Dominant Expression of Type III Hyperlipoproteinemia
Pathophysiological Insights Derived from the Structural and Kinetic Characteristics of ApoE-1(Lys^146→Glu)

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Abstract

Type III hyperlipoproteinemia is characterized by delayed chylomicron and VLDL remnant catabolism and is associated with homozygosity for the apoE-2 allele. We have identified a kindred in which heterozygosity for an apoE mutant, apoE-1(Lys^146→Glu), is dominantly associated with the expression of type III hyperlipoproteinemia. DNA sequence analysis of the mutant apoE gene revealed a single-point mutation that resulted in the substitution of glutamic acid (GAG) for lysine (AAG) at residue 146 in the proposed receptor-binding domain of apoE. The pathophysiological effect of this mutation was investigated in vivo by kinetic studies in the patient and six normal subjects, and in vitro by binding studies of apoE-1(Lys^146→Glu) to LDL receptors on human fibroblasts and to heparin. The kinetic studies revealed that apoE-1(Lys^146→Glu) was catabolized significantly slower than apoE-3 in normals (P < 0.005). In the proband, the plasma residence times of both apoEs were substantially longer and the production rate of total apoE was about two times higher than in the control subjects. ApoE-1(Lys^146→Glu) was defective in interacting with LDL receptors, and its ability to displace LDL in an in vitro assay was reduced to 7.7% compared with apoE-3. The affinity of apoE-1(Lys^146→Glu) to heparin was also markedly reduced compared with both apoE-2(Arg^135→Cys) and apoE-3. These abnormal in vitro binding characteristics and the altered in vivo metabolism of apoE-1(Lys^146→Glu) are proposed to result in the functional dominance of this mutation in the affected kindred. (J. Clin. Invest. 1995. 96:1100–1107.) Key words: atherosclerosis · dysbetalipoproteinemia · in vivo kinetics · low density lipoprotein (apoB, apoE) receptor binding · heparin binding

Introduction

Human apoE is a constituent of several classes of plasma lipoproteins and plays an important role in modulating the catabolism of chylomicron and VLDL remnants by binding to LDL (apoB, apoE) and putative apoE (remanent) receptors on cell membranes. DNA sequence analysis of the apoE-2 allele revealed a single-point mutation that resulted in the substitution of glutamic acid (GAG) for lysine (AAG) at residue 146 in the proposed receptor-binding domain of apoE. The pathophysiological effect of this mutation was investigated in vivo by kinetic studies in the patient and six normal subjects, and in vitro by binding studies of apoE-1(Lys^146→Glu) to LDL receptors on human fibroblasts and to heparin. The kinetic studies revealed that apoE-1(Lys^146→Glu) was catabolized significantly slower than apoE-3 in normals (P < 0.005). In the proband, the plasma residence times of both apoEs were substantially longer and the production rate of total apoE was about two times higher than in the control subjects. ApoE-1(Lys^146→Glu) was defective in interacting with LDL receptors, and its ability to displace LDL in an in vitro assay was reduced to 7.7% compared with apoE-3. The affinity of apoE-1(Lys^146→Glu) to heparin was also markedly reduced compared with both apoE-2(Arg^135→Cys) and apoE-3. These abnormal in vitro binding characteristics and the altered in vivo metabolism of apoE-1(Lys^146→Glu) are proposed to result in the functional dominance of this mutation in the affected kindred. (J. Clin. Invest. 1995. 96:1100–1107.) Key words: atherosclerosis · dysbetalipoproteinemia · in vivo kinetics · low density lipoprotein (apoB, apoE) receptor binding · heparin binding

Methods

Patient characterization

The heterozygous proband of the apoE-1(Lys^146→Glu) kindred has been described in detail previously (23). Briefly, he is a 50-y-old white male

1. Abbreviations used in this paper: HLP, hyperlipoproteinemia; RT, residence time.
with a history of hyperlipidemia and coronary artery disease. Physical findings include palmar xanthomas, tuberous xanthomas on his extremities, and xanthelasma. At the time of the study, he was without lipid-lowering medications, and his plasma total cholesterol value was 404 mg/dl with a VLDL cholesterol of 284 mg/dl (Table 1). His total plasma apoE concentration was 50 mg/dl, his apoE-1(Lys146→Glu) level being 36.5 mg/dl and his apoE-3 level 13.5 mg/dl. The ratio of apoE-1 to apoE-3 was determined by an isoelectric focusing gel of the proband's plasma, followed by immunostaining with an apoE antibody and subsequent densitometric scanning (23).

**DNA isolation**
Genomic DNA was purified from white blood cells using an automated nucleic acid extractor (model 340A; Applied Biosystems Inc., Foster City, CA).

**Enzymatic amplification of genomic DNA**
1 μg each of genomic DNA from a control and the apoE-1(Lys146→Glu)/apoE-3 subject was amplified with 5 U of Taq DNA polymerase (Perkin-Elmer Corp., Norwalk, CT) as described (21), using an automated DNA thermal cycler (Perkin-Elmer Corp.). The reaction mixture was subjected to 30 cycles of denaturation at 94°C for 1 min, annealing at 62°C for 1 min, and polymerization at 72°C for 1 min.

**DNA sequence analysis**
DNA amplified by the automated polymerase chain reaction was digested with the restriction endonucleases HindIII and EcoRI (New England BioLabs, Beverly, MA), according to the manufacturer's recommendations. Fragments were isolated from low melting point agarose (Bethesda Research Laboratories, Gaithersburg, MD) and subcloned into M13 vectors mp18 and mp19 (Bethesda Research Laboratories; reference 26). Sequencing was performed with the dideoxynucleotide chain termination method (27) using T7 DNA polymerase (Sequenase; United States Biochemical Corp., Cleveland, OH).

**In vivo kinetic studies**
Study subjects. In addition to the patient described above, six healthy young adults, three males and three females, aged 19–23, were studied as controls. The six subjects had an apoE-3/3 phenotype, normal lipid levels for age and sex, normal fasting glucose concentrations, normal kidney, liver, and endocrine functions, and were taking no medications (Table 1). All normal volunteers and the proband from the Harrisburg kindred were studied as in-patients at the Clinical Center of the National Institutes of Health and gave informed consent to the study protocol, which was approved by the Institute Review Board of the National Heart, Lung, and Blood Institute.

apoE isolation. apoE-1(Lys146→Glu) was isolated from the proband, whereas apoE-2(Arg188→Cys) and apoE-3 were purified from individuals homozygous for the respective alleles. VLDL was obtained by preparative ultracentrifugation (28, 29) and delipidated with chloroform/methanol (2:1). Heparin affinity and Sephacryl S-200 gel permeation chromatography were used to separate apoE from the other apo (30). Briefly, resolubilized VLDL was applied to a 2.5×48-cm column packed with heparin-Sepharose in 5 M urea, 25 mM NaCl, and 2 mM sodium phosphate (pH 7.7). Using a linear NaCl gradient (25–220 mM) at 4°C, apoE-1(Lys146→Glu) was recovered at a concentration of 50 mM NaCl, while apoE-3 was eluted with 100 mM NaCl. Finally, samples were dialyzed against 10 mM ammonium bicarbonate.

**Iodination.** apoE was iodinated by the iodine monochloride method (31), with previously described modifications (32). The amount of iodine monochloride added was calculated to result in less than 1 mol iodine introduced per 2 mol of protein. The iodination efficiency ranged from 15 to 20%. To remove potentially damaged apoE, radiolabeled apoE was reassociated with autologous plasma by incubation at 37°C for 30 min. The plasma sample was adjusted to a density of 1.21 grams/ml with potassium bromide. The supernatant was isolated by ultracentrifugation and dialyzed at 4°C against PBS. The recovery of radioactivity in the 1.21-gm/ml top fraction was 70% of the starting activity. Assuming that recovery of radioactivity corresponded to the recovery of apoE mass, the specific activities could be estimated with −7,500 cpm/mg. HSA was then added to a final concentration of 1%. The solution was sterilized by filtration through a 0.22-μm filter (Millipore Corp., Bedford, MA), and testing was performed for pyrogens and sterility before injection.

**Study protocol.** The study subjects were placed on a defined isoenergy diet (42% carbohydrate, 42% fat, 16% protein, 200 mg cholesterol/1,000 kcal, ratio polyunsaturated/saturated fat 0.2) 3 days before the start of the study. 1 d before injection, the subjects began to receive 900 mg potassium iodide per day given in equal doses three times a day with meals; this was continued through the end of the study. After the injection of 111-I-apoE-3 and 113-I-apoE-1(Lys146→Glu), blood samples were obtained after 10 min, at 1, 3, 6, 12, 18, and 24 h, and every day until day 7. The exact amount of apoE injected was not calculated but was estimated to be <1% of the total pool size and thus did not significantly affect steady state conditions. Samples were collected into tubes containing 0.1% Na2EDTA, kept at 4°C, and plasma was obtained by low-speed centrifugation at 2,000 rpm for 30 min. Sodium azide (0.05%) and aprotinin (200 Kallikrein inhibitor units/ml; Boehringer Mannheim Biochemicals, Indianapolis, IN) were added to the plasma. Radioactivity was quantitated in a gamma counter (model Autogamma 5260; Packard Instruments Co., Downers Grove, IL).

**Analytical methods.** apoE phenotypes were determined by isoelectric-focusing gel electrophoresis (33), and apoE levels were quantitated by radioimmunossay (34). Plasma cholesterol and triglyceride concentrations were determined on an Abbott VPSS analyzer (Abbott Laboratories, North Chicago, IL). HDL cholesterol was measured after dextran sulfate precipitation of plasma (35). The other lipid and lipoprotein analyses were performed as published (36). The residence time (RT) was determined from the area under the plasma decay curves using the computer-assisted curve-fitting program SAAM 29 (37). The fractional

<table>
<thead>
<tr>
<th>Study Group</th>
<th>Age</th>
<th>Sex</th>
<th>Weight</th>
<th>apoE</th>
<th>Cholesterol</th>
<th>Triglycerides</th>
<th>VLDL cholesterol</th>
<th>LDL cholesterol</th>
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<tr>
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<td>F</td>
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<td>180</td>
<td>109</td>
<td>29</td>
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<td>39</td>
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<tr>
<td></td>
<td>19</td>
<td>F</td>
<td>71.7</td>
<td>3/3</td>
<td>145</td>
<td>103</td>
<td>30</td>
<td>57</td>
<td>58</td>
</tr>
<tr>
<td></td>
<td>21</td>
<td>M</td>
<td>70.9</td>
<td>3/3</td>
<td>145</td>
<td>83</td>
<td>19</td>
<td>84</td>
<td>42</td>
</tr>
<tr>
<td></td>
<td>20</td>
<td>F</td>
<td>57.8</td>
<td>3/3</td>
<td>187</td>
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<td>164</td>
<td>40</td>
<td>15</td>
<td>101</td>
<td>48</td>
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<td>22</td>
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<td>81.5</td>
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<td>106</td>
<td>12</td>
<td>129</td>
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<td>Proband</td>
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<td>1/3</td>
<td>404</td>
<td>578</td>
<td>284</td>
<td>84</td>
<td>39</td>
</tr>
</tbody>
</table>

**Table I. Characteristics of the Study Subjects**
catabolic rate is the reciprocal of the RT. Production rates were calculated according to the following formula: production rate = (apoE concentration × plasma volume)/(RT × weight). The plasma volume was determined by dividing the total radioactivity injected by the radioactivity per unit volume in a sample obtained 10 min after injection. Intergroup comparisons were performed using Student’s t test.

In vitro binding studies

LDL displacement by apoE. To determine the LDL receptor-binding affinity of apoE-2, apoE-3, and the mutant apoE-1(Lys144→Glu), their ability to displace radioidinated LDL from receptors on normal human fibroblasts was quantitated. Normal fibroblasts were isolated and grown as previously described (38). On day 0, cells were placed in 24-well plates (Costar Corp., Cambridge, MA) and maintained in growth medium (Eagle’s minimal essential medium, 2 mM glutamine, 10% v/v FCS). On day 6, the FCS was replaced with 10% lipoprotein-depleted serum to upregulate LDL receptors. 48 h later the cells were chilled to 4°C for 20 min. The medium was replaced with medium containing 2.5 μg protein/ml 125I-LDL and apoE liposomes in concentrations of 1, 2, 4, and 8 μg/ml (apoE-1(Lys144→Glu), apoE-2) or 0.1, 0.5, 1, 2, and 4 μg/ml (apoE-3), and the cells were incubated for an additional 4 h at 4°C. The supernatant was collected, and the cells were washed and lysed with 0.1 N sodium hydroxide. Cell-associated radioactivity was quantitated by TCA precipitation of an aliquot of the lysed cells. Cell protein content was determined, and cell-associated LDL was determined per microgram of cell protein (39). The competition data were used to determine the apoE liposome concentration required to displace 50% of bound LDL. The inhibition of LDL binding by normal apoE-3 was defined as 100%, and the relative inhibition by apoE-1(Lys144→Glu) and apoE-2 was calculated (40).

The ligands used were prepared as follows: LDL in 10 M glycine (pH 10) was iodinated with iodine-125 (1 mCi/mg LDL) by the iodine monochloride method as modified for lipoproteins (41). Specific activity ranged from 100 to 150 cpm/ng LDL, containing <4% labeled lipid as determined by Folch extraction. ApoE-phospholipid complexes (apoE liposomes) were prepared as described by Hovinga et al. (42).

Heparin binding. Heparin-Sepharose CL 6B and Sepharose CL 6B were obtained from Pharmacia LKB Biotechnology Inc. (Piscataway, NJ) and prepared as recommended by the manufacturer. Heparin-binding assays were performed under non-denaturing and denaturing conditions.

For binding of radiolabeled apoE under non-denaturing conditions, 4 mg of heparin-Sepharose or 4 mg Sepharose was incubated for 4 h at 4°C in 200 μl 10 mM Tris-HCl, pH 7.5, 50 mM NaCl, and 1% BSA containing radioidinated apoE liposomes (1 μg protein/ml). ApoE-1(Lys144→Glu), apoE-2, and apoE-3 were radiiodinated (specific activity, 1,714–1,900 dpm/ng) and packed into liposomes as described earlier. After incubation of the gel with the respective apoE, the gel matrix was sedimented by centrifugation at 10,000 g. The radioactivity in 100-μl aliquots of the top and bottom fractions was quantitated, and the amount of gel-bound ligand was determined by subtracting the radioactivity of the top aliquot from that of the bottom aliquot. Heparin binding was corrected for nonspecific Sepharose binding by subtracting Sepharose gel-bound radioactivity and expressed as nanograms of apoE bound per microgram of heparin. The mass of heparin in heparin-Sepharose was quantitated by uronic acid determination (43).

For binding under denaturing conditions, the assay was modified as follows: the incubation buffer consisted of 5 M urea, 2 mM NaH2PO4, pH 8.0, 50 mM NaCl, and 1% BSA. Radiolabeled apoE-1(Lys144→Glu), apoE-2, and apoE-3 (1 μg protein/ml) were used as ligands.

Results

Structural analysis. To determine the underlying mutation in apo E-1 Patient, genomic DNAs from the proband and from a control subject with the phenotype apoE-3/3 were amplified by PCR using apoE-specific primers. Sequence analysis revealed a single base exchange of a G for an A at position 254 within exon 4 of the apoE gene, which results in the substitution of lysine (AAG) by glutamic acid (GAG) at amino acid position 146 in the proposed receptor-binding domain of apoE. The other allele was found to be normal, thus establishing that the patient is a heterozygote for the Σ-1(Lys146→Glu) allele.

In vivo kinetic studies. Six normal subjects with an apoE-3/3 phenotype and the proband participated in the in vivo kinetic study. The lipid and lipoprotein profiles of the participants are shown in Table I. Radiolabeled 125I-apoE-1(Lys146→Glu) and 131I-apoE-3 were injected simultaneously, and blood samples were obtained at intervals through day 7. Steady state conditions during the study were demonstrated by multiple determinations of plasma apoE levels (data not shown). Radioactivity was quantitated at each time point, plasma decay curves were plotted, and residence times were calculated using computer-assisted curve fitting. The catabolism of apoE-1(Lys146→Glu) and apoE-3 in one of the normal subjects is illustrated in Fig. 1 A, representative for the six controls studied. ApoE-1-(Lys146→Glu) was catabolized significantly slower (RT, 0.91 d) than apoE-3 (RT, 0.66 d) in all normal subjects (P < 0.005). The proband, in contrast, catabolized both apoE-1(Lys146→Glu) (RT, 3.76 d) and normal apoE-3 (RT, 2.23 d) much slower than did the controls (Fig. 1 B). The kinetic parameters of radiolabeled apoE-1(Lys146→Glu) and apoE-3 are summarized...
in Table II. The mean RT for apo E-3 in the normal subjects was 0.66±0.07 d, with the production rate at 3.09±0.75 mg/kg × d, which agrees with previous turnover studies of apoE-3 (32). In the proband, RTs were greatly prolonged, with 3.76 d for apo E-1 (Lys<sup>146</sup>-Glu) and 2.32 d for apoE-3. Moreover, the proband’s production rate for total apoE was about two times higher than in controls, with 6.2 mg/kg × d. The increased apoE levels in the proband were therefore due to both a decreased fractional catabolic rate and an increased production rate.

The lipoprotein distribution of <sup>125</sup>I-apoE-1 (Lys<sup>146</sup>-Glu) and <sup>131</sup>I-apoE-3, analyzed 10 min after injection by sequential ultracentrifugation of plasma samples of two normal subjects and of the proband, was very similar, as shown in Table III. However, in contrast to the controls, the majority of apoE-1 (Lys<sup>146</sup>-Glu) and apoE-3 accumulated in the VLDL subfraction of the proband.

Table III. Lipoprotein Distribution of ApoE-1 (Lys<sup>146</sup> → Glu) and ApoE-3

<table>
<thead>
<tr>
<th>Lipoprotein</th>
<th>Subject 1</th>
<th>Subject 2</th>
<th>Proband</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>E-1</td>
<td>E-3</td>
<td>E-1</td>
</tr>
<tr>
<td>VLDL</td>
<td>28.2</td>
<td>26.4</td>
<td>39.2</td>
</tr>
<tr>
<td>LDL</td>
<td>4.5</td>
<td>4.0</td>
<td>7.2</td>
</tr>
<tr>
<td>LDL</td>
<td>10.8</td>
<td>10.9</td>
<td>8.1</td>
</tr>
<tr>
<td>HDL</td>
<td>29.3</td>
<td>36.3</td>
<td>21.1</td>
</tr>
<tr>
<td>LPDS</td>
<td>27.2</td>
<td>24.3</td>
<td>25.3</td>
</tr>
</tbody>
</table>

Plasma lipoproteins of two normal subjects (1 and 2) and the proband were separated by sequential ultracentrifugation of a plasma sample at 10 min after injection of <sup>125</sup>I-apoE-1<sup>146</sup> and <sup>131</sup>I-apoE-3. Figures given are percentages.

Fibroblast binding. To assess the LDL receptor affinity of apoE-1 (Lys<sup>146</sup>-Glu) as compared with apoE-2 and apoE-3, the ability of the different apoE isoproteins to displace LDL from LDL receptors on normal human fibroblasts was determined. ApoE liposomes were used for these in vitro competition assays to exclude apoB-LDL receptor interactions. Normal human fibroblasts were incubated with 2.5 µg/ml LDL and increasing concentrations of liposomes containing either apoE-1 (Lys<sup>146</sup>-Glu), apoE-2, or apoE-3. The apoE liposome concentration required to displace 50% LDL (IC<sub>50</sub>) was calculated, and the receptor affinities of apoE-1 (Lys<sup>146</sup>-Glu) and apoE-2 were expressed relative to normal binding of apoE-3. Binding of apoE-3 was defined as 100%. Both apoE-1 (Lys<sup>146</sup>-Glu) and apoE-2 were comparably defective in their ability to displace LDL from receptors on normal human fibroblasts (apoE-1 (Lys<sup>146</sup>-Glu): 7.7±0.8%; apoE-2: 8.5±1.0%, Table IV).

Heparin binding. To investigate further the interaction of apoE-1 (Lys<sup>146</sup>-Glu) and apoE-2 (Arg<sup>128</sup>-Cys) with polyionic binding sites similar to the LDL (apoB, apoE) receptor, heparan was used as a model for negatively charged cell surface proteoglycans. Under nondenaturing conditions, the amount of apoE-1 (Lys<sup>146</sup>-Glu) and apoE-2 bound to heparin-Sepharose was decreased to ~10 and 60%, respectively, compared with the amount of apoE-3 bound (Fig. 2 A). Denaturation of the apoEs with 5 M urea restored the ability of apoE-2 to bind to heparin-Sepharose, while apoE-1 (Lys<sup>146</sup>-Glu) remained defective (Fig. 2 B).

Discussion

We have previously described a kindred in which heterozygosity for an apoE mutant, apoE-1 (Lys<sup>146</sup>-Glu), is dominantly associated with the expression of type III HLP (23). In the present study, we investigated this mutant apoE by sequence
analysis, in vivo kinetic studies, and in vitro binding studies. Analysis of the proband's apoE gene revealed a single base exchange of a G for an A at position 254 within exon 4, resulting in a lysine (AAG) to glutamic acid (GAG) substitution at amino acid position 146 in the proposed receptor binding domain of the apoE molecule. The other allele encoded the normal apoE-3. These results agree with previously performed amino acid analyses of the mutant apoE (24).

To assess the metabolic effect of this mutation, an in vivo kinetic study was performed by simultaneously injecting 131I- apoE-1 (Lys146-Glu) and 131I-apoE-3 into the proband and into six normal volunteers (Table II). In the control subjects, apoE-1 (Lys146-Glu) was catabolized significantly slower than was apoE-3; in the proband, both residence times were markedly prolonged. These results demonstrate that apoE-1 (Lys146-Glu) not only has an impaired in vivo metabolism but also interferes with the catabolism of apoE-3, thus leading to a greatly prolonged residence time of the normal apoE protein in the patient. In addition, there was an approximately twofold increase in the production rate of apoE-1 (Lys146-Glu) compared with its normal counterpart (Table II). In the patient, therefore, the ~10 times elevated plasma apoE levels were due to both a decreased fractional catabolic rate and an increase in apoE production. Previously performed kinetic studies of apoE-2- (Arg158-Cys) also showed a decreased fractional rate for apoE-2 (9) as an experimental correlate of impaired function. However, the defect of apoE-1 appears more severe, because it results in dominant manifestation of disease as opposed to the recessive mode observed for apoE-2. One possible explanation is that the LDL receptor-binding affinity of apoE-1 (Lys146-Glu) is lower than that of apoE-2. However, this is not the case. ApoE-2 and apoE-1 (Lys146-Glu) were both shown to possess 98% of normal receptor-binding activity in an in vitro competition assay using apoE liposomes. Thus, differential binding of apoE-1 and apoE-2 to the LDL receptor binding does not discriminate between dominant and variable manifestation of type III HLP. This conclusion is supported by the data recently published by Moriyama et al. (25). His group reported that apoE-1 (Lys146-Glu), which was isolated from a Japanese proband and complexed with dimyristoylphosphatidylcholine, bound to the LDL receptor with less than 10% of the affinity of normal apoE-3. With the exception of apoE-2 (Arg158-Cys), which has a receptor-binding activity as low as 1% (10), apoE-1 (Lys146-Glu) is the most binding-defective naturally occurring apoE mutant described. In comparison, the binding activities of apoE-3 (Cys122→Arg,Arg142→Cys), apoE-3 (duplication of residues 120–126), and apoE-2 (Lys146-Glu) were ~20, 25, and 35–40%, respectively, compared with apoE-3 (13, 18, 44). The latter mutant demonstrates the importance not only of the position of the mutated residue, but also of the charge of the newly introduced amino acid on LDL receptor-binding activity. The substitution of the basic residue 146, lysine, by the polar amino acid glutamine in apoE-2 (Lys146-Glu) and by the negatively charged glutamic acid in apoE-1 (Lys146-Glu) occurs in the center of the proposed receptor-binding domain of apoE, which contains six positively charged amino acids. A three dimensional crystallographic analysis of this domain revealed that residue 146, lysine, is directed toward the aqueous phase and is not involved in forming salt bridges within the apoE molecule, thus being available for intermolecular interactions with acidic residues of the LDL receptor (45). A substitution at this position is expected to adversely affect binding by directly altering the ionic interactions between ligand and receptor.

Based on the fact that several apoEs are present on a single lipoprotein particle, multiple interactions between apoE molecules and its receptors may be necessary (oligomeric ligand-receptor interaction). This concept is supported by data from Nykjaer et al. (46), who showed that it is the dimeric form of lipoprotein lipase that mediates binding of β-VLDL to LRP. Thus the binding-defective apoE mutant would prevent the formation of the required number of ionic interactions of the exposed apoE on the particle surface with its receptor, even in the presence of normal apoE-3 (Fig. 3). This would explain why the proband from the Harrisburg kindred catalyzed both apoE-1 (Lys146-Glu) and apoE-3 much slower than did the normal controls, resulting in the accumulation in plasma of particles containing both mutant and normal apoE. The especially slow removal from circulation of the mutant apoE and its twofold increased production rate and secretion, compared with the normal counterpart, explains the predominance of plasma of apoE-1 (Lys146-Glu) over normal apoE-3 at a ratio of approximately 3:1 and the increased pool size of lipoproteins with low receptor affinity. In addition, apoE-1 (Lys146-Glu) may associate preferentially with chylomicrons and VL LDL. The enrichment of VL LDL particles with binding-defective apoE-3 (Cys122→Arg,Arg142→Cys) and apoE-3 (duplication of residues 120–126) has already been proposed to promote the expression of type III HLP (44, 47). Our data, however, indicate that apoE-1 (Lys146-Glu) and apoE-3 distribute equally well to all lipoprotein classes in both normal subjects and in the proband (Table III). The predominant association of both apoE with the VL LDL subfraction of the proband is explained by the high levels of VL LDL accumulating in the patient’s plasma due to the delayed catabolism of apoE-1 (Lys146-Glu). Since VL DLs readily acquire added apoE, the enrichment of these particles with apoE appears to be a symptom of type III HLP rather than a consequence of an increased affinity of the mutant apoE for triglyceride-rich lipoproteins.

To investigate further LDL receptor-independent ionic interactions of apoE with cell membranes, we analyzed the binding of apoE-3, apoE-2 (Arg158-Cys), and apoE-1 (Lys146-Glu) to heparin as a model for cell surface proteoglycans. There are two heparin-binding sites in the apoE molecule (48, 49), and one is contained within the region of the molecule implicated in LDL receptor binding (amino acids 135–150). The replacement of residue 142, arginine, by cysteine in apoE-3 (Cys122→Arg,Arg142→Cys) has already been shown to result in a weaker affinity for heparin as measured by salt gradient elution (44). We could show that the binding of apoE-1 (Lys146-Glu) was reduced to 10% of apoE-3, whereas the affinity of apoE-2 was

| Table IV. LDL Receptor Binding of ApoE Liposomes |
|----------------|----------------|----------------|
| ApoE isopein   | IC50 (ng/ml)±SEM* | Percentage of activity |
| ApoE-1(Lys146-Glu) | 887±87         | 7.7±0.7         |
| ApoE-2         | 825±96         | 8.4±1.0         |
| ApoE-3         | 69±7           | 100±11          |

Competition data were obtained by incubating normal human fibroblasts with 2.5 μg/ml 125I-LDL with increasing concentrations of apoE liposomes for 4 h at 4°C. * IC50, concentration of apoE required to displace 50% 125I-LDL.
59% of normal. Unfolding of the protein under denaturing conditions resulted in a normalization of the heparin binding of apoE-2, while apoE-1 (Lys\(^{146}\)
\(-\text{Glu}\)) remained defective. Thus, it is not a change in the three-dimensional conformation of apoE-1 (Lys\(^{146}\)
\(-\text{Glu}\)) but rather the charge shift in the linear amino acid sequence that seems responsible for its low affinity to heparin.

This finding agrees with previously performed chemical modification experiments demonstrating the importance of positively charged arginine and lysine residues in the binding of apoB and apoE to heparin (50). It is interesting that the opposite appears to be true for apoE-2 (Arg\(^{150}\)
\(-\text{Cys}\)). This result corroborates earlier studies which demonstrated that the severe binding defect of apoE-2 is due mainly to conformational changes (44, 51).

The physiological relevance of the in vivo binding of apoE to heparin is unknown. However, evidence from in vitro experiments indicates that cell surface proteoglycans, including heparin, could have a receptor-like function for both apoE- (52, 53) and apoB- (54) containing lipoproteins. To analogize to the fibroblast growth factor family (55–57) these lipoproteins could interact with heparin or heparin-like heparan sulfate proteoglycans on the cell surface and, in a second step, be presented to LDL or apoE (remnant) receptors, which in turn mediate the cellular uptake of these particles (53, 58). In addition, binding of apoE to heparan sulfate proteoglycans could bring triglyceride- and apoE-rich particles into close spatial relationship with lipolytic enzymes (hepatic lipase, reference 59, lipoprotein lipase: reference 60), thereby enhancing hydrolysis of triglycerides as well as cellular binding and uptake of these lipoproteins. Nascent apoE may also become bound to proteoglycans inside the cell and then be transported to the membrane, where they serve as an apoE reservoir for the uptake of lipoproteins (61, 62). Defective proteoglycan binding of apoE may thus result in ineffective retardation of apoE on the cell surface with an apparently higher rate of secretion. In our kinetic study, the production rate of the mutant apoE-1 was indeed increased twofold, compared with apoE-3, which would be in accordance with the proposed mechanism. However, direct measurements of transcriptional or secretional levels could not be done because a liver biopsy would have been required.

At least two other apoE mutants associated with dominant expression of type III HLP, apoE-2 (Lys\(^{146}\)
\(-\text{Gln}\)) and apoE-4 (Glu\(^{15}\)
\(-\text{Lys},\text{Arg}\(^{145}\)
\(-\text{Cys}\)), are also impaired in their binding

Figure 3. Schematic model of the proposed oligomeric ligand-receptor interactions prevented by apoE-1 (Lys\(^{146}\)
\(-\text{Glu}\)). (Top) Normal binding of remnant particles to cells may involve two steps. First apoE mediates binding of the particle to heparan sulfate proteoglycans (HSPG) by an interaction of the positive residues in the receptor-binding region of apoE with negative charges in the polysulfated heparan side chains. This sequestration of the particle on the cell surface may facilitate step 2: the binding to lipoprotein receptors (LR) before receptor-mediated uptake (58, 62). Again multiple interactions with negatively charged residues in the ligand-binding repeats (■) of the lipoprotein receptor may be required. (Bottom) The presence of apoE-1 (Lys\(^{146}\)
\(-\text{Glu}\)) may prevent the formation of a sufficient number of ionic interactions, even in the heterozygous state. The charge of residue 146 may play a crucial role in this process and is indicated as plus (Lys\(^{146}\)) or minus (Glu\(^{146}\)) sign. The missing of oligomeric ligand-receptor interactions could represent one important factor in the dominant expression of type III HLP.
to heparin. Ji et al. (63) recently showed that defective binding
of apoE variants to heparan sulfate proteoglycans paralleled
defective remnant binding, mediated by the respective apoE vari-
ante in vitro. It was concluded from this study that defective
binding of apoE to proteoglycans may correlate with the expres-
sion of type III HLP. Following these data, together with the
presented results of our in vivo kinetic studies, heparin binding,
in addition to plasma apoE levels (22), may be used to differen-
tiate between recessive and dominant apoE mutants.

In conclusion, in vitro binding assays and in vivo turnover
demonstrate that the dominant expression of type III HLP
associated with apoE-1(Lys146→Glu) is due to (a) an
overexpression of the mutated allele, (b) an impaired binding
of the mutant apoE to LDL receptors, and (c) a slow removal
from the plasma compartment of particles containing both mu-
 tant and normal apoE. Defective binding of apoE-1(Lys146→
Glu) to heparin or heparin-like proteoglycans may play a sig-
ificant role in the manifestation of this disease. Our combined
data indicate that dominantly transmitted type III HLP is a
multifactorial disorder of apoE metabolism.

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