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Ambulatory monitoring of the cardiovascular system: the role of Pulse Wave Velocity

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1. Introduction

Currently the leading cause of mortality in western countries, cardiovascular diseases (CVD) are largely responsible for the ever increasing costs of healthcare systems. During the last decade it was believed that the best trade-off between quality and costs of a healthcare system would pass through the promotion of healthy lifestyles, the early diagnosis of CVD, and the implantation of home-based rehabilitation programs. Hence, the development of novel healthcare structures will irremediably require the availability of techniques allowing the monitoring of patients' health status at their homes. Unfortunately, the ambulatory monitoring of the cardiovascular vital parameters has not evolved as required to reach this aim.

To be exploitable in the long term, ambulatory monitors must of course provide reliable vital information, but even more important, they must be comfortable and inconspicuous: whoever has experimented wearing an ambulatory blood pressure monitor (ABPM) for 24 hours understands what cumbersomeness means. Surprisingly, even if intermittent and obtrusive, ABPM prevails nowadays as the single available method to assess a vascular-related index at home. Hence, a clear demand to biomedical engineers arises: healthcare actors and patients require the development of new monitoring techniques allowing the non-invasive, unobtrusive, automatic and continuous assessment of the cardiac and vascular health status. Concerning cardiac health status, the ambulatory measurement of the electrical and mechanical activities of the heart is already possible by the joint analysis of the electrocardiogram, the phonocardiogram and the impedancardiogram. But surprisingly, little has been proposed so far for the ambulatory monitoring of vascular-related parameters.

Because several studies have recently highlighted the important role that arterial stiffness plays in the development of CVD, and since central stiffness has been shown to be the best independent predictor of both cardiovascular and all-cause mortality, one might suggest stiffness to be the missing vascular-related parameter in ambulatory cardiovascular

monitoring. However, the only available technique for measuring arterial stiffness non-invasively so far is the so-called Pulse Wave Velocity (PWV). In this chapter we will see that the state of the art in PWV assessment is not compatible with the requirements of ambulatory monitoring. The goal of our work is thus to examine the limitations of the current techniques, and to explore the introduction of new approaches that might allow PWV to be established as the new gold-standard of vascular health in ambulatory monitoring.

This chapter is organized as follows: in Section 2 we introduce the phenomenon of pulse propagation through the arterial tree. In section 3 we provide a large review on the clinical relevance of aortic stiffness and its surrogate, PWV. In Section 4 we perform an updated analysis of the currently existing techniques available for the non-invasive assessment of PWV. Section 5 describes a novel approach to the measurement of PWV based on a non-obtrusive and unsupervised beat-to-beat detection of pressure pulses at the sternum. Finally, Section 6 reviews the historic and current trends on the use of PWV as a non-obtrusive surrogate for arterial blood pressure.

2. The genesis and propagation of pressure pulses in the arterial tree

In cardiovascular research and clinical practice, PWV refers to the velocity of a pressure pulse that propagates through the arterial tree. In particular, we are interested in those pressure pulses generated during left ventricular ejection: at the opening of the aortic valve, the sudden rise of aortic pressure is absorbed by the elastic aorta walls. Subsequently a pulse wave naturally propagates along the aorta exchanging energy between the aortic wall and the aortic blood flow (Figure 1). At each arterial bifurcation, a fraction of the energy is transmitted to the following arteries, while a portion is reflected backwards. Note that one can easily palpate the arrival of arterial pressure pulses at any superficial artery, such as the temporal, carotid or radial artery: already in the year 1500, traditional chinese medicine performed clinical diagnosis by palpating the arrival of pressure pulses at the radial artery (King et al., 2002). But why do clinicians nowadays get interested on the velocity of such pulses, and especially in the aorta? The reason is that the velocity of propagation of aortic pressure pulses depends on the elastic and geometric properties of the aortic wall. We will show later that while arterial stiffness is difficult to measure non-invasively, PWV is nowadays available *in vivo* to clinicians. Hence, the PWV parameter is an easily-accessible potential surrogate for the constitutive properties of the arterial walls.

In order to provide a better understanding of the biomechanics of pulse propagation, we describe here the commonly accepted model of pulse propagation: the Moens-Korteweg equation. For a complete derivation of the model see (Nichols & O'Rourke, 2005). This model assumes an artery to be a straight circular tube with thin elastic walls, and assumes it being filled with an inviscid, homogeneous and incompressible fluid. Under these hypotheses the velocity of a pressure pulse propagating through the arterial wall is predicted to be:

$$PWV^2 = Eh / d\rho \quad (1)$$

where E stands for the elasticity of the wall (Young's modulus), h for its thickness, D for its diameter and ρ corresponds to the density of the fluid. Even if this model is only a rough

approximation of reality, it provides an intuitive insight on the propagation phenomenon in arteries and, in particular, it predicts that, the stiffer the artery (increased E), the faster a pressure pulse will propagate through it. Therefore, for large elastic arteries such the aorta where the thickness to diameter ratio (h / D) is almost invariable, PWV is expected to carry relevant information related to arterial stiffness.

3. Clinical relevance of Pulse Wave Velocity as a marker of arterial stiffness

We already demonstrated that, from a biomechanical point of view, the velocity of propagation of pressure pulses in large arteries is a surrogate indicator of arterial stiffness. Due to the recent commercialization of semi-automatic devices performing routine measurements of PWV, numerous studies investigating the clinical relevance of arterial stiffness have been conducted during the last decade (Asmar, 1999). In this section we review the most prominent conclusions of these studies. An additional review is given by (Mitchel, 2009).

Cardiovascular disease is the leading cause of morbidity and mortality in western countries and is associated with changes in the arterial structure and function. In particular, arterial stiffening has a central role in the development of such diseases. Nowadays, aortic PWV is considered the gold standard for the assessment of arterial stiffness and is one of the most robust parameters for the prediction of cardiovascular events. Because the structure of the arterial wall differs between the central (elastic) and the peripheral (muscular) arteries, several PWV values are encountered along the arterial tree, with increasing stiffness when moving to the periphery. Because carotid-to-femoral PWV is considered as the standard measurement of aortic arterial stiffness, we will refer to it as simply PWV. In the following we review the most important factors influencing PWV, then we justify the need for a reliable PWV monitoring: on one hand we analyse the pathophysiological consequences of increased arterial stiffness and, on the other hand we highlight the clinical relevance of PWV as an independent marker of cardiovascular risk.

Fig. 1. Genesis of pressure pulses: after the opening of the aortic valve the pulse propagates through the aorta exchanging energy between the aortic wall and the blood flow. Adapted with permission from (Laurent & Cockcroft, 2008).

Fig. 2. The dependency of PWV with age for central elastic arteries (dashed line) and peripheral muscular arteries (continuous line). Adapted from (Avolio et al., 1985).

Major determinants of PWV under normal conditions

Before elucidating the role that PWV plays in the generation and diagnosis of pathological situations, it is necessary to understand which are its determinant factors under normal conditions. It is currently accepted that the four major determinants of PWV are age, blood pressure, gender and heart rate.

Age affects the wall properties of central elastic arteries (aorta, carotid, iliac) in a different manner than in muscular arteries (brachial, radial, femoral, popliteal). With increasing age the pulsatile strain breaks the elastic fibers, which are replaced by collagen (Faber & Oller-Hou, 1952). These changes in the arterial structure lead to increased arterial stiffness, and consequently to increased central PWV (Figure 2). On the other hand, there is only little alteration of distensibility of the muscular, *i.e.*, distal, arteries with age (Avolio, 1983; Avolio, 1985; Nichols et al., 2008). This fact supports the use of generalized transfer functions to calculate the central aortic pressure wave from the radial pressure wave in adults of all ages, as will be described in Section 4 (Nichols, 2005).

Arterial blood pressure is also a major determinant of PWV. Increased blood pressure is associated with increased arterial stiffness and vice versa. Ejection of blood into the aorta generates a pressure wave that travels along the whole arterial vascular tree. A reflected wave that travels backwards to the ascending aorta is principally generated in the small peripheral resistance arterioles. With increasing arterial stiffness both the forward and the reflected waves propagate more rapidly along the vessels. Consequently, instead of reaching back the aorta during the diastole, the reflected pulse wave reaches it during the systole. This results in an increase of aortic pressure during systole and reduced pressure during diastole, thus leading to an increase of the so-called Pulsatile Pressure (PP) parameter (Figure 3). Asmar (Asmar et al., 2005) studied large untreated populations of normotensive and hypertensive subjects and found that the two major determinants of PWV were age and systolic blood pressure in both groups. This result confirms the close interdependence between systolic blood pressure and arterial stiffness.

Concerning gender, studies in children revealed no gender difference in PWV, whereas in young and middle age, healthy adult men displayed higher PWV values compared to women (London et al., 1995; Sonesson et al., 1993). Indeed premenopausal women show lower carotid-radial PWV values than age-matched men, but carotid-femoral PWV is found to be similar. Once women become postmenopausal, PWV values become similar to those of age-matched men (London, 1995).

Fig. 3. Consequences of increased arterial stiffness on central blood pressure: increase of systolic and decrease of diastolic central pressures. Pulsatile Pressure is defined as the difference of both pressure amplitudes. PP_n stands for PP under normal conditions and PP_s stands for PP under stiff conditions.

Heart rate is related to PWV through two independent mechanisms. Firstly, heart rate influences PWV because of the frequency-dependant viscoelasticity of the arterial wall: if heart rate increases, the time allowed to the vessels to distend is reduced, resulting in an increased rigidity of the arterial wall. Hence, increasing rate is associated with increasing arterial stiffness. In a recent study, (Benetos et al., 2002) showed that particularly in hypertensive patients increased heart rate was one of the major determinants of accelerated progression of arterial stiffness. Secondly, heart rate is related to PWV through the influence of the sympathetic nervous system: sympathetic activation is associated with increased stiffness of the arteries (Boutouyrie et al., 1994) due to an increase in heart rate, blood pressure and smooth muscle cells tonus.

Why keep arterial stiffness under control?

Up to this point we simply outlined that increased arterial stiffness appears to be normally associated to factors such as aging and blood pressure, among others. As natural as it seems, one might then wonder, why do we need to keep arterial stiffness under controlled (low) values? We will answer this question backwards: what would happen if we did not do so? In other words, we are interested in understanding the pathophysiological consequences of increased arterial stiffness.

Firstly we describe the role of arterial stiffness in the development of endothelial dysfunction. Endothelial dysfunction is the first step in the development of atherosclerosis and plays a central role in the clinical emergence and progression of atherosclerotic vascular disease (Figure 4). The endothelium plays not only an important role in atherogenesis but also in the functional regulation of arterial compliance since endothelial cells release a number of vasoactive mediators such as the vasodilator nitric oxide (NO) and the vasoconstrictor endothelin. The complex interplay between endothelial function and arterial stiffness leads to a vicious cycle of events, as illustrated in Figure 4 (Dart & Kingwell, 2001).

Fig. 4. Vicious circle of events resulting from endothelial dysfunction and augmented arterial stiffness.

Increased arterial stiffness is also an important determinant of myocardium and coronary perfusions. In Figure 3 we already described the mechanism through which increasing arterial stiffness leads to augmented central PP, *i.e.*, the difference between systolic and diastolic aortic pressures. The increase in central systolic pressure is thus associated with an increased afterload, which if persistent, promotes the development of left ventricular (LV) hypertrophy, an independent cardiovascular risk factor (Bouthier et al., 1985; Toprak et al., 2009). Conversely, the decrease in central diastolic pressure compromises myocardial blood supply, particularly in patients with coronary artery stenosis. However, the increased LV-mass induced by the augmented afterload will require an increased oxygen supply. Therefore, a mismatch between oxygen demand and supply may occur, leading to myocardial ischemia, LV diastolic and later systolic dysfunction. The full mechanism is illustrated in Figure 5.

Finally, the widening of central PP induced by increasing arterial stiffness may affect the vascular bed of several end-organs, particularly of brain and kidney. Because both organs are continually and passively perfused at high-volume flow throughout systole and diastole, and because their vascular resistance is very low, pulsations of pressure and flow are directly transmitted to the relatively unprotected vascular bed. By contrast, other organs if exposed to increased PP may protect themselves by vasoconstriction (O'Rourke & Safar, 2005). This unique situation predisposes the brain and kidney to earlier micro- and macrovascular injuries (Laurent et al, 2003; Henskens et al, 2008; Fesler et al., 2007).

Relevance of PWV in clinical conditions

We already described the factors that modify arterial stiffness in normal conditions. We also reviewed the consequences of an increase of arterial stiffness to endothelial function, coronary perfusion and possible damages to heart muscle, brain and kidneys. We are interested in reviewing now the broad uses of PWV as an independent cardiovascular risk factor and its interaction with the others classical risk factors such as arterial hypertension, diabetes mellitus, and dyslipidemia. The independent predictive value of PWV for cardiovascular and all-cause mortality is finally underlined.

Fig. 5. Effects of increased arterial stiffness on the myocardium and its function.

Structural arterial abnormalities are already observed at an early stage of hypertension. Changes in the structure of the arterial wall, particularly of the matrix and the three-dimensional organization of the smooth muscle cells, have an important impact in determining arterial stiffness. Studies of white-coat hypertension (Glen et al., 1996) and borderline hypertension (Girerd et al., 1989) showed higher values of PWV compared to controls. Moreover, for a similar blood pressure, PWV was higher in patients than in controls, suggesting that the increased PWV was not only due to the elevated blood pressure but also to some structural changes of the arterial wall. As already mentioned, increased arterial stiffness leads to increased central systolic blood pressure, augmented afterload and ultimately left ventricular hypertrophy (Figure 5), which is itself a major cardiovascular risk factor (Bouthier et al., 1985; Lorell et al., 2000). Arterial stiffness and its associated augmented PWV is now recognized as an independent marker of cardiovascular risk (Willum-Hansen et al., 2006; Laurent et al., 2001) especially in hypertensive patients (Mancia et al., 2007).

Diabetes mellitus is one of the major cardiovascular risk factors and has been associated with premature atherosclerosis. There are numerous studies showing that both patients suffering from type 1 diabetes (van Ittersum et al., 2004) and type 2 diabetes (Cruickshank et al., 2002; Schram et al., 2004) have an increased arterial stiffness compared to controls. The increase in arterial stiffening in patients with type 1 and type 2 diabetes mellitus is evident even before clinical micro- and macrovascular complications occur (Giannattasio et al., 1999; Ravikumar et al., 2002), being already present at the stage of impaired glucose tolerance (Henry et al., 2003). Moreover, as in hypertensive patients, increased aortic PWV is identified as an independent predictor of mortality in diabetics (Cruickshank et al., 2002). The increase in arterial stiffness in patients suffering from diabetes mellitus is multifactorial (Creafer et al., 2003) and is associated with structural (Airaksinen et al., 1993) (extracellular matrix), functional (endothelium dysfunction) and metabolic (increased oxidative stress, decreased nitric oxide bioavailability) alterations. The most important mechanism seems to be the glycation of the extracellular matrix with the formation of advanced glycation end-products (AGEs): hyperglycemia favors AGEs formation which is responsible for the altered collagen content of the arterial wall (Airaksinen et al., 1993). A new class of drugs called "AGE breakers" is able to decrease the numbers of collagen cross-links and improve arterial stiffness in both diabetic rats (Wolffenbuttel et al., 1998) and humans (Kass et al., 2001).

Fig. 6. Changes in mean BP (solid circles) and aortic PWV (open circles) of patients with end-stage renal disease for survivors and non-survivors: despite achievement of target BP non-survivors showed no improvement or even an increase in PWV, demonstrating on the one hand the presence of a pressure-independent component of PWV, and on the other hand, the relevance of PWV as an independent predictor for mortality. Adapted from (Guerin et al., 2001).

The association between lipids and arterial stiffness has been studied since the seventies, but the results are so far controversial. In patients suffering from coronary artery disease (CAD), an association between increased arterial stiffness and higher LDL has been proved (Cameron et al., 1995). On the other hand, in the general population the results regarding the relationship between LDL and arterial stiffness are controversial and some studies have reported a lack of association between total cholesterol and arterial stiffness (Dart et al., 2004).

Acute smoking is associated with increased arterial stiffness in healthy individuals and several patients subgroups, including normotensive, hypertensive and CAD. Studies on the chronic effects of smoking demonstrated contradictory results. However, the largest studies showed that chronic cigarette smoking was associated with increased PWV both in normotensive and hypertensive subjects (Liang et al., 2001; Jatoi et al., 2007).

Arterial hypertension and arterial stiffness induce the same end-organ damages such as coronary artery disease (CAD), cerebrovascular disease (CVD), peripheral artery disease (PAD) and chronic kidney disease (CKD) (Mancia et al., 2002). Many studies showed an association between increased PWV and the severity of CAD (Hirai et al., 1989; Giannattasio et al., 2007), CVD (Laurent et al., 2003; Henskens et al., 2008; Mattace-Raso et al., 2006), PAD (van Popele et al., 2001) and CKD (London et al., 1996; Shinohara et al., 2004).

Beyond its predictive value of morbidity, aortic stiffness appears to be relevant because of its independent predictive value for all-cause and cardiovascular mortality, in patients with arterial hypertension (Laurent et al., 2001), with type-2 diabetes (Cruickshank et al., 2002),

with CKD (Blacher et al., 1999), older age (Mattace-Raso et al., 2006; Meaume et al., 2001; Sutton-Tyrrell et al., 2005) and even in the general population (Willum-Hansen et al., 2006). Figure 6 demonstrates PWV to be a blood-pressure-independent cardiovascular risk factor for patients with end-stage renal disease.

Hence, if it is nowadays accepted (Nilsson et al., 2009) that arterial stiffness and PWV may be regarded as a “global” risk factor reflecting the vascular damage provoked by the different classical risk factors and time, how can we explain its limited use in clinical practice? The main reason seems to be its difficulty to measure. While blood pressure and heart rate are at present easily automatically measured, reliable PWV measurements still require complex recent equipments and, even worse, they require the continuous presence of a skilled well-trained operator.

4. Measuring aortic Pulse Wave Velocity *in vivo*

In the preceding sections we pointed out the need of including a vascular-related parameter into ambulatory monitoring, and we highlighted the clinical relevance of PWV as a surrogate measurement of arterial stiffness. In this section we analyse the strategies and devices that have been so far developed to measure PWV *in vivo*. Although in some cases these techniques rely on rather simplistic physiologic and anatomic approximations, their commercialization has triggered the interest in the diagnostic and prognostic uses of PWV (Boutouyrie et al., 2009). For the sake of clearness, Table 1 summarizes the different approaches described in this section.

In general, given an arterial segment of length D , we define its PWV as:

$$\text{PWV} = D / \text{PTT} \quad (2)$$

where PTT is the so-called Pulse Transit Time, *i.e.*, the time that a pressure pulse will require to travel through the whole segment. Formally PTT is defined as:

$$\text{PTT} = \text{PAT}_d - \text{PAT}_p \quad (3)$$

where PAT_p corresponds to the arrival time of the pressure pulse at the proximal (closer to the heart) extremity of the artery, and PAT_d corresponds to the arrival time of the pressure pulse at its distal (distant to the heart) extremity.

In particular, concerning the aorta, we define PWV as the average velocity of a systolic pressure pulse travelling from the aortic valve (proximal point) to the iliac bifurcation (distal point), as Figure 7 illustrates. Note that this definition concerns the propagation of the pulse through anatomically rather different aortic segments, namely the ascending aorta, the aortic arch and the descending aorta. Accordingly, we re-define aortic PWV as:

$$\text{PWV} = (D_{\text{asc}} + D_{\text{arch}} + D_{\text{desc}}) / \text{PTT}_a \quad (4)$$

Fig. 7. Aortic PWV is defined as the average velocity of a pressure pulse when travelling from the aortic valve, through the aortic arc until it reaches the iliac bifurcation.

Hence, the *in vivo* determination of aortic PWV is a two-step problem: first one needs to detect the arrival times of a pressure pulse at both the ascending aorta and the iliac bifurcation, and secondly one needs to precisely measure the distance travelled by the pulses.

A first group of aortic PWV measurement methods corresponds to those approaches that measure transit times in the aorta in a straight-forward fashion, that is, without relying in any model-based consideration. Because the aorta is not easily accessible by neither optical nor mechanical means, the strategy is to detect the arrival of a pressure pulse at two substitute arterial sites, remaining as close as possible to the aorta (Asmar et al., 1995). Starting from the aorta and moving to the periphery, the first arteries that are accessible are the common carotid arteries (at each side of the neck) and the common femoral arteries (at the upper part of both thighs, near the pelvis). This family of devices assumes thus the carotid-to-femoral transit time to be the best surrogate of the aortic transit time. Currently four commercial automatic devices based on this assumption are available: the Complior (Artech Medical, Paris, France), the Vicorder (Skidmore Medical, Bristol, UK), the SphygmoCor (AtCor Medical, New South Wales, Australia), and the PulsePen (DiaTecne, Milano, Italy). While Complior simultaneously records the arrival of a pressure pulse at the carotid and femoral arteries by means of two pressure sensors (Figure 8), SphygmoCor and PulsePen require performing the two measurements sequentially by means of a single hand-

held tonometer. A simultaneously recorded ECG supports the post-processing of the data obtained from both measurements (Figure 9). It has been suggested that because measurements are not performed on the same systolic pressure pulses, the SphygmoCor might introduce artifactual PTT variability (Rajzer et al., 2008). Unfortunately, there is so far no consensus on whether the transit times obtained by Complior and SphygmoCor display significant differences (Millasseau et al., 2005; Rajzer et al., 2008). Concerning the estimation of the travelled distance D , each manufacturer provides different and inconsistent recommendations on how to derive D from superficial morphological measurements with a tape (Rajzer et al., 2008). Regrettably Complior, SphygmoCor and PulsePen require the constant presence of a skilled operator who manually localizes the carotid and femoral arteries and holds the pressure sensors during the examination.

Fig. 8. Pulse transit time (PTT) as measured by Complior. The arrival time of a pressure pulse is simultaneously detected on the carotid and femoral artery. Complior implements as well a correlation-based PTT estimation.

Fig. 9. Pulse transit time (PTT) as measured by SphygmoCor. The delay between the R-Wave on the ECG and the arrival time of a pressure pulse is sequentially measured on the carotid and the femoral arteries. Both measurements are further combined to obtain a single PTT value.

Fig. 10. Time to reflection (T_r) is defined as the arrival time of a pressure pulse that has been reflected in the arterial tree and travels back towards the heart. This example illustrates an important shortening of T_r for a male adult when performing a handgrip effort. During the sustained handgrip, mean arterial pressure is augmented, increasing the stiffness of the aorta and thus aortic PWV. Consequently, the reflected pulse reaches the aortic valve prematurely: T_r is shifted to the left in the bottom pressure pulse.

A second group of devices estimate aortic transit time based on wave reflection theory (Segers et al., 2009). It is generally accepted (Westerhof et al., 2005) that any discontinuity on the arterial tree encountered by a pressure pulse traveling from the heart to the periphery (downstream) will create a reflected wave on the opposite direction (upstream). Main reflection sites in humans are high-resistance arterioles and major arterial branching points. In particular, the iliac bifurcation at the distal extremity of the descending aorta has empirically been shown to be a main source of pulse reflections (Latham et al., 1985). Consequently, a pulse pressure generated at the aortic valve is expected to propagate downstream through the aorta, to reflect at the iliac bifurcation and to propagate upstream towards the heart, reaching its initial point after T_r seconds (Figure 10). Commonly depicted as Time to Reflection, T_r is related to the aortic length (D) and the aortic pulse wave velocity as:

$$T_r = 2 D / \text{PWV} \quad (5)$$

Even though the concept of a unique and discrete reflection point in the arterial tree is not widely accepted and is currently the source of fervent discussions (Nichols, 2009), PWV values derived from the time to reflection method have been shown to be at least positively correlated to PWV measured by Complior, $r=0.69$ (Baulmann et al., 2008) and $r=0.36$ (Rajzer et al., 2008).

Fig. 11. Example of aortic pressure pulse, radial pressure pulse and the generalized transfer function that relates them. Adapted from (Chen et al., 1997).

Obviously a main issue is how to record aortic pressure pulses non-invasively (Hirata et al., 2006). Two approaches have been proposed so far. A first device, Arteriograph (TensioMed, Budapest, Hungary), records a sequence of pressure pulses at the upper arm by inflating a brachial cuff above systolic pressure, typically 35 mmHg. The brachial pressure waveform is then simply assumed to be a surrogate of the aortic one. Regardless of its manifest lack of methodological formalism, Arteriograph is so far the unique fully automatic and unsupervised commercial available device. Similarly, some recent studies aim at analyzing pressure pulses recorded at the finger to obtain similar results (Millasseau et al., 2006). A second device, SphygmoCor (AtCor Medical, New South Wales, Australia), records pressure pulses at the radial artery by a hand-held tonometer and then estimates an associated aortic pressure pulse by applying a generalized transfer function. In brief, the generalized transfer function approach relies on a series of empirical studies conducted during the 90s in which it was proven that the relationship between aortic and radial pressure pulses is consistent among subjects and unaffected even by aging and drug action (O'Rourke, 2009). Consequently, transfer functions provide a method for universally estimating aortic pressure pulses from radial artery measurements in a non-invasive fashion. Figure 11 illustrates the modulus and phase of the widely accepted aortic-to-radial general transfer function (Chen, et al. 1997). Large population studies (Gallagher et al., 2004) and numerical models of the arterial tree (Karamanoglu et al., 1995) have shown that the generalized transfer function is indeed consistently unchanged for frequencies below 5 Hz.

A third group of approaches comprises those developments based on the R-wave-gated pulse transit time. In brief, this technique exploits the strength of the ECG signal on the human body, and assumes its R-wave to trigger the genesis of pressure pulses in the aorta, at time T_{R-wave} . Then, by detecting the arrival time of a pressure pulse on a distal location (PAT_d) one calculates:

$$PTT_{R-wave} = PAT_d - T_{R-wave} \quad (6)$$

Unfortunately the physiological hypothesis relating PTT_{R-wave} to PWV neglects the effects of cardiac isovolumetric contraction: indeed, after the onset of the ventricle depolarization (R-Wave in the ECG) left ventricles start contracting while the aortic valve remains closed. It is only when the left ventricle pressure exceeds the aortic one, that the aortic valve opens and generates the aortic pressure pulse. The introduced delay is commonly known as Pre-Ejection Period (PEP) and depends on physiological variables such as cardiac preload, central arterial pressure, and cardiac contractility (Li & Belz, 1993). Hence, PTT_{R-wave} is to be corrected for the delay introduced by PEP as proposed in (Payne et al., 2006):

$$PTT'_{R-wave} = PAT_d - (T_{R-wave} + PEP) \quad (7)$$

Several strategies to assess PEP non-invasively are nowadays available, mainly based on the joint analysis of the ECG (Berntson et al., 2004) and either an impedance cardiogram or a phono-cardiogram (Lababidi et al., 1970; DeMarzo & Lang, 1996; Ahlström 2008). Nevertheless, even obviating the PEP correction, PTT_{R-wave} has been shown to be correlated with PWV ($r=0.37$) (Abassade & Baudouy, 2002) and systolic blood pressure ($r=0.64$) (Payne et al. 2006). Concerning the distal detection of the pressure pulse arrival time (PAT_d), different approaches have been proposed so far. We describe here the most relevant ones. Novacor (Cedex, France) commercializes an ambulatory method to monitor PWV based on a fully automatic auscultatory approach: the so-called Qkd index. Qkd is defined as the time interval between the R-Wave on the ECG and the second Korotkoff sound detected on an inflated brachial cuff. The device is currently being used to evaluate long-term evolution of systemic sclerosis in large population studies (Constans et al., 2007). A different technology, photo-plethysmography, is probably the approach that has given rise to the largest number of research developments and studies in the field (Naschitz et al., 2005). Being non-obtrusive and cheap, this technology consists in illuminating a human perfused tissue with an infrared light source and to analyse the changes in absorption due to arterial pulsatility (Allen, 2007). Each time a pressure pulse reaches the illuminated region, the absorption of light is increased due to a redistribution of volumes in the arterial and capillary beds. The analysis of temporal series of light absorption then allows the detection of the arrival of the pressure pulse. Regrettably, to obtain reliable photo-plethysmographic signals is not a simple task and, so far, only those body locations displaying very rich capillary beds have been exploited: namely the finger tips or phalanxes (Smith et al., 1999; Fung et al. 2004; Schwartz, 2004; Muehlsteff et al., 2006, Banet, 2009), the toes (Sharwood-smith et al., 2006; Nitzan et al., 2001) and the ear lobe (Franchi et al, 1996). Undoubtedly, the listed locations correspond to the classical placement of probes for pulse oximetry, or SpO_2 , in clinical practice (Webster, 1997). It is to be highlighted that recent studies have investigated the feasibility of performing pulse oximetry at innovative regions such as the sternum (Vetter et al., 2009). To reduce the cumbersomeness of measuring ECG has also been the aim of recent researches: a capacitively-coupled ECG mounted on a chair has been recently proposed to monitor PTT_{R-wave} in computer users (Kim et al., 2006).

Finally, an emerging non-invasive technique remains to be cited, although its implantation in ambulatory monitoring seems nowadays unfeasible: the phase-contrast MR imaging (PCMRI) (Lotz et al., 2002). PCMRI opens the possibility to perform local measurements of PWV for any given segment of the aorta, by simply defining two regions of interest on the

image: a proximal and a distal region. By analysing the evolution of the regional blood flow velocity in each region, one determines the arrival times (PTT_p and PTT_d) of the pressure pulse. Because the distance between both aortic regions (D) can now be precisely measured, this approach is expected to provide highly accurate regional aortic PWV measurements. PCMRI was already introduced in the 90s (Mohiaddin et al., 1989), but the recent advances in MRI capturing rates seem to be encouraging the apparition of new studies (Boese et al., 2000; Gang et al., 2004; Laffon et al., 2004; Giri et al., 2007; Butlin et al., 2008). Fitting in the same category, some studies have been published on the assessment of PAT_d by means of ultrasound Doppler probes (Baguet et al. 2003; Meinders et al., 2001; Jiang et al., 2008).

Note that we have intentionally skipped from our analysis some works that have been performed on the tracking of pressure pulses artificially induced to the arterial wall by mechanical oscillators (Nichols & O'Rourke, 2005). Similarly, we have excluded those works based on the analysis of pressure-diameter and flow-diameter measurements (Westerhof et al., 2005).

Segments of the arterial tree	Method	Measurements of PTT and D	AMB	COM
Carotid to Femoral PTT (simultaneous)	PTT is measured by two pressure sensors placed over the carotid and femoral arteries. D is estimated from superficial morphologic measurements.	No	Complior Vicorder
	Carotid to Femoral PTT (sequential)	PTT is measured by a single pressure sensor placed sequentially over the carotid and femoral arteries. ECG is used for synchronization purposes. D is estimated from superficial morphologic measurements.	No	SphygmoCor PulsePen
Time to reflection, from brachial pressure pulse	PTT is measured by extracting Tr from the brachial pressure pulse recorded by a brachial obtrusive cuff. D is estimated from superficial morphologic measurements.	Yes	Arteriograph
	Time to reflection, from radial pressure pulse (generalized transfer function)	The aortic pressure pulse is estimated by applying a generalized transfer function to a radial pressure pulse recorded by a handheld tonometer. PTT is measured from the associated Tr . D is estimated from superficial morphologic measurements.	No	SphygmoCor

ECG to brachial pulse transfer time	PTT is approximated as the delay between the R-Wave at the ECG, and the arrival of the pressure pulse at the brachial artery, recorded by a brachial obtrusive cuff. <i>D</i> is estimated from superficial morphologic measurements	Yes	NovaCor
ViSi	PTT is approximated as the delay between the R-Wave at the ECG, and the arrival of the pressure pulse at the digital artery, recorded by photo-plethysmography. <i>D</i> is estimated from superficial morphologic measurements	Yes	-
MR Imaging of aortic blood flow	PTT is measured by detecting the arrival of the pressure pulse at two or more different aortic sites, associated to different regions of interest in the PCMR images. <i>D</i> is accurately determined from the images.	No	-
Sequential Doppler measurements of aortic blood flow	PTT is measured by detecting the arrival of the pressure pulse at two or more different aortic sites, by performing ECG-gated Doppler measurements. <i>D</i> is estimated from superficial morphologic measurements	No	-

Table 1. Summary of most relevant approaches to measure aortic PWV. Detailed descriptions are available on the text. PTT stands for Pulse Transit Time, *D* for distance, AMB for ambulatory compatibility, and COMM for commercial devices.

Determination of Pulse Arrival Times

Up to this point we assumed that detecting the arrival time of a pressure pulse at a certain aortic site was an obvious operation. Yet, clinical experience has shown that this is not the case: given a pressure pulse recorded either by tonometry, photo-plethysmography or any other measurement technique, it is not straight-forward to objectively define its Pulse Arrival Time, or PAT (Chiu et al., 1991; Solà et al., 2009). In the past, originally based on the analysis of pressure pulses obtained from cardiac catheterization, PAT was proposed to be estimated by identifying a collection of characteristic points (Chiu et al., 1991). Simply stated, a characteristic point is a typical feature that is expected to be found in any pressure pulse waveform. In particular one is interested in those features describing the position of the wavefront of a pulse. The justification is rather simple: on one hand the wavefront is the most patent representative feature of the arrival time of a pulse (Chiu et al., 1991), and on

the other hand it is expected to be free of deformations created by reflected waves, thus maintaining its identity while propagating through the arterial tree. Conversely, any other feature of the pressure pulse waveform cannot be assigned an identity in a straight-forward manner (Westerhof et al., 2005).

Fig. 12. Characteristic points encountered on a pressure pulse (bold curve) according to state-of-the-art definitions. Time zero corresponds to the R-Wave of a simultaneously recorded ECG.

Hence, the state-of-the-art extraction of characteristic points relies on the morphologic analysis of the wavefront of pressure pulses. The analysis is commonly based on empirically-determined rules, as illustrated in Figure 12. For the sake of completeness, we briefly describe them: the foot of a pressure pulse (FOOT) is defined as the last minimum of the pressure waveform before the beginning of its upstroke. In (Chiu et al., 1991) an iterative threshold-and-slope technique to robustly detect FOOT was proposed. The partial amplitude on the rising edge of the pulse (PARE) is defined as the location at which the pressure pulse reaches a certain percentage of its foot-to-peak amplitude. The maximum of the pressure pulse (MAX) is defined as the time at which the pressure pulse reaches its maximum amplitude. The maximum of the first derivative (D1) is defined as the location of the steepest rise of the pressure pulse. The first derivative is commonly computed using the central difference algorithm in order to reduce noise influences (Mathews & Fink, 2004). The maximum of the second derivative (D2) is defined as the location of the maximum inflection

point of the pressure pulse at its anacrotic phase. Finally, the intersecting tangent (TAN) is defined as the intersection of a tangent line to the steepest segment of the upstroke, and a tangent line to the foot of the pressure pulse. Nowadays TAN is the characteristic point commonly implemented in commercial devices such as SphygmoCor.

Fig. 13. Six seconds of simulated photo-plethysmographic signals, corresponding to different signal-to-noise scenarios. Noise model according to (Hayes & Smith, 1998).

Fig. 14. Given a pressure pulse, the state-of-the-art strategy consists on determining its Pulse Arrival Time (PAT) by identifying a characteristic point on its waveform. The novel parametric approach consists in modeling the whole pulse wavefront, and to extract from this model a PAT-related index. While highly correlated ($r=0.99$), the robustness to noise is five times higher.

Unfortunately, when targeting ambulatory applications, one must consider that the recording of pressure pulses is severely affected by measurement and artefact noises, with signal to noise ratios (SNR) reaching values below 10dB. In order to illustrate the influence of such noises on the waveform analysis of pressure pulses, Figure 13 displays a series of

simulated photo-plethysmographic recordings to which noises of different amplitudes have been introduced (Hayes & Smith, 1998). Unquestionably, the straight-forward identification of state-of-the-art characteristic points under such noisy conditions leads to erratic and unrepeatable results (Solà et al., 2009).

A common strategy to reduce the influence of noise in the determination of PAT is the pre-processing of the raw recorded data. Commercial PWV devices mostly rely on the so-called ensemble averaging approach (Hurtwitz et al. 1990), that is, by assuming the source of pressure pulses (*i.e.* the heart) to be statistically independent from the source of noise, the averaging of N consecutive pressure pulses is expected to increase the signal-to-noise ratio by a factor of $\sqrt{N}$. However, the main drawback of such an approach is that while increasing N , one eliminates any information concerning short-term cardiovascular regulation: by instance, the Complior device requires averaging at least 10 heart cycles, blurring thus any respiration-related information contained into PWV. To overcome the smoothing effects of ensemble averaging some authors have explored the use of innovative pre-processing strategies based either on ICA denoising (Foo, 2008), neighbor PCA denoising (Xu et al., 2009), sub-band frequency decomposition (Okada et al., 1986), or ECG-assisted projection on local principal frequency components (Vetter et al., 2009). The main limitation is that, in order to preserve information on the original arrival time, any pre-processing operator applied to the raw pressure pulse signals must be designed to control any source of (phase) distortions.

According to this principle we have recently proposed a novel PAT estimation approach (Solà et al., 2009): instead of initially de-noising the raw pressure pulse, we design a robust PAT detector that minimizes the need of pre-processing, and thus works on a real beat-to-beat basis (Figure 14). The so-called parametric PAT estimation relies on the analysis of the whole wavefront of the pressure pulse, rather than searching for a punctual feature on it. The approach consists on initially fitting a parametric model to the pressure pulse wavefront and then on obtaining arrival time information from the parameters of the model. The correlation analysis performed on more than 200 hours of photo-plethysmographic data has shown that the new parametric approach highly correlates with the state-of-the-art characteristic points D1 ($r=0.99$) and TAN ($r=0.96$) when hyperbolic models were used. Concerning the robustness to noise, the parametric approach has been shown to improve the temporal resolution of PAT estimations by at least a five-fold factor (Solà et al., 2009).

5. Towards the ambulatory monitoring of Pulse Wave Velocity

The review of existing technologies in Table 1 highlights the current lack of approaches allowing the ambulatory non-obtrusive monitoring of aortic PWV. So far, the only commercial devices that might be considered as being ambulatory-compatible rely on the use of brachial cuffs, and hence require a pneumatic inflation each time a measurement is to be performed (Arteriograph and NovaCor). Other devices (Complior, SphygmoCor and PulsePen) are limited to clinical uses because they require the supervision of a well-trained operator. Moreover in ambulatory scenarios one must additionally consider the important role of hydrostatic pressure: while in supine position variations of hydrostatic pressure through the arterial tree are negligible, at standing or sitting positions the pressure gradient

from the iliac bifurcation to the aortic arch can reach values of almost 60 mmHg, *i.e.* about 75 mmHg per meter of altitude difference (Westerhof et al., 2005). Therefore, because PWV is affected by blood pressure to a high degree, changes in patient position would severely affect PWV measurements in setups such as Complior, SphygmoCor or PulsePen. Although a few methods for compensating for punctual hydrostatic pressure changes have been proposed in the past in the field of ambulatory blood pressure monitoring (Hiroyuki Ota & Kenji Taniguchi, 1998; McCombie et al., 2006), these cannot be applied to PWV for two reasons. Firstly, because contrary to blood pressure, PWV is not a point measurement but a distributed one, depicting propagation properties of a whole segment of the arterial tree. Secondly, because there is no one-to-one relationship between pressure and PWV changes, different unknown factors playing important roles as depicted by the Moens-Korteweg model in Equation 1.

In conclusion, there is a lack of methods that provide aortic PWV measurements automatically, continuously and in a non-obtrusive way, while remaining unaffected by changes in body position. A novel approach fulfilling these requirements is currently under investigation at CSEM, based on the continuous measurement of transit times of a pressure pulse when travelling from the Aortic Valve to the Sternum, the so-called av2sPTT. We describe now the benefits of introducing such an approach in ambulatory monitoring.

From a metrological perspective, the av2sPTT parameter is prone to be assessed continuously and non-obtrusively, a possible measurement setup being a textile harness mounted on the thorax. In particular we are working on a harness that integrates two dry ECG electrodes, a phono-cardiograph and a multi-channel photo-plethysmograph. While the joint analysis of the ECG and the phono-cardiogram provides information on the opening of the aortic valve (Alhstrom, 2008), the ECG-supported processing of the multi-channel photo-plethysmograph provides information on the arrival of the pressure pulse at the sternum (Vetter et al., 2009). Hence, PTT values are obtained through Equation 7. Note that for assessing av2sPTT none of the implemented sensing technologies requires the inflation of any cuff, and thus the approach remains fully non-obtrusive.

From a physiological perspective, the clinical relevance of the av2sPTT parameter is supported by a simple anatomical model of the arterial tree (Figure 15). In Table 2 we have detailed the arterial segments through which a pulse pressure propagates before reaching the sternum, together with the expected delays introduced at each segment. Typical PWV values have been obtained from (Nichols & O'Rourke, 2005) and (Acar et al., 1991). According to the model, the timing information contained into the av2sPTT parameter is expected to be 85% related to large vessels (aortic and carotid) and only 15% related to conduit arteries (internal thoracic artery). In other words, the arrival time of a pressure pulse at the sternum is mainly expected to be determined by the propagation through large vessels, and only minimally affected by secondary muscular arteries.

The simple anatomical model in Table 2 foresees that the av2sPTT parameter should be a good surrogate for arterial PTT, the incidence of central elastic arteries in the total measured PTT being of 85%. The statistical consistence of such an approach has not been validated yet,

and it is currently under investigation: the results of a validation study will be published in the future.

Fig. 15. 3D model of arterial segments involved in the measurement of av2sPTT: the pressure pulse propagates through the ascending aorta, the aortic arch, the brachiocephalic trunk and the internal thoracic artery. 3D model courtesy and copyright from Primal Pictures Ltd.

Segments of the arterial tree	Typical PWV	Typical length	Typical delay	Expected incidence
Ascending aorta and aortic arch	4.8 m/s	7 cm	15 ms	37%
Brachiocephalic trunk	5.1 m/s	10 cm	20 ms	49%
Internal thoracic artery (mammary)	8.5 m/s	5 cm	6 ms	15%
Overall av2sPTT	5.3 m/s	22 cm	41 ms	100%

Table 2. Propagation delays for the different arterial segments involved in the av2sPTT parameter.

In the following, we illustrate the potential use of the av2sPTT parameter as a method for the unsupervised continuous non-obtrusive monitoring of aortic PWV. The experiment consisted in sixty minutes of continuous recording on a healthy male subject on whom we simultaneously assessed the av2sPTT parameter and a carotid-to-radial PTT, obtained by a Complior device (Artech Medical, Paris, France). Beat-to-beat blood pressure was measured as well by a PortaPres device (FMS, Amsterdam, The Netherlands). During the experiment, three types of stresses were induced to the subject aiming at increasing his aortic stiffness,

and thus to decrease the measured PTT values. The stresses were, by chronological order: an arithmetic task stress, a sustained handgrip test and a cold stress test. Finally, the influence of muscular arteries on the av2sPTT parameter was tested by the oral administration of 250µg isosorbide-dinitrate (UCB-Pharma, Bulle, Switzerland). Isosorbide-dinitrate is expected to create a dilatation of conduit arteries, and thus to augment av2sPTT because of the propagation through the internal thoracic artery. As depicted in Figure 16, av2sPTT was successfully decreased by all three stresses, in accordance with the Complior measurements. As one would expect (Payne et al., 2006), the decrease of PTT coincided with a marked increase of blood pressure. After administering isosorbide-dinitrate, both Complior and av2sPTT detected an increase of PTT even if systolic blood pressure remained unchanged. Overall, a correlation coefficient of $r=0.89$ was encountered between the $n=9$ Complior measurements and the synchronous av2sPTT estimations. Although this experiment confirmed the hypothesis underlying the av2sPTT measurement principle, larger populations are required to validate the approach.

In conclusion, a novel method of assessing aortic PWV has been proposed that can be measured continuously (beat-by-beat), automatically (*i.e.* unsupervised) and non-obtrusively, thus paving the way towards the ambulatory measurement of PWV. Even more, since the measurement site is located at the sternum, perturbations due to hydrostatic pressure changes at different body positions have been minimized. Although high correlation with aortic PWV has been demonstrated from a theoretical perspective, statistically consistent data on the reliability of the approach is still missing.

6. Pulse Wave Velocity as a surrogate of mean arterial pressure

During the last 20 years a lot has been written on the use of PWV as a surrogate of arterial blood pressure (BP). The current lack of non-obtrusive devices for measuring BP has probably awakened the interest on this particular use of the PWV parameter. Yet, no commercial device has so far been released on the market. We present here the basics of this approach and we analyse the obstacles that researches are currently facing.

Historically, the dependency of PWV with respect to blood pressure was already observed in 1905 by Frank (Frank, 1905), although the underlying mechanism was not fully understood until half a century later (Hughes et al., 1979). We already described that PWV depends on arterial stiffness, and we introduced a mathematical model for this relationship: the Moens-Korteweg equation (Equation 1). Accordingly, the greater the stiffness of an artery, the faster a pressure pulse will propagate through it. Assume now that we increase the transmural pressure of the arterial wall, for instance by increasing systemic blood pressure. Because of the elastic properties of the wall, the artery will increase in diameter and decrease in thickness, while becoming stiffer. Additionally, at a certain point the recruitment of collagen fibers will start and will enhance the stiffness in a highly non-linear way (Nichols & O'Rourke, 2005). Hence, the stiffness of the arterial wall will depict a strong dependence to its transmural pressure. Putting now the two puzzle pieces together, we conclude that an increase of blood pressure will raise arterial stiffness and thus, increase PWV. Unfortunately, the relationship blood pressure - arterial stiffness - PWV is not unique, and several other parameters play important roles, as illustrated by Figure 17.

Fig. 16. Example of sixty minutes of non-invasive cardiovascular monitoring performed on a healthy male subject. The upper figure plots the evolution of systolic blood pressure as recorded by PortaPres. The lower figure shows continuous av2sPTT measurements (bold line) and simultaneous Carotid-to-Radial Complior measurements (black dots). Aortic Pulse Transit Time was altered during the experiment through different stress tasks.

Fig. 17. Under controlled situations, increasing the transmurial pressure (P) of an artery increases its stiffness (E), and consequently augments its PWV. Unfortunately other factors such as arterial muscular tonus and geometric modifications influence PWV as well.

A prominent study published in 2006 analyzed the dependency of PTT with respect to blood pressure for 12 subjects to whom vasoactive drugs were administered (Payne et al., 2006). The goal of the experiment was to quantify how the predictive capacities of PTT were degraded by drug alterations of arterial stiffness. All subjects and all drug conditions confounded, Payne still found that PTT was correlated with Diastolic Blood Pressure (DBP) with $r=0.64$. However, when “speaking in mmHg” this correlation appeared to be insufficient. Assuming that one would generate a calibration function for each subject in order to use PTT measurements to predict DBP, one would obtain a BP measurement device classed as “grade D” according to the British Society of Hypertension (BSH) (O’Brien et al., 2001), the cumulative percentage of readings falling within 5 mmHg, 10 mmHg and 15 mmHg of actual DBP values being 44%, 66% and 73% respectively. Consequently, following the BSH directives, a PTT-based blood pressure monitor should not be recommended.

Nevertheless, the dependency of PWV versus blood pressure has still been shown to be sufficiently dominant to be statistically exploitable under standard conditions (Poon & Zhang, 2005; Meigas et al., 2006; Foo et al., 2006). In this sense, and especially because of the expected clinical and commercial impact of a non-obtrusive BP monitor, promising research works and developments have been released in the past years. We now review the most relevant ones.

A good illustrative development of the state-of-the-art is that of (Chen et al., 2000): assuming that the factors that may interfere with the relationship blood pressure - PWV have slow time dynamics, *i.e.* slower than actual blood pressure changes, Chen investigated the use of a hybrid BP monitor: while a brachial oscillometric cuff regularly inflated to perform reference BP measurements, continuous PTT measurements were performed through a non-obtrusive ECG-fingertip approach. Chen then proposed to use the PTT series to interpolate the BP intermittent readings, beat-by-beat. Such a setup still demanded the use of brachial cuffs, but was a first step towards the non-obtrusive beat-to-beat BP monitoring. With this setup, Chen analyzed 42 hours of data on 20 subjects and obtained BSH cumulative percentages of 38.8%, 97.8% and 99.4%. Note that Chen did not account for PEP changes in his measurements, and that therefore, his results might have been improved.

Hence, PWV-based methods rely nowadays on the mapping of measured PTT values (in ms) to estimated BP values (in mmHg) through some initial or intermittent calibrations performed by oscillometric brachial cuffs (Stephoe et al., 1976). Several different calibration strategies and techniques have been described so far, mainly based on the use of neural networks (Barschdorff et al., 2000), linear regressions (Park et al., 2005; Kim et al., 2007; Poon & Zhang, 2008), model-based functions (Chen et al. 2003; Yan & Zhang, 2007) or hydrostatic-induced changes (McCombie et al., 2007).

Fig. 18. When estimating BP from PWV measurements, we suggest accounting for changes of arterial geometry and stiffness in order to reduce the frequency of calibrations (Sola et al., 2008). The proposed model requires introducing Cardiac Output (CO) as an additional parameter.

In 2008 we proposed a novel approach aiming at reducing the frequency of these calibrations (Sola et al., 2008). Assuming that the major source of calibration modifications are the changes of diameter and stiffness of conduit arteries (Madero, et al., 2003), we proposed a technique that compensates for these changes without necessitating additional calibrations. In order to do so, the technique requires the continuous measurement of both PWV and Cardiac Output (CO), and assumes a Moens-Korteweg-like model of pulse wave propagation together with a Poiseuille's model of fluidic resistance (Figure 18). The most likely BP value is then computed by solving:

$$BP = \operatorname{argmin}_{BP} d(PWV_m, PWV_p(CO, BP)) \quad (8)$$

that is, by finding the BP value that makes the measured PWV_m to be as close as possible to the PWV_p predicted by the model (given BP and CO). A distance metric (d) between two series of PWV measurements needs to be previously defined. Although the approach has been shown to reliably provide BP measurements even during induced vasoconstriction (Sola et al., 2008), its consistency in large population studies remains to be demonstrated.

In conclusion, the use of PWV as a surrogate for blood pressure appears to be justified under controlled conditions, that is, when the effects of vasomotion can be neglected. Unfortunately, the ambulatory monitoring of PWV currently passes through the detection of pressure pulses at distal sites such as the radial artery, or the fingertip. Because such measurement setups involve the propagation of pressure pulses through conduit arteries, the effects of vasomotion are, at least, unfavorable. The ideal solution would pass through the monitoring of PWV of proximal arteries only. However, as we demonstrated, there is currently no available technology capable of providing ambulatory very-proximal (*i.e.* aortic) PWV measures. The development of novel technologies as the herein presented aortic valve-to-sternum PTT (Figure 15) might provide great chances for the implantation of PWV into the field of portable BP monitoring.

7. Conclusion and outlook

The continuous ambulatory monitoring of cardiac and vascular functions is a clear requirement for the deployment of the new generation of healthcare structures. Unfortunately, there is so far no technology available for the non-obtrusive measurement of vascular-related health status. The recent spread of automatic devices for the measurement of aortic PWV has promoted numerous research studies on the clinical uses of PWV. Widely accepted as a surrogate for arterial stiffness, PWV appears to be an independent global marker of vascular damage risk. In this chapter we reviewed the major clinical findings and we analyzed the developed technologies. The results suggest that PWV could be promoted as a candidate to fill in the gap of vascular-related health indexes. However, the state-of-the-art in PWV measurement is currently inappropriate for the ambulatory monitoring of arterial stiffness. Therefore, we propose a novel aortic PWV measurement approach that can be integrated into a chest belt and that fulfils the requirements of continuity (beat-by-beat) and non-obtrusiveness. The strategy relies on measuring the transit time of pressure pulses from the opening of the aortic valve to their arrival at the sternum capillaries. In contrast with existing techniques, the new approach does not require the presence of skilled operators, and is expected to minimize the influences of hydrostatic pressure interferences due to postural changes.

Independently of the actual measurement approach, aortic PWV might in the near future play a central role in the diagnosis and follow up of cardiovascular diseases. It is in this context that the coordinated work of biomedical engineers and clinicians will determine to which extent PWV will be able to penetrate into clinical and medical practice.

8. References

- Abassade, P. & Baudouy, Y. (2002). Relationship between arterial distensibility and left ventricular function in the timing of Korotkoff sounds (QkD internal). An ambulatory pressure monitoring and echocardiographic study. *American Journal of Hypertension*, 15, 4, (2002) 67A
- Acar, C.; Jebara, V. A.; Portoghèse, M.; Fontaliran, F.; Dervanian, P.; Chachques, J. C., Meininger, V. & Carpentier, A. (1991). Comparative anatomy and histology of the radial artery and the internal thoracic artery. *Surg Radiol Anat*, 13, (1991) 283-288
- Ahlström, C. (2008). *Nonlinear Phonocardiographic Signal Processing*, Linköpings Universitet, 978-91-7393-947-8, Linköping, Sweden
- Airaksinen, K.E.; Salmela, P.I.; Linnaluoto, M.K.; Ikaheimo, M.J.; Ahola, K & Ryhanen, L.J. (1993) Diminished arterial elasticity in diabetes: association with fluorescent advanced glycosylation end products in collagen. *Cardiovasc Res*, 27 (1993) 942-945
- Allen, J. (2007). Photoplethysmography and its application in clinical physiological measurement. *Physiol Meas*, 28, (2007) R1-R39
- Asmar, R. (1999). *Arterial Stiffness and Pulse Wave Velocity. Clinical Applications*, ELSEVIER, 2-84299-148-6
- Asmar, R.; Beneros, A.; Topouchian, J.; Laurent, P.; Pannier, B. Brisac, A.; Target, R. & Levy, B. (1995). Assessment of Arterial Distensibility by Automatic Pulse Wave Velocity Measurement. *Hypertension*, 26, 3, (1995) 485-490

- Avolio, A.P.; Deng, F. Q.; Li, W. Q. et al. (1985) Effects of aging on arterial distensibility in populations with high and low prevalence of hypertension: comparison between urban and rural communities in China. *Circulation*, 71 (1985) 202-210
- Avolio, AP; Chen, SG; Wang, R.P.; Zhang, C.L.; Li, M.F. & O'Rourke, M.F. (1983) Effects of aging on changing arterial compliance and left ventricular load in a northern Chinese urban community. *Circulation*, 68, (1983) 50-58
- Baguet, J.; Kingwell, B. A.; Dart, A. L; Shaw, J.; et al. (2003). Analysis of the regional pulse wave velocity by Doppler: methodology and reproducibility. *J Human Hypertension*, 17, (2003) 407-412
- Banet, M. (2009) ViSi: Body-Worn System for Monitoring Continuous, Non-Invasive Blood Pressure cNIBP. *Proceedings of DARPA Workshop on Continuous, Non-Invasive Monitoring of Blood Pressure (CNIMBP)*, 2009, Coronado
- Barschdorff, D.; Erig, M. & Trowitzsch E. (2000). Noninvasive continuous blood pressure determination, *Proceedings XVI IMEKO World Congress*, 2000, Wien
- Baulmann, J.; Schillings, U.; Rickert, S.; Uen, S.; Düsing, R.; Cziraki, R.; Illyes, M. & Mengden, T. (2008). A new oscillometric method for assessment of arterial stiffness: comparison with tonometric and piezo-electronic methods. *Journal of Hypertension*, 26, 3, (2008) 523-528
- Benetos, A.; Adamopoulos, C.; Bureau, J. M. et al. (2002) Determinants of accelerated progression of arterial stiffness in normotensive subjects and in treated hypertensive subjects over a 6-year period. *Circulation*, 105 (2002) 1202-1207
- Berntson, G. G.; Lozano, D. L., Chen Y. J. & Cacioppo J. T. (2004). Where to Q in PEP. *Psychophysiology*, 41, 2, (2004) 333-337
- Blacher, J.; Guerin, A.P.; Pannier, B.; Marchais, S.J.; Safar, M.E. & London, G.M. (1999) Impact of aortic stiffness on survival in end-stage renal disease. *Circulation* 99 (1999) 2434-2439
- Boese, J. M.; Bock, M.; Bahner, M. L.; Albers, J. & Schad, L. R. (2000) In vivo Validation of Aortic Compliance Estimation by MR Pulse Wave Velocity Measurement, *Proc. Intl. Soc. Mag. Reson. Med.*, 8 (2000) 357
- Bouthier, J.D.; De Luca, N.; Safar, M.E. & Simon, A.C. (1985) Cardiac hypertrophy and arterial distensibility in essential hypertension. *Am Heart J*, 109 (1985) 1345-52.
- Boutouyrie, P.; Briet, M.; Collin, C.; Vermeersch, S. & Pannier, B. (2009). Assessment of pulse wave velocity. *Artery Research*, 3, (2009) 3-8
- Boutouyrie, P.; Lacolley, P.; Girerd, X.; Beck, L.; Safar, M. & Laurent S. (1994) Sympathetic activation decreases medium-sized arterial compliance in humans. *Am J Physiol*, 267 (1994) H1368-H1376
- Butlin, M.; Hickson, S.; Graves, M. J.; McEniery, C. M; et al. (2008). Determining pulse wave velocity using MRI: a comparison and repeatability of results using seven transit time algorithms. *Artery Research*, 2, 3, (2008) 99
- Cameron, J.D.; Jennings, G.L. & Dart, A.M. (1995) The relationship between arterial compliance, age, blood pressure and serum lipid levels. *J Hypertens*, 13 (1995) 1718-1723
- Chen, C. H.; Nevo, E.; Fetcs, B.; Pak, P. H.; Yin, F. C.; Maughan, W. L. & Kass, D. A.(1997). Estimation of central aortic pressure waveform by mathematical transformation of radial tonometry pressure. Validation of generalized transfer function. *Circulation*, 95, 7, (1997) 1827-1836

- Chen, W.; Kobayashi, T.; Ichikawa, S.; Takeuchi, Y. & Togawa, T. (2000). Continuous estimation of systolic blood pressure using the pulse arrival time and intermittent calibration. *Medical & Biological Engineering & Computing*, 38, (2000) 569-574
- Chen, Y.; Li, L.; Hershler, C. & Dill, R. P. (2003). Continuous non-invasive blood pressure monitoring method and apparatus. *US Patent 6,599,251*, July 2003
- Chiu, Y. C.; Arand, P. W.; Shroff, S. G.; Feldman, T. Et al. (1991). Determination of pulse wave velocities with computerized algorithms. *Am Heart J*, 121, 5, (1991) 1460-1470
- Constans, J.; Germain, C.; Gossec, P. Taillard, J. et al. (2007). Arterial stiffness predicts severe progression in systemic sclerosis: the ERAMS study. *J Hypertension*, 25, (2007) 1900-1906
- Creager, M.A.; Luscher, T.F.; Cosentino, F. & Beckman, J.A. (2003) Diabetes and vascular disease: pathophysiology, clinical consequences, and medical therapy: Part I. *Circulation*, 108 (2003) 1527-32
- Cruickshank, K.; Riste, L.; Anderson, S.G.; Wright, J.S.; Dunn, G. & Gosling, R.G. (2002) Aortic pulse-wave velocity and its relationship to mortality in diabetes and glucose intolerance: an integrated index of vascular function? *Circulation*, 106 (2002) 2085-90
- Dart, A.M. & Kingwell, B. A. (2001) Pulse pressure--a review of mechanisms and clinical relevance. *J Am Coll Cardiol*, 37 (2001) 975-984
- Dart, A.M.; Gatzka, C.D.; Cameron, J.D., et al. (2004) Large artery stiffness is not related to plasma cholesterol in older subjects with hypertension. *Arterioscler Thromb Vasc Bio*, 24 (2004) 962-968
- DeMarzo, A. P. & Lang, R. M. (1996). A new algorithm for improved detection of aortic valve opening by impedance cardiography. *Computers in Cardiology*, 8, 11, (1996) 373-376
- Faber M. & Oller-Hou G. (1952). The human aorta. V. Collagen and elastin in the normal and hypertensive aorta. *Acta Pathol Microbiol Scand*, 31, (1952) 377-82.
- Fesler, P.; Safar, M.E.; du Cailar, G.; Ribstein, J. & Mimran, A. (2007) Pulse pressure is an independent determinant of renal function decline during treatment of essential hypertension. *J Hypertens*, 25 (2007) 1915-1920
- Foo, J. Y. A. (2008). Use of Independent Component Analysis to Reduce Motion Artifact in Pulse Transit Time Measurement. *Signal Processing Letters*, 15, (2008) 124-126
- Foo, J. Y. A.; Lim, C. S. & Wang, P. (2006). Evaluation of blood pressure changes using vascular transit time. *Physiol. Meas.*, 17, (2006) 685-694
- Franchi, D.; Bedini, R.; Manfredini, F.; Berti, S.; Palagi, G.; Ghione, S. & Ripoly, A. (1996). Blood pressure evaluation based on arterial pulse wave velocity. *Computers in Cardiology*, 8, 11, (1996) 397-400
- Frank, O. (1905) Der Puls in den arterien, *Zeitschrift für Biologie*, 45 (1905) 441-553
- Fung, P.; Dumont, G.; Ries, C.; Mott, C. & Ansermino, M. (2004). Continuous Noninvasive Blood Pressure Measurement by Pulse Transit Time, *Proceedings of the 26th Annual International Conference of the IEEE EMBS*, pp. 738-741, 0-7803-8439-3, San Francisco, September 2004, IEEE
- Gallagher, D.; Adj, A. & O'Rourke, M. F. (2004). Validation of the transfer function technique for generating central from peripheral upper limb pressure waveform. *Am J Hypertens*, 17, 11, (2004) 1059-1067

- Gang, G.; Mark, P.; Cockshott, P., Foster, J., et al. (2004). Measurement of Pulse Wave Velocity using Magnetic Resonance Imaging, *Proceedings of the 26th Annual International Conference of the IEEE EMBS*, pp. 3684-3687, 0-7803-8439-3, San Francisco, September 2004, IEEE
- Giannattasio, C.; Capra, A.; Facchetti, R. et al. (2007) Relationship between arterial distensibility and coronary atherosclerosis in angina patients. *J Hypertens*, 25 (2007) 593-598.
- Giannattasio, C.; Failla, M.; Piperno, A. et al. (1999) Early impairment of large artery structure and function in type I diabetes mellitus. *Diabetologia*, 42 (1999) 987-994
- Girerd, X.; Chanudet, X.; Larroque, P.; Clement, R.; Laloux, B. & Safar, M. (1989) Early arterial modifications in young patients with borderline hypertension. *J Hypertens Suppl*, 7 (1989) S45-S47.
- Giri, S.S.; Ding, Y.; Nishikima, Y.; Pedraza-Toscano, et al. (2007). Automated and Accurate Measurement of Aortic Pulse Wave Velocity Using Magnetic Resonance Imaging. *Computuvers in Cardiology*, 34, (2007) 661-664
- Glen, S.K.; Elliott, H.L.; Curzio, J.L.; Lees, K.R. & Reid J.L. (1996) White-coat hypertension as a cause of cardiovascular dysfunction. *Lancet*, 348 (1996) 654-657
- Guerin, A. P.; Blacher, J.; Pannier, B.; Marchais, S. J.; Safar, M. E. & London, G. (2001). Impact of Aortic Stiffness Attenuation on Survival of Patients in End-Stage Renal Failure. *Circulation*, 103, (2001) 987-992
- Hayes, M. J. & Smith, P. R. (1998). Artifact reduction in photoplethysmography. *Applied Optics*, 37, 31, (1998) 7437-7446
- Henry, R.M.; Kostense, P.J.; Spijkerman, A.M. et al. (2003) Arterial stiffness increases with deteriorating glucose tolerance status: the Hoorn Study. *Circulation* 107 (2003) 2089-95
- Henskens, L.H.; Kroon, A.A.; van Oostenbrugge, R.J., et al. (2008) Increased aortic pulse wave velocity is associated with silent cerebral small-vessel disease in hypertensive patients. *Hypertension*, 52 (2008) 1120-1126
- Hirai, T.; Sasayama, S.; Kawasaki, T. & Yagi, S. (1989) Stiffness of systemic arteries in patients with myocardial infarction. A noninvasive method to predict severity of coronary atherosclerosis. *Circulation*, 80 (1989) 78-86
- Hirata, K.; Kawakami, M. & O'Rourke M. F. (2006) Pulse Wave Analysis and Pulse Wave Velocity – A review of Blood Pressure Interpretation 100 Years After Korotkov, *Circ J*, 70 (2006) 1231-1239
- Hiroiyuki Ota, H.; Kenji Taniguchi, K.; (1998). Electronic blood pressure meter with posture detector. *US Patent 5,778,879*, July 1998
- Hughes, D. J.; Babbs, C. F.; Geddes, L. A. & Bourland, J. D. (1979) Measurement of Young's Modulus of Elasticity of the Canine Aorta with Ultrasound, *Ultrasonic Imaging*, 1 (1979) 356-367
- Hurtwitz, B. E.; Shyu, L. Y.; Reddy, S P.; Shneiderman, N. & Nagel, J. H (1990). Coherent ensemble averaging techniques for impedance cardiography, *Proceedings of Third Annual IEEE Symposium on Computer-Based medical Systems*, 2009
- Jatoi, N.A.; Jerrard-Dunne, P.; Feely, J. & Mahmud, A. (2007) Impact of smoking and smoking cessation on arterial stiffness and aortic wave reflection in hypertension. *Hypertension*, 49 (2007) 981-5

- Jiang, B.; Liu, B.; McNeill, K. L. & Chowienczyk, P. J. (2008) Measurement of Pulse Wave Velocity Using Pulse Wave Doppler Ultrasound: Comparison with Arterial Tonometry, *Ultrasound in Med. & Biol.*, 34, 3 (2008) 509-512
- Karamanoglu, M.; Gallegher, D. E.; Avolio, A. P. & O'Rourke, M. F. (1995). Pressure wave propagation in a multibranched model of the human upper limb. *Am J Physiol*, 269, 4, (1995) H1363-1369
- Kass, D.A.; Shapiro, E.P.; Kawaguchi, M. et al. (2002) Improved arterial compliance by a novel advanced glycation end-product crosslink breaker. *Circulation*, 104 (2002) 1464-1470
- Kim, J. S.; Kim, K. K.; Lim Y. G. & Park K. S. (2007) Two unconstrained methods for blood pressure monitoring in a ubiquitous home healthcare, *Proceedings of the Fifth International Conference Biomedical Engineering*, pp. 293-296, 978-0-88986-648-5, Innsbruck, February 2007
- Kim, J.; Park, J.; Kim, K.; Chee, Y.; Lim, Y. & Park, K. (2006). Development of a Noninvasive Blood Pressure System for Somputer Users. *Telemedicine and e-Health*, 13, 1, (2006) 57-64
- King, E.; Cobbin, D.; Walsh, S. & Ryan, D. (2002). The Reliable Measurement of Radial Pulse Characteristics. *Acupuncture in Medicine*, 20, 4, (2002) 150-159
- Lababidi, Z.; Ehmke, D. A.; Durnin, R. E.; Leaverton, P. E. & Lauer, R. M. (1970). The First Derivative Thoracic Impedance Cardiogram. *Circulation*, 41, (1970) 651-658
- Laffon, E.; Marthan, R.; Mantaudon, M.; Latrabe, V.; Laurent F. & Ducassou, D. (2005) Feasibility of aortic pulse pressure and pressure wave velocity MRI measurement in young adults, *J. Magn. Reson. Imaging*, 21, 1 (2005) 53-58
- Latham, R.D.; Westerhof, N.; Sipkema, P. et al. (1985). Regional wave travel and reflections along the human aorta: a study with six simultaneous micromonometric pressures. *Circulation*, 72, 3, (1985) 1257-1269
- Laurent, S. & Cockcroft, J. (2008). *Central aortic blood pressure*, Elsevier, 978-2-84299-943-8
- Laurent, S.; Boutouyrie, P.; Asmar, R. et al. (2001) Aortic stiffness is an independent predictor of all-cause and cardiovascular mortality in hypertensive patients. *Hypertension*, 37 (2001) 1236-41
- Laurent, S.; Katsahian, S.; Fassot, C. et al. (2003) Aortic stiffness is an independent predictor of fatal stroke in essential hypertension. *Stroke*, 34 (2003) 1203-1206
- Li, Q. & Belz, G. G. (1993). Systolic time intervals in clinical pharmacology. *Eur J Clin Pharmacol*, 44, (1993) 415-421
- Liang, Y.L.; Shiel, L.M.; Teede, H. et al. (2001) Effects of Blood Pressure, Smoking, and Their Interaction on Carotid Artery Structure and Function. *Hypertension*, 37 (2001) 6-11
- London, G.M.; Guerin, A.P.; Marchais, S.J. et al. (1996) Cardiac and arterial interactions in end-stage renal disease. *Kidney Int*, 50 (1996) 600-8
- London, G.M.; Guerin, A.P.; Pannier, B.; Marchais, S.J. & Stimpel, M. (1995) Influence of sex on arterial hemodynamics and blood pressure. Role of body height. *Hypertension*, 26 (1995) 514-519
- Lorell, B.H. & Carabello, B.A. (2000) Left ventricular hypertrophy: pathogenesis, detection, and prognosis. *Circulation*, 102 (2000) 470-9
- Lotz, J.; Meier, C.; Leppert, A. & Galanski, M. (2002). Cardiovascular flow measurement with phase-contrast MR imaging: basic facts and implementation. *Radiographics*, 22, 3, (2002) 651-671

- Madero, R.; Lawrence, H. & Sai, K. (2003). Continuous, non-invasive technique for measuring blood pressure using impedance plethysmography. *European Patent Office, EP 1 344 4891 A1*, February 2003
- Mancia, G.; De Backer, G.; Dominiczak, A. et al. (2007) Guidelines for the Management of Arterial Hypertension: The Task Force for the Management of Arterial Hypertension of the European Society of Hypertension (ESH) and of the European Society of Cardiology (ESC). *J Hypertens*, 25 (2007) 1105-1187
- Mathews, J. H & Fink, K. D. (1998). *Numerical Methods Using MATLAB*, Prentice Hall, 978-0132700429
- Mattace-Raso, F.U.; van der Cammen, T.J.; Hofman, A., et al. (2006) Arterial stiffness and risk of coronary heart disease and stroke: the Rotterdam Study. *Circulation*, 113 (2006) 657-663
- McCombie, D. B.; Reisner, A. T. & Asada, H. H. (2006). Adaptive blood pressure estimation from wearable PPG sensors using peripheral pulse wave velocity measurement and multi-channel blind identification of local artery dynamics, *Proceedings of the 28th Annual International Conference of the IEEE EMBS*, pp. 3521-3524, 1-4244-0033-3, New York City, September 2006, IEEE
- McCombie, D. B.; Shaltis, P. A.; Reisner, A. T. & Asada, H. H. (2007). Adaptive hydrostatic blood pressure calibration: Development of a wearable, autonomous pulse wave velocity blood pressure monitor, *Proceedings of the 29th Annual International Conference of the IEEE EMBS*, pp. 370-373, 1-4244-0788-5, Lyon, August 2007, IEEE
- Meaume, S.; Benetos, A.; Henry, O.F.; Rudnichi, A. & Safar, M.E. (2001) Aortic pulse wave velocity predicts cardiovascular mortality in subjects >70 years of age. *Arterioscler Thromb Vasc Biol*, 21 (2001) 2046-2050
- Meigas, K.; Lass, J.; Karai, D.; Kattai, R. & Kaik, J. (2006). Pulse Wave Velocity in Continuous Blood Pressure Measurements, *Proceedings of the Worl's Congress on Medical Physics and Biomedical Engineering*, pp. 626-629, 978-3-540-36839-7, Seoul, September 2006, Springer
- Meinders, J. M.; Kornet, L.; Brands, P.J. & Hoeks, A. P. (2001) Assessment of local pulse wave velocity in arteries using 2D distension waveforms, *Ultrason Imaging*, 23, 4 (2001) 199-215
- Millasseau, S. C.; Ritter, J. M.; Takazawa, K. & Chowienczyk, P. J. (2006). Contour analysis of the photoplethysmographic pulse measured at the finger. *Journal of Hypertension*, 24, 8, (2006) 1149-1456
- Millasseau, S. C.; Stewart, A. D.; Patel, S. J.; Redwood, S. R. & Chowienczyk, P. J. (2005). Evaluation of carotid-femoral pulse wave velocity: influence of timing algorithm and heart rate. *Hypertension*, 45, 2, (2005) 222-226
- Mitchell, G. F. (2009) Arterial stiffness and wave reflection: Biomarkers of cardiovascular risk, *Artery Research*, 2 (2009) 56-64
- Mohiaddin, R. H.; Longmore, D. B. (1989). MRI studies of atherosclerotic vascular disease: structural evaluation and physiological measurements. *British Medical Bulletin*, 45, (1989) 968-990
- Muehlsteff, J.; Aubert, X. L. & Schuett, M. (2006). Cuffless Estimation of Systolic Blood Pressure for Short Effort Bicycle Tests: The Prominent Role of the Pre-Ejection Period, *Proceedings of the 28th Annual International Conference of the IEEE EMBS*, pp. 5088-5092, 1-4244-0033-3, New York City, September 2006, IEEE

- Naschitz, J. E.; Bezobchuk, S.; Mussafia-Priselac, R.; Sundick, S.; et al. (2004). Pulse transit time by R-wave-gated infrared photoplethysmography: review of the literature and personal experience. *J Clin Monit Comput*, 18, 5-6, (2004) 333-342
- Nichols W.W. (2005) Clinical measurement of arterial stiffness obtained from noninvasive pressure waveforms. *Am J Hypertens*, 18 (2005) 3S-10S
- Nichols, W. W. & O'Rourke, M. F (2005). *McDonald's blood flow in arteries*, Hodder Arnold, 0 340 80941 8
- Nichols, W. W. (2009). Aortic Pulse Wave Velocity, Reflection Site Distance, and Augmentation Index. *Hypertension*, 53, 1, (2009) e9
- Nichols, W. W.; Denardo, S. J.; Wilkinson, I. B.; McEniery, C. M.; Cockcroft, J. & O'Rourke, M. F. (2008) Effects of Arterial Stiffness, Pulse Wave Velocity, and Wave Reflections on the Central Aortic Pressure Waveform, *J Clin Hypertens*, 10 (2008) 295-303
- Nilsson, P. M.; Boutouyrie, P. & Laurent, S. (2009) Vascular Aging : A Tale of EVA and ADAM in Cardiovascular Risk Assessment and Prevention, *Hypertension*, 54 (2009) 3-10
- Nitzan, M.; Khanokh, B. & Slovik, Y. (2002). The difference in pulse transit time to the toe and finger measured by photoplethysmography. *Physiol Meas*, 23, (2002) 85-93
- O'Brien, E.; Waeber, B.; Parati, G.; Staessen, J. & Myers M. G. (2001) Blood pressure measuring devices: recommendations of the European Society of Hypertension. *BMJ*, 322 (2001) 531-536
- O'Rourke M. F. (2009). Time domain analysis of the arterial pulse in clinical medicine. *Med Biol Eng Comp*, 47, 2, (2009) 119-129
- Okada, M.; Kimura, S. & Okada, M. (1986) Estimation of arterial pulse wave velocities in the frequency domain: method and clinical considerations, *Medical & Biological Engineering & Computing*, 24, 3 (1986) 255-260
- O'Rourke, M.F. & Safar, M.E. (2005) Relationship between aortic stiffening and microvascular disease in brain and kidney: cause and logic of therapy. *Hypertension*, 46 (2005) 200-204
- Park E. K.; Cho B. H.; Park, S. H.; Lee, J. Y.; Lee, J. S.; Kim, I. Y. & Kim S. I (2005). Continuous measurement of systolic blood pressure using the PTT and other parameters, *Proceedings of the 27th Annual International Conference of the IEEE EMBS*, pp. 3555-3558, 0-7803-8740-6, Shanghai, September 2005, IEEE
- Payne, R. A.; Symeonides, C. N.; Webb, D. J. & Maxwell, S. R. (2006). Pulse transit time measured from the ECG: an unreliable marker of beat-to-beat blood pressure. *J Appl Physiol*, 100, 1, (2006) 136-141
- Poon, C. C. Y. & Zhang Y. T. (2005). Cuff-less and Noninvasive Measurement of Arterial Blood Pressure by Pulse Transit Time, *Proceedings of the 27th Annual International Conference of the IEEE EMBS*, pp. 5877-5880, 0-7803-8740-6, Shanghai, September 2005, IEEE
- Poon, C. Y. & Zhang, Y. T. (2008). The Beat-to-Beat Relationship between Pulse Transit time and Systolic Blood Pressure, *Proceedings of the 5th International Conference on Information Technology and Application in Biomedicine*, pp. 342-343, 978-1-4244-2255-5, Shenzhen, May 2008, IEEE
- Rajzer, M.; Wojciechowska, W.; Kloczek, M.; Palka, I.; Brzozowska, B. & Kawecka-Jaszcz, K. (2008). Comparison of aortic pulse wave velocity measured by three techniques: Complior, SphygmoCor and Arteriograph. *Journal of Hypertension*, 26, 10, (2008) 2001-2007

- Ravikumar, R.; Deepa, R.; Shanthirani, C. & Mohan, V. (2002) Comparison of carotid intima-media thickness, arterial stiffness, and brachial artery flow mediated dilatation in diabetic and nondiabetic subjects (The Chennai Urban Population Study [CUPS-9]). *Am J Cardiol*, 90 (2002) 702-7
- Schram, M.T.; Henry, R.M.; van Dijk, R.A. et al. (2004) Increased central artery stiffness in impaired glucose metabolism and type 2 diabetes: the Hoorn Study. *Hypertension*, 43 (2004) 176-81
- Schwartz, D. J. (2005). The pulse transit time arousal index in obstructive sleep apnea before and CPAP. *Sleep Medicine*, 6, (2005) 199-2003
- Segers, P.; Kips, J.; Trachet, B.; Swillens, A.; Vermeersch, S.; Mahieu, D.; Rietzschel, E.; Buyzere, M. D. & Bortel, L. V. (2009) Limitations and pitfalls of non-invasive measurement of arterial pressure wave reflections and pulse wave velocity, *Artery Research*, 3, 2 (2009) 79-88
- Sharwood-Smith, G.; Bruce, J. & Drummond, G. (2005). Assessment of pulse transit time to indicate cardiovascular changes during obstetric spinal anaesthesia. *Br J Anaesthesia*, 96, 1, (2006) 100-105
- Shinohara, K.; Shoji, T.; Tsujimoto, Y. et al. (1999) Arterial stiffness in predialysis patients with uremia. *Kidney Int*, 65 (1999) 936-43
- Smith, R. P.; Argod, J.; Pépin, J. & Lévy, A. (1999). Pulse transit time: an appraisal of potential clinical applications. *Thorax*, 54, (1999) 452-457
- Solà, J.; Chételat, O. & Luprano, J. (2008) Continuous Monitoring of Coordinated Cardiovascular Responses, *Proceedings of the 30th Annual International Conference of the IEEE EMBS*, pp. 1423-1426, 978-1-4244-1814-5, Vancouver, August 2008, IEEE
- Solà, J.; Vetter, R.; Renevey P.; Chételat, O.; Sartori, C. & Rimoldi, S. F. (2009) Parametric estimation of Pulse Arrival Time: a robust approach to Pulse Wave Velocity, *Physiol. Meas.*, 30 (2009) 603-615
- Sonesson, B.; Hansen, F.; Stale, H. & Lanne, T. (1993) Compliance and diameter in the human abdominal aorta--the influence of age and sex. *Eur J Vasc Surg*, 7 (1993) 690-697.
- Stephoe, A.; Smulyan, H. & Gribbin B. (1976) Pulse Wave Velocity and Blood Pressure Change: Calibration and Applications. *Psychophysiology*, 13, 5 (1976) 488-493.
- Sutton-Tyrrell, K.; Najjar, S.S.; Boudreau, R.M. et al. (2005) Elevated aortic pulse wave velocity, a marker of arterial stiffness, predicts cardiovascular events in well-functioning older adults. *Circulation*, 111 (2005) 3384-90
- Toprak, A.; Reddy, J.; Chen, W.; Srinivasan, S. & Berenson, G. (2009) Relation of pulse pressure and arterial stiffness to concentric left ventricular hypertrophy in young men (from the Bogalusa Heart Study). *Am J Cardiol*, 103 (2009) 978-984.
- van Ittersum, F.J.; Schram, M.T.; van der Heijden-Spek, J.J. et al. (2004) Autonomic nervous function, arterial stiffness and blood pressure in patients with Type I diabetes mellitus and normal urinary albumin excretion. *J Hum Hypertens*, 18 (2004) 761-8
- van Popele, N.M.; Grobbee, D.E.; Bots, M.L. et al. (2001) Association between arterial stiffness and atherosclerosis: the Rotterdam Study. *Stroke*, 32 (2001) 454-460
- Vetter, R.; Rossini, L.; Ridolfi, A.; Solà, J.; Chételat, O.; Correvon, M. & Krauss, J. (2009). Frequency domain SpO₂ estimation based on multichannel photoplethysmographic measurements at the sternum, to appear in *Proceedings of Medical Physics and Biomedical Engineering World Congress*, Munich, September 2009
- Webster, J. G. (1997). *Design of pulse oximeters*, CRC Press, 978-0-750-30467-2

- Westerhof, N.; Stergiopulos, N. & Noble, M. (2005). *Snapshots of Hemodynamics*, Springer, 978-0-387-23345-1
- Willum-Hansen, T.; Staessen, J.A.; Torp-Pedersen, C. et al. (2006) Prognostic value of aortic pulse wave velocity as index of arterial stiffness in the general population. *Circulation*, 113 (2006) 664-670
- Wolffenbuttel B.H.; Boulanger, C.M.; Crijns, F.R., et al. (1998) Breakers of advanced glycation end products restore large artery properties in experimental diabetes. *Proc Natl Acad Sci USA*, 95 (1998) 4630-4634
- Xu, P.; Bergsneider, M. & Hu, X. (2009) Pulse onset detection using neighbor pulse-based signal enhancement, *Medical Engineering & Physics*, 31 (2009) 337-345
- Yan, Y. S. & Zhang, Y. T. (2007). A Novel Calibration Method for Noninvasive Blood Pressure Measurement Using Pulse Transit Time, *Proceedings of the 4th IEEE-EMBS International Summer School and Symposium on Medical Devices and Biosensors*, Cambridge, August 2007

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