Apolipoprotein B100 Metabolism in Autosomal-Dominant Hypercholesterolemia Related to Mutations in PCSK9

Khadija Ouguerrram, Maud Chetiveaux, Yassine Zair, Philippe Costet, Marianne Abifadel, Mathilde Varret, Catherine Boileau, Thierry Magot, Michel Krempf

Objective—We have reported further heterogeneity in familial autosomal-dominant hypercholesterolemia (FH) related to mutation in proprotein convertase subtilisin/kexin type 9 (PCSK9) gene previously named neural apoptosis regulated convertase 1 (Narc1). Our aim was to define the metabolic bases of this new form of hypercholesterolemia.

Methods and Results—In vivo kinetics of apolipoprotein B100-containing lipoproteins using a 14-hour primed constant infusion of [3H]leucine was conducted in 2 subjects carrying the mutation S127R in PCSK9, controls subjects, and FH subjects with known mutations on the low-density lipoprotein (LDL) receptor gene (LDL-R). Apo B100 production, catabolism, and transfer rates were estimated from very LDL (VLDL), intermediate-density lipoprotein (IDL), and LDL tracer enrichments by compartmental analysis. PCSK9 mutation dramatically increased the production rate of apolipoprotein B100 (3-fold) compared with controls or LDL-R mutated subjects, related to direct overproduction of VLDL (3-fold), IDL (3-fold), and LDL (5-fold). The 2 subjects also showed a decrease in VLDL and IDL conversion (10% to 30% of the controls). LDL fractional catabolic rate was slightly decreased (by 30%) compared with controls but still higher than LDL-R–mutated subjects.

Conclusion—These results showed that the effect of the S127R mutation of PCSK9 on plasma cholesterol homeostasis is mainly related to an overproduction of apolipoprotein B100. (Arterioscler Thromb Vasc Biol. 2004;24:1-6.)

Key Words: PCSK9 hypercholesterolemia apolipoprotein B100 kinetic analysis modeling

Received December 23, 2003; revision accepted April 8, 2004.
Correspondence to Pr M. Krempf, Centre de Recherche en Nutrition Humaine, INSERM U 539, Hotel Dieu, 44093 Nantes cedex 1, Paris, France.
E-mail mkrempf@sante.univ-nantes.fr
© 2004 American Heart Association, Inc.
Arterioscler Thromb Vasc Biol. is available at http://www.atvhaha.org

DOI: 10.1161/01.ATV.0000133684.77013.88
details. These 2 affected subjects, a mother (S1 II9) and her son (S2 II10), participated in the kinetic study after they gave their informed consent. Neither had used any hypolipidemic drugs for at least 3 weeks before the study. Concentration of Lp(a) obtained from plasma frozen samples was <10 mg/dL for the 2 studied subjects. Eleven controls and 5 heterozygous FH patients previously studied by the same methods were used for comparison. The study was approved by the ethical committee of Nantes University Hospital.

**Kinetic Studies**

The endogenous labeling of apoB100 was performed by constant infusion of [1H],-leucine. Subjects fasted overnight before the study and remained fasting during the entire procedure. Each patient received intravenously a prime of 10 μmol·kg⁻¹·tracer·h⁻¹, immediately followed by a constant tracer infusion (10 μmol·kg⁻¹·h⁻¹) for 14 hours. Venous blood samples were drawn into EDTA tubes (Venoject, Paris, France) at baseline and at 15, 30, and 45 minutes, and 1, 1.5, 2, and 2.5 hours, and then hourly until 14 hours had passed. Sodium azide, an inhibitor of bacterial growth, and Pefabloc SC (Interchim, Montluçon, France), a protease inhibitor, were added to blood samples at a final concentration of 1.5 and 0.5 mmol/L, respectively.

**Analytical Procedures**

Isolation by ultracentrifugation and measurement of enrichment of lipoprotein containing apoB100. VLDL1 (Sf 400 to 600) and VLDL2 (Sf 60 to 20) were separated by density gradient ultracentrifugation using a swinging bucket rotor at 40 000 revolutions/min for 24 hours at 10°C (RPS 40T, Hitachi). IDL (1.006 < Sf < 1.020 g/mL) and LDL (1.020 < Sf < 1.063 g/mL) were separated by standard sequential ultracentrifugation methods using a fixed-angle rotor at 40 000 revolutions/min for 22 hours at 10°C (CP70, Hitachi).

Isolation and measurement of leucine enrichment in apoB100 have been described previously. Briefly, apoB100 in lipoproteins was isolated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis. Apolipoprotein bands were dried under a vacuum and then hydrolyzed. The amino acids were purified by cation exchange chromatography, then esterified and derivatized. Electron-impact gas chromatography, then esterified and derivatized. Electron-impact gas chromatography was performed on a 5891A gas chromatograph connected with a 5971A quadrupole mass spectrometer. The isotopic ratio was determined by selected ion monitoring at m/z 282 and 285. Calculations of apoB100 kinetic parameters were based on the tracer-to-trace mass ratio.

**Isolation of Lipoproteins by Fast Protein Liquid Chromatography**

Lipoprotein isolation was performed as described by Chetiveaux et al. Briefly, 200 μL of plasma were injected into a MV-7 multi-injection loop and separation was performed on 2 superose 6 HR 10/30 columns in series at flow rate of elution at 0.35 mL/min; 0.35 mL was collected for each fraction and the entire profiles was completed within 105 minutes. The system was controlled by FPLC DIRECTOR software (Amersham Pharmacia Biotech Inc).

**Measurements of Lipids and ApoB100**

All measurements were realized at 4 different sampling times on plasma and once on fast protein liquid chromatography (FPLC) fractions. Cholesterol and TG levels were measured using commercially available enzymatic kits (Biomérieux, Marcy l’Etoile, France). ApoB100 concentrations were obtained in VLDL, IDL, and LDL by combining selective precipitation and mass spectrometry and by immunonephelometry (Biomérieux) on FPLC fractions. The percentage recovery of cholesterol, TGs, and apoB100 after centrifugation was higher than 85%.

**Modeling**

Kinetic analysis of tracer-to-tracer ratios was achieved using computer software for simulation, analysis, and modeling (SAAMII). This model (Figure 1) has previously been used and published for controls and hypertriglyceridemic subjects. Briefly, in this model, heterogeneity in VLDL is represented by large VLDL (VLDL1), small VLDL (VLDL2); and a shunt between VLDL and LDL. Data are obtained by chemical analysis for VLDL1 and VLDL2. For VLDL1 fraction, supplementary heterogeneity was introduced in the model by 2 compartments: the first one, named VLDL1a and linked to VLDL2 by delipidation cascade, and the second, named VLDL1 remnants (VLDL1R), related to a direct removal of VLDL. This was performed to get a better fit of the data. ApoB-100 enters into plasma through VLDL secretion and direct production of IDL and LDL.

ApoB-100 direct removal occurs from VLDL1a (k(20,10), VLDL2 (k(30,20), VLDL1R (k(20,30)), IDL (k(10,30)), and LDL (k(40,10)). ApoB-100 transfer to higher density lipoproteins occurs by delipidation for VLDL1a (k(10,10), VLDL2 (k(30,10), k(20,30)), IDL (k(10,30)), and LDL (k(40,10)). The use of more complex models did not provide significant improvement in the fit from F-test and Akaike information criterion. A forcing function determined with SAAMII program through interpolation between experimental data and corresponding to the time course of plasma-leucine enrichment was used to drive the appearance of leucine tracer into apoB100 of the different lipoprotein fractions.

For comparison between the 3 groups (PCSK9-mutated patients, heterozygous FH, and controls) the VLDL1a, VLDL2, and VLDL1R data were presented as VLDL delipidation rate and VLDL fractional catabolic rate (FCR), which represents the sum of delipidation and direct removal rate. The VLDL conversion rate was calculated as VLDL2 delipidation flux divided by total VLDL mass. The VLDL direct removal was calculated as a sum of VLDL1a, VLDL2, and VLDL1R direct removal divided by total VLDL mass. The apoB100 production rate (PR) in mg/kg per day represents the product of FCR and pool size of apoB100 in lipoprotein fractions assuming plasma volume equal to 4.5% of body weight.

**Results**

Patients mutated for PCSK9 had higher plasma TC and apoB100 than controls (Table 1). Plasma lipids (cholesterol, TGs) profiles from FPLC lipoprotein are shown in Figure 2 for S1 and S2 and for 1 representative control and 1 representative FH. Compared with control, apoB100-rich lipoproteins from S1 and S2 exhibited higher cholesterol concentrations, although they remained lower than in heterozygous FH subjects. VLDL and IDL profiles were also different compared with controls (Table 2). They were more enriched in cholesterol than in TG. The TC/TG ratio was high.
TABLE 1. Characteristics of Controls, Patients With PCSK9 Mutation (S1 and S2), and Heterozygous Familial Hypercholesterolemic Patients

<table>
<thead>
<tr>
<th>Age</th>
<th>Sex</th>
<th>BMI</th>
<th>TC</th>
<th>TG</th>
<th>LDL-C</th>
<th>HDL-C</th>
<th>ApoB</th>
</tr>
</thead>
<tbody>
<tr>
<td>S1</td>
<td>43</td>
<td>F</td>
<td>19</td>
<td>292</td>
<td>75</td>
<td>241</td>
<td>36</td>
</tr>
<tr>
<td>S2</td>
<td>23</td>
<td>M</td>
<td>22</td>
<td>372</td>
<td>180</td>
<td>301</td>
<td>40</td>
</tr>
<tr>
<td>FH</td>
<td>53</td>
<td>2F/3M</td>
<td>24.7</td>
<td>440</td>
<td>150</td>
<td>345</td>
<td>37</td>
</tr>
<tr>
<td>SD</td>
<td>9</td>
<td>1F/11M</td>
<td>26.3</td>
<td>170</td>
<td>80</td>
<td>82</td>
<td>49.8</td>
</tr>
</tbody>
</table>

FH indicates familial hypercholesterolemic; TC, total cholesterol; TG, Triglycerides; ApoB, Apolipoprotein B100.

TABLE 2. Lipoprotein Composition in Different Subjects

<table>
<thead>
<tr>
<th></th>
<th>VLDL</th>
<th>IDL</th>
<th>LDL</th>
<th>HDL</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>TC</td>
<td>TG</td>
<td>ApoB</td>
<td>TC</td>
</tr>
<tr>
<td>S1</td>
<td>9.1</td>
<td>27.4</td>
<td>7.7</td>
<td>5.9</td>
</tr>
<tr>
<td>S2</td>
<td>23.3</td>
<td>77.5</td>
<td>17.3</td>
<td>6.6</td>
</tr>
<tr>
<td>FH</td>
<td>80</td>
<td>74</td>
<td>5.9</td>
<td>46</td>
</tr>
<tr>
<td>SD</td>
<td>9</td>
<td>10</td>
<td>4</td>
<td>8</td>
</tr>
<tr>
<td>Controls</td>
<td>4.5</td>
<td>23</td>
<td>2.7</td>
<td>3.0</td>
</tr>
<tr>
<td>SD</td>
<td>2.6</td>
<td>10</td>
<td>1</td>
<td>1.8</td>
</tr>
</tbody>
</table>

Data are in mg/dl.
TC indicates total cholesterol; TG, Triglycerides; ApoB, Apolipoprotein B100.

Discussion
Kinetics of VLDL, IDL, and LDL apoB100 were performed in 2 hypercholesterolemic subjects carrying the mutation S127R in PCSK9 and were compared with control and heterozygous FH subjects with identified mutation on LDL-
Hypercholesterolemia in these 2 subjects was the result of a dramatically apoB100-containing lipoprotein overproduction accompanied by a slight decrease in their removal. The delayed clearance observed for VLDL and IDL was a consequence of a decreased delipidation rate.

Although only 2 subjects from the same family carrying the S127R mutation were studied, the results are similar and consistent enough to be reported and used for the design of research plans on NARC-1 and hypercholesterolemia. Both subjects showed a strong increase of the direct production of VLDL, IDL, and LDL. The high VLDL release was mainly related to VLDL2 production (data not shown) and did not affect TGs pool size. These finding are different from those reported in patients with LDL receptor or apolipoprotein B100 defect. In the former, a strong reduction of catabolism and a slight overproduction of apoB100-containing lipoproteins were reported. A delayed conversion of VLDL to IDL and IDL to LDL was also reported in these subjects. Familial-deficiency apolipoprotein B100 heterozygous patients were characterized by a lower LDL catabolism and direct production rate and a low conversion rate from IDL to LDL compared with controls.4,25,26

The overproduction of VLDL, IDL, and LDL observed in the present study was already reported in other familial hypercholesterolemia,3 but the magnitude of this phenomena is dramatically higher in the patients with the PSCK9 mutation. It was already shown that substrate availability, especially cholesterol ester mass, modulates hepatic secretion of lipoproteins.27 The response of the liver to increased delivery of lipid is increased secretion rather than decreased uptake.27 However, there are still controversies and in cultured hepatoma cells starved of lipoproteins, an enhancement of apoB mRNA was observed and was reversed by an addition of LDL to the culture medium,28 as we have observed in FH patients after LDL apheresis.17 A strong relationship was also recently reported between LDL-R receptor activity and apoB100 degradation in mice hepatocytes.29 Some studies using statins have shown that the inhibition on apoB100 synthesis is dependent on the magnitude of the inhibition of endogenous cholesterol synthesis.30 In this complex regulation, we can speculate that PCSK9 could increase the degradation of newly synthesized apoB100. It is well known that a large amount of novel synthesized apoB100 is not secreted but is degraded within the cell and that apoB100 concentration is mainly regulated after transcription.15,31 Numerous stud-

![Figure 3. Time course of enrichment of VLDL1 (●), VLDL2 (○), IDL (□), and LDL (+) apolipoprotein B100 in a representative control subject (A) and in S1 (B) and S2 (C) subjects during [2H3]-leucine perfusion.](http://atvb.ahajournals.org/)

**TABLE 3. ApoB Kinetic Parameters of Controls, Patients With PCSK9 Mutation (S1 And S2), and Heterozygous Familial Hypercholesterolemic Patients**

<table>
<thead>
<tr>
<th></th>
<th>VLDL</th>
<th></th>
<th>IDL</th>
<th></th>
<th>LDL</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>TPR</td>
<td>FCR</td>
<td>TPR</td>
<td>PRd</td>
<td>FCR</td>
</tr>
<tr>
<td>S1</td>
<td>29.94</td>
<td>9.50</td>
<td>1.39</td>
<td>8.42</td>
<td>4.0</td>
</tr>
<tr>
<td>S2</td>
<td>31.28</td>
<td>4.46</td>
<td>0.29</td>
<td>6.23</td>
<td>4.21</td>
</tr>
<tr>
<td>FH</td>
<td>15.50</td>
<td>6.32</td>
<td>4.12</td>
<td>11.50</td>
<td>6.62</td>
</tr>
<tr>
<td>SD</td>
<td>8.70</td>
<td>3.20</td>
<td>2.48</td>
<td>7.20</td>
<td>3.21</td>
</tr>
<tr>
<td>controls</td>
<td>10.42</td>
<td>9.20</td>
<td>7.93</td>
<td>11.30</td>
<td>1.44</td>
</tr>
<tr>
<td>SD</td>
<td>3.70</td>
<td>4.43</td>
<td>3.62</td>
<td>3.38</td>
<td>1.01</td>
</tr>
</tbody>
</table>

FCR indicates fractional catabolic rate (day⁻¹); TPR, total production rate (mg/kg/day); PRd, direct production rate (mg/kg/day).
ies$^{29-35}$ have shown that several situations like hormonal and nutritional states or apoB100 gene mutations can alter apoB100 rate of secretion by changing its intracellular degradation. The activity of PCSK9 could be critical for this pathway and its regulation, but the mechanisms related to this mutation on cholesterol homeostasis and apoB100 secretion need further investigation.

A lower conversion rate of VLDL and IDL compared with controls and heterozygous FH patients was also observed in the 2 subjects. It is demonstrated that in normal subjects, most apoB100 LDL comes from VLDL lipolysis.$^{16,17}$ The contribution of VLDL and IDL to LDL formation accounted for almost 28% in these subjects versus 75% in the controls, and the direct synthesis was the principal pathway for LDL production. In FH heterozygous patients,$^{3,4,17}$ LDL direct secretion was equal to the formation from IDL, whereas in the familial deficiency apolipoprotein B$^{4}$ the synthesis from IDL was the major source (80%). The composition of nascent VLDL, mainly TG content, may govern their subsequent metabolic behavior in plasma. TG-rich particles undergo rapid lipolysis via lipoprotein lipase.$^{36}$ The VLDL composition of the 2 subjects showed a higher cholesterol to TGs ratio. As demonstrated by Schreier et al,$^{37}$ this ratio is negatively correlated with the affinity of lipoprotein lipase. Thus, these enriched cholesterol particles are probably not well-recognized by lipoprotein lipase and then make little contribution to LDL production. Our kinetic data and the composition of lipoproteins found in the 2 subjects are in good agreement with this hypothesis. The mechanisms involved in the elevated cholesteryl ester-to-TGs ratio. As demonstrated by Schreier et al,$^{37}$ this ratio is negatively correlated with the affinity of lipoprotein lipase. Thus, these enriched cholesterol particles are probably not well-recognized by lipoprotein lipase and then make little contribution to LDL production. Our kinetic data and the composition of lipoproteins found in the 2 subjects are in good agreement with this hypothesis. The mechanisms involved in the elevated cholesteryl ester-to-TGs ratio in the familial hypercholesterolemia are critical for their technical assistance.

Acknowledgments

This work was supported by La direction de la recherche clinique of Nantes University Hospital and INSERM (contrat Progrès). We thank P. Maugère and C. Levalleig for their technical assistance.

References


Apolipoprotein B100 Metabolism in Autosomal-Dominant Hypercholesterolemia Related to Mutations in PCSK9

Khadija Ouguerram, Maud Chetiveaux, Yassine Zair, Philippe Costet, Marianne Abifadel, Mathilde Varret, Catherine Boileau, Thierry Magot and Michel Krempf

Arterioscler Thromb Vasc Biol. published online May 27, 2004;
Arteriosclerosis, Thrombosis, and Vascular Biology is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2004 American Heart Association, Inc. All rights reserved.
Print ISSN: 1079-5642. Online ISSN: 1524-4636

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://atvb.ahajournals.org/content/early/2004/05/27/01.ATV.0000133684.77013.88.citation

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in Arteriosclerosis, Thrombosis, and Vascular Biology can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to Arteriosclerosis, Thrombosis, and Vascular Biology is online at:
http://atvb.ahajournals.org//subscriptions/