to an unknown ligand. In rat, CD53 antigen is detected on several mature cell types of the hematopoietic system, including macrophages, monocytes, granulocytes, leukocytes, and B and T cells, as well as osteoblasts and osteoclasts. In rat macrophages we have found that the CD53 protein also stimulates the production of nitric oxide (submitted for publication), which is implicated in septic and hemorrhagic shocks. Preliminary data based on somatic cell hybrids indicated that the human CD53 gene is located on chromosome 1. To confirm these data and to determine precisely the location of human *CD53* gene in chromosome 1, we have used fluorescence *in situ* hybridization (FISH). Using a rat OX-44 cDNA probe, a human genomic clone, hROX44, containing a 17-kb-long insert was isolated and demonstrated to include the complete human *OX-44/CD53* gene. FISH was performed as previously reported. Briefly, peripheral blood lymphocytes were obtained from a human male by Lymphoprep gradient centrifugation and stimulated with phytohemagglutinin for 72 h. Colchicine (0.2 μ g/ml) was added to the culture and kept for 1 h. The X-hROX44 genomic clone was labeled by nick-translation with digoxigenin-11-dUTP. After 1 h of suppression hybridization at 37°C, hybridization was performed in 50% formamide containing 10% dextran sulfate, 2 $\times$ SSC, and 50 mM phosphate, pH 7.0, 2 ng of labeled probe, and a 500-fold excess of both *Cot1* DNA (fast reassociation fraction of human genomic DNA) and sonicated human placental DNA at 37°C overnight. Washings and detection with TRITC-conjugated antibodies were performed as described. Chromosomes were counterstained with 75 ng/ml of 4',6-diamino-2-phenylindole in antifade medium. After the fluorescence microscopy, GTG-banding was performed as described. A total of 35 metaphase spreads were analyzed for the presence of fluorescent spots; 89% of them showed hybridization signals on the pericentromeric region of the short arm of chromosome 1, in one (39%) or both (61%) chromatids (Fig. 1). Hybridization was highly specific, and no other chromosomes showed spots systematically. This nonrandom distribution was analyzed after GTG-banding of metaphase spreads to determine the precise location of the *CD53* gene (Figs. 1A and 1B). The results of this analysis are summarized in Fig. 2; 62 and 19% of the hybridization spots were localized on the 1p13 and 1p21 bands, respectively.

From these data we conclude that the human *CD53* gene maps to 1p13, proximal to 1p21.

FIG. 1. Fluorescence *in situ* hybridization on human chromosomes with a digoxigenin-labeled 17-kb-long genomic DNA probe of the human *CD53* gene. Arrows indicate the specific site of hybridization. (A) Rhodamine detection of digoxigenin-labeled probe and DAPI counterstaining in human mitotic cells and GTG-banding of the same metaphase spread. (B) Examples of chromosome 1 from other cells.

Chromosomal region 1p13 contains some other lymphoid genes like *CD2* and its ligand *CD58* (12). Whether the three lymphoid genes are linked, forming a cluster, is currently unknown. Interestingly, *CD53* and *CD2* antigens coprecipitate with antibodies against *CD53* in rat T lymphocytes and natural killer cells, suggesting that they are associated on the cell membrane (3). Furthermore, in these cells the response to antigenic stimulation of *CD2*⁺ cells is increased severalfold if they are *CD53*⁺ (3). These observations raise the possibility that these two genes might have a coordinated expression in T-cells. This interaction and the pattern of OX-44/*CD53* expression at different stages of hematopoietic development and in very different mature cell types make the regulation of *CD53* gene transcription an interesting problem. Chromosomal translocations specific for leukemias have been very instrumental in the discovery of genes implicated in oncogenesis. There are some translocations involving the 1p13 region in different types of tumors (7).

Among them is t(1;22) (p13;q13), which is detected in 70% of the cases of M7 acute megakaryocytic leukemia in young children and has a fatal prognosis (6). Because of the possible effects of *CD53* on cell proliferation, it might be implicated in the oncogenic phenotype. Two *RAS-related* genes (10), *NRAS* and *RAP1A*, and other genes, such as *NGFB* and *TSH*, which might have effects on the growth properties of a cell, also map to 1p13. Which genes are affected by the t(1;22) (p13;q13) translocation is not known, but a better characterization of this region should help to clarify this point.

FIG. 2. Assignment of the *CD53* gene to 1p13. The idiogram of human chromosome 1 schematically indicates the intrachromosomal

ACKNOWLEDGMENTS:

This work has been supported by grants from Fundación Ramón Areces and CICYT (SAL90-0209 and SAL91-0043) to P.A.L. and FIS (92-0889) to S.R.C.