The fundamental role of inflammation in atherosclerosis is well established through animal and autopsy studies, which have promoted the development of imaging approaches for in vivo assessment of plaque inflammation. Among many biological targets implicated in plaque inflammation, macrophages and neovessels have gained broad interest because they are abundant and detectable by clinical imaging modalities such as 18F-fluorodeoxyglucose (FDG) positron emission tomography (PET) and dynamic contrast-enhanced (DCE) MRI. Increased metabolic activity measured as target-to-background ratio (TBR) is thought to indicate enhanced macrophage recruitment and activation. In contrast, the transfer constant ($K_{\text{trans}}$) in DCE-MRI is considered to reflect microvessel density and permeability. As integral components of the inflammatory process, previous studies have suggested that cellular events, including macrophage recruitment and activation, and vascular adaptation, including angiogenesis and increased permeability, could affect each other during atherogenesis.

We studied their relationship using in vivo imaging.

**Methods**

**Study Sample**

All investigations were approved by the institutional review board. From March 2011 to June 2013, 46 patients with carotid plaque (thickness $\geq 2$ mm) by ultrasound were enrolled with informed consent. Indications for ultrasound referrals included ischemic stroke, transient ischemic attack, coronary artery disease, and hypertension. Subjects underwent DCE-MRI followed by 18F-FDG PET/computed tomography (CT) within 2 weeks. Bilateral carotids were imaged, of which 25 had a history of transient ischemic attack ($n=2$) or ischemic stroke ($n=23$) appropriate to the distribution of carotid plaque inflammation by inflammatory cells.

**Results**

Overall, the correlation between target-to-background ratio and $K_{\text{trans}}$ was weak and marginal ($r=0.22$; $P=0.068$). They were correlated in the symptomatic plaques ($r=0.59$; $P=0.006$) but not in the asymptomatic plaques ($r=0.07$; $P=0.625$; $P=0.033$ for difference in $r$). Neither target-to-background ratio nor $K_{\text{trans}}$ was significantly higher in the symptomatic plaques, but both showed an inverse relationship with time since last neurological event ($r=-0.94$ and $-0.69$ for target-to-background ratio and $K_{\text{trans}}$, respectively).

**Conclusions**

The correlation between 18F-fluorodeoxyglucose positron emission tomography and dynamic contrast-enhanced MRI measurements varied with clinical conditions, pointing to a complex interplay between macrophages and neovessels under different pathophysiological conditions. The moderate correlation shown only in symptomatic plaques indicates the presence of acute plaque inflammation with increased metabolic activity and cytokine production by inflammatory cells.

**Key Words:** atherosclerosis ■ carotid arteries ■ magnetic resonance imaging ■ positron emission tomography
artery within the past 4 months. The ipsilateral carotid plaque was considered the culprit based on clinical workup to rule out other common causes and defined as symptomatic.

Dynamic Contrast-Enhanced MRI
A previously published DCE-MRI protocol was used. Images were analyzed by 2 readers (X.Z. and H.C.) who followed a previously described approach to obtain plaque $K_{trans}$ while blinded to clinical information and PET/CT results (Figure 1).5

18F-Fluorodeoxyglucose Positron Emission Tomography/Computed Tomography
PET/CT was performed on a dedicated PET/CT scanner (Biograph Truepoint, Siemens). Images were analyzed on a workstation (SynGo Somaris/5 software, Siemens) by 2 readers (Z.G. and C.L.) who were blinded to clinical information and DCE-MRI results. The approach described by Rudd et al9 was followed to obtain maximum TBR across slices that match MRI coverage (Figure 1).

Statistical Analysis
Spearman correlation coefficient ($r$) was used to assess correlation between variables. Correlation coefficients of mutually exclusive subgroups were compared using Fisher $z$ transformation.9 Mann–Whitney test was used to compare TBR and $K_{trans}$ between symptomatic and asymptomatic plaques.

Results
After excluding 5 subjects because of poor MR image quality, 41 subjects (34 men; aged 65±12 years) contributed 68 carotid arteries with atherosclerotic plaque (thickness >2 mm), of which 20 (29.4%) were symptomatic as a result of recent ischemic events (median: 29 days; range: 4–120 days).

The correlation between TBR and $K_{trans}$ was overall weak and marginal ($r=0.22; P=0.068$). Although no systemic condition modified their relationship, symptomatic status seemed to be a critical factor (Table). The 2 measurements were correlated in the symptomatic plaques but not in the asymptomatic plaques ($r=0.59$ versus $r=0.07; P=0.033$ for difference in $r$; Figure 2).

Table. Correlation Between TBR and $K_{trans}$ Under Different Clinical Conditions

<table>
<thead>
<tr>
<th>Clinical Variables</th>
<th>$r$</th>
<th>$P$ Value</th>
<th>$P$ Value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female (n=14)</td>
<td>0.18</td>
<td>0.53</td>
<td>0.80</td>
</tr>
<tr>
<td>Male (n=54)</td>
<td>0.26</td>
<td>0.06</td>
<td></td>
</tr>
<tr>
<td>Symptomatic</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>+(n=20)</td>
<td>0.59</td>
<td>0.006</td>
<td>0.033</td>
</tr>
<tr>
<td>-(n=48)</td>
<td>0.07</td>
<td>0.625</td>
<td></td>
</tr>
<tr>
<td>Coronary artery disease</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>+(n=39)</td>
<td>0.31</td>
<td>0.054</td>
<td>0.90</td>
</tr>
<tr>
<td>-(n=29)</td>
<td>0.08</td>
<td>0.68</td>
<td></td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>+(n=17)</td>
<td>0.04</td>
<td>0.87</td>
<td>0.38</td>
</tr>
<tr>
<td>-(n=51)</td>
<td>0.24</td>
<td>0.08</td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>+(n=52)</td>
<td>0.17</td>
<td>0.24</td>
<td>0.50</td>
</tr>
<tr>
<td>-(n=16)</td>
<td>0.42</td>
<td>0.10</td>
<td></td>
</tr>
<tr>
<td>Hypercholesterolemia</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>+(n=18)</td>
<td>0.20</td>
<td>0.43</td>
<td>0.80</td>
</tr>
<tr>
<td>-(n=50)</td>
<td>0.27</td>
<td>0.06</td>
<td></td>
</tr>
<tr>
<td>Smoker</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>+(n=35)</td>
<td>0.23</td>
<td>0.19</td>
<td>0.35</td>
</tr>
<tr>
<td>-(n=33)</td>
<td>0.20</td>
<td>0.26</td>
<td></td>
</tr>
</tbody>
</table>

*Comparison of correlation coefficients between 2 mutually exclusive subgroups.
Neither TBR nor $K^{\text{trans}}$ was significantly higher in the symptomatic plaques compared with the asymptomatic ones. Nonetheless, both showed an inverse relationship with time since last ischemic event ($r=-0.94$ and $-0.69$ for TBR and $K^{\text{trans}}$, respectively) in the symptomatic group.

Discussion

There is generally a hypothesized link between plaque macrophages and neovascularization. However, the nature and strength of their relationship are vague. Neovessels serve as a major entry route for inflammatory cells, and activated macrophages in turn may promote angiogenesis by secreting vascular endothelial growth factor and others alike. Such a relationship is complicated by the identification of macrophage subpopulations with diverse phenotypes and functions in atherogenesis. Although M1 macrophages may outnumber M2 macrophages in established lesions, it is suggested that the latter is involved more in angiogenesis. It is also equivocal whether improved perfusion via angiogenesis enhances (bringing more tracers) or suppresses (alleviating hypoxia) FDG uptake. The weak and marginal correlation between TBR and $K^{\text{trans}}$ reflects the many influential factors on macrophages and neovessels, as well as their imaging read-outs, which implies that the correlation could vary under different pathophysiological conditions.

A moderate correlation was noted in symptomatic plaques but not in asymptomatic ones. A prominent phenomenon in symptomatic plaques is the exacerbation of plaque inflammation, especially in the acute phase. The recruitment and activation of inflammatory cells lead to a parallel increase in proinflammatory cytokines that may have mediated the correlation between $^{18}$F-FDG PET and DCE-MRI signals seen in symptomatic plaques. We attribute the correlation to cytokines that influence microvascular permeability rather than microvessel density because the postevent decrease of $K^{\text{trans}}$ was rapid and paralleled the decrease in TBR, which could not be easily explained by regression in neovascularization.

In contrast to the traditional view of atherosclerosis as a chronic inflammatory process, our findings add in vivo evidence to the presence of acute plaque inflammation concurrently with clinical events, which was characterized by increased metabolic activity and cytokine production by inflammatory cells. Whether it is responsible for or just a consequence of clinical events remains to be investigated.

Conclusions

Our data from a heterogeneous population showed that the correlation between $^{18}$F-FDG PET and DCE-MRI measurements varied with clinical conditions, pointing to a complex interplay between macrophages and neovessels under different pathophysiological conditions. The moderate correlation shown only in symptomatic plaques indicates the presence of acute plaque inflammation and its clinical relevance.

Sources of Funding

This work was supported by Army Medical Research Funds of China (11BJZ19) and National Natural Science Foundation of China (81371540).

Disclosures

Dr Yuan has been a consultant for Philips Healthcare. The other authors report no conflicts.

References


Varying Correlation Between $^{18}$F-Fluorodeoxyglucose Positron Emission Tomography and Dynamic Contrast-Enhanced MRI in Carotid Atherosclerosis: Implications for Plaque Inflammation

Juan Wang, Hongbin Liu, Jie Sun, Hao Xue, Leixing Xie, Shengyuan Yu, Changzai Liang, Xu Han, Zhiwei Guan, Liqun Wei, Chun Yuan, Xihai Zhao and Huijun Chen

*Stroke*. 2014;45:1842-1845; originally published online May 1, 2014;
doi: 10.1161/STROKEAHA.114.005147

*Stroke* is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2014 American Heart Association, Inc. All rights reserved.
Print ISSN: 0039-2499. Online ISSN: 1524-4628

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://stroke.ahajournals.org/content/45/6/1842

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in *Stroke* 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 *Stroke* is online at:
http://stroke.ahajournals.org/subscriptions/