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## Substitution of Alanine<sup>543</sup> with a Threonine Residue at the Carboxy Terminal End of the $\beta$ -Chain Is Associated with Thermolabile Hexosaminidase B in a Jewish Family of Oriental Ancestry

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**Thermolabile forms of the lysosomal enzyme  $\beta$ -hexosaminidase B (Hex B), likely to result from different genetic defects, have been described. Ten individuals in five generations of a family of Oriental Jewish ancestry were identified biochemically as carriers of a thermolabile Hex B form. The  $\beta$ -chain thermolability was found to be associated with the presence of a G  $\rightarrow$  A transition at nucleotide 1627 of the HEXB gene causing the substitution of Ala<sup>543</sup> with a threonine. The same mutation was detected in an unrelated Oriental Jew whose Hex B was heat labile. Since thermolabile Hex B has been shown to occur more frequently among Jews of Oriental origin, the Ala<sup>543</sup>  $\rightarrow$  Thr mutation may be the common mutation associated with  $\beta$ -chain thermolability in this ethnic group.** © 1995 Academic Press, Inc.

Two major forms of the lysosomal enzyme  $\beta$ -hexosaminidase ( $\beta$ -Hex, E.C. 3.2.1.52), differing in their physicochemical properties and substrate specificity are normally found in human tissues. Hex A, the enzyme responsible for the catabolism of ganglioside G<sub>M2</sub>, is a heat labile heterodimer composed of  $\alpha$  and  $\beta$  subunits coded by separate genes (HEXA and HEXB, respectively), while Hex B is a heat stable homopolymer composed exclusively of  $\beta$ -chains. Serum contains a third major form, Hex I, which is also composed of  $\beta$ -chains. Deficiency of Hex A activity caused by mutations in the HEXA and HEXB genes results in a group of neurodegenerative disorders collectively known as G<sub>M2</sub>-gangliosidoses (1). For the diagnosis of G<sub>M2</sub>-gangliosidosis and for carrier identification the relative levels of Hex A and B in tissues and body fluids are measured by a simple heat inactivation assay which takes advantage of the different thermal stability of the two enzyme

forms (1). This assay, however, becomes unreliable if Hex B is no longer thermostable. There have been several reports in the literature describing the existence of thermolabile forms of the  $\beta$ -chain (2–8). Depending upon the case, the thermolabile  $\beta$ -chain was found to form both Hex A and Hex B, albeit characterized by increased thermolability (5, 8), to associate only with the  $\alpha$ -chain resulting in the total absence of Hex B (Hexosaminidase Paris variant) (2), or to lose its ability to associate with the  $\alpha$ -chain, resulting in deficiency of Hex A activity (3, 4, 7, 9). These observations suggest that the  $\beta$ -chain thermolability is the result of more than one genetic defect.

Studies in the Jewish population have shown that a thermolabile Hex B form is found with the highest frequency in Jews of Oriental origin (10, 11). In this paper we report the biochemical characterization of a thermolabile Hex B form found in a family of Oriental Jewish descent and the identification of a mutation in exon 14 of the HEXB gene likely to be associated with the  $\beta$ -chain thermolability.

### MATERIALS AND METHODS

Five generations of a family of Oriental Jewish descent (Iran) carrying the thermolabile Hex B trait were studied (see the pedigree in Fig. 1A). The proband, individual IV-4 in the pedigree, was originally identified as a potential carrier of a thermolabile  $\beta$ -chain through a community screening program for Tay-Sachs disease carriers. By the standard heat inactivation assay he was found to have only 9–12% residual heat stable  $\beta$ -Hex activity in serum and leukocytes. We have also studied another unrelated Jewish individual of Middle Eastern ori-



bilon S (Millipore, Bedford, MA) and hybridized overnight at 42°C to the following <sup>32</sup>P-labeled oligonucleotides: 5' GAATAGCTGCACAACCT (normal probe) and 5' GAATAGCTACACAACCT (mutant probe). The filters were then washed in 6× SSC, 0.1% SDS, at 50°C for 10 min and autoradiographed.

Site-directed mutagenesis of the  $\beta$ -chain cDNA was performed using the mutagenic oligonucleotide 5' GAGGTTGTGTAGCTATTCC as described (15). The wild-type and the mutant  $\beta$ -chain cDNA were subsequently subcloned into the eukariotic expression vector pSVL (15) and the resulting constructs purified by CsCl-ethidium bromide isopycnic centrifugation. Each construct (5  $\mu$ g) was transfected into COS-7 cells using the calcium phosphate coprecipitation method (16). As an internal control for efficiency, 2  $\mu$ g of the indicator plasmid pBLCAT2 containing the chloramphenicol acetyl transferase gene (CAT) (17, 18) was included in each transfection experiment. After 48 h, the cells were harvested and the  $\beta$ -Hex and CAT activity (19) was determined in cell extracts. In some experiments, 20  $\mu$ M leupeptin was added to the culture medium after transfection (20).

## RESULTS

### *Biochemical Characterization of $\beta$ -Hex*

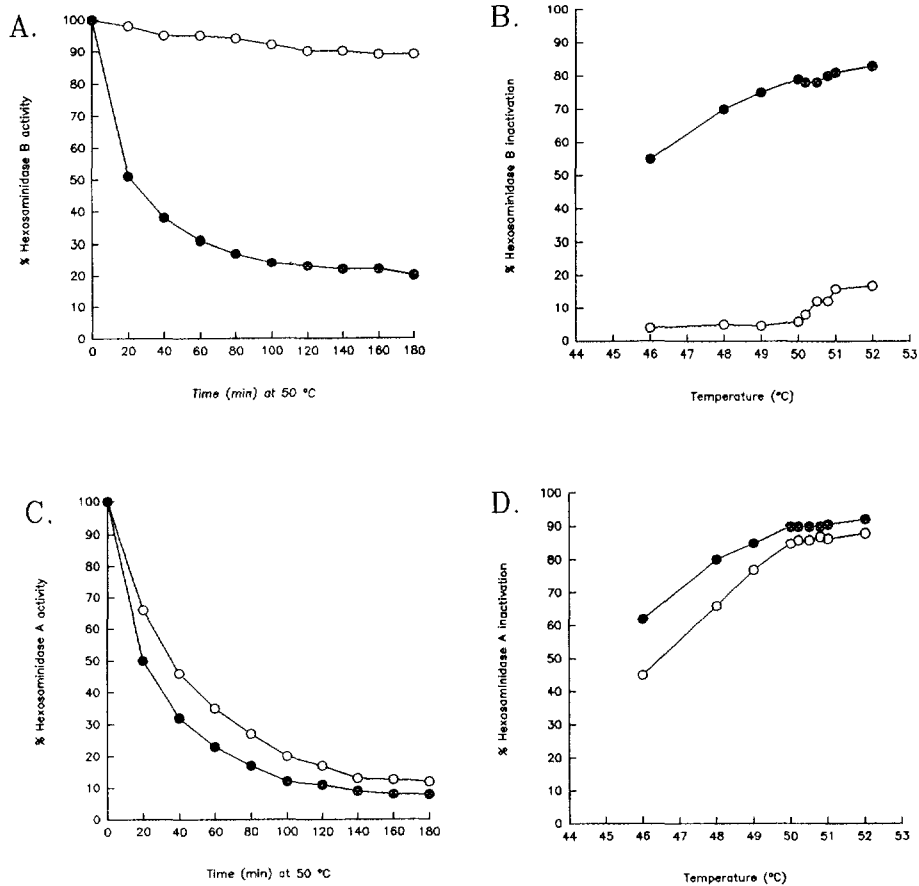
In heat inactivation assays the serum, leukocytes, and cultured fibroblasts of individual IV-4 (see pedigree in Fig. 1A) were found to have reduced levels of Hex B. Separation of the  $\beta$ -Hex forms by ion-exchange chromatography showed that the levels of Hex B in the fibroblasts of individual IV-4 were similar to those found in the fibroblasts of a control individual, suggesting that the apparently reduced level of heat stable  $\beta$ -Hex in IV-4 was due to the presence of a thermolabile Hex B. Thermal stability studies using the Hex B fraction isolated by ion-exchange chromatography showed that at 50°C the enzyme from IV-4 lost 50% of its activity in the first 20 min. This was followed by a slower inactivation phase so that after 180 min only 25% of the original activity was heat stable. Under the same conditions, the control Hex B was stable (Fig. 2A). Heat inactivation assays at different temperatures from 46 to 52°C confirmed the intrinsic thermolability of the Hex B from IV-4 (Fig. 2B). Similar results were obtained with Hex I (results not shown). The Hex A isolated from IV-4 also appeared to be slightly more thermolabile than the control Hex A at 50°C (Fig. 2C). This became more evident when the stability of

Hex A was studied as a function of the temperature. As shown in Fig. 2D, at 46°C the control Hex A lost about 40% of its activity in 2 h, while under the same conditions the Hex A isolated from IV-4 lost 60% of the original activity.

We then studied five generations of the family of IV-4 as seen in Fig. 1A and were able to identify nine additional individuals who appeared to have inherited the thermolabile Hex B trait (see Fig. 1A). These included the oldest member of the family (I-1), a 104-year-old man. The  $\beta$ -Hex elution profiles obtained by ion-exchange chromatography of the sera of all the family members tested were essentially identical, again suggesting that those individuals who had reduced levels of Hex B in heat inactivation assays actually had normal amounts of the enzyme. As shown in Table 1, the Hex B fraction isolated from the sera of III-1, IV-1, IV-3, IV-6, and IV-8 was stable after 180 min at 50°C, while the enzyme isolated from the remaining family members was thermolabile and lost, depending upon the individual, about 55 to 75% of the original activity. Similar results were obtained with the Hex I fraction (Table 1). The biochemical results also suggested that the 10 individuals with heat labile Hex B must be heterozygous for the allele causing  $\beta$ -chain thermolability, otherwise no residual thermostable Hex B activity would have been found after 2 h at 50°C.

### *Molecular Analysis of the $\beta$ -Chain Coding Sequences*

To understand the molecular basis of the Hex B thermolability, we sequenced the  $\beta$ -chain cDNA amplified from the mRNA isolated from the fibroblasts of IV-4. A G  $\rightarrow$  A transition at position 1627 of the coding sequence was the only base alteration found in the entire  $\beta$ -chain cDNA. Sequence analysis of PCR-amplified genomic sequences containing exon 14 confirmed the presence of the G<sup>1627</sup>  $\rightarrow$  A mutation in IV-4 (Fig. 3). This mutation was found in about half of the clones analyzed, which was consistent with the biochemical data suggesting a carrier status for the thermolabile allele. As a result of the mutation, Ala<sup>543</sup>, which is located at the C-terminal end of the enzyme, is substituted by a threonine. Analysis of the Ala<sup>543</sup>  $\rightarrow$  Thr mutation in the HEXB gene of the other family members showed that this mutation was carried only by those individuals whose Hex B was thermolabile (see Figs. 1A and 1B). The same Ala<sup>543</sup>  $\rightarrow$  Thr mutation was also found in another Middle Eastern Jewish individual unrelated to the family described above, whose bio-



**FIG. 2.** Thermal inactivation (A) and thermal stability at different temperatures (B) of Hex B isolated from the fibroblasts of IV-4 (●) and of a control individual (○). The Hex B fraction isolated by ion-exchange chromatography was incubated for the indicated periods of time at 50°C (A) or for 2 h at the indicated temperatures (B) and the residual activity measured with MUG. (C and D) Same as A and B, except that the experiments were performed with the Hex A fraction isolated from the fibroblasts of IV-4 (●) and of a control individual (○).

chemical data had suggested that he may also be a carrier of heat labile  $\beta$ -chain (Fig. 4).

A mutant  $\beta$ -chain cDNA containing the  $G^{1627} \rightarrow A$  mutation was prepared by site-directed mutagenesis and cloned into the eukariotic expression vector pSVL. Transfection of two independent mutant constructs in COS-7 cells did not result in the expression of any Hex B activity over background. Under the same conditions, the Hex B activity in COS cells transfected with the wild-type  $\beta$ -chain was on average 13-fold over the activity of mock-transfected cells. Cotransfection of each mutant construct separately with the plasmid containing the wild-type cDNA resulted in the expression of thermostable Hex B activity only. Changes in the amount of DNA transfected and in the method of transfection as well as addition of 20  $\mu$ M leupeptin to the medium after transfection (20) did not result in expression of any mutant Hex B activity. Cotransfection of the mutant

$\beta$ -chain cDNA with the  $\alpha$ -chain cDNA also failed to result in increased Hex A activity. The amount of CAT activity expressed in the cells transfected with the mutant constructs and with the wild-type plasmid was comparable, indicating that the lack of expression of the mutant enzyme was not due to a poor efficiency of transfection. Also sequence analysis of the entire mutant cDNA did not reveal the presence of any other alteration which could explain the lack of expression in COS cells.

## DISCUSSION

We have identified 10 individuals belonging to five generations of a Jewish family of Middle Eastern origin in which the allele conferring heat lability to the Hex  $\beta$ -chain had segregated. The biochemical results indicated that they must be heterozygotes for the heat labile  $\beta$ -chain allele because of the pres-



