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Combined Pancreas and Kidney Transplantation Normalizes Protein Metabolism in Insulin-dependent Diabetic-Uremic Patients

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Abstract

In order to assess the combined and separate effects of pancreas and kidney transplantation on whole-body protein metabolism, 9 insulin-dependent diabetic-uremic patients (IDDUP), 14 patients after combined kidney-pancreas transplantation (K-P-Tx), and 6 insulin-dependent diabetic patients with isolated kidney transplant (K-Tx), were studied in the basal postabsorptive state and during euglycemic hyperinsulinemia (study 1). [1-14C]Leucine infusion and indirect calorimetry were utilized to assess leucine metabolism. The subjects were studied again with combined infusion of insulin and amino acids, given to mimic postprandial amino acid levels (study 2). In the basal state, IDDUP demonstrated with respect to normal subjects (CON): (a) higher free-insulin concentration (17.6±2.8 vs. 6.8±1.1 μU/ml, P < 0.01) (107±17 vs. 41±7 PM); (b) reduced plasma leucine (92±9 vs. 124±2 μM, P < 0.05), branched chain amino acids (BCAA) (297±34 vs. 416±10 μM, P < 0.05), endogenous leucine flux (ELF) (28.7±0.8 vs. 39.5±0.7 μmol·m⁻²·min⁻¹, P < 0.01), and nonoxidative leucine disposal (NOLD) (20.7±0.2 vs. 32.0±0.7 μmol·m⁻²·min⁻¹, P < 0.01; (c) similar leucine oxidation (LO) (8.0±0.1 vs. 7.5±0.1 μmol·m⁻²·min⁻¹; P = NS). Both K-P-Tx and K-Tx patients showed a complete normalization of plasma leucine (116±5 and 107±9 μM), ELF (38.1±0.1 and 38.5±0.9 μmol·m⁻²·min⁻¹), and NOLD (28.3±0.6 and 31.0±1.3 μmol·m⁻²·min⁻¹) (P = NS vs. CON). During hyperinsulinemia (study 1), IDDUP showed a defective decrease of leucine (42% vs. 53%; P < 0.05), BCAA (38% vs. 47%, P < 0.05), ELF (28% vs. 33%, P < 0.05), and LO (0% vs. 32%, P < 0.05) with respect to CON. Isolated kidney transplant reverted the defective inhibition of ELF (34%, P = NS vs. CON) of IDDUP, but not the inhibition of LO (18%, P < 0.05 vs. CON) by insulin. Combined kidney and pancreas transplantation normalized all kinetic parameters of insulin-mediated protein turnover. During combined hyperinsulinemia and hyperaminoacidemia (study 2), IDDUP showed a defective stimulation of NOLD (27.9±0.7 vs. 36.1±0.8 μmol·m⁻²·min⁻¹, P < 0.01 compared to CON), which was normalized by transplantation (44.3±0.8 μmol·m⁻²·min⁻¹). (J. Clin. Invest. 1994. 93:1948–1958.)

Key words: diabetes mellitus · uremia · kidney transplant · kidney-pancreas transplant · protein turnover

Introduction

Both diabetes mellitus and chronic uremia determine profound alterations of whole-body protein metabolism (1–3). In the fasting state, poorly controlled diabetic patients are characterized by negative nitrogen balance, increased blood amino acid concentrations, and increased protein turnover and oxidation rates, as assessed by commonly employed isotope techniques (4–6). In contrast, patients with chronic uremia usually present reduced blood amino acid concentrations and reduced whole-body leucine turnover and oxidation rates (7). The result of these metabolic alterations is a negative nitrogen balance. Furthermore, in the insulin-stimulated condition, both diabetes mellitus (4) and chronic uremia (8) are characterized by insulin resistance with respect to glucose metabolism, whereas the insulin effect on protein metabolism is normal. This suggests that the defect in insulin action is at postreceptorial sites in both diseases.

Isolated kidney and combined kidney-pancreas transplantation constitute valid therapeutic approaches in the treatment of insulin-dependent diabetic patients who develop end-stage renal failure. Previous reports already emphasized the importance of co-transplanting the pancreas along with the kidney, in order to obtain a near normalization of glucose metabolism (9–15). The effect of kidney-pancreas transplantation on whole-body protein metabolism is still unknown. An additional problem is constituted by the need to administer chronic immunosuppressive therapy in diabetic patients after kidney and pancreas transplantation. The specific aims of this work are (a) to assess the effect of isolated kidney transplant and combined kidney-pancreas transplantation on basal leucine turnover and oxidation rates (which are an estimate of both protein catabolism and protein synthesis), (b) to assess the effect of acute hyperinsulinemia on leucine/protein metabolism before and after isolated kidney and combined kidney-pancreas transplantation, (c) to study the effect of combined hyperinsulinemia/hyperaminoacidemia on protein synthesis before and after kidney-pancreas transplantation, (d) to determine the effect of chronic immunosuppressive therapy per se on leucine/protein metabolism in patients with posterior chronic uveitis (localized ocular disease without systemic disease). Our results demonstrate that combined kidney-pancreas transplantation completely reverts the alterations of amino acid/protein metabolism in diabetic-uremic patients.

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Table I. Clinical and Laboratory Data of the Five Study Groups

<table>
<thead>
<tr>
<th>Number</th>
<th>Group 1 (IDDUP)</th>
<th>Group 2 (KP-Tx)</th>
<th>Group 3 (K-Tx)</th>
<th>Group 4 (CU)</th>
<th>Group 5 (CON)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yr)</td>
<td>35±4</td>
<td>36±2</td>
<td>34±5</td>
<td>31±4</td>
<td>32±5</td>
</tr>
<tr>
<td>Duration of diabetes (yr)</td>
<td>20±4</td>
<td>22±3</td>
<td>19±4</td>
<td>—</td>
<td>—</td>
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<tr>
<td>Blood pressure (mmHg)</td>
<td>166±3/97±2*</td>
<td>148±3/87±1</td>
<td>150±5/88±6</td>
<td>128±2/84±3</td>
<td>121±4/82±3</td>
</tr>
<tr>
<td>Blood pH (pH units)</td>
<td>7.32±0.04</td>
<td>7.38±0.02</td>
<td>7.36±0.03</td>
<td>7.40±0.02</td>
<td>7.40±0.02</td>
</tr>
<tr>
<td>Plasma glucose (mg/dl)</td>
<td>133±4/9.0§</td>
<td>95±4/7.2</td>
<td>118±83/6§</td>
<td>86±4/1.4</td>
<td>89±3.2</td>
</tr>
<tr>
<td>Insulin dose (U/d)</td>
<td>44±3</td>
<td>—</td>
<td>48±4</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>HbA1C (%)</td>
<td>9.2±0.3§</td>
<td>5.9±0.2</td>
<td>8.7±0.4§</td>
<td>5.6±0.2</td>
<td>5.2±0.1</td>
</tr>
<tr>
<td>Plasma urea (mg/dl)</td>
<td>86±8/23.5§</td>
<td>21.5±0.5</td>
<td>34.7±0.3</td>
<td>24.3±2.5</td>
<td>15.6±0.2</td>
</tr>
<tr>
<td>Plasma creatinine (mg/dl)</td>
<td>10.2±1.1§</td>
<td>1.1±0.2</td>
<td>1.3±0.1</td>
<td>1.1±0.1</td>
<td>0.8±0.2</td>
</tr>
<tr>
<td>Nitrogen excretion (mg/min)</td>
<td>11.5±0.8*</td>
<td>7.0±0.2</td>
<td>8.4±0.7</td>
<td>8.0±0.3</td>
<td>8.1±0.5</td>
</tr>
<tr>
<td>Plasma-free insulin (μU/ml)</td>
<td>17.8±2.8§</td>
<td>14.8±1.6§</td>
<td>6.2±1.8</td>
<td>9.1±1.0</td>
<td>6.8±1.1</td>
</tr>
<tr>
<td>Plasma C-peptide (ng/ml)</td>
<td>0.06±0.01§</td>
<td>2.89±0.32*</td>
<td>0.05±0.02§</td>
<td>2.51±0.14*</td>
<td>1.70±0.20</td>
</tr>
<tr>
<td>Plasma growth hormone (ng/ml)</td>
<td>8.93±2.12§</td>
<td>2.02±0.30*</td>
<td>1.35±0.16*</td>
<td>0.47±0.14</td>
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<tr>
<td>Plasma glucagon (pg/ml)</td>
<td>152±24*</td>
<td>127±19</td>
<td>162±7*</td>
<td>120±14</td>
<td>68±7</td>
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<tr>
<td>Plasma cortisol (ng/ml)</td>
<td>154±17§</td>
<td>65±14</td>
<td>52±2</td>
<td>28±4*</td>
<td>71±7</td>
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<tr>
<td>Prednisone dose (mg/d)</td>
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<td>10±2</td>
<td>10±1</td>
<td>15±3</td>
<td>—</td>
</tr>
<tr>
<td>Cyclosporine dose (mg/kg per d)</td>
<td>—</td>
<td>5±1</td>
<td>6±1</td>
<td>5±2</td>
<td>—</td>
</tr>
<tr>
<td>Azathioprine dose (mg/kg per d)</td>
<td>—</td>
<td>1.0±0.2</td>
<td>1.1±0.1</td>
<td>1.1±0.1</td>
<td>—</td>
</tr>
</tbody>
</table>

Conversion factors to SI units: glucose = 0.05551, urea = 0.357, creatinine = 88.4, free-insulin = 6.0, C-peptide = 0.331, growth hormone = 1.0, glucagon = 1.0, cortisol = 2.759.

* P < 0.05 vs. CON; † P < 0.01 vs. CON; ‡ P < 0.01 vs. KP-Tx; § P < 0.01 vs. CU; ¶ P < 0.01 vs. K-Tx.

Methods

Subjects and experimental protocol

9 insulin-dependent diabetic-uremic patients (IDDUP); body mass index [BMI] 22.1±2.0; five males, four females), 14 patients after combined kidney-pancreas transplantation (KP-Tx; BMI 22.1±2.4; eight males, six females), 6 insulin-dependent diabetic patients after isolated kidney transplant (K-Tx; BMI 22.3±2.1; three males, three females), 5 patients with chronic uretis (CU) on the same immunosuppressive therapy as transplanted patients (BMI 23.0±2.2; three males, two females), and 8 normal healthy controls (CON; BMI 23.8±2.9; five males, three females) were studied. All transplants were from cadaveric donors. 8 patients received a segmental and 6 patients received a total pancreas transplant. The patients were admitted to the Metabolic Unit of San Raffaele Hospital at 8:00 a.m., at least 24 h before the study. All subjects underwent a 180-min euglycemic hyperinsulinemic (40 mU·m-2·min-1) clamp, combined with indirect calorimetry as previously described (13). All IDDP, 9 patients after KP-Tx, and all K-Tx, CON, and CU subjects underwent a [1-14C]leucine infusion to assess whole-body leucine turnover and oxidation. In the remaining KP-Tx patients (5) a euglycemic insulin clamp without tracer infusion was performed. Therefore, only data on cold substrate concentration are available for those subjects. All the subjects followed an isocaloric diet containing at least 250 g of carbohydrates and 70–90 g of proteins per day in 2 wk preceding the study. The day before the study the fat-free mass was assessed in all subjects by means of whole-body bio-electrical impedance as previously described (16, 17). In IDDP the fat-free mass was assessed at the end of the hemodialytic session (17). IDDP were dialyzing 3 d/wk; the study was usually done within 48 h from the last hemodialytic session. All subjects were fully informed of the possible risks of the study and gave their consent. The experimental protocol was approved by the Institutional Ethical Committee. Table I summarizes the clinical and laboratory data of the five study groups. All values in the table refer to blood samples drawn the morning of study 1.

After an overnight fast, at 6:00 a.m. a polyethylene catheter (18 G, Terumo, Leuven, Belgium) was inserted into an antecubital vein for the infusion of all test substances. A second catheter was inserted retrogradely into a wrist vein and advanced to the dorsum of the hand for blood sampling. The hand was placed in an heated box (70°C) to ensure arterIALIZATION of the venous blood (18). A prime-continuous infusion of [1-14C]leucine was administered (16 µCi as a bolus injection, followed by 0.20 µCi/min as a continuous infusion; Amersham Corp., Arlington Heights, IL; aqueous solution containing 2% ethanol), in combination with a priming dose of NaH14CO3 (2.3 µCi, Amersham, sodium salt) to prime the body bicarbonate pool (4, 19). After 2 h of isotope equilibration, samples were drawn at 10-min intervals from 120 to 180 min for the determination of basal leucine and α-ketoisocaproic acid (α-KIC) specific activities, as well as hormone and substrate concentrations.

Continuous indirect calorimetry was started 90 min after the tracer infusion was initiated, and continued throughout the study. Expired air samples were collected at 15-min intervals and bubbled through a CO2 trapping solution (hyamine hydroxide/absolute ethanol/0.1% phenolphthalein, 3:5:1) as described previously (4, 19). The solution was titrated to trap 1 mmol CO2 per 3 ml of solution. The 14C radioactivity was subsequently determined with a β scintillation counter (Packard Instrument Co. Inc., Meriden, CT) and the expired 14CO2 specific activity was calculated. Total 14CO2 expired per minute was determined by multiplying the 14C specific activity by the total CO2 production, determined as described below. The experimental protocol is depicted in Fig. 1.

1. Abbreviations used in this paper: BCAA, branched chain amino acids; BMI, body mass index; CON, normal subjects; CU, chronic uretis (patients); ELF, endogenous leucine flux; IDDUP, insulin-dependent diabetic uremic patients; K-Tx, kidney-transplanted (patients); similarly, KP-Tx, kidney/pancreas-transplanted (patients); a-KIC, α-ketoisocaproic acid; LO, leucine oxidation; NLB, net leucine balance; NOLD, nonoxidative leucine disposal.
Study 1: insulin clamp. All the subjects participated in this study protocol. After a 180-min equilibration period an insulin clamp was performed as previously described (13). Briefly a prime-continuous infusion of crystalline human insulin (Actrapid HM, Novo Nordisk, Copenhagen, Denmark) was administered at the rate of 40 mU · m⁻² · min⁻¹ to achieve and maintain an increment in plasma insulin concentration of ~75 μU/ml. The plasma glucose concentration was maintained at the basal level by measuring it at 5-min intervals and by periodic adjustment of a 20% glucose infusion based on a negative feedback principle.

Study 2: insulin clamp with amino acid infusion. Six IDDUP, seven KP-Tx patients, four K-Tx patients, four CU patients, and five CON were studied with a repeat insulin clamp. The insulin clamp was performed as described above with one exception. A balanced amino acid solution was used at the start of the insulin infusion and continued throughout the study as previously described (4). The rate of amino acid infusion averaged 0.98±0.11 ml · m⁻² · min⁻¹ (cold leucine infusion rate of 51±6 μmol · m⁻² · min⁻¹) in all study groups. These infusion rates were chosen to simulate the postprandial plasma amino acid concentrations. The aim was to obtain the same increment with respect to the basal levels in all groups.

Respiratory exchange measurements. In all studies, total CO₂ production rate and O₂ consumption rate were measured by continuous indirect calorimetry as reported previously (13), by means of Sensor-Medics instrumentation, Model Horizon, Anaheim, CA. Nonprotein respiratory quotient was measured using the Lusk’s table as reviewed and discussed by Simonson and DeFronzo (20). Nitrogen excretion rate in the IDDUP was quantified as previously described (13).

Analytical determinations and calculations

Plasma leucine concentration and specific activity were determined using an HPLC system (System Gold, Beckman Instruments, Inc., Palo Alto, CA) constituted by a programmable solvent module (model 126AA) and a programmable detector module, model 168 as previously described (4). Briefly, to precipitate plasma proteins 50 mg of sulfosalycilic acid (dry powder) was added to 1.0 ml of plasma and a 100-μl aliquot of the supernatant was analyzed in duplicate for amino acid concentration. 1 ml of the remaining supernatant was placed in duplicate on a cation exchange resin column (Dowex 50 G, Bio-Rad Laboratories, Richmond, CA) and the free amino acid fraction was eluted with 4 N HCl, dehydrated, and reconstituted in water. 10 ml of Picofluor 15 scintillation fluid (Packard) was added to each vial and 14C radioactivity was measured in a scintillation counter. The recovery of [1-14C]leucine added to plasma was 98±2%. The interassay and intraassay variations for the determination of [1-14C]leucine specific activity were 3±1% and 4±2%, respectively. >98% of the radioactivity collected in the amino acid fraction was in the leucine peak after separation by ion exchange chromatography (21). Plasma α-KIC specific activity was measured as previously described (4, 22). Briefly, 1 ml of plasma was placed in duplicate on a Dowex 50 G cation exchange resin column (Bio-Rad Laboratories) and the free α-ketoacid fraction was eluted with 4 ml of 0.01 N HCl in 50 ml cell culture tubes. 35 ml of methylene chloride was added and after shaking vigorously for 1 min the tubes were centrifuged for 5 min at 2,000 rpm to extract the free α-ketoacid fraction from the plasma. After decanting the supernatant the α-ketoacids were extracted in 350 μl of 0.2 M NaH₂PO₄ at pH 7.0. After a brief centrifugation 200 μl of the resultant supernatant was injected into an HPLC system. The system utilized a C₁₈ reverse-phase column (Altex-Ultralhpe ODS, 0.6 × 25 cm; Beckman, Instruments, Inc., Palo Alto, CA) that was eluted with 5% acetonitrile in 0.2 M H₂PO₄ buffer (pH 7.4) at the rate of 1.4 ml/min. Absorbance of α-KIC was monitored at 206 nm. Radioactivity eluting with the α-KIC peak was measured by liquid scintillation β-counting (4). All calculations were made during the last hour of the tracer equilibration and of the insulin clamp period when steady-state conditions for leucine and α-KIC specific activities and concentrations existed. Data were analyzed using a stochastic model as previously described (4, 23). The model generates the following equations in which the total leucine turnover or flux equals: Q = S + C = B + I, where S is the rate of leucine incorporation into protein (nonoxidative leucine disposal); C is the rate of leucine oxidation; B is the rate of leucine release from protein (endogenous leucine appearance); and I is the rate of exogenous leucine input. The rate of leucine turnover (Q) is calculated as follows: Q = F/leucine specific activity, where F is the infusion rate of [1-14C]leucine (in dpm/min) and leu specific activity is the specific radioactivity of leucine in the plasma compartment under conditions of equilibrium. The leucine oxidation rate (C) is calculated as follows: C = O/(K · leucine specific activity), where O is the rate of appearance of 14CO₂ in the expired air (in dpm/min) and K is a correction factor (0.81) which takes into account the incomplete recovery of labeled 14CO₂ from the bicarbonate pool. In particular, the 0.81 factor accounts for the fraction of 14CO₂ released into the body’s bicarbonate pool from the oxidation of [1-14C]-leucine, but not recovered in expired air (24, 25). An estimate of the rate of leucine incorporation into protein (S) can be calculated as follows: S = Q – C. An estimate of the rate of leucine release into plasma space from endogenous protein (B) can be calculated as follows: B = Q – I.
When the subjects are in the postabsorptive state, the leucine intake (I) equals O and B = Q. When amino acids are being infused intravenously, the rate of exogenous leucine infusion (I) must be subtracted from the total leucine flux (Q) in order to calculate the rate of endogenous leucine release from proteins. To calculate the rates of leucine turnover and oxidation, we used the plasma leucine specific activity. Recent publications have suggested that the plasma specific activity of the transaminated product of leucine, the α-KIC, may provide a better indicator of the specific activity in the intracellular mixing pool (26, 27). Therefore, estimates of leucine kinetics were also carried out using the plasma α-KIC specific activity.

Plasma glucose concentrations were determined by a differential-pH method previously described (28). Methods for determining plasma free insulin (13), glucagon (13), cortisol (13), growth hormone (13), and epinephrine (29) concentrations have been previously described. Nitrogen excretion rate was assessed as previously described (13).

**Statistical analysis**

All values are expressed as the mean±SE. Comparisons between the basal and the insulin-stimulated states within a group were performed with the Student's t test for paired data. Comparisons among the various groups were performed with the ANOVA and the Student's t test (30).

**Results**

**Plasma glucose, hemoglobin (Hb) A₁c, hormone concentrations, and nitrogen excretion rate** *(Tables I–III)*. In the basal state, plasma glucose concentration was higher in IDDP and K-Tx respect to KP-Tx, CU, and CON *(P < 0.01)*. In contrast, during the last hour of studies 1 and 2, it did not differ significantly among the various study groups (Tables I and III and Fig. 1). HbA₁c concentration was increased in IDDP and K-Tx with respect to the other groups (Table I). Postabsorptive free-IRI concentration was increased in the basal state in IDDP and K-Tx with respect to CU and CON *(P < 0.01)*. During studies 1 and 2, the increment of free-insulin was similar in the five groups (Table II, Fig. 1). Basal glucagon concentration was higher in IDDP and K-Tx with respect to KP-Tx, CU, and CON *(P < 0.05)* (Table I). During both studies 1 and 2, plasma glucagon concentration did not change significantly from the basal state, in any of the groups (Table II). Basal growth hormone concentration was increased *(P < 0.01)* in IDDP (8.93±2.12 vs. KP-Tx (2.02±0.30), K-Tx (1.35±0.16), CU (0.47±0.14), and CON (0.51±0.07). Growth hormone concentration increased significantly with respect to basal in KP-Tx and K-Tx *(P < 0.01)* during euglycemic hyperinsulinemia (Table II). Cortisol concentration was higher in IDDP vs. all groups *(P < 0.01)* (Table II). Plasma epinephrine concentration was increased in the basal state in IDDP (1.96±0.320 pm) vs. all groups *(P < 0.01)* (Table II).

In the basal state, plasma FFA were similar in IDDP (0.589±0.08), KP-Tx (0.521±0.06), K-Tx (0.681±0.04), CU (0.610±0.01), and CON (0.592±0.01 mM) (Table III). During euglycemic hyperinsulinemia the FFA concentration decreased of a similar value in IDDP (0.398±0.01) respect to KP-Tx (0.358±0.01), K-Tx (0.524±0.02), CU (0.457±0.1), and CON (0.449±0.2 mM).

The nitrogen excretion rate was increased in IDDP with respect to KP-Tx, K-Tx, and CON *(P < 0.01)*.

**Fat-free mass.** In IDDP *(BW = 62.1±3.2 kg)* the fat-free mass *(73.7±2.9%)* was slightly reduced compared to KP-Tx *(77.2±2.1%)*, K-Tx *(74.2±1.8%)*, CU *(74.4±2.1%)*, and CON *(75.4±2.3%)*. However, no statistically significant difference was detected among any of the groups.

**Plasma amino acid, leucine, and α-KIC concentrations and specific activities in the postabsorptive state in the five study groups** *(Table III, Fig. 1)*. In IDDP, the fasting branched chain amino acid (BCAA) *(297±34 μM)*, and leucine concentrations *(92±9 μM)* were all lower *(P < 0.05)* than those in control subjects *(CON: 416±10 and 124±2 μM)* for BCAA and leucine, respectively (Table II). The fasting α-KIC concentration *(19±3 vs. 28±2 μM)* was also lower *(P < 0.05)* with respect to CON. After KP-Tx and in insulin-dependent diabetic patients after isolated K-Tx, the basal plasma BCAA *(340±14, and 311±11 μM)*, respectively increased with respect to IDDP, still being significantly lower than normals *(P < 0.05)*. Basal BCAA in CU *(459±32 μM)* was similar to those in CON *(416±10 μM)*. Basal plasma leucine *(116±5, 107±9, and 121±8 μM)*, and basal plasma α-KIC *(23±1, 24±2, and 35±4 μM)* in KP-Tx, K-Tx, and CU, respectively, did not differ from those in CON (Table II). Basal plasma leucine, *(3.61±0.52 dpmmol)* and α-KIC *(2.44±0.71 dpmmol)* specific activities in IDDP were significantly higher compared with those of CON *(1.94±0.01 and 1.27±0.02 dpmmol)*, respectively *(P < 0.05)*, as well as in KP-Tx, K-Tx, and CU, respectively (plasma leucine *(2.93±0.24, 2.63±0.48, and 2.91±0.03 dpmmol)* and α-KIC *(1.97±0.11, 1.77±0.22, and 1.79±0.08 dpmmol)*). Plasma phenylalanine concentration was not statistically different among the five study groups (Table III). Plasma alanine concentration was higher in IDDP and CU vs. Groups KP-Tx, K-Tx, and CON subjects *(P < 0.05)*. A steady-state plasma plateau was achieved for the leucine and α-KIC specific activities during the last hour of the equilibration period in all study groups.

**Plasma amino acid, leucine, and α-KIC concentration and specific activities during study 1 (hyperinsulinemia) and study 2 (combined hyperinsulinemia/hyperaminoacidemia)** *(Table III, Fig. 1)*. During study 1, IDDP demonstrated a decrease in all measured plasma amino acid concentrations with the notable exception of alanine. The BCAA concentration decreased from *(297±34 to 182±25, from 340±14 to 151±9, from 311±11 to 133±24, from 459±32 to 264±28, and from 416±10 to 221±13 μM)* in IDDP, KP-Tx, K-Tx, CU, and CON, respectively. Plasma leucine concentration decreased from *(92±9 to 53±6, from 116±5 to 42±5, from 107±9 to 43±4, from 121±8 to 59±4, and from 124±2 to 58±2 μM)* in IDDP, KP-Tx, K-Tx, CU, and CON, respectively. Plasma α-KIC concentration decreased from *(19±3 to 12±2, from 23±1 to 14±1, from 24±2 to 12±1, from 35±4 to 19±2, and from 28±2 to 14±2 μM)* in IDDP, KP-Tx, K-Tx, CU, and CON, respectively. The absolute decrease of plasma leucine concentration from baseline *(39±6 vs. 66±4 μM)* was significantly less *(P < 0.01)* in IDDP compared to CON (Table III). In the patients studied after KP-Tx, the absolute decreases in BCAA *(189±36 vs. 195±20 μM), leucine *(74±11 vs. 66±4 μM), and α-KIC (9±1 vs. 14±2 μM)* concentrations were nearly identical to those observed in CON (Fig. 1, Table III). A steady-state plateau of plasma leucine and α-KIC concentrations and specific activities was achieved in all study subjects during the last hour of insulin infusion.

The plasma BCAA *(487±58, 640±84, 598±92, 602±87, and 683±69 μM)*, leucine *(157±22, 220±8, 207±23, 209±17, and 215±5 μM)*, and α-KIC *(23±2, 23±1, 29±2, 30±3, and 27±3 μM)* concentrations were similar in IDDP, KP-Tx, K-Tx, and CON.
Table II. Blood Hormone Concentrations in the Basal State, during Euglycemic Hyperinsulinemia, and during Combined Hyperinsulinemia/Hyperaminoacidemia in the Five Study Groups

<table>
<thead>
<tr>
<th></th>
<th>Group 1 (IDDUP)</th>
<th>Group 2 (KP-Tx)</th>
<th>Group 3 (K-Tx)</th>
<th>Group 4 (CU)</th>
<th>Group 5 (CON)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Basal</td>
<td>Insulin</td>
<td>Insulin + AA</td>
<td>Basal</td>
<td>Insulin</td>
</tr>
<tr>
<td>Free-IRI (μU/ml)</td>
<td>17.8±2.8*</td>
<td>77.6±6.1*</td>
<td>79.6±4.3*</td>
<td>14.8±1.6*</td>
<td>85.2±5.0*</td>
</tr>
<tr>
<td>C-peptide (ng/ml)</td>
<td>0.06±0.01*</td>
<td>0.04±0.01*</td>
<td>0.04±0.01*</td>
<td>2.89±0.32**</td>
<td>2.67±0.04**</td>
</tr>
<tr>
<td>Glucagon (pg/ml)</td>
<td>152±24**</td>
<td>129±23**</td>
<td>130±9</td>
<td>127±19</td>
<td>105±17</td>
</tr>
<tr>
<td>Growth hormone (ng/ml)</td>
<td>8.93±2.12*</td>
<td>12.21±2.71*</td>
<td>9.11±1.96</td>
<td>2.02±0.30**</td>
<td>7.63±2.32**</td>
</tr>
<tr>
<td>Cortisol (ng/ml)</td>
<td>154±17**</td>
<td>210±27**</td>
<td>169±21</td>
<td>65±14</td>
<td>69±15</td>
</tr>
<tr>
<td>Epinephrine (pM)</td>
<td>1961±320**</td>
<td>—</td>
<td>—</td>
<td>1350±285</td>
<td>—</td>
</tr>
</tbody>
</table>

Conversion factors to SI units: Insulin = 6.0, C-peptide = 0.331, growth hormone = 1.0, glucagon = 1.0, cortisol = 2.759.

* P < 0.01 vs. CON; † P < 0.01 vs. CU; ‡ P < 0.01 vs. K-Tx; § P < 0.01 vs. basal; ‡‡ P < 0.05 vs. KP-Tx; ** P < 0.05 vs. CON; †† P < 0.05 vs. basal.

Table III. Blood Metabolites Concentration and Plasma Leucine and αKIC Specific Activities, in the Basal State, during Euglycemic Hyperinsulinemia, and during Combined Hyperinsulinemia/Hyperaminoacidemia in the Five Study Groups

<table>
<thead>
<tr>
<th></th>
<th>Group 1 (IDDUP)</th>
<th>Group 2 (KP-Tx)</th>
<th>Group 3 (K-Tx)</th>
<th>Group 4 (CU)</th>
<th>Group 5 (CON)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Basal</td>
<td>Insulin</td>
<td>Insulin + AA</td>
<td>Basal</td>
<td>Insulin</td>
</tr>
<tr>
<td>Plasma glucose (mg/dl)</td>
<td>133.4±9.0*</td>
<td>115.2±5.4*</td>
<td>120.6±7.2*</td>
<td>95.4±7.2</td>
<td>90.1±2.7</td>
</tr>
<tr>
<td>Plasma FFA (mM)</td>
<td>0.58±0.08</td>
<td>0.19±0.01</td>
<td>0.137±0.03‡‡</td>
<td>0.521±0.06</td>
<td>0.163±0.01‡‡</td>
</tr>
<tr>
<td>Plasma leucine (μM)</td>
<td>92±5**</td>
<td>53±6†</td>
<td>157±22**</td>
<td>116±5</td>
<td>42±5†</td>
</tr>
<tr>
<td>Plasma αKIC (μM)</td>
<td>19±3**</td>
<td>12±2</td>
<td>23±2‡</td>
<td>23±1</td>
<td>14±3‡</td>
</tr>
<tr>
<td>Plasma leucine sp act (dpm/nmol)</td>
<td>3.61±0.52**</td>
<td>5.01±0.58*</td>
<td>1.71±0.11**</td>
<td>2.93±0.24</td>
<td>5.88±0.54‡‡</td>
</tr>
<tr>
<td>Plasma α-KIC sp act (dpm/nmol)</td>
<td>2.44±0.71**</td>
<td>3.32±0.69*</td>
<td>1.66±0.19**</td>
<td>1.97±0.11</td>
<td>3.77±0.27**</td>
</tr>
<tr>
<td>Plasma BCAA (μM)</td>
<td>297±34**</td>
<td>182±25**</td>
<td>487±58**</td>
<td>340±14**</td>
<td>151±9**</td>
</tr>
<tr>
<td>Plasma alanine (μM)</td>
<td>347±44**</td>
<td>286±29</td>
<td>395±45**</td>
<td>237±17</td>
<td>220±8</td>
</tr>
<tr>
<td>Plasma phenylalanine (μM)</td>
<td>43±4</td>
<td>27±3</td>
<td>94±4‡</td>
<td>57±1</td>
<td>32±1</td>
</tr>
</tbody>
</table>

Conversion factors to SI units: glucose = 0.55551. Abbreviation: AA, amino acid.

* P < 0.01 vs. CON; † P < 0.01 vs. CU; ‡ P < 0.01 vs. K-Tx; § P < 0.01 vs. basal; ‡‡ P < 0.05 vs. KP-Tx; ** P < 0.05 vs. CON; †† P < 0.05 vs. basal; ††† P < 0.05 vs. KP-Tx.
Tx, Cu, and CON during the last hour of study 2. The plasma
leucine and α-KIC specific activities, respectively, were signifi-
cantly higher ($P < 0.05$) in IDDUP (1.71±0.11 and 1.66±0.19
dpm/nmol) with respect to KP-Tx (1.46±0.17 and 1.31±0.14
dpm/nmol), K-Tx (1.33±0.31 and 0.96±0.11 dpm/nmol),
Cu (1.34±0.28 and 0.88±0.23 dpm/nmol), and CON
(1.34±0.12 and 1.09±0.09 dpm/nmol).

**Total leucine flux.** In the postabsorptive state, total leucine
flux averaged 28.7±0.8 μmol·m⁻²·min⁻¹ in IDDUP, a signifi-
cantly lower value ($P < 0.01$) compared to KP-Tx (38.1±0.1),
K-Tx (38.5±0.9), Cu (38.0±0.4), and CON (39.5±0.7
μmol·m⁻²·min⁻¹).

During study 1 total leucine flux decreased less ($P < 0.01$)
in IDDUP (from 28.7±0.8 to 20.5±0.7) compared to KP-Tx,
K-Tx, Cu, and CON (from 38.1±0.1 to 18.9±0.4, 38.5±0.9 to
25.1±0.7, 38.0±0.4 to 18.8±0.5, and 39.5±0.7 to 26.0±0.3
μmol·m⁻²·min⁻¹, respectively). During study 2, IDDUP
showed a 56% increase of total leucine flux (28.7±0.8 to
44.9±3.1 μmol·m⁻²·min⁻¹). This increase was similar to that
in KP-Tx (from 42.8±1.7 to 84.4±2.6), K-Tx (from 37.8±1.1
to 75.7±8.2), Cu (37.9±0.5 to 78.5±7.9), and CON
(41.5±0.6 to 69.4±3.1 μmol·m⁻²·min⁻¹).

**Endogenous leucine flux (ELF) (protein catabolism) (Table
IV).** The endogenous leucine flux (which provides a mea-
sure of protein degradation) is equal to the total leucine flux in
the basal state as well as in the studies performed without
amino acid infusion.

In study 2, the combination of hyperinsulinemia and hy-
peraminoacidemia caused a similar absolute decline of ELF in
IDDUP (28.7±0.8 to 4.8±0.5 μmol·m⁻²·min⁻¹), KP-Tx
(42.8±1.7 to 20.5±0.5), K-Tx (37.8±1.1 to 19.9±0.9), Cu
(37.9±0.5 to 22.3±1.1), and CON (41.5±0.6 to 18.9±0.9
μmol·m⁻²·min⁻¹).

**Leucine oxidation (LO) (Table IV).** In the postabsorptive
state, leucine oxidation was similar in IDDUP (8.0±0.1
μmol·m⁻²·min⁻¹), KP-Tx (9.8±0.3), K-Tx (7.5±0.3), Cu
(6.6±0.4), and CON (7.5±0.1 μmol·m⁻²·min⁻¹). During
study 1 both IDDUP (from 8.0±0.1 to 8.5±0.3 μmol·m⁻²·
min⁻¹) and K-Tx (from 7.5±0.3 to 6.1±0.4 μmol·m⁻²·
min⁻¹) demonstrated a defective suppression of leucine oxidation
when compared to KP-Tx (from 9.8±0.3 to 4.6±0.2), Cu
(from 6.6±0.4 to 3.1±0.1), and CON (from 7.5±0.1 to
5.1±0.1 μmol·m⁻²·min⁻¹). During study 2 the leucine oxidation
increased less in IDDUP (from 5.0±0.4 to 17.0±0.5
μmol·m⁻²·min⁻¹) with respect to KP-Tx (6.0±0.8 to
40.1±0.4), K-Tx (6.9±0.9 to 35.6±1.2), Cu (7.0±1.1 to
33.2±1.0), and CON (7.3±0.2 to 33.3±0.2 μmol·m⁻²·
min⁻¹).

**Nonoxidative leucine disposal (protein synthesis) (NOLD)
(Table IV).** In the postabsorptive state the NOLD (an index of
protein synthesis) was significantly reduced in IDDUP
(20.7±0.2 μmol·m⁻²·min⁻¹) with respect to normal controls
(32.0±0.7 μmol·m⁻²·min⁻¹; $P < 0.01$). KP-Tx (28.3±0.6
μmol·m⁻²·min⁻¹), K-Tx (31.0±1.3 μmol·m⁻²·min⁻¹), and
Cu (31.4±0.1 μmol·m⁻²·min⁻¹) demonstrated no difference
in NOLD with respect to normals ($P = NS$). During the insulin
clamp (study 1), NOLD decreased less in IDDUP (20.7±0.2 to
12.1±0.4) than in KP-Tx (28.3±0.6 to 14.3±0.3), K-Tx
(31.0±1.3 to 19.0±0.9), Cu (31.4±0.1 to 15.7±0.5), and

* $P < 0.01$ vs. CON; $P < 0.05$ vs. KP-Tx; $P < 0.05$ vs. K-Tx.
CON (32.0±0.7 to 21.5±0.6). During study 2, NOLD rose significantly above the basal rate in KP-Tx (from 34.1±1.1 to 44.3±0.8), K-Tx (30.9±1.2 to 40.1±2.7), CU (30.5±1.9 to 45.3±2.9), and CON (from 31.2±0.8 to 36.1±0.8), while in IDDP (from 24.1±1.1 to 27.9±0.7; *P = NS*) it did not show any significant change from the basal value.

Net leucine balance. In the basal state all study groups demonstrated a similar negative flux of leucine into and out of proteins, averaging −8.0±0.1, −9.8±0.3, −7.5±0.3, −6.6±0.4, and −7.5±0.1 μmol·m⁻²·min⁻¹ in IDDP, KP-Tx, K-Tx, CU, and CON, respectively. During study 1, the net leucine balance did not change from the basal value in IDDP (−8.5±0.3 μmol·m⁻²·min⁻¹), while it became less negative in KP-Tx (−4.6±0.2), K-Tx (−6.1±0.4), CU (−3.1±0.1), and CON (−5.1±0.1) compared with the value in the basal state. During study 2, the combined hyperaminoacidemia/hyperinsulinemia induced a similar positive net leucine balance in IDDP (from −5.0±0.4 to 23.1±0.9 μmol·m⁻²·min⁻¹), KP-Tx (from −6.0±0.8 to 23.8±1.9), K-Tx (−6.9±0.9 to 20.2±2.0), CU (−7.0±1.1 to 23.0±2.1), and CON (from −7.3±0.4 to 17.2±1.6 μmol·m⁻²·min⁻¹).

Leucine turnover based on KIC specific activity (Tables IV and V). Because it has been suggested that plasma KIC specific activity may provide a more accurate index of the intracellular precursor pool involved in leucine turnover, all calculations were repeated utilizing the steady-state plasma KIC specific activity achieved during the last hour of the basal and experimental periods. Table IV shows that the qualitative changes in leucine turnover (LO, ELF, and NOLD) calculated with the plasma KIC specific activity (reciprocal pool model), closely parallel those determined with the plasma leucine specific activity (primary pool model) under the present experimental conditions. During combined hyperinsulinemia/hyperaminoacidemia the NOLD was defectively stimulated in IDDP (from 30.5±2.2 to 28.6±0.5 μmol·m⁻²·min⁻¹), while it showed a similar stimulation in KP-Tx, K-Tx, CU, and CON (49.4±2.1, 54.2±5.4, 60.7±5.1 and 54.4±1.6, respectively).

Glucose metabolism. During the 120–180 min of study 1 in IDDP, the glucose infusion rate required to maintain euglycemia was 3.4±1.6 mg·kg⁻¹·min⁻¹, significantly less (*P < 0.01*) than the other groups. After KP-Tx and after K-Tx, the glucose infusion rate required to maintain euglycemia increased to 6.0±0.8 and 4.2±0.9 mg·kg⁻¹·min⁻¹, respectively. CU patients’ insulin-stimulated glucose metabolism was still defective with respect to CON (normals) (6.2±0.2 vs. 7.5±0.3 mg·kg⁻¹·min⁻¹).

During study 2 the glucose infusion rates in IDDP, KP-Tx, K-Tx, CU, and CON were 3.2±1.2, 6.3±1.5, 5.9±0.9, 6.5±0.6, and 7.1±0.4 mg·kg⁻¹·min⁻¹, respectively. These rates were similar to those observed when the insulin clamp study was performed without amino acid infusion (*P = NS* vs. study 1 for all groups).

**Discussion**

The overall incidence of diabetic nephropathy in the diabetic population is 35% in Western countries (31). All patients who develop proteinuria enter a hemodialysis program or get a kidney transplant within 5–10 years (32). Isolated kidney transplantation is still the first choice surgical approach for type I uremic-diabetic patients (33). One of the major drawbacks of isolated kidney transplant is the need to continue life-long therapy with prednisone (10 mg·d⁻¹), cyclosporine (5 mg·kg⁻¹·d⁻¹), and azathioprine (1 mg·kg⁻¹·d⁻¹) in insulin-dependent diabetic patients (13, 34). During the last two decades the improvement of both patient and organ survival has made it possible to extend pancreas transplant, alone or in combination with the kidney to a greater number of patients (34). In fact, both segmental and total pancreas transplantation are capable of maintaining long-term insulin independ-

**Table V. Metabolic and Kinetic Parameters of K-Tx and CU during Combined Hyperinsulinemia-Hyperaminoacidemia**

<table>
<thead>
<tr>
<th></th>
<th>K-Tx</th>
<th>CU</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Bas</td>
<td>Stu</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Patients (n)</td>
<td>4</td>
<td>4</td>
</tr>
<tr>
<td>HbA1c (%)</td>
<td>7.8±0.5#</td>
<td>5.8±0.4*</td>
</tr>
<tr>
<td>Plasma glucose (mg/dl)</td>
<td>120.7±10.8*</td>
<td>84.7±3.6*</td>
</tr>
<tr>
<td>Plasma leucine (μM)</td>
<td>104±8</td>
<td>110±9</td>
</tr>
<tr>
<td></td>
<td></td>
<td>209±17*</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ELFbas</td>
<td>37.8±1.1</td>
<td>19.9±0.9#</td>
</tr>
<tr>
<td>ELFtot</td>
<td>50.4±2.6</td>
<td>39.3±5.4#</td>
</tr>
<tr>
<td>LObas</td>
<td>6.9±0.9</td>
<td>35.6±1.2#</td>
</tr>
<tr>
<td>LOWt</td>
<td>9.9±1.4</td>
<td>50.9±3.6#</td>
</tr>
<tr>
<td>NOLDbas</td>
<td>30.9±1.2</td>
<td>40.1±2.7#</td>
</tr>
<tr>
<td>NOLDtot</td>
<td>44.1±2.9</td>
<td>54.2±5.4#</td>
</tr>
<tr>
<td>NLABas</td>
<td>−6.0±1.9</td>
<td>20.2±2.0#</td>
</tr>
<tr>
<td>NLABt</td>
<td>−9.9±1.4</td>
<td>−7.0±1.1</td>
</tr>
<tr>
<td></td>
<td>−10.6±1.9</td>
<td>14.9±0.8*</td>
</tr>
</tbody>
</table>

μmol·m⁻²·min⁻¹

Data are expressed using both the leu and α-KIC specific activities.

* *P < 0.01* compared to CU.  † *P < 0.01* compared to basal.
dence (1–10 years) (13, 35). In addition, previous studies have demonstrated that transplantation of the pancreas along with the kidney has a better impact on the quality of life of insulin-dependent diabetic patients compared with transplantation of the kidney alone (36, 37).

In the present study we used labeled leucine to assess basal and insulin-stimulated whole body protein metabolism in IDDUP before and after isolated kidney transplant or combined pancreas and kidney transplant. Our data demonstrate that the combination of diabetes mellitus and uremia causes a serious impairment of the basal and insulin-stimulated protein metabolism, that kidney transplant only partially reverses the protein/metabolic alterations of IDDUP, and finally, that the combination of the replacement of the kidney and the pancreas leads to a near normalization of whole-body protein metabolism, despite the need of chronic immunosuppressive therapy.

The nitrogen excretion rate was significantly higher in the basal state in IDDUP compared to the other groups. This indicated a negative nitrogen balance before transplantation which was reverted by combined kidney and pancreas transplantation (Table I). The protein content of the diet was comparable in IDDUP, in K-Tx alone, and in KP-Tx. Therefore, the difference in nitrogen excretion rate was not attributable to a difference in diet. Transplanted kidney may present a relapse of diabetic nephropathy. In such cases, in order to prevent the progression of renal damage a low-protein diet should be administered (38). A recent report by Brodsky et al. (39) has shown that a low protein diet may cause an impairment of leucine metabolism and a worsening of nitrogen balance and nutritional parameters. Therefore, a possible advantage of performing the transplant of the pancreas along with the kidney is that it is not necessary to give a protein restricted diet to IDDUP patients after the transplant and during chronic prednisone administration.

In a previous report we showed that IDDM in poor metabolic control is characterized by increased basal ELF (proteolysis), LO, and NOLD (an index of protein synthesis) (4). The amelioration of the metabolic control by optimizing conventional insulin therapy normalized basal ELF, NOLD, and LO in IDDM subjects (4). The basic defect in protein metabolism in type 1 diabetes mellitus would therefore seem to be a primary increase in proteolysis. In addition, in IDDM, the stimulation of protein synthesis by combined hyperinsulinemia/hyperaminoacidemia is similar to that in normals. In contrast, we have previously shown that patients with chronic uremia are characterized by reduced amino acid concentrations and reduced proteolysis (ELF), LO, and NOLD (8). In the same study, the primary defect in chronic uremia was found to be at the level of protein synthesis, while insulin action on proteolysis was found to be normal. Previous reports had shown that hemodialytic therapy (40) and chronic ambulatory peritoneal dialysis (41) are not capable of normalizing protein turnover and insulin action in uremic patients. Possible explanations of the particularly low ELF and NOLD in postabsorptive IDDUP when compared to controls include the following: (a) increased insulin levels with respect to normals (8), (b) increased growth hormone concentration (Table II), and (c) increased epinephrine concentration. In particular both high insulin (4, 6) and growth hormone (42) concentrations have been linked to the low leucine fluxes. We have previously shown that epinephrine infusion in normals causes a mild decline of amino acid concentrations (43). From the present results it seems that IDDUP patients demonstrate intracellular defects localized both at the proteolytic and proteosynthetic pathways. This explains the striking negative nitrogen balance in such patients.

Apparently both the anti-proteolytic and the proteosynthetic defects are acquired. Insulin deficiency is the primary cause of increase of proteolysis in fasting IDDM (4); when IDDM patients develop uremia a new defect in protein synthesis may be selectively determined. During exogenous insulin infusion (study 1) the inhibition of ELF is defective in IDDUP compared to the other groups indicating an insulin resistance with respect to the proteolytic pathway, only when diabetes mellitus and uremia are combined. During study 2 the stimulation of protein synthesis was defective in IDDUP compared to all the other groups. Therefore, the higher nitrogen excretion rate present in IDDUP is due partially to a defective inhibition of proteolysis and partially to a defective stimulation of the protein synthetic pathway by insulin. The combination of both defects makes more amino acid nitrogen available for the intracellular oxidative processes.

Patients after successful kidney transplant have to continue conventional insulin therapy with the additional problem of chronic immunosuppressive drugs. Furthermore, Nair et al. (44) have recently shown that hyperglucagonemia in combination with insulin deficiency (the hormone-metabolic condition in diabetic patients after isolated kidney transplant) accelerates protein catabolism. In such patients (K-Tx) the basal ELF is similar or increased with respect to normals depending upon the degree of insulinization. Patients with successful pancreas and kidney transplant are insulin independent and are capable of overcoming the alterations determined by the steroid therapy on glucose metabolism by means of basal and post-prandial hyperinsulinemia (9, 10, 12, 35, 45). During study 1 (insulin infusion) the defect of endogenous leucine flux (proteolysis) of IDDUP is reverted in K-Tx. In contrast, insulin-induced inhibition of the leucine oxidative pathway is still defective in K-Tx. Therefore, isolated kidney transplant may revert most of the intracellular metabolic pathways, with the notable exception of insulin-stimulated suppression of LO. This finding may be due to a direct glucocorticoid activation of muscle branched-chain a-ketoacid dehydrogenase (46). The fact that K-Tx have an absolute value of NOLD similar to CON and higher than KP-Tx during hyperinsulinemia may be explained as follows: (a) patients with isolated kidney transplant are generally insulin resistant as regards whole body leucine metabolism, although the impaired insulin action reaches statistical significance only for LO; in contrast, patients with combined pancreas and kidney transplantation apparently have an increased insulin sensitivity in the protein metabolic pathways; (b) considering the variation from basal instead of the absolute value of NOLD, only in IDDUP the decrease is significantly lower than CON; (c) in order to correct for the fall in amino acids, the best assessment of protein synthetic capacity is obtained in a condition of combined hyperinsulinemia/hyperaminoacidemia (4).

In our 14 patients after combined kidney-pancreas transplantation, we found a clear-cut dissociation between insulin effects on glucose and protein metabolism. In fact, after combined transplantation glucose metabolism is impaired, while
insulin-stimulated protein metabolism is normal. We had already found this in IDDM (4) and in patients with chronic renal failure (8). The presence of a dissociation between glucose and protein metabolism indicates a postreceptor defect in insulin action (47). From a clinical standpoint this finding supports the fact that the alterations of protein metabolism are easier to correct than the alterations of glucose metabolism by intensified insulin therapy, and by pancreas transplantation itself, in IDDM.

When the pancreas was transplanted along with the kidney a complete reversal of all defects in basal and insulin-stimulated protein metabolism was obtained. In particular, stimulation of NOLD after combined hyperinsulinemia/hyperaminocidemia was reduced in the diabetic-uremic group compared to controls. This alteration was completely normalized by the combined kidney and pancreas transplantation, indicating that the defect of the protein synthetic pathway is also reversible. Interestingly, also the transplantation of kidney alone was capable to revert the defect in protein synthetic rate during hyperaminocidemia.

The increased basal insulin concentration present in the kidney-pancreas transplanted group may be determined by increased secretion or reduced clearance of the molecule. Previous reports by Diem et al. (11), Osei et al. (48), and Blackman et al. (49), indicate a reduction of insulin clearance by the liver as the leading cause of basal hyperinsulinemia in KP-Tx. A reduction of insulin clearance is also present in diabetic-uremic patients (50) as a consequence of impaired renal function. An additional cause of increased basal insulin concentration is the denervation of the transplanted pancreas. In recent works by ourselves (29) and by Blackman et al. (49), it has been shown that the denervated pancreas has an increased basal insulin secretion rate, along with a defective insulin-insulin autoinhibition. Therefore, a strict linkage exists between the basal insulin concentration and the basal concentrations and leucine flux of all study groups. This is underlined by a close statistical correlation between the basal insulin level and the basal endogenous leucine flux (r = -0.64; P < 0.05) in normal volunteers, in patients with CU and in patients after KP-Tx. This finding indicates that the basal proteolysis is mainly influenced by the insulin concentration, independently of the group of subjects considered. In contrast, when the insulin and ELF of patients of groups 1 and 3 (who were on exogenous insulin administration) were plotted, no correlation was found (r = 0.01; P = NS).

Hyperinsulinemia has been elsewhere shown to cause deleterious effects on glucose, lipid, and lipoprotein metabolism (51–53). However, Homan et al. (54) have shown in the dog that peripheral insulin delivery (causing hyperinsulinemia) does not induce any major defect on glucose, protein and palmitate metabolism. Although the question of whether hyperinsulinemia is metabolically deleterious is still unanswered, our results support the hypothesis that pancreas transplantation can completely correct the post-receptor alterations induced by diabetes and uremia on protein metabolism.

It is also important to note that the hyperinsulinemia of pancreas transplanted patients is only in the systemic circulation and not in the portal blood. Actually, pancreas transplant recipients might also have a portal hypoinsulinemia when compared to normals. In addition, the systemic hyperinsulinea after transplantation is for no matter worse than iatrogenic hyperinsulinemia of the pre-transplant condition.

Prednisone (55, 56) cyclosporin A (57), and azathioprine (58) have deleterious effects on whole-body and muscle protein metabolism. In particular, steroids have been shown to mainly stimulate protein breakdown and amino acid oxidation and to directly antagonize insulin action at the muscle site; cyclosporin and azathioprine have been shown to inhibit, with different mechanisms, intracellular protein synthesis. Since all organ transplantations require a long-term administration of these drugs, caution should be used when an organ has to be transplanted in a recipient who might suffer particularly from the effect of these drugs. An important example is found in children and adolescents who have to undergo isolated kidney transplantation (59, 60). On the basis of our results the low-dose chronic immunosuppressive therapy does not appear to have a major effect on protein metabolism in IDDM patients or in patients with chronic uveitis. In fact, also during combined hyperinsulinemia/hyperaminocidemia, CU patients showed a normal stimulation of protein synthetic rate. To note that the results obtained in patients with chronic uveitis are also relevant for other organ transplantations, such as liver and heart, and indicate that the immunosuppressive therapy at the doses used after a transplant does not significantly affect whole-body protein homeostasis (59, 60). A possible counteraction of hyperinsulinemia might explain the lack of deleterious effects of chronic immunosuppression in pancreas transplanted patients. In fact, previous work has shown a close correlation between height velocity and basal insulin concentration in diabetic children (61, 62) postulating a major role of chronic low-dose hyperinsulinemia on the stimulation of protein synthesis.

The basal epinephrine concentration of our transplanted patients is slightly higher than previously published work (12). This is probably due to the different analytical technique utilized by us (HPLC vs. radioenzymatic).

In conclusion, we have shown that the alterations of basal and insulin-stimulated protein metabolism found in diabetic-uremic patients can be completely reverted by combined kidney-pancreas transplantation despite the chronic immunosuppressive therapy. The present data indicates the usefulness of transplantation of the pancreas along with the kidney in diabetic-uremic patients, especially in all clinical and pathophysiological conditions characterized by a higher rate of whole-body protein turnover.

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References

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