Genetic spectrum and clinical correlates of somatic mutations in aldosterone-producing adenoma.

This is the author’s manuscript

Original Citation:
Genetic spectrum and clinical correlates of somatic mutations in aldosterone-producing adenoma. / Fernandes-Rosa FL;Williams TA;Riester A;Steichen O;Beuschlein F;Boulkroun S;Strom TM;Monticone S;Amar L;Matchi T;Mantero F;Cicala MV;Quinkler M;Fallo F;Alloolio B;Bernini G;Maccario M;Giacchetti G;Jeunemaitre X;Mulatero P;Reincke M;Zennaro MC. - In: HYPERTENSION. - ISSN 0194-911X. - STAMPA. - 64:2(2014), pp. 354-361.

Availability:
This version is available http://hdl.handle.net/2318/153168 since

Published version:
DOI:10.1161/HYPERTENSIONAHA.114.03419

Terms of use:
Open Access
Anyone can freely access the full text of works made available as "Open Access". Works made available under a Creative Commons license can be used according to the terms and conditions of said license. Use of all other works requires consent of the right holder (author or publisher) if not exempted from copyright protection by the applicable law.

(Article begins on next page)
This is the author's final version of the contribution published as:

Fernandes-Rosa FL;Williams TA;Riester A;Steichen O;Beuschlein F;Boulkroun S;Strom TM;Monticone S;Amar L;Meatchi T;Mantero F;Cicala MV;Quinkler M;Fallo F;Allolio B;Bernini G;Maccario M;Giacchetti G;Jeunemaitre X;Mulatero P;Reincke M;Zennaro MC. Genetic spectrum and clinical correlates of somatic mutations in aldosterone-producing adenoma.. HYPERTENSION. 64 (2) pp: 354-361.
DOI: 10.1161/HYPERTENSIONAHA.114.03419

The publisher's version is available at:
http://hyper.ahajournals.org/cgi/doi/10.1161/HYPERTENSIONAHA.114.03419

When citing, please refer to the published version.

Link to this full text:
http://hdl.handle.net/2318/153168
Genetic Spectrum and Clinical Correlates of Somatic Mutations in Aldosterone-Producing Adenoma

Fabio Luiz Fernandes-Rosa, Tracy Ann Williams, Anna Riester, Olivier Steichen, Felix Beuschlein, Sheerazed Boukloum, Tim M. Strom, Silvia Monticone, Laurence Amar, Tchao Meatchi, Franco Mantero, Maria-Verena Cicala, Marcus Quinkler, Francesco Fallo, Bruno Allolio, Giampaolo Bernini, Mauro Maccario, Gilberta Giacchetti, Xavier Jeunemaitre, Paolo Mulatero, Martin Reincke and Maria-Christina Zennaro

Abstract

Primary aldosteronism is the most common form of secondary hypertension. Somatic mutations in KCNJ5, ATP1A1, ATP2B3, and CACNA1D have been described in aldosterone-producing adenomas (APAs). Our aim was to investigate the prevalence of somatic mutations in these genes in unselected patients with APA (n=474), collected through the European Network for the Study of Adrenal Tumors. Correlations with clinical and biochemical parameters were first analyzed in a subset of 199 patients from a single center and then replicated in 2 additional centers. Somatic heterozygous KCNJ5 mutations were present in 38% (180/474) of APAs, whereas ATP1A1 mutations were found in 5.3% (25/474) and ATP2B3 mutations in 1.7% (8/474) of APAs. Previously reported somatic CACNA1D mutations as well as 10 novel CACNA1D mutations were identified in 44 of 474 (9.3%) APAs. There was no difference in the cellular composition of APAs or in CYP11B2, CYP11B1, KCNJ5, CACNA1D, or ATP1A1 gene expression in APAs across genotypes. Patients with KCNJ5 mutations were more frequently female, diagnosed younger, and with higher minimal plasma potassium concentrations compared with CACNA1D mutation carriers or noncarriers. CACNA1D mutations were associated with smaller adenomas. These associations were largely dependent on the population structure of the different centers. In conclusion, recurrent somatic mutations were identified in 54% of APAs. Young women with APAs are more likely to be KCNJ5 mutation carriers; identification of specific characteristics or surrogate biomarkers of mutation status may lead to targeted treatment options.

Introduction

Arterial hypertension is a major cardiovascular risk factor that affects 10% to 40% of the adult population in industrialized countries. Detection of secondary forms of hypertension is particularly important because it allows the targeted management of the underlying disease. Primary aldosteronism (PA) is the most common form of secondary hypertension with an estimated prevalence of ≈10% in referred patients and 4% in primary care but as high as 20% in patients with resistant hypertension.1-5 PA is caused by the excessive production of aldosterone from the adrenal cortex, resulting in hypertension associated with high plasma aldosterone levels, low plasma renin activity, and varying degrees of hypokalemia and metabolic alkalosis. Long-term consequences include an increased risk of myocardial infarction, stroke, and atrial fibrillation.6,7 Aldosterone-producing adenoma (APA) and bilateral adrenal hyperplasia (also known as idiopathic hyperaldosteronism) are the most frequent subtypes of PA together accounting for ≈95% of all cases.8

In recent years, major advances in the knowledge of the genetic basis of APA development have been made. The activation of the calcium signaling pathway is central for the regulation of aldosterone production, in particular by increasing the expression of CYP11B2, encoding aldosterone synthase. Zona glomerulosa (ZG) cells are sensitive to changes in extracellular potassium concentration (K⁺). The ZG cell membrane is selectively permeable to a wide range of K⁺ concentrations, and the depolarization of these cells leads to calcium (Ca²⁺) entry through voltage-dependent calcium channels.9 Somatic and inherited mutations of KCNJ5, coding for the potassium channel GIRK4, have been implicated in the formation of APA and in familial hyperaldosteronism type 3.10,11 More recently, somatic mutations in ATP1A1 (coding for the α1 subunit of the Na⁺/K⁺-ATPase) and ATP2B3 (coding for the plasma membrane Ca²⁺-ATPase, type 3) were identified in APA.12 Mutations in KCNJ5 and ATP1A1 affect adrenal ZG cell membrane potential and intracellular ionic homeostasis, with chronic depolarization leading to opening of voltage-dependent calcium channels and activation of calcium signaling and aldosterone production.10,12,13 Finally, somatic
mutations in the CACNA1D encoding the Ca$_{\alpha}$1.3 channel (calcium channel, voltage dependent, L type, α-1d subunit) have been identified in APA. The altered residues are located in particular segments bordering the channel pore.$^{14,15}$ These changes result in channel activation and opening at lower voltages, leading, like ATP2B3 mutations, to increased intracellular calcium concentrations.

A large collection of APA samples, endowed with comprehensive clinical and biological information, collected through reference centers is organized within the European Network for the Study of Adrenal Tumors (ENS@T). The aim of the present study was to establish the prevalence of KCNJ5, CACNA1D, ATP1A1, and ATP2B3 mutations in this European cohort and to explore the clinical and biological correlates of the mutation status across the spectrum of APA.

Subjects and Methods

An expanded Methods section is available in the online-only Data Supplement.

Patients

Patients with PA were recruited between 1994 and 2012 from 9 different centers belonging to the ENS@T APA working group (http://www.ENSAT.org). An extended phenotype was available for patients from Paris, Munich, and Torino.$^{11,12,16,17}$ Case detection and subtype identification of PA were performed according to institutional and Endocrine Society guidelines.$^{18-22}$ In patients diagnosed with PA, a thin slice computed tomographic scan or MRI of the adrenal or an adrenal venous sampling (AVS) was performed to differentiate between unilateral and bilateral aldosterone hypersecretion. Baseline clinical and biochemical characteristics of the patients are indicated in Table S1 in the online-only Data Supplement. Further details are available in the online-only Data Supplement.

DNA Isolation and Sequencing

Details are available in the online-only Data Supplement.

Pathological Analysis

Details are available in the online-only Data Supplement.

Statistical Analyses

Quantitative variables are reported as medians and interquartile range and compared with the Kruskal–Wallis (3 groups) or Mann–Whitney tests (2 groups). Categorical variables are reported as percentages and compared with Fisher exact test. Because of the low number of patients with mutations in ATP1A1 and ATP2B3, extended phenotypes in patients from Paris were only compared between patients with KCNJ5 or CACNA1D mutations or no mutations. A P value <0.05 was considered significant for global comparisons between the 3 groups; significant differences were further explored with 3 post hoc pairwise comparisons, for which a P value <0.05/3 was considered significant. We tried to replicate the significant pairwise differences in patients from the other 2 centers with large cohorts (Munich and Torino), first in each center and then by overall comparisons stratified by center (stratified linear regression for quantitative variables and Mantel–Haenszel test for categorical variables).

Results

Spectrum of Somatic Mutations in APA
Hot spot regions for mutation in KCNJ5, CACNA1D, ATP1A1, and ATP2B3 were successfully sequenced in 474 somatic APA DNA samples from 9 different centers. Genetic screening of mutations in KCNJ5, ATP1A1, and ATP2B3 was partially described in previous studies (online-only Data Supplement).11,12,17 In addition, whole exome sequencing was performed in 25 paired somatic and germline DNA samples. Somatic mutations were found in 54.2% of APAs with 38.0% of KCNJ5 mutations, 9.3% of CACNA1D mutations, 5.3% of ATP1A1 mutations, and 1.7% of ATP2B3 mutations (Table 1). There was no difference in the distribution of mutations across different centers (P=0.21). From the 25 patients who underwent exome sequencing, somatic mutations affecting one of the KCNJ5, CACNA1D, ATP1A1, and ATP2B3 genes were identified in 11 patients. The somatic mutations were screened in paired germline DNA, and no mutation was observed.

**Table 1.** Type and Frequency of Somatic Mutations in Aldosterone-Producing Adenomas Across Different Centers

<table>
<thead>
<tr>
<th>Center</th>
<th>No. With Successful Sequencing</th>
<th>KCNJ5 Mutations (%)</th>
<th>CACNA1D Mutations (%)</th>
<th>ATP1A1 Mutations (%)</th>
<th>ATP2B3 Mutations (%)</th>
<th>Total (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Paris</td>
<td>199</td>
<td>74 (37.2)</td>
<td>27 (13.6)</td>
<td>9 (4.5)</td>
<td>3 (1.5)</td>
<td>113 (56.7)</td>
</tr>
<tr>
<td>München</td>
<td>93</td>
<td>32 (34.4)</td>
<td>7 (7.5)</td>
<td>7 (7.5)</td>
<td>4 (4.3)</td>
<td>50 (53.7)</td>
</tr>
<tr>
<td>Torino</td>
<td>73</td>
<td>29 (39.7)</td>
<td>4 (5.4)</td>
<td>5 (6.8)</td>
<td>1 (1.3)</td>
<td>39 (53.4)</td>
</tr>
<tr>
<td>Padova A</td>
<td>37</td>
<td>14 (37.8)</td>
<td>2 (5.4)</td>
<td>1 (2.7)</td>
<td>0</td>
<td>17 (45.9)</td>
</tr>
<tr>
<td>Berlin</td>
<td>23</td>
<td>11 (47.8)</td>
<td>1 (4.3)</td>
<td>1 (4.3)</td>
<td>0</td>
<td>13 (56.5)</td>
</tr>
<tr>
<td>Ancona</td>
<td>21</td>
<td>10 (47.6)</td>
<td>2 (9.5)</td>
<td>2 (9.5)</td>
<td>0</td>
<td>14 (66.6)</td>
</tr>
<tr>
<td>Padova B</td>
<td>11</td>
<td>3 (27.2)</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>3 (27.2)</td>
</tr>
<tr>
<td>Würzburg</td>
<td>9</td>
<td>4 (44.4)</td>
<td>1 (11.1)</td>
<td>0</td>
<td>0</td>
<td>5 (55.5)</td>
</tr>
<tr>
<td>Pisa</td>
<td>8</td>
<td>3 (37.5)</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>3 (37.5)</td>
</tr>
<tr>
<td>Total</td>
<td>474</td>
<td>180 (38.0)</td>
<td>44 (9.3)</td>
<td>25 (5.3)</td>
<td>8 (1.7)</td>
<td>257 (54.2)</td>
</tr>
</tbody>
</table>

Somatic KCNJ5 mutations were found in 180 APA samples (Table S3). The recurrent mutations p.Gly151Arg (c.451G>A or c.415G>C) and p.Leu168Arg (c.503T>G) accounted for 62.7% (113/180) and 36.1% (65/180) of KCNJ5 mutations, respectively. There was no difference in the frequency of p.Gly151Arg and p.Leu168Arg across different centers (P=0.33). The cohort also included 2 patients harboring 2 additional previously described somatic mutations (p.Thr158Ala/c.472A>G and p.Trp126Arg/c.376T>C).17

CACNA1D mutations were found in 44 APAs from 7 centers. Nineteen CACNA1D mutations were identified; these mutations are located in exons 6, 8A, 8B, 14, 16, 23, 27, and 32 (Table S4). Among these mutations, we identified 9 previously described CACNA1D mutations14,15 (please note that mutations p.Phe747Leu and p.Ile750Met described here and in Azizan et al14 correspond to p.Phe767Leu and p.Ile770Met in Scholl et al15 because of the use of different reference sequences as indicated in Table S4). The mutations p.Gly403Arg (c.1207G>C), p.Phe747Leu (c.2239T>C), and p.Val1338Met (c.4012 G>A) were the most prevalent, being found in 8, 6, and 6 APAs, respectively. Six novel CACNA1D mutations were identified: p.Tyr741Cys (c.2222A>G) and p.Ile750Phe (c.2248A>T) located in exon 16; p.Val979Asp (c.2936T>A), p.Val981Asn (c.2943G>C), p.Ala998Val (c.2993G>T), and p.Ala998Ile (c.2992-2993GC>AT) located in exon 23. Four additional new mutations were identified in exons not previously reported to carry somatic CACNA1D mutations, with whole exome sequencing: p.Leu655Pro (c.1964T>C) and p.Ser652Leu (c.1955C>T) located in exon 14 and p.Val1151Phe (c.3451G>T) and p.Ile1152Asn (c.3455T>A) located in exon 27 (Figure 1). The new mutations are located in regions conserved across different species (Figure S1). All but 2 novel mutations were present in single samples; the mutations p.Ala998Ile and p.Val1151Phe were present in 3 and 2 samples, respectively.
Table 2. Extended Phenotype of Patients From Paris With Different Mutation Status

<table>
<thead>
<tr>
<th>Clinical and Biological Parameters</th>
<th>No Mutation (A) (n=86)</th>
<th>KCNJ5 (B) (n=74)</th>
<th>CACNA1D (C) (n=27)</th>
<th>ATP1A1 (n=9)</th>
<th>ATP2B3 (n=3)</th>
<th>P Value Overall</th>
<th>A vs B</th>
<th>A vs C</th>
<th>B vs C</th>
</tr>
</thead>
<tbody>
<tr>
<td>Females</td>
<td>45%</td>
<td>72%</td>
<td>33%</td>
<td>11%</td>
<td>67%</td>
<td>&lt;0.001</td>
<td>0.001</td>
<td>0.37</td>
<td>0.001</td>
</tr>
<tr>
<td>Age at HTN diagnosis, y</td>
<td>39.5 (31–45)</td>
<td>36.5 (28–43)</td>
<td>39 (35–44)</td>
<td>40 (35–46)</td>
<td>38 (38–38)</td>
<td>0.33</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age at PA diagnosis, y</td>
<td>45.5 (40–51)</td>
<td>40 (35–45)</td>
<td>46 (39–51)</td>
<td>50 (43–53)</td>
<td>44 (42–53)</td>
<td>0.001</td>
<td>&lt;0.001</td>
<td>0.95</td>
<td>0.007</td>
</tr>
<tr>
<td>Lowest plasma K, mmol/L</td>
<td>3.0 (2.7–3.3)</td>
<td>3.3 (3.0–3.6)</td>
<td>3.0 (2.8–3.3)</td>
<td>3.1 (2.5–3.2)</td>
<td>1.9 (1.7–3.3)</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>0.69</td>
<td>0.009</td>
</tr>
<tr>
<td>Systolic BP, mm Hg</td>
<td>144.5 (134–158)</td>
<td>138 (131–153)</td>
<td>150 (138–164)</td>
<td>146 (138–151)</td>
<td>162 (157–167)</td>
<td>0.05</td>
<td>0.08</td>
<td>0.24</td>
<td>0.03</td>
</tr>
<tr>
<td>Diastolic BP, mm Hg</td>
<td>90 (85–97)</td>
<td>90 (80–99)</td>
<td>92 (85–95)</td>
<td>88 (84–95)</td>
<td>107 (81–111)</td>
<td>0.51</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Treatment score, n</td>
<td>2 (1–3)</td>
<td>2 (1–3)</td>
<td>2 (1–3)</td>
<td>2 (1–2)</td>
<td>3 (2–3)</td>
<td>0.52</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>eGFR, mL/min per 1.73 m²</td>
<td>92 (82–113.5)</td>
<td>94.5 (84–114)</td>
<td>91 (69–103)</td>
<td>94 (74–107.5)</td>
<td>92 (92–92)</td>
<td>0.45</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Urinary aldosterone, mmol/24 h</td>
<td>93.5 (64–147)</td>
<td>78 (63–112)</td>
<td>115 (75–142)</td>
<td>124 (80–223)</td>
<td>330 (59–757)</td>
<td>0.15</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Plasma aldosterone, pmol/L</td>
<td>812 (511–1176)</td>
<td>827 (626–1148)</td>
<td>685 (520–1012)</td>
<td>1102 (894–2157)</td>
<td>2193 (1249–9576)</td>
<td>0.27</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Plasma rennin, mU/L</td>
<td>1.7 (1.0–3.1)</td>
<td>1.7 (1.0–2.7)</td>
<td>2.9 (1.2–4.7)</td>
<td>1.9 (1.3–5.2)</td>
<td>1.0 (1.0–8.7)</td>
<td>0.16</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ARR, pmol/mU</td>
<td>149 (92–203)</td>
<td>165 (122–230)</td>
<td>126 (95–195)</td>
<td>218 (179–431)</td>
<td>252 (250–1915)</td>
<td>0.10</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>AVS lateralization ratio</td>
<td>16 (10–35)</td>
<td>23 (13–45)</td>
<td>10 (7–32)</td>
<td>19 (13–58)</td>
<td>53 (6–99)</td>
<td>0.03</td>
<td>0.05</td>
<td>0.19</td>
<td>0.02</td>
</tr>
<tr>
<td>Solitary adenoma</td>
<td>79%</td>
<td>92%</td>
<td>81%</td>
<td>89%</td>
<td>100%</td>
<td>0.06</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Largest adenoma, mm</td>
<td>12 (10–20)</td>
<td>16 (13–20)</td>
<td>9 (8–12)</td>
<td>10 (10–18)</td>
<td>20 (12–25)</td>
<td>&lt;0.001</td>
<td>0.02</td>
<td>0.002</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>SBP at FU, mm Hg</td>
<td>127 (118–135)</td>
<td>125 (117–134)</td>
<td>126 (122–135)</td>
<td>133 (124–134)</td>
<td>136.5 (133–140)</td>
<td>0.62</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diastolic BP at FU, mm Hg</td>
<td>82 (75–87)</td>
<td>82 (76.5–88)</td>
<td>82 (76–87)</td>
<td>86 (79–89)</td>
<td>87 (84–90)</td>
<td>0.77</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Treatment score at FU, n</td>
<td>0 (0–1)</td>
<td>0 (0–1)</td>
<td>1 (0–2)</td>
<td>1 (0–2)</td>
<td>0 (0–0)</td>
<td>0.10</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>K at FU, mmol/L</td>
<td>4.2 (3.9–4.4)</td>
<td>4.1 (3.9–4.5)</td>
<td>4.2 (4.0–4.3)</td>
<td>4.5 (4.1–4.7)</td>
<td>4.6 (4.0–5.2)</td>
<td>0.87</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ARR at FU, pmol/mU</td>
<td>9.3 (4.8–16.1)</td>
<td>9.8 (6.6–19.7)</td>
<td>9.6 (4.1–18.1)</td>
<td>5.9 (2.3–6.7)</td>
<td>...</td>
<td>0.63</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Adjusted change in SBP, mm Hg*</td>
<td>26 (16–38)</td>
<td>26 (13–35)</td>
<td>25 (17–37)</td>
<td>21 (10–30)</td>
<td>45 (44–46)</td>
<td>0.58</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hypertension cure</td>
<td>56%</td>
<td>58%</td>
<td>32%</td>
<td>44%</td>
<td>50%</td>
<td>0.08</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Significant BP improvement†</td>
<td>85%</td>
<td>78%</td>
<td>76%</td>
<td>78%</td>
<td>100%</td>
<td>0.45</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Statistical analysis has been done on groups A, B, and C only. Characteristics of ATP1A1 and ATP2B3 mutation carriers are presented for completeness but were not included in statistical comparisons because of small sample sizes. Categorical data reported as number (percentage) and compared with Fisher exact test; quantitative data reported as median (interquartile range) and compared with the Kruskal–Wallis test (overall P, significant if <0.05) or Mann–Whitney test (pairwise P, significant if <0.05/3). ARR indicates aldosterone-to-renin ratio; AVS, adrenal venous sampling; BP, blood pressure; eGFR, estimated glomerular filtration rate; FU, follow-up; HTN, hypertension; K, potassium; PA, primary aldosteronism; and SBP, systolic blood pressure. * Change in SBP adjusted for treatment score by ANCOVA. † Hypertension cure or adjusted change in SBP >20 mm Hg.

ATP1A1 mutations were found in 25 APAs from 6 centers (Table S5). The mutation p.Leu104Arg (c.311T>G) identified in 17 APAs is the most prevalent. Another 3 mutations were observed, p.Phe100_Leu104del (c.299_313delTCTCAATGTTACTGT) in 5 samples, p.Val332Gly (c.995T>G) in 2 samples, and p.Gly99Arg (c.295G>A) in 1 sample. Novel somatic ATP1A1 mutations were not identified. Somatic ATP2B3 mutations were found in 8 APAs (Table S6). Three different in frame deletion mutations were observed affecting the region comprised between residues 424 and 427. The deletion p.Leu425_val426del (c.1272‐1277delGCTGGT or c.1273‐1278delCTGGTC), identified in 5 APAs, was the most prevalent. The mutation p.Val426_val427del (c.1277_1282delTCGTGG) was present in 1 sample. A new mutation p.Val424_leu425del (c.1270‐1275del GTGCTG) was identified in 2 patients from the Paris cohort.

Correlation of Mutation Status With Morphological and Molecular Characteristics of APA

To verify the hypothesis that mutations in different genes are associated to a specific cellular composition of APA, we analyzed the cellular composition of 78 APAs from the Paris cohort. APAs are generally composed of zona fasciculata (ZF)‐like cells or ZG‐like cells in variable proportions; sometimes, smaller eosinophilic cells resembling cells from the zona reticularis are also observed. ZF cells are large cells with pale‐staining nuclei and acidophilic and clear cytoplasm with abundant lipid droplets (Figure S2A). ZG cells are small cells with basophilic nuclei and clear cytoplasm with few intracellular fat droplets (Figure S2B). Seventy‐two percent of APAs were composed of a majority (>50%) of ZF‐like cells. Overall, ZF‐like cells represented the main cellular component of APA: ZF‐like cells median, 70 (50–90); ZG‐like cells median, 10
(10–35); P<0.001 (Figure S2C). Analysis of the cellular composition of APA according to the mutation status did not show any difference in the percentage of ZF-like and ZG-like cells across groups (Figure S2D and S2E).

To verify the hypothesis that the mutation status is associated to different levels of CYP11B2 or CYP11B1 expression, we retrieved CYP11B2 and CYP11B1 mRNA expression from a transcriptome study performed on 92 APAs with complete genotypes. This analysis included 45 APAs with KCNJ5 mutations, 10 APAs with CACNA1D mutations, 5 APAs with ATP1A1 mutations, and 31 APAs without mutations. There was no difference in CYP11B2 or CYP11B1 mRNA expression as a function of the mutation status (Figure 2A and 2B). In addition, we also investigated KCNJ5, CACNA1D, and ATP1A1 mRNA expression according to the mutation status. No difference in gene expression was observed across groups with different mutations (Figure 2C–2E).

**Figure 2.**


Combined information on mutation status, CYP11B2 and CYP11B1 mRNA expression, and cellular composition was available for 31 APAs. There was no correlation between CYP11B2 expression and cellular composition in the different groups of APA carrying KCNJ5, CACNA1D, or no mutations (Figure S3A and S3B). Similarly, no correlation was observed between CYP11B1 expression and cellular composition in APAs carrying KCNJ5 or no mutations; although APAs carrying CACNA1D mutations showed a weak correlation of CYP11B1 expression with the percentage of ZG-like and ZF-like cells, no definitive conclusions can be drawn because of the small numbers in this group (Figure S3C and S3D).

**Genotype–Phenotype Correlations**

The comparison of patients from Paris with KCNJ5 or CACNA1D1 mutations or no mutation showed that patients with KCNJ5 mutations were more often females, were diagnosed younger, and had higher minimal plasma potassium concentrations than patients from the 2 other groups (Table 2). Patients with CACNA1D1 mutations had smaller adenomas compared with those from the 2 other groups. Overall, significant differences between the 3 groups were also observed for systolic blood pressure and AVS lateralization.
ratio, but pairwise differences were not statistically significant. There was no association between the mutation status and preoperative plasma aldosterone or renin levels, the aldosterone to renin ratio, or the number of medications taken before surgery. There was also no association with postoperative blood pressure outcome as measured by blood pressure and treatment score at follow-up, cure, or improvement of hypertension.

In replication analyses, patients with KCNJ5 mutations were significantly more often females than those with no mutation in the Munich cohort (69% versus 21%; Fisher P<0.001) but not in the Torino cohort (56% versus 31%; P=0.07), resulting in a significant overall difference (P=0.001 with the Mantel–Haenszel test controlling for center). Median age at diagnosis was lower in patients with KCNJ5 mutations compared with those with no mutation in the Munich cohort (47 versus 58 years; P<0.001) but not in the Torino cohort (50 versus 51 years; P=0.50), resulting in a nonsignificant overall difference (P=0.61 with stratified linear regression controlling for center). Median minimal plasma potassium was similar in patients with KCNJ5 mutations or no mutation in both cohorts from Munich (2.8 versus 2.9 mmol/L; P=0.08) and Torino (3.0 versus 3.0 mmol/L; P=0.40), resulting in a nonsignificant overall difference (P=0.08). There was no difference in median adrenocortical size between patients with CACNA1D mutations or KCNJ5 mutations or no mutation in the 2 replication cohorts and overall.

**Discussion**

Here, we report the prevalence of recurrent somatic mutations in KCNJ5, ATP1A1, ATP2B3, and CACNA1D in a large cohort of 474 patients recruited through ENS@T and the analysis of association of the mutation status with clinical and biological parameters of the disease. CACNA1D mutations were the second most prevalent in 474 APAs enrolled in this study with a frequency of 13.6% in the Paris cohort. This is much higher than the previously described prevalence of CACNA1D in 5% to 7.8% of APAs. CACNA1D codes for the pore-forming Ca1.3 subunit of the L-type voltage-gated calcium channel. Ca1.3 contains 4 repeated domains (I–IV), each with 6 transmembrane segments (S1–S6) and a membrane-associated loop between S5 and S6. S5, S6, and the interposed loop line the channel pore. In previous studies, CACNA1D mutations were identified in genomic regions encoding transmembrane S6 segments (p.Gly403Arg, p.Ile750Met, p.Phe747Leu) known to form the channel activation gate, or in regions coding for the transmembrane S4 segment involved in voltage sensing (p.Arg990His) and the cytoplasmic S4–S5 linker coupling the voltage-sensing domain to the pore (p.Pro1336Arg, p.259Asp). We identified 10 new CACNA1D mutations, including mutations in exons different from those described in previous studies. These mutations are located in regions encoding transmembrane segments S4, S5, and S6 and cytoplasmic segments S4 and S5. These regions are predicted to be involved in voltage sensing and may increase the intracellular concentration of Ca2+. Although in vitro studies would be necessary to define the exact consequences of mutations located in this region on channel function, they are likely to alter channel function in a manner similar to that described previously. Importantly, the identification of new mutations in different regions of Ca1.3 implies the necessity of a large genotyping of CACNA1D in APAs, including exons 14 and 27.

Although there was no statistical difference in the distribution of the mutations across the different centers participating in this study, CACNA1D mutations were highest in the Paris cohort, whereas ATP2B3 mutations were highest in the Munich cohort. These differences may depend on the different characteristics of the populations investigated in these studies, in particular with regard to sex, age at diagnosis of PA, severity of the disease, in terms of aldosterone production and blood pressure levels, and size of adenoma. Given the large retrospective nature of the Paris cohort including samples diagnosed before introduction of systematic AVS for diagnostic workup, we were able to further explore the occurrence of somatic mutations depending on the diagnostic procedure for subtype identification. The distribution of mutations was significantly different depending on whether AVS or computed tomography was used (P=0.002; Table S7), suggesting that smaller adenomas harboring a CACNA1D mutation may be missed at diagnosis in the absence of AVS. Remarkably, mutations in ATP2B3 were not identified in recent studies investigating somatic mutations in APA by whole exome sequencing, indicating that these mutations may be specifically associated with a subtype of APA. Because of the small number of patients
with ATPase mutations in each single cohort, analysis of clinical and biological correlates could not be performed in our study design.

APAs present a marked histological heterogeneity, being composed of cells with characteristics of different zones of the adrenal cortex.\textsuperscript{14,25} Based on these observations, some authors have proposed that there may be more than 1 type of APA and that their cellular origins might be different. Some studies have identified a higher proportion of ZF-like cells in APA harboring KCNJ5 mutations and higher proportion of ZG-like cells in APA harboring CACNA1D and ATP1A1 mutations.\textsuperscript{14,25} In our study, histological analysis of 78 APAs did not reveal any difference in the proportion of ZF or ZG-like cells across groups with different mutation status, the majority of APAs being mainly composed of ZF-like cells. Heterogeneity between samples from different centers may explain absent replication of previously described correlations between mutation status and morphological characteristics of the APA.\textsuperscript{14,25} Although CYP11B2 expression seemed somewhat lower in APA carrying KCNJ5 mutations compared with CACNA1D mutations, similar to previous reports,\textsuperscript{17,25} it was not statistically different across mutation groups when taking into account nonmutated tumor samples and independent of the cellular phenotype (Figure S3A and S3B). This supports previous data by our group showing that despite their ZF-like cellular phenotype, APAs are composed of cells expressing ZG markers such as CYP11B2, Dab2, and CD56.\textsuperscript{26}

In this study, we have explored the clinical and biological correlates of somatic mutations in APA. Given the large individual sample sizes available from 3 different centers, we have performed an exploratory study in the Paris cohort followed by replication analyses in the Torino and Munich cohorts. This approach was chosen to obtain strong correlates that were independent of recruitment biases relative to single centers. In the Paris cohort, we confirmed the previously described association of KCNJ5 mutations with female sex and a younger age of diagnosis of PA.\textsuperscript{11} KCNJ5 mutations were also associated with higher minimal recorded potassium. These results are slightly different from our previous observations in a group of 309 patients, in which KCNJ5 mutations were associated with higher preoperative plasma aldosterone levels in a grouped analysis including all centers.\textsuperscript{11} Because the centers participating in the 2 studies are not completely overlapping, different results may reflect a strong individual contribution by 1 single center to the associations reported previously. In addition, the group without mutations analyzed previously included patients carrying CACNA1D and ATPase mutations, which were unknown at that time.

In the Paris cohort, adenoma size was associated to the mutation status. APAs with CACNA1D mutations were smaller than APAs without mutation or with KCNJ5 mutations. These observations partially confirm results described previously that compared KCNJ5-mutant APAs versus CACNA1D- and ATP1A1-mutant APAs,\textsuperscript{14} in which APAs with CACNA1D mutations or ATP1A1 mutations were smaller than APAs with KCNJ5 mutations.

The associations observed in the Paris cohort were partially replicated for age and sex, but not for minimal recorded potassium concentration, systolic blood pressure, or adenoma size. Again, these results likely reflect the different population characteristics observed among the 3 largest cohorts. While patients from the Munich cohort are more frequently males, older, and with lower minimal recorded potassium, patients from the Torino cohort present higher preoperative blood pressure, plasma aldosterone and aldosterone-to-renin ratio, and larger adenoma (Table S1). Patients from the Paris cohort are significantly younger than the patients from the other 2 cohorts. Although guidelines for the management of PA have standardized screening procedures and treatment among specialized centers,\textsuperscript{22} some patients analyzed in the present study were diagnosed before the publication and implementation of the guidelines in each center. In particular, AVS, which represents the gold standard method for subtype identification in PA, has been implemented only recently for the systematic screening of patients with PA in all centers. On the contrary, differences in the type of reference center recruiting the patients (hypertension versus endocrinology) and the criteria for patients’ referral (hypokalemia or resistant hypertension) may explain the differences in the population structure, frequency of mutations, and clinical correlations observed among centers.
**Perspectives**

*KCNJ5, CACNA1D, ATP1A1,* and *ATP2B3* mutations were identified in >50% of APAs in the ENS@T cohort. The mutation status of APA was associated to specific clinical and biochemical characteristics of PA, but this association was largely dependent on the population structure. There was marked variability in clinical and biochemical characteristics of patients with PA across centers, possibly reflecting different referral procedures, ethnic background, and the retrospective study design. Nevertheless, young women are more likely to be *KCNJ5* mutation carriers. It will be particularly relevant in the future to identify additional features or surrogate biomarkers associated with the mutation status for assessing the usefulness of stratifying patients for targeted treatment before surgery involving molecules such as verapamil specifically blocking mutated *KCNJ5* potassium channels or L-type calcium channel blockers specifically inhibiting Ca$_{1.3}$ in patients carrying *CACNA1D* mutations.

**References**


