In vivo P-glycoprotein function before and after epilepsy surgery

ABSTRACT

Objectives: To study the functional activity of the multidrug efflux transporter P-glycoprotein (Pgp) at the blood-brain barrier of patients with temporal lobe epilepsy using [R]-[11C]verapamil (VPM)-PET before and after temporal lobe surgery to assess whether postoperative changes in seizure frequency and antiepileptic drug load are associated with changes in Pgp function.

Methods: Seven patients with drug-resistant temporal lobe epilepsy underwent VPM-PET scans pre- and postsurgery. Patients were followed up for a median of 6 years (range 4–7) after surgery. Pgp immunoreactivity in surgically resected hippocampal specimens was determined with immunohistochemistry.

Results: Optimal surgical outcome, defined as seizure freedom and withdrawal of antiepileptic drugs, was associated with higher temporal lobe Pgp function before surgery, higher Pgp-positive staining in surgically resected hippocampal specimens, and reduction in global Pgp function postoperatively, compared with nonoptimal surgery outcome.

Conclusions: The data from our pilot study suggest that Pgp overactivity in epilepsy is dynamic, and complete seizure control and elimination of antiepileptic medication is associated with reversal of overactivity, although these findings will require confirmation in a larger patient cohort.

GLOSSARY

AED = antiepileptic drug; Pgp = P-glycoprotein; TL = temporal lobe; TLE = temporal lobe epilepsy; VOI = volume of interest; VPM = [R]-[11C]verapamil.

Approximately 30% of patients with epilepsy have inadequate seizure control despite taking antiepileptic drugs (AEDs). Surgery with removal of the anterior and mesial temporal lobe (TL) structures may result in favorable seizure outcome in 60% to 75% of patients, but long-term seizure freedom is achieved in only 50% with most of those patients still taking medication. Biomarkers predicting outcome of epilepsy surgery would be of high clinical interest.

According to the transporter hypothesis of drug resistance, regional overactivity of multidrug efflux transporters, such as P-glycoprotein (Pgp), impedes access of substrate AEDs to their brain target sites, thereby rendering them ineffective. The hypothesis is well supported in some animal models of epilepsy. However, relevant in vivo human data are sparse. PET with the radiolabeled Pgp substrate [R]-[11C]verapamil (VPM) can be used to noninvasively measure cerebral Pgp function. Two previous studies in patients with drug-resistant TL epilepsy (TLE) showed a trend for increased Pgp function in the ipsilateral TL compared with the contralateral side, and a significant bilateral increase in TL Pgp function compared with seizure-free patients. It remains unclear

*These authors contributed equally to this work.
whether Pgp overactivity is static or changes with disease progression, and to what extent it depends on seizure activity or AED load.9

In the present study, we examined patients with TLE using VPM-PET before and after TL surgery in order to (1) assess whether postoperative changes in seizure frequency and AED load are associated with changes in Pgp function, and (2) correlate in vivo measurements of Pgp function with ex vivo immunohistochemistry. We hypothesized that optimal outcome after surgery would be associated with a reduction in Pgp activity.

METHODS Standard protocol approvals, registrations, and patient consents. The study was approved by the Ethics Committee of the Medical University of Vienna and was conducted as a pilot study. Written informed consent was obtained from all patients.

Patients. Seven patients (2 females; median age at time of first PET scan: 50 years, range 33–54) with drug-resistant TLE10 were included in this study after surgical treatment with either selective amygdalohippocampectomy (n = 4) or anteromedial temporal lobectomy (n = 3). Patients were recruited from a group of 9 patients studied preoperatively with VPM-PET; 2 previous patients were not included because one patient was not available for follow-up after surgery and the other had no arterial input function for presurgery PET. A second VPM-PET scan was performed a median of 42 months (range 17–54) after surgery (see table for details). Surgery outcome in individual patients was ranked according to (1) complete seizure freedom after surgery, (2) the type and number of seizures during follow-up period after surgery, and (3) intake of AEDs (figure e-1 on the Neurology® Web site at Neurology.org).

PET and MRI. Dynamic VPM-PET scans were acquired on an Advance PET scanner (GE Medical Systems, Waukesha, WI) as described before,7 with arterial blood sampled throughout. Radiolabeled metabolites of VPM were measured in discrete arterial blood samples using a previously described solid-phase extraction assay.7 For all patients, T1-weighted MRIs were acquired before and after surgery with an Achieva 3.0T scanner (Philips Medical Systems, Best, the Netherlands). The PET and MRIs were processed as described previously.7 PET data analysis was performed using the Hammersmith 3-dimensional maximum probability atlas.7 Six TL volumes of interest (VOIs) (amygdala, parahippocampal gyrus, anterior temporal lobe, middle and inferior temporal gyrus, superior temporal gyrus, and posterior temporal lobe) and 2 extratemporal VOIs (superior parietal gyrus and cerebellum) were chosen for analysis. The hippocampus itself could not be analyzed because of spillover of radioactivity from the adjacent choroid plexus, which showed high VPM uptake.

Kinetic modeling of PET data was performed by using an arterial input function corrected for polar radiolabeled metabolites of VPM and a 1-tissue 2-rate constant compartment model to derive the transfer rate constant $K_1$ (mL/min/cm$^3$) of VPM from plasma into brain as an outcome parameter of Pgp function, with low $K_1$ indicating high Pgp function.7 To minimize the influence of radiolabeled metabolites of VPM on $K_1$ estimates, only the first 10 minutes of the PET data were considered for kinetic modeling.6

Immunohistochemistry. Brain tissue (hippocampus for all patients, and anterior-mesial parts of the TL of 3 patients) was removed during epilepsy surgery and fixed in 10% neutral buffered formalin immediately after surgery. For immunohistochemical staining, brain sections were prepared and probed with antibodies recognizing Pgp (clones, JSB1 1:400, C219 1:80, CA94 1:1,000; Alexis Biochemicals, Lausen, Switzerland) followed by quantitative image assessment as published.4,11

Statistical analyses. Statistical testing was performed using Prism 5.0 software (GraphPad Software, La Jolla, CA). Data are presented as mean ± SD for PET data and median (range) for demographic data. Friedman test with Dunn post hoc test was used when multiple groups were compared, and Wilcoxon matched-pairs signed rank test was used when 2 groups were compared. Correlations were assessed by calculating Spearman rank correlation coefficients ($r$). A value of $p < 0.05$ was considered statistically significant.

RESULTS Preoperative Pgp activity. In all patients, VPM-$K_1$ values from preoperative VPM-PET scans were lower in regions pathophysiologically connected to the epileptic focus (amygdala: 0.032 ± 0.011; superior temporal gyrus: 0.037 ± 0.012) compared with a reference region considered not to be involved in epileptogenic pathways (cerebellum: 0.045 ± 0.012) (figure 1A). Moreover, VPM-$K_1$ was lower in amygdala than in posterior TL (0.043 ± 0.011). There were no differences between TL VPM-$K_1$ values ipsilaterally or contralaterally to the seizure focus.

Correlation of preoperative Pgp activity with Pgp expression. Pgp immunoreactivity in surgically resected hippocampal specimens was determined using Pgp-specific antibodies (figure 1B). The higher the Pgp immunoreactivity in surgically resected hippocampal specimens, the lower were the ipsilateral VPM-$K_1$ values for anterior TL ($r = −0.821$, $p = 0.034$; figure 1C) and superior temporal gyrus ($r = −0.857$, $p = 0.024$).

Postoperative Pgp activity. Postoperative VPM-$K_1$ values were lower in superior temporal gyrus (0.033 ± 0.007) compared with cerebellum (0.041 ± 0.008) and the posterior TL (0.041 ± 0.009; figure 1D). The AED load was reduced in all patients, with 2 patients coming off all AEDs, but this did not affect peripheral metabolism of VPM; i.e., fractions of polar radiolabeled metabolites of VPM in plasma at 10 minutes after injection were similar before (0.140 ± 0.039) and after surgery (0.136 ± 0.042, $p = 0.938$).

Correlation of pre- and postoperative Pgp activity with postoperative outcome. Patients with better outcome had lower mean presurgical VPM-$K_1$ values in ipsilateral TL (higher Pgp function) than patients with poorer outcome ($r = 0.837$, $p = 0.024$) (figure 2A). Patients with better outcome had high areas of Pgp immunopositive labeling in surgically resected hippocampal specimens, while patients with poorer outcome had...
low Pgp immunoreactivity ($r = -0.782, p = 0.048$; figure 2B). In the 3 patients with best postoperative outcome (patients 1 and 6: seizure-free and without medication; patient 3: seizure-free and with very low AED load), VPM-$K_1$ values increased relative to preoperative PET scans in all brain VOIs studied, whereas $K_1$ values decreased in patients with worst outcome (figure 2C).

**DISCUSSION**

In a longitudinal pilot VPM-PET study in patients with drug-resistant TLE before and after TL surgery with a median follow-up period after surgery of 6 years (range 4–7), we detected a “normalization” of Pgp activity in patients with long-lasting seizure freedom postoperatively. Our preoperative in vivo PET findings are supported by direct correlation with Pgp immunoreactivity in surgical specimens in the same patients. Furthermore, high Pgp activity in epileptogenic pathways preoperatively was associated with better postoperative outcome. Our longitudinal PET data suggest that Pgp overactivity is dynamic rather than static, and seizure freedom and reduction in AED load are associated with reversal of upregulation.

In line with the transporter hypothesis of drug resistance, we found significantly lower VPM-$K_1$ values in some TL brain regions close to the presumed epileptic focus (amygdala, superior temporal gyrus) as compared with a reference region outside the TL (cerebellum) or with a region more distant from the focus (posterior TL), pointing to Pgp overactivity in epileptogenic brain tissue. Consistent with previous findings, we show that Pgp overactivity was not restricted to the ipsilateral TL, but also extended to the contralateral side involved in epileptogenic pathways.

After epilepsy surgery, patients with optimal surgery outcome defined as seizure freedom and withdrawal of AEDs had global VPM-$K_1$ increases relative to presurgery PET scans, suggestive of decreased Pgp function. This “return to normal” after successful epilepsy surgery is consistent with our recent report of higher VPM-$K_1$ values in pharmacosensitive compared with drug-refractory patients and with findings in postmortem brain tissue, which showed almost no Pgp overexpression in the sclerotic hippocampus of patients with epilepsy who had entered terminal remission before death.

A previous immunohistochemistry study showed higher Pgp expression in the TL white matter of patients with postoperative seizure relapse compared with postoperatively seizure-free patients, in keeping with our finding of increased Pgp activity postoperatively in patients with poorer outcome. Our preoperative in vivo PET data and ex vivo

### Table: Clinical, EEG, and MRI data for patients with drug-resistant TLE who underwent surgery

<table>
<thead>
<tr>
<th>Patient Number</th>
<th>Age, y</th>
<th>Sex</th>
<th>Syndrome</th>
<th>MRI, EEG</th>
<th>Age at onset of epilepsy, y</th>
<th>Average seizure frequency per year before surgery</th>
<th>Interval last seizure to presurgery PET, d</th>
<th>Time interval presurgery PET to surgery, mo</th>
<th>Time interval surgery to postsurgery PET, mo</th>
<th>No. of seizures since surgery</th>
<th>Follow-up period after surgery, mo</th>
<th>Outcome rank</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>37</td>
<td>M</td>
<td>RmTLE</td>
<td>RHA, RTL</td>
<td>3</td>
<td>12</td>
<td>7</td>
<td>3</td>
<td>34</td>
<td>0</td>
<td>84</td>
<td>1</td>
</tr>
<tr>
<td>2</td>
<td>50</td>
<td>M</td>
<td>LmTLE</td>
<td>LHA, LTL</td>
<td>11</td>
<td>36</td>
<td>15</td>
<td>2</td>
<td>10</td>
<td>42</td>
<td>0</td>
<td>5</td>
</tr>
<tr>
<td>3</td>
<td>52</td>
<td>F</td>
<td>LmTLE</td>
<td>LHA, BTL</td>
<td>3</td>
<td>12</td>
<td>12</td>
<td>3</td>
<td>34</td>
<td>17</td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>4</td>
<td>50</td>
<td>M</td>
<td>RmTLE</td>
<td>RHA, RTL</td>
<td>34</td>
<td>144</td>
<td>1</td>
<td>37</td>
<td>9</td>
<td>45</td>
<td>0</td>
<td>5</td>
</tr>
<tr>
<td>5</td>
<td>50</td>
<td>M</td>
<td>LmTLE</td>
<td>LHA, BTL</td>
<td>6</td>
<td>36</td>
<td>90</td>
<td>14</td>
<td>36</td>
<td>31</td>
<td>3</td>
<td>6</td>
</tr>
<tr>
<td>6</td>
<td>50</td>
<td>M</td>
<td>LmTLE</td>
<td>LHA, BTL</td>
<td>0</td>
<td>36</td>
<td>360</td>
<td>4</td>
<td>36</td>
<td>31</td>
<td>3</td>
<td>6</td>
</tr>
<tr>
<td>7</td>
<td>41</td>
<td>M</td>
<td>LmTLE</td>
<td>LHA, LTL</td>
<td>0</td>
<td>36</td>
<td>360</td>
<td>4</td>
<td>36</td>
<td>31</td>
<td>3</td>
<td>6</td>
</tr>
</tbody>
</table>

**Abbreviations:** AED = antiepileptic drug; BTL = bitemporal; CBZ = carbamazepine; LEV = levetiracetam; LHA = left hippocampal atrophy; LTL = left temporal lobe; OXC = oxcarbazepine; PHT = phenytoin; RHA = right hippocampal atrophy; RmTLE = right mesial temporal lobe epilepsy; RTL = right temporal lobe.

**Outcome ranking modified from Wieser’s outcome classification (see figure e-1).**

**Patients 2, 3, 5, and 6 had amygdalohippocampectomy and patients 1, 4, and 7 had anteromedial temporal lobectomy.**
immunohistochemistry, however, suggest that Pgp overactivity in epileptogenic pathways and hippocampal overexpression are associated with optimal outcome after TL surgery. This discrepancy in predicting postoperative outcome might be explained by methodologic differences: we quantified hippocampal Pgp immunoreactivity, whereas Kwan et al.\textsuperscript{12} used semiquantitative approaches and found differences in Pgp immunoreactivity in TL white matter, but also a trend toward higher neuronal Pgp expression in postoperatively seizure-free patients. The latter finding is in keeping with our in vivo PET finding of Pgp overactivity in TL gray matter as a predictor of optimal outcome.

Our pilot study was not designed to assess the predictive ability of VPM-PET as biomarker of surgery outcome; this should be tested prospectively in a larger cohort, and integrated with other known risk factors for less favorable outcome, such as psychiatric comorbidities and incompleteness of resection. Moreover, our study cannot distinguish between the effects of seizure cessation from dose decrease or discontinuation of AEDs on postoperative Pgp function. Our cohort was too small to statistically compare subgroups of patients, e.g., seizure-free (n = 4) vs not seizure-free (n = 3) patients, or seizure-free patients without AEDs (n = 2) vs seizure-free patients on AEDs (n = 2), with meaningful results. Alternatively, one could conduct a prospective study to leave the patients on their medication for the first year after surgery and to acquire 2 separate postsurgery PET scans, before and after the change in medication. However, our data suggest that AED exposure also affects Pgp activity, because our outcome ranking mainly based on seizure burden after surgery correlates with AED load. Another limitation of this study is that we were not able to obtain VPM-$K_1$ values in hippocampus, which would require the availability of a PET scanner with higher spatial resolution and advanced image reconstruction to overcome the spill-over from the high VPM uptake in the adjacent choroid plexus.\textsuperscript{8}

Our study cannot address whether Pgp overactivity is the cause of drug resistance in TLE, and thus “surgical” removal of this overactivity contributed to seizure freedom, or whether upregulation is a marker of seizure burden rather than a cause.\textsuperscript{9} In both scenarios, Pgp overactivity may reduce target concentrations of AEDs and thereby possibly contribute to drug resistance. Our study demonstrates that Pgp overactivity is not static, and thus could be therapeutically targeted by drugs that inhibit or downregulate Pgp to possibly prevent development of, or bring about reversal of, drug resistance mediated by Pgp.

Our data confirm previous findings that VPM-PET reflects Pgp function in vivo in patients with TLE, and
that cerebral Pgp overactivity is associated with drug resistance in TLE. Our postoperative VPM-PET data extend previous findings, indicating a “normalization” of Pgp function in those patients who benefited most from surgery. Our findings also suggest that Pgp overactivity in the TL on preoperative VPM-PET might be indicative of optimal postoperative outcome.

AUTHOR CONTRIBUTIONS
Dr. Bauer: study concept or design, acquisition of data, analysis or interpretation of data, drafting/revising the manuscript for content. Dr. Karch: analysis or interpretation of data, statistical analysis, drafting/revising the manuscript for content. Dr. Zeitlinger: study concept and design, drafting/revising the manuscript for content. Dr. Liu: acquisition of data, analysis or interpretation of data, drafting/revising the manuscript for content. Dr. Koepp, Dr. Asselin, Dr. Sisodiya, and Dr. Hainfellner: analysis or interpretation of data, drafting/revising the manuscript for content. Dr. Wadsak and Dr. Mitterhauser: drafting/revising the manuscript for acquisition of data. Dr. Müller and Dr. Pataia: study concept or design, analysis or interpretation of data, drafting/revising the manuscript for content. Dr. Langer: study concept or design, acquisition of data, analysis or interpretation of data, drafting/revising the manuscript for content.

ACKNOWLEDGMENT
The authors thank the staff of the PET center at the Department of Biomedical Imaging and Image-guided Therapy (Medical University of Vienna) and Research Nurse Edith Lackner (Department of Clinical Pharmacology, Medical University of Vienna) for their support in performing this study.

STUDY FUNDING
Supported by the European Community’s Seventh Framework Program (grant 201380, to M.J.K., S.M.S., M.-C.A., M. Müller, and O.L.), the Austrian Science Fund (FWF) (grant F 3513-B20, to M. Müller and O.L., and KLI 139-B00, to M. Müller and O.L.), and the Department of Health’s National Institute for Health Research Biomedical Research Centres funding scheme (to M.J.K. and S.M.S.).

DISCLOSURE
M. Bauer, R. Karch, M. Zeitlinger, and J. Liu report no disclosures relevant to the manuscript. M. Koepp is funded by the European Community’s Seventh Framework Program (grant 201380, expired). Dr. Koepp served on a scientific advisory board of GE Healthcare, and has received honoraria for lectures from UCB Pharma, Eisai, Novartis, and Desitin. M. Asselin is funded by the European Community’s Seventh Framework Program (grant 201380, expired). S. Sisodiya served on a scientific advisory board for UCB Pharma, and has received institutional support or honoraria for lectures from UCB Pharma, GSK, and Eisai. J. Hainfellner, W. Wadsak, and M. Mitterhauser report no disclosures relevant to the manuscript. M. Müller is funded by the European Community’s Seventh Framework Program (grant 201380, expired) and the Austrian Science Fund (FWF) (grants F 3513-B20 and KLI 139-B00). E. Pataraia reports no disclosures relevant to the manuscript. O. Langer is funded by the European Community’s Seventh Framework Program (grant 201380, expired) and the Austrian Science Fund (FWF) (grants F 3513-B20 and KLI 139-B00). Go to Neurology.org for full disclosures.

Received December 20, 2013. Accepted in final form July 11, 2014.

REFERENCES


Save These Dates for AAN CME Opportunities!

Mark these dates on your calendar for exciting continuing education conferences by the American Academy of Neurology. Learn more at AAN.com/conferences.

AAN Fall Conference
- October 31-November 2, 2014, Las Vegas, NV, The Cosmopolitan of Las Vegas

AAN Annual Meeting
- April 18-25, 2015, Washington, DC, Walter E. Washington Convention Center

BrainPAC

BrainPAC is the American Academy of Neurology’s federal political action committee.

- Since its inception, nearly 3,000 AAN members have contributed $1,500,000 to BrainPAC.

- BrainPAC will contribute $600,000 to candidates running for Congress in 2014.

- During the 2012 congressional campaign, 89 percent of candidates supported by BrainPAC won their elections.

BrainPAC supports both Democrats and Republicans who support issues important to the practice of neurology and the care of patients with neurologic conditions. US AAN members are invited to learn more at BrainPAC.org.
In vivo P-glycoprotein function before and after epilepsy surgery
Martin Bauer, Rudolf Karch, Markus Zeitlinger, et al.
Neurology 2014;83;1326-1331 Published Online before print September 3, 2014
DOI 10.1212/WNL.0000000000000858

This information is current as of September 3, 2014

Updated Information & Services
including high resolution figures, can be found at:
http://www.neurology.org/content/83/15/1326.full.html

Supplementary Material
Supplementary material can be found at:
http://www.neurology.org/content/suppl/2014/09/03/WNL.0000000000000858.DC1.html

References
This article cites 13 articles, 1 of which you can access for free at:
http://www.neurology.org/content/83/15/1326.full.html##ref-list-1

Subspecialty Collections
This article, along with others on similar topics, appears in the following collection(s):
All Epilepsy/Seizures
http://www.neurology.org/cgi/collection/all_epilepsy_seizures
Epilepsy surgery
http://www.neurology.org/cgi/collection/epilepsy_surgery_
Functional neuroimaging
http://www.neurology.org/cgi/collection/functional_neuroimaging
PET
http://www.neurology.org/cgi/collection/pet
PET in epilepsy
http://www.neurology.org/cgi/collection/pet_in_epilepsy

Permissions & Licensing
Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at:
http://www.neurology.org/misc/about.xhtml#permissions

Reprints
Information about ordering reprints can be found online:
http://www.neurology.org/misc/addir.xhtml#reprintsus

Neurology® is the official journal of the American Academy of Neurology. Published continuously since 1951, it is now a weekly with 48 issues per year. Copyright © 2014 American Academy of Neurology. All rights reserved. Print ISSN: 0028-3878. Online ISSN: 1526-632X.