Arterial Stiffness and Pharmacological Interventions – The TRanscend Arterial stiffness Substudy (TRANS study)

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Abstract: The degree of arterial stiffness is correlated with the risk of cardiovascular diseases and it is a powerful predictor for morbidity and mortality. Studies have shown that arterial stiffness reduction is associated with an improvement in survival. Reduction of arterial stiffness by pharmacological drugs varies according to the drugs and doses used and duration of treatment. This effect on the arteries differs among the various classes of drugs and among individual drugs in the same class. Quantification of the stiffness and other properties of the arterial wall can be used to monitor the responses to therapy in individuals with hypertension and other cardiovascular diseases. These measures can then be used as surrogate markers for the risk of clinical events. Inhibition of the renin-angiotensin system (RAS) is associated with an important decrease in cardiovascular risk. Findings from clinical trials support the hypothesis that the protective effects of RAS inhibition are partly independent from blood pressure reduction and related to several mechanisms including vascular protective effects. The aim of the TRanscend Arterial stiffness Substudy (TRANS) is to assess the effect of an angiotensin II receptor blocker (ARB), telmisartan, on the arterial stiffness in a subgroup of patients from the Telmisartan Randomized Assessment Study in a CE intolerant subjects with cardiovascular Disease (TRANSCEND) trial. The TRANSCEND trial is an international, multicenter, randomized double blind placebo controlled trial of telmisartan that enrolled patients at high risk for cardiovascular events. Some clinical baseline data of the TRANS substudy are reported. When completed, the results of the TRANS substudy will show whether the beneficial effects of treatment with telmisartan on cardiovascular outcome may be related to an improvement in arterial stiffness.

Keywords: arterial stiffness, cardiovascular prevention, ARBs, telmisartan, pulse wave velocity, antihypertensive

Introduction

The degree of arterial stiffness, obtained in various populations, has been found to be a powerful independent marker of vascular target organ damage and an independent prognostic predictor for cardiovascular morbidity, as well as cardiovascular and all-cause mortality (Blacher et al 1999; Laurent et al 2001, 2003; Meaume et al 2001; Boutouyrie et al 2002; Cruickshank et al 2002; Dernellis et al 2005; Shokawa et al 2005; Sutton-Tyrrell et al 2005; Mattace-Raso et al 2006; Willum-Hansen et al 2006). Measuring pulse wave velocity (PWV) to assess arterial stiffness is a simple and reproducible method. The underlying principles and technique of this method have been described in detail previously (Asmar 1999). Several experimental studies have shown that PWV is related to the arterial wall structure, function, geometry and endothelium functions (Asmar 1999). Validation studies have shown that automatic measurements of PWV are simple, non-invasive, accurate, and reproducible (Asmar et al 1995; Van Bortel et al 2002; Laurent et al 2006), making this technique a convenient, sensitive and useful tool in physiological and pharmacological studies.
Basic pharmacological concepts of arterial stiffness

Several important points serve to better understand the effects of pharmacological intervention on arterial stiffness.

The arterial site

Atherosclerosis, arterial abnormalities, and their progression vary in different arterial sites. Arteries are heterogenous in structure and the arterial site has to be considered in assessment of the pharmacological treatment (Asmar 1999). The impact of a given pharmacological agent may differ on the various components of the arterial wall (elastin, collagen, muscle) according to its pharmacodynamic properties. It is logical to assume that the arterial effects of a given drug administered at a given dose and period of time may differ according to the arterial site, which may be more elastic (aorta, carotid) or more muscular (radial) arteries (Topouchian et al 1999). Figure 1 shows an example of the different effects on the arterial sites produced by the same antihypertensive drug in the same patients (Asmar 1999; Topouchian et al 1999).

Duration of treatment

Since several mechanisms may be involved in producing reductions in arterial stiffness with a given treatment, assessment of arterial stiffness has to distinguish between the effects of acute, short-term, or long-term chronic treatments. For example, after acute administration of an antihypertensive drug, improvement of arterial stiffness is principally related to functional or mechanical mechanisms such as reduction of distension pressure, reduction of smooth muscle tone, enhancement of endothelial functions, whereas after long-term chronic treatment, additional mechanisms can be involved, eg, changes in the arterial geometry and structure, reduction in degree of fibrosis, increase in elastin/collagen ratio, remodeling of the arterial wall (Asmar 1999; Laurent et al 2002). Experts agree that assessment of arterial stiffness after a long-term treatment period should be preferred because of the underlying pathophysiological mechanisms involved and because acute effects may not predict long-term efficacy.

Drug doses

In assessing the efficacy of a drug, one needs to consider the dose/effect relationship. This relationship may differ from other properties of the drug such as the dose/effect of its antihypertensive properties (Asmar 1999; Laurent et al 2002). A clear dissociation between the antihypertensive effect and the reduction of arterial stiffness by angiotensin converting enzyme (ACE) inhibitors has been reported. There is evidence that the effect on arterial wall properties can be seen at higher doses of ACE inhibitors than with the doses required for effective blood pressure (BP) reduction in

Figure 1 Change in arterial distensibility after antihypertensive treatment in three arterial sites: abdominal aorta, carotid artery and brachial artery. Significant site effect was observed:

- **Carotid artery**
  - Before: 1.6 ± 0.7
  - After: 2.2 ± 0.8

- **Brachial artery**
  - Before: 0.7 ± 0.3
  - After: 1.3 ± 0.6

- **Abdominal aorta**
  - Before: 0.9 ± 0.3
  - After: 1.3 ± 0.3

(Derived from Topouchian et al 1999)
hypertensive patients (Asmar et al 1992a, b). These findings, independent of the drug effects on BP, have been recently confirmed by the results of large clinical studies on patients at high cardiovascular risk. In these studies, positive results on arterial stiffness have been observed with high doses of ACE inhibitors (Yusuf et al 2000; PROGRESS Collaborative Group 2001; Fox et al 2003).

**Effect of antihypertensive agents on arterial stiffness**

Several pharmacological studies have evaluated the effects of antihypertensive drugs on arterial stiffness. Table 1 shows the effects on PWV of different antihypertensive drug classes, administered double blind, either short-term (<28 days) or long-term (≥28 days) (Asmar 1999; Asmar et al 2002; Rajzer et al 2003; White et al 2003; Handbook of Hypertension 2006). During long-term treatment, improvements in central and peripheral arterial stiffness have been found with ACE inhibitors. Less marked improvement has been reported with angiotensin II receptor blockers. Beta-blockers have exhibited variable results according to the particular drugs used. With calcium channel blockers, the results are more complex. At the aortic level, all calcium channel blockers have shown a significant reduction in arterial stiffness, in parallel with BP reduction, but at the peripheral level, arterial stiffness reduction was less evident. Diuretics have shown no significant effects. Therefore, antihypertensive treatment is associated with variable effects on arterial stiffness, due not only to BP reduction but also duration of treatment. More specific long-term studies are needed.

Arterial stiffness or distensibility, evaluated by PWV or other measurements, depends on BP level. Therefore, any blood pressure decrease (decrease of the distension pressure) is theoretically associated with a decrease of PWV. For this reason, analysis of PWV changes according to BP change is important. Different situations have been described in the literature (Asmar 1999), the most frequent of which are:

- decrease of PWV and BP
- decrease of PWV and small changes of BP (this may reflect an independent arterial effect of the drug)
- decrease of BP with unchanged PWV. This may reflect a relatively “harmful” impact on the arterial wall by other mechanisms such as increasing sympathetic tone (hydralazine) or heart rate, or other mechanisms related to the impact of the drug concerned.

Taking into consideration these aspects, analysis of PWV changes should be discussed according to other hemodynamic changes such as BP, heart rate, and peripheral resistance.

**Future perspectives**

Activation of the renin-angiotensin system (RAS) has been implicated in the pathogenesis of a wide variety of diseases affecting the cardiovascular system. Clinical trials have shown that inhibition of the RAS, particularly by ACE inhibitors, is associated with important decreases in cardiovascular risk in a broad range of patients (Yusuf et al 2000; PROGRESS Collaborative Group 2001; Fox et al 2003; HOPE/HOPE-TOO Study Investigators 2005). More recently, angiotensin II receptor blockers (ARBs) have been reported to reduce cardiovascular events in different populations, such as patients with diabetes, hypertension, or heart failure (Lacourcière et al 1998; Neutel et al 1998; Dalhöf et al 2002; Young et al 2004). Both pathophysiological studies and clinical trials support the hypothesis that the protective effects of RAS inhibition observed with ACE inhibitors and ARBs are, at least partly, independent from their blood-pressure-lowering effects. Consistent improvement in arterial stiffness seems to be observed with some classes of drugs that target the RAS. Whether this effect on arterial stiffness may be observed with other drugs, such as ARBs, in different populations needs clarification.

There are several observations regarding ACE inhibitors (eg, adverse effects and lack of full RAS blockade during

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**Table I Effects of antihypertensive drugs on pulse wave velocity**

<table>
<thead>
<tr>
<th></th>
<th>Short term treatment &lt;28 days</th>
<th>Long term treatment ≥28 days</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Aorta</td>
<td>Arm/Leg</td>
</tr>
<tr>
<td>Vasodilators</td>
<td>=</td>
<td>=</td>
</tr>
<tr>
<td>Beta-blockers</td>
<td>/ =</td>
<td></td>
</tr>
<tr>
<td>Diuretics</td>
<td>=</td>
<td>/ =</td>
</tr>
<tr>
<td>Calcium antagonists</td>
<td>=</td>
<td></td>
</tr>
<tr>
<td>ACE inhibitors</td>
<td>/ =</td>
<td></td>
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<tr>
<td>ARBs</td>
<td>/ =</td>
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</tbody>
</table>

**Abbreviations:** ACE, angiotensin converting enzymes; ARB, angiotensin II receptor blocker
chronic treatment) and ARBs (powerful blocking effect on angiotensin II receptors) that can be discussed. While the relationship between reduction in arterial stiffness and decreased cardiovascular risk has been shown with ACE inhibitors, this relationship is not clear with ARBs. Thus, it is important to assess the arterial effect of ARBs after long-term chronic treatment in high risk patients.

Telmisartan is an ARB with a specific pharmacokinetic profile (Lacourcière et al 1998; Neutel et al 1998). Several studies have investigated the effects of telmisartan on end-organ damage and demonstrated beneficial results. Results of a pilot crossover, placebo-controlled study showed that telmisartan reduced arterial stiffness in hypertensive patients with type 2 diabetes (Asmar et al 2002). The TRANSCEND study (Telmisartan Randomized Assessment Study in ACE iNtolerant subjects with cardiovascular Disease) seeks to determine whether the ARB, telmisartan, is superior to placebo in high risk patients who are intolerant to ACE inhibitors, in reducing the risk of cardiovascular death, myocardial infarction, stroke, and hospitalization for heart failure. Details of the TRANSCEND trial have been published previously elsewhere (Lacourcière et al 1998; Neutel et al 1998; Teo et al 2004; Asmar 2006). The TRanscend Arterial stiffNess Substudy (TRANS) is designed to determine the effect of telmisartan on arterial stiffness. This will assess whether the expected results of the TRANSCEND trial might be explained by the vascular protective effects of telmisartan and estimate whether the efficacy of telmisartan on cardiovascular outcomes may be related to an improvement in arterial stiffness.

Methods
Study design
The TRANSCEND study was an international, multicenter, randomized, double-blind, placebo-controlled clinical trial (Teo et al 2004). The primary objective of the TRANSCEND study was to determine if telmisartan 80 mg daily was superior to placebo in reducing the composite endpoint of cardiovascular death, myocardial infarction, stroke, or hospitalization for heart failure. In this substudy (TRANS), assessment of arterial stiffness was performed initially during the run-in period before randomization, after 6 months of randomized treatment and yearly for 3 years. The primary objective of the TRANS study was to investigate whether the lower incidence of cardiovascular events with telmisartan was correlated with an improvement in arterial stiffness. Secondary objectives were to compare early changes in arterial stiffness between the treatment groups, to estimate the prognostic factors relating to changes in arterial stiffness, and to determine if an early change in arterial stiffness might be a prognostic factor for cardiovascular events.

Patients
Patient eligibility, inclusion, and exclusion criteria of the TRANSCEND study have been described elsewhere (Teo et al 2004). Approximately 6000 high risk patients were enrolled.

Specific non-inclusion criteria for the TRANS study were: known significant peripheral vascular disease with proximal artery stenosis or limb amputation, history of vascular surgery at the level of the carotid artery, femoral artery, or on the aorta, body mass index >40 kg/m², and atrial fibrillation. The purpose of these additional exclusion criteria was to avoid conditions that may impair the quality and reliability of PWV measurements.

Assessment of arterial stiffness
The Complior® device (Artech Medical, Paris, France) was used to record PWV in this study because of its reproducibility (Asmar 1999) and simplicity of use. These properties were important in this large international multicenter study, with many sites obtaining recordings and a centralized assessment in a core-laboratory (Asmar et al 2001). This device also allowed a valid measurement of the velocity of the same pulse wave recorded simultaneously in different sites of the arterial tree. Basic principles of PWV measurements have been described elsewhere (Asmar 1999). The pressure pulse generated by ventricular ejection is propagated along the arterial tree at a speed determined by the geometric and elastic properties of the arterial wall. PWV is calculated from measurements of pulse transit time and the distance traveled by the pulse between two recording sites, according to the following formula: PWV (m/s) = distance (m)/transit time (s). All measurements are recorded on the right side of the body with the patient lying at ease in the supine position.

Blood pressure measurement
Blood pressure (BP) was measured before PWV recording and according to standard published guidelines (O’Brien et al 2005). In order to minimize its variability the following recommendations were followed: patients in supine position, their arms bared, and supported at heart level. Patients refrained from smoking or ingesting caffeine during the 30 minutes preceding the measurement. Measurements began after at least 5 minutes of rest. The appropriate cuff size was used to ensure accurate measurement.
Logistics and organization
A total of 34 centers (from 12 countries) were selected; all were provided with Complior devices. All investigators participated in specific training sessions on PWV assessment organized by the study core laboratory. In order to limit inter-center variability of the PWV measurements, and to ensure data homogeneity, a pre-study assessment was carried out as part of the investigators’ certification to participate in the study.

Data analysis
Demographic and baseline characteristics
Demographic and baseline characteristics, risk factors, and relevant clinical variables (eg, cardiovascular history and medication), current status of diabetes, current medical conditions, and concurrent medications were summarized by treatment group and overall inclusion.

Efficacy analysis
Primary efficacy criterion
The primary efficacy parameter was the change from baseline, after 3 years of treatment, in the aortic stiffness as assessed by carotid-femoral PWV. The change from baseline was defined as the mean value at the end of the treatment period minus the mean value at baseline. The change from baseline was compared between the treatment groups using a t-test. If the data did not conform to the assumptions of normality, then a non-parametric test was used. Multiple regression analysis was used to explore BP changes and baseline characteristics. The same analysis was performed with the change from baseline after 6 months of treatment. In a second step, repeated measures of analysis of variance with PWV at 6 months, 1 year, 2 years and 3 years of treatment and between treatment groups was performed.

Secondary efficacy criteria
Regression analyses were performed to estimate factors related to carotid-femoral PWV at baseline. Only known factors or factors significantly related to PWV in simple regression analysis were included in a stepwise selection multiple regression analysis. If the treatment effect on PWV was not constant throughout the trial, regression analyses were performed for changes after 6 months of treatment and for changes between 2 and 3 years of treatment.

Results
Logistics and organization
Among the centers participating in the TRANSCEND trial, 34 centers from 12 countries participated in the TRANS study. All of these centers were provided with Complior devices. Investigators from all centers participated in specific training sessions and received their certification by the end of 2003. Twenty-six of the 34 centers participated actively by including patients in the TRANS study. Table 2 shows the numbers of centers and patients enrolled per center.

Study population
Recruitment for the TRANS study took place from November 2003 to June 2004. During this period 296 patients from 26 centers were enrolled. Table 2 shows their geographic distribution. At randomization, 17 patients were excluded because their PWV recordings were judged (n = 17) to be of insufficient quality by the PWV Core laboratory. Therefore a total of 279 patients were included in the TRANS study. Their main baseline clinical characteristics are shown in Table 3.

Table 3 Baseline clinical characteristics of the TRANS study population

<table>
<thead>
<tr>
<th>Gender</th>
<th>N = 279</th>
<th>Mean ± SD</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>67 ± 7</td>
<td>43–88</td>
<td></td>
</tr>
<tr>
<td>M: 160 (57.3%)</td>
<td>F: 119 (42.6%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Height (cm)</td>
<td>166 ± 9</td>
<td>140–191</td>
<td></td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>77 ± 13</td>
<td>41–123</td>
<td></td>
</tr>
<tr>
<td>Systolic BP (mmHg)</td>
<td>139 ± 17</td>
<td>95–200</td>
<td></td>
</tr>
<tr>
<td>Diastolic BP (mmHg)</td>
<td>80 ± 10</td>
<td>51–112</td>
<td></td>
</tr>
<tr>
<td>Heart rate (beat/min)</td>
<td>66 ± 11</td>
<td>45–111</td>
<td></td>
</tr>
<tr>
<td>Carotid-femoral PWV (m/s)</td>
<td>10.0 ± 2.6</td>
<td>4.7–21.4</td>
<td></td>
</tr>
<tr>
<td>Carotid-radial PWV (m/s)</td>
<td>9.1 ± 1.4</td>
<td>5.3–13.6</td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: BP, blood pressure; PWV, pulse wave velocity.
and pulse wave velocity (carotid-femoral = 10.0 ± 2.6 m/s; carotid-radial = 9.1 ± 1.4 m/s). These relatively “normal” values may be related to the baseline treatments received by this population at high cardiovascular risk.

Tables 4 and 5 show that patients included in the TRANS sub-study are at high cardiovascular risk, as intended for the TRANSCEND study. Results show that about 68% of patients had previous coronary artery disease, more than 30% had experienced previous stroke, 13% had high-risk diabetes with evidence of end organ damage and about 4% experienced recent Transient Ischemic Attacks (TIA). Details of biochemical results are shown in Table 5.

**Conclusion**

The TRANS arterial stiffness sub-study of the TRANSCEND trial assessed the effect of telmisartan on arterial stiffness in patients at high cardiovascular risk. The results of this sub-study should be helpful in the interpretation of the results of both the TRANSCEND and the parallel ONgoing Telmisartan Alone and in combination with Ramipril Global Endpoint Trial (ONTARGET) trials. In fact, among the several mechanisms supporting the cardioprotective effects of RAS inhibition, improvement of arterial stiffness is of major importance. Moreover, the TRANS substudy will provide information about direct assessment of the arterial wall which constitutes the major target and the site of treatment of cardiovascular disease.

**References**


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**Table 4** Diagnosis and previous diseases of the TRANS study population

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>N</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coronary artery disease</td>
<td>190</td>
<td>68</td>
</tr>
<tr>
<td>Peripheral arterial disease</td>
<td>18</td>
<td>6</td>
</tr>
<tr>
<td>Previous stroke</td>
<td>87</td>
<td>31</td>
</tr>
<tr>
<td>TIA (&gt;7 days and &lt;1 year)</td>
<td>11</td>
<td>4</td>
</tr>
<tr>
<td>High-risk diabetes</td>
<td>35</td>
<td>13</td>
</tr>
</tbody>
</table>

**Abbreviations:** TIA, transient ischemic attack.

**Table 5** Biochemical results of the TRANS study population

<table>
<thead>
<tr>
<th>Biochemical Results</th>
<th>N</th>
<th>Mean</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Creatinine (mmol/L)</td>
<td>280</td>
<td>91.8</td>
<td>23.8</td>
</tr>
<tr>
<td>Potassium (mmol/L)</td>
<td>280</td>
<td>4.4</td>
<td>0.4</td>
</tr>
<tr>
<td>Sodium (mmol/L)</td>
<td>280</td>
<td>140.7</td>
<td>4.1</td>
</tr>
<tr>
<td>Urea (mmol/L)</td>
<td>279</td>
<td>6.8</td>
<td>3.3</td>
</tr>
<tr>
<td>Plasma glucose (mmol/L)</td>
<td>280</td>
<td>6.5</td>
<td>2.4</td>
</tr>
<tr>
<td>Cholesterol (mmol/L)</td>
<td>280</td>
<td>5.2</td>
<td>1.2</td>
</tr>
<tr>
<td>Triglycerides (mmol/L)</td>
<td>280</td>
<td>1.9</td>
<td>1.4</td>
</tr>
<tr>
<td>HDL (mmol/L)</td>
<td>279</td>
<td>1.3</td>
<td>0.3</td>
</tr>
<tr>
<td>LDL (mmol/L)</td>
<td>277</td>
<td>3.1</td>
<td>1.0</td>
</tr>
</tbody>
</table>

**Abbreviations:** HDL, high density lipoprotein; LDL, low density lipoprotein.
Arterial stiffness in high risk patients


Coronary artery disease in women: a review on prevention, pathophysiology, diagnosis, and treatment

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Abstract: Despite numerous studies on women's cardiac health throughout the past decade, the number of female deaths caused by cardiovascular disease still rises and remains the leading cause of death in women in most areas of the world. Novel studies have demonstrated that cardiovascular disease, and more specifically coronary artery disease presentations in women, are different than those in men. In addition, pathology and pathophysiology of the disease present significant gender differences, which leads to difficulties concerning diagnosis, treatment and outcome of the female population. The reason for this disparity is all steps for female cardiovascular disease evaluation, treatment and prevention are not well elucidated; and an area for future research. This review brings together the most recent studies published in the field of coronary artery disease in women and points out new directions for future investigation on some of the important issues.

Keywords: coronary artery disease, women, risk factors, prevention, diagnosis, treatment.

Introduction
The first female-specific recommendations for preventive cardiology were published in 1999 (Mosca et al 1999). Even though research in the treatment of cardiovascular disease (CVD) had advanced in many areas, it remains the leading cause of death in women in most parts of the world. Studies have shown that 500 thousand women die of CVD every year in the United States, somewhat near one death every minute (American Heart Association 2003). Such index exceeds not only the number of deaths in men, but also the next seven causes of death in women combined, and more importantly, coronary artery disease (CAD) is believed to be the major cause responsible for these deaths (American Heart Association 2003). Over a quarter of a million deaths per year are attributed to CAD alone in the United States (Merz et al 2004). Although already high, these figures are expected to rise even more during the next decades, due to an increase of diabetes and obesity, as well as the aging of the world population (Merz et al 2004).

Even though women have a higher frequency of chest pain/angina than men, the incidence of obstructive CAD in the female population is lower when compared with men with similar symptoms (Kenedy et al 1982; Diamond et al 1983; Merz et al 1999). In addition, it would appear that young women with obstructive CAD have a worse prognosis after acute myocardial infarction (AMI), whereas older women in similar circumstances often present with larger number of comorbidities that adversely influence the outcome, when compared to men (Coronado et al 1997). Women with acute coronary syndromes (ACS) are also less likely to receive rapid effective diagnosis and treatment than are men (Ayanian and Epstein 1991; Maynard et al 1996; Pope et al 2000).

Regarding the North American population, the Women's Ischemic Syndrome Evaluation (WISE) study workshop (Hayes et al 2004; Maseri 2004; Nabel et al 2004; Pepine et al 2004; Shaw et al 2004; Waters et al 2004) from the National Heart, Lung and...