

Diving Into Research of Biomedical Engineering in Scuba Diving

Tobias Cibis, *Student Member, IEEE*, Alistair McEwan, *Member, IEEE*, Arne Sieber, Bjoern Eskofier, *Member, IEEE*, John Lippmann, Karl Friedl, *Member, IEEE*, and Michael Bennett

Abstract—The physiologic response of the human body to different environments is a complex phenomenon to ensure survival. Immersion and compressed gas diving, together trigger a set of responses. Monitoring those responses in real-time may increase our understanding of these and help to develop safety procedures and equipment. This review outlines diving physiology and diseases and identifies physiological parameters worthy of monitoring. Subsequently, we have investigated technological approaches matched to those in order to evaluate their capability for underwater application. We focused on wearable biomedical monitoring technologies, or those which could be transformed to wearables. We have also reviewed current safety devices, including dive computers and their underlying decompression models and algorithms. The review outlines the necessity for biomedical monitoring in scuba diving and should encourage research and development of new methods to increase diving safety.

Index Terms—Decompression Sickness, Decompression Algorithms, Diving Medicine, Dive Computer, Medical Engineering, Physiological Monitoring, Scuba Diving, Wearables.

I. INTRODUCTION

Diving is practiced in many settings including recreational, commercial and military diving [1]. All variations carry physical risks related to the underwater environment and can result in negative consequences manifesting in different diving-related diseases, ranging from immediate abnormalities such as decompression illness [2], [3] to long-term or late effects such as neurological impairment [4]– [6]. Despite the rarity of lethal injuries, the risks associated with diving suggest the need to develop a device capable of real-time monitoring and analysis of bio-signals while diving and during diving injury assessment [7]. The development of underwater monitoring technology may result in safety equipment for divers that could be used for both real-time monitoring and data acquisition for later analysis, as well as increased underwater working time for commercial and military divers [1]. Evaluated data may improve the fundamental understanding of the pathophysiology in diving because physiological changes are measured as

they occur.

In diving medicine, the physical effects of immersion and pressure elicit physiological responses. These responses have been well-described and are closely associated with the development of a number of diving-related diseases [8]. In relation to the preeminent physiological parameters, matching monitoring approaches are investigated. On the basis that invasive methods of monitoring are impractical for scuba diving, we assessed each technology for the ability to be adapted for the use as a wearable, including electrocardiography, blood pressure and volume shift technologies.

Common dive technologies currently in use include dive computers that calculate decompression obligations in real-time using various decompression tables, models and algorithms [9]. Diving computers are the state of the art tool for divers to monitor their depth and time in order to reduce the chance of decompression injuries [10]. The core function of any dive computer is the operating decompression models / algorithms used. These models calculate decompression schedules to step divers safely to the surface while avoiding critical supersaturation of various tissues with inert gas.

Sophisticated methods for monitoring the physiological status of divers while in the water is a topical subject. While recent articles have highlighted interest in this area [11] – [15], our aim is to comprehensively review the relationship between technical approaches to wearable monitoring during immersion and the prevention and diagnosis of diving-related pathology.

II. DIVING MEDICINE

Diving and hyperbaric medicine [16] deals with diseases and injuries related to high ambient pressure, hyperbaric oxygen and immersion. Fatalities and injuries are reported by the Divers Alert Network (DAN) Annual Diving Reports [17]. In the latest report from 2015, DAN presents some analyses of scuba dive fatalities involving U.S. citizens, predominantly in the USA or Caribbean (Fig. 1). Most fatalities occurred in male divers aged 50-59 years. Investigations of experience level and diving activity revealed that most fatalities occurred when divers were inexperienced and the deaths occurred during recreational diving (65%). Although drowning is the most commonly attributed cause of death, this is sometimes secondary to disabling agents such as cardiac arrhythmias or myocardial infarctions, particularly in older divers. A discussion of monitoring that might prevent cardiac death in general is outside the scope of this review but may become applicable

Corresponding author: Tobias Cibis; e-mail: tobias.cibis@fau.de

T. Cibis and B. Eskofier were with the Pattern Recognition Lab, Digital Sports Group, Friedrich-Alexander University, Erlangen-Nürnberg, Germany

T. Cibis and A. McEwan were with the Department of Electrical Engineering, The University of Sydney, Sydney, NSW, Australia.

A. Sieber was with the SEABEAR Diving Technology, Leoban, Austria.

K. Friedl was with the Department of Neurology, University of California, San Francisco, USA.

J. Lippmann was with the Divers Alert Network Asia-Pacific, Melbourne, Victoria, Australia.

M. Bennett was with the Department of Diving and Hyperbaric Medicine, Prince of Wales Clinical School, UNSW, Randwick, Sydney, Australia.

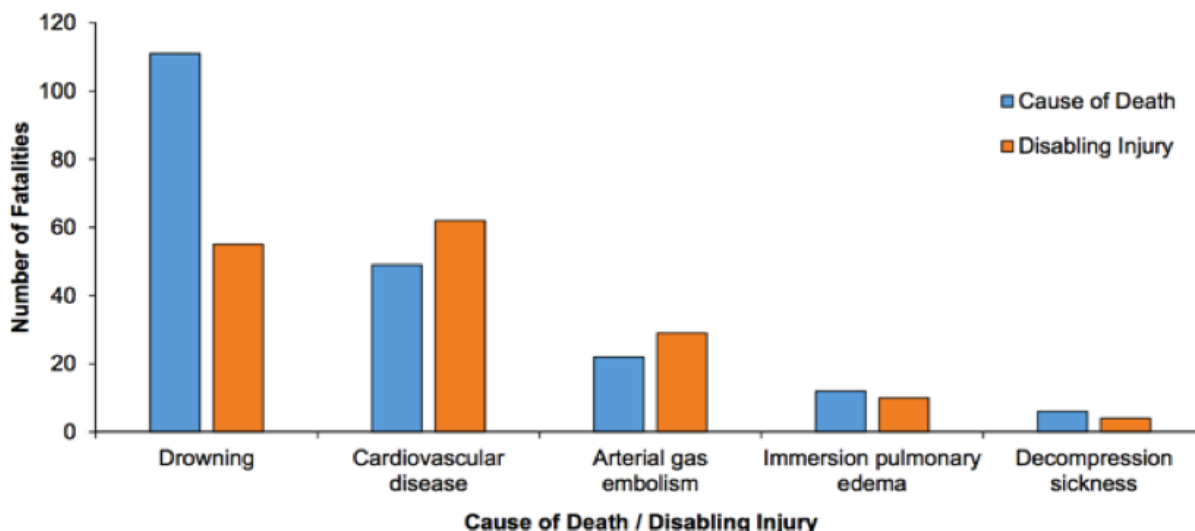


Fig. 1. Divers Alert Network investigation of diving fatalities in the USA from the Annual Diving Report 2015 [17]. Drowning and pre-existing cardiovascular diseases are the two major causes of death. However, drowning has also been considered as a result of other existing diseases or injuries.

in divers in the future. The pattern of fatality indicates the need to target injury monitoring and prevention including improved diver education.

A. DIVING PHYSIOLOGY

The human body is adapted to the terrestrial environment, performing best at 1 atmosphere absolute (ATM). Unsurprisingly, significant physiological changes occur when the body is immersed and subjected to increased pressure. The 'diving reflex' is the initial physiological response to immersion and is most prominent in cold water and exposure of the face [5]. Initially there is a restriction of blood flow to the peripheries (peripheral arterial vasoconstriction) and a reflex bradycardia [16]. There is also venoconstriction secondary to hydrostatic forces and combined with cooling peripheries this results in shunting of blood into central veins [18]. The net result of increased venous return, increased stroke volume and the reflex bradycardia is a modest increase in cardiac output [19]. The increased venous return also stimulates atrial receptors to release atrial natriuretic factors (ANF) in response to stretching which, in addition to an associated decrease of anti-diuretic hormone (ADH), results in unopposed diuresis [8]. High ambient pressure also impacts respiratory mechanical factors [20]. When the diver is upright (particularly head out of the water), the higher pressure on the chest wall compared to the mouth will result in compression of the thoracic structures and require an increased negative pressure generation in the lungs to achieve inspiration. This further encourages increased venous return to the heart and exacerbates the hemodynamic changes.

B. DIVING DISEASES AND INJURIES

1) **DROWNING**: Drowning, defined as a process of experiencing respiratory impairment from submersion in liquid [21], is the fourth commonest cause of injury-related death

accounting about half a million deaths annually worldwide [22]. During this period the victim is unable to ventilate the lungs, resulting in oxygen depletion and carbon dioxide retention. The victim will become hypercarbic, hypoxemic and acidotic and will frequently inhale water leading to death [23]. Drowning often occurs as a consequence of a primary cause associated with a reduced level of consciousness and the loss of upper airway reflexes [24].

2) **CARDIAC DISEASES**: Sudden cardiac death while immersed is mostly a result of myocardial infarction or a cardiac dysrhythmia [25]. Diving induces a series of stresses (exercise, cold, breath-holding, diving reflexes) which may impact on cardiac function. These factors can be exacerbated by a reduction of blood volume secondary to the diuresis induced by immersion, tachycardia or bradycardia, hypertension and increased cardiac work [26]. Undoubtedly, pre-existing cardiac disease increases the risk of sudden death. While the presence of myocarditis, hypertrophic cardiomyopathy or genetic disorders predispose young subjects to cardiac death, cardiac disorders in older subjects are more often associated with hypertension, dysrhythmia or triggering events [27]. Minor cardiac conditions may preface major problems in divers.

3) **BAROTRAUMA**: Describes injuries to the body as a result of changes in barometric pressure [8]. Rapid changes in ambient pressure can cause serious damage to several body systems where gas is contained within tissue. Most commonly barotrauma affects the middle-ear, lungs and respiratory sinuses.

4) **GAS TOXICITY**: Oxygen toxicity and nitrogen narcosis are hazards caused by inspired gases during scuba diving. Oxygen toxicity [28] occurs when breathing an elevated partial pressure of oxygen and depends on the pressure degree and exposure time. It is considered to result from reactive oxygen species (sometimes called 'free radicals'). When present in sufficient concentration, these species can overwhelm the available antioxidant systems and result in temporary or per-

manent tissue injury [29]. While the lungs may be affected over a longer period of continuous exposure, the more grave danger for divers is central neurological oxygen toxicity which can manifest as a grand mal seizure without warning.

Nitrogen narcosis [30] is a result of the ability of nitrogen to produce narcotic effects similar to volatile anesthetic agents. With an increasing partial pressure of nitrogen in the central nervous system, the diver will be increasingly impaired and ultimately lose consciousness. The narcotic effect is probably due to an impairment of cell membrane function. While oxygen effects on the lungs and other tissues may persist for some time, nitrogen narcosis is completely reversible when ambient pressure decreases.

5) **HYPOTHERMIA**: Water conducts heat 25 times better than air. Determined divers may drop their core temperature to a point of mild hypothermia (33–35°C). This causes shivering, marked diuresis and numbness from peripheral vasoconstriction. Lethargy may cause poor decision-making with further cooling causing incoordination, unconsciousness and drowning [31].

C. DECOMPRESSION ILLNESS

Decompression illness (DCI) is the umbrella term to describe health problems caused by the effect of gas (usually nitrogen) in the blood, the tissues and organ systems as a result of exposure to changing ambient pressures. The term includes Decompression Sickness (DCS) where problems derive from gas dissolved within the tissues and later liberated on decompression and arterial gas embolism (AGE) (Fig. 2), [32]. AGE often results from pulmonary over-distension and the subsequent entry of breathing gas in pulmonary vasculature. However, it is also associated with previously dissolved inert gas from the tissues that forms bubbles in the venous blood, passes through the great veins to the right heart and crosses through an intracardiac or pulmonary vascular abnormality to the systemic arterial system [33].

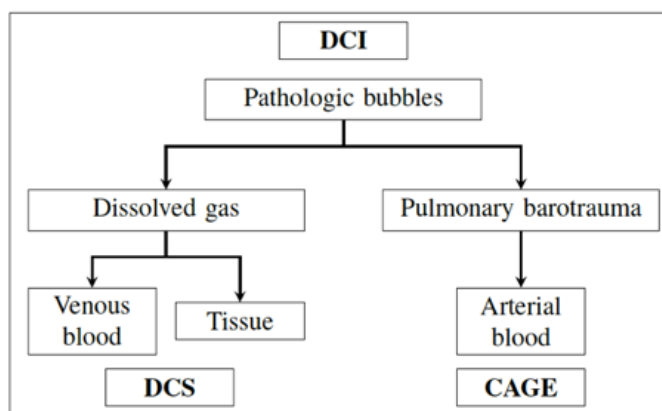


Fig. 2. Definition of bubble related diving diseases. The umbrella term Decompression Illness (DCI) includes Decompression Sickness (DCS) and Cerebral Arterial Gas Embolism (CAGE).

DCS is predominantly caused by bubble formation from dissolved gas absorbed during scuba diving. These bubbles

form during or after reduction in ambient pressure on ascent. Divers absorb inert gas while breathing at an increased ambient pressure [34], [35]. During decompression the ambient pressure falls, prompting the elimination of dissolved gas down a pressure gradient. This implies the tissue will contain more dissolved gas than it would when in an equilibrium state with the ambient pressure (the tissue is 'supersaturated'). If the level of supersaturation is high enough, the gas will come out of solution and form bubbles. Bubbles can result in a wide variety of symptoms depending on the location, volume and persistence of the gas. These include cardio-pulmonary, cerebral, inner ear, spinal cutaneous and 'mild syndromes' [36].

III. DIVE COMPUTER, DECOMPRESSION TABLES AND ALGORITHMS

A. DIVE TABLES

Diving tables were developed to protect divers from DCS [37]. These tables define stepwise ascent protocols for minimizing bubble formation over a range of duration and depth exposure. All diving tables predict the uptake and elimination of inert gas during a dive and the subsequent decompression. The aim is to manipulate the decompression in order to limit the supersaturation of tissue to a level that can be tolerated without developing DCS. Tables may be entirely based on an algorithm or the result of an algorithm modified by human trial exposures, or by theoretical preferences of the designer.

B. DIVE COMPUTER

Dive computers provide real-time information about depth and duration. The primary functions of these computers are to calculate the risk of dangerous supersaturation in the tissue and avoid this point by stepping divers safely from the pressure exposure back to the surface. Many utilize a number of decompression algorithms in order to calculate a perceived amount of inert gas present in a range of theoretical tissues within the body [10]. Comparing the calculated saturation of each tissue will identify the 'critical' tissue controlling ascent. The critical tissue is that which determines when and for how long a decompression stop is required in order to avoid a dangerous supersaturation that is likely to result in clinical DCS. Some dive computers also measure physiological information, such as breathing gas consumption or heart rate [10].

The ascent schedules depend on the implemented decompression model / algorithm [38], [39]. Despite a variety of existing models and algorithms, manufacturers of dive computers often do not provide detailed information about the software. A standard method of dive computer validation has not been established, although the need has been identified [40]. In most cases, manufacturers do not have significant hard data to confidently justify claims of safety from a functional point of view, biasing their algorithms on theoretical tissue compartment behavior [40]. Several studies have nevertheless attempted to evaluate dive computers functionality from a safety perspective.

Azzopardi et al. [41] investigated the technical specifications of 47 different dive computers. They compared whether dive

computer performance matched the technical specification described in the device manual. Some computers did not reflect the capabilities as described in their manuals. Their review emphasized the necessity for the development of standardized validation procedures.

Angelini et al. [42] investigated six dive computers during an identical set of exposures. All devices were pressurized simultaneously in a hyperbaric chamber to a maximum pressure of 8 ATM (70 msw) in three different contexts: a no-decompression stop dive, a dive requiring formal decompression stops and a set of repetitive dives. The data displayed on each computer was compared to data sets prepared in previously performed computer simulations. The results suggested all computers proposed different decompression schedules with substantial variation in decompression stop depths and durations. The authors concluded that the devices operated differently and a central data set for accurate evaluation is missing.

Most dive computer still accept a 2% DCS probability, representing a gap between latest research and current application [36]. Dive injuries still occur, even if divers stay within the suggested schedules of their dive computers [17]. DAN Asia-Pacific records (unpublished) indicate that approximately 80% of divers who had contacted them with symptoms of suspected DCS had been diving within the limits of their dive computers. This suggests that divers should be conservative when following the advice of their computer. Where possible, it may be wise to set the computer into a more conservative mode to reduce DCS risks.

C. DECOMPRESSION ALGORITHM

Decompression algorithms are derived from models, data and ascent schedules applied for underwater operation [45]. Complex equations delineate decompression schedules managing dissolved and free gas in blood and tissue (taking real-time depth and duration measurements into account) to stage divers as safely as possible to the surface. The equations are governed by a number of different factors [37] representing biological systems and medias and physical and chemical mechanisms [46] – [50].

In 1908, Haldane, Boycott and Damant [33] proposed that body tissues will hold gas in a supersaturated state. They determined tissue perfusion was the limiting factor in inert gas uptake and concluded tissues can tolerate a halving of ambient pressure before the onset of DCS symptoms. This approach was accepted for many years and practiced widely – resulting in a considerable reduced incidence of serious DCS. However, the overall incidence remained unacceptably high as diving techniques and equipment permitted longer and deeper exposures.

In 1965, Goodman and Workman proposed decompression models which extended Haldanes' supersaturation concept [51], [52]. They introduced the concept of an 'M-value' – defined as the maximum value of inert gas pressure (absolute) that a hypothetical tissue compartment can tolerate without presenting overt symptoms of DCS [53]. Based on the M-values for a range of such hypothetical tissues, decompression

schedules were modeled to stage divers to the surface with greater safety.

Further developments in the physics of phase separation, bubble behavior, and gas transfer mechanisms culminated in dissolved-free phase theories [46] – [56]. Unlike the supersaturation models, phase theories also consider free gas in addition to inert gas. Commonly implemented phase models are the varying permeability model (VPM) and the reduced gradient bubble model (RGBM).

The VPM was constructed using a critical bubble volume hypothesis [56]. This model assumes that microbubbles will grow after they exceed a critical cutoff radius that is dependent upon the level of supersaturation in the tissue. The VPM estimates acceptable pressure gradients that aim to minimize the total volume of bubbles. The aim of this model is to keep the total bubble volume less than a pre-defined limit volume to keep bubbles in the microbubble stage and not excite them into growth.

The RGBM approach [37] employs a phase volume constraint across a total dive profile. The model is parameterized by biological, chemical and physical factors [37]. Bubbles are assumed to expand and contract during diving. The material properties of these bubbles dictate the response to pressure changes, gas diffusion and growth. The estimation of the volume constraint is defined in terms of a bubble phase function, (Φ), dependent on the number of gas bubbles stimulated into growth by decompression, a supersaturation gradient, seed diffusion and Bolye's expansion-contraction law. The RGBM stages decompression iteratively ensuring that Φ does not exceed pre-defined limitation. It is unknown if any dive computers incorporate the full RGBM.

These are theoretical models and neither of these have been validated using human trials [43]. Possibly because of the large number of different factors, the equations are still not able to predict bubble behavior accurately [57]. As a result, the implementation of physiological parameters in decompression models has been very limited.

Gutvik et al. [58] proposed a first attempt at decompression modeling including physiological parameter in the Copernicus model [60]. They introduced a mathematical model using continuous heart rate measurements to calculate customized decompression schedules [59]. The heart rate was assumed responsive to cardiac workload and to provide information on the blood flow to body tissue, and thus to changes in the rate of uptake and elimination of inert gas. The aim was to combine this information with depth and time to control bubble growth. Despite operating on the assumption of static parameters [58], the model is designed to give a quantitative description of the physiological parameters significantly affecting decompression. The model's accuracy has yet to be validated with large data sets.

The most recent approaches in decompression research involve the potential role for changes in serum microparticles, inflammatory markers and genetic expressions with hyperbaric exposure. The diving environment triggers a cellular stress response which results in inflammation and the generation of microparticles both of which can be linked to the appearance of intravascular bubbles [61], [62]. The analysis of gene

activity after diving indicates that some sets of genes respond to hyperbaric stress and predispose an individual to DCS. It has been noted in the field that some individuals are more likely to suffer DCS than others, and if this is so, there may in part be some genetic predisposition. It is also possible that in the future, genetic information may be used as biomarkers for early DCS recognition [61] – [63].

IV. MEDICAL MONITORING TECHNOLOGY FOR DIVING

The consideration of diving physiology and diving-related diseases above highlights the need to monitor physiological parameters. In particular, the hemodynamic changes on diving along with the potential of assessing inert gas uptake and elimination, indicate that measuring cardiac parameters such as blood pressure, heart rate, cardiac volumes and output are of major interest. These parameters are already monitored in a wide range of application in marine mammals by sensors and data loggers transplanted into the mammals [64] – [68]. The potential to extend these measurements to humans using wearable technologies, would greatly enhance diving physiological research in general and the management of decompression in particular [68]. Monitoring physiological parameters will reflect an individual's response to diving, rather than relying on a general model, and thereby may increase diving safety [69]. To date, wearable means to measure and evaluate the onset of DCI have not emerged, although there has been some related laboratory and animal work published in recent years [67], [69], [70] – [72]. These methodologies could be converted to wearable approaches to satisfy the need for real-time monitoring [65]. For example, Evgendis et al. [73] introduced an impedance measurement concept to detect bubbles using advanced time and frequency analysis for correlation with bubble characteristics. Such an approach has the potential to determine the onset of DCS in real-time.

A. WEARABLES

Wearable devices are increasingly used over a wide range of sporting and medical applications. These devices are able to analyze vital and environmental events and display the results to consumers. The usefulness of these devices is enhanced by modern processors, micro-controllers and battery technology. This innovative field offers ways to develop wearables with multiple sensors acquiring biomedical signals simultaneously, leading to a better analysis of physiological changes and the relation between different physiological effects.

Work is performed on a number of related fronts. Physiological parameter monitoring in an extreme situation has been described by Garbino et al. [13]. They investigated physiological changes during a stratospheric jump. The subject was equipped with a device capturing ECG, respiration rate and three-axis acceleration. The device stored data on a microSD card and transmitted data in real-time. Fei et al. [12] has proposed an embedded biomedical sensor system for astronauts. The system is capable of monitoring several parameters in parallel for space walks, including sensors for temperature, galvanic skin resistance, plethysmography, oxygen saturation and heart rate.

They achieved parallel data analysis by developing hardware components for each sensor. Data acquisition and processing modules were applied inside the space suit and connected to the wrist display via wires. Seabrook et al. [74] investigated mobile platforms, such as Apple iOS and Android, for their capacity for medical applications while Laurino et al. [75] have developed a telemonitoring system for commercial divers that is applicable in hyperbaric chambers. The system acquires vital parameters and communicates via internet connection with external monitors outside the chamber. Sieber et al. [11] has presented instrumentation for physiological monitoring during scuba diving using an underwater data controller, containing an 8-bit micro-controller and a 3x16 character display. Companies are continually developing smaller, faster and more powerful components with capabilities for measuring different scenarios simultaneously. At the time of writing however, multi-sensor platforms with real-time processing capability during scuba diving are not available.

B. IMPEDANCEMETRY

Electrodes typically acquire high input impedances in biomedical measurements. During submersion, however, electrodes are exposed to a high conductivity medium water, compared to the poorly conductive air. Water is assumed as a low-impedance $Z_0 (= 10 \Omega)$ parallel to a high electrode-skin interface impedance $Z_{IF} (= 10 k\Omega)$. The water impedance significantly reduces the interface-input impedance of the amplifier from kilo-ohms to ohms, making biopotential measurements impossible using standard equipment. To solve this problem, hydrophobic-sealed electrodes are needed.

The current practice is to seal standard Ag/AgCl electrodes with adhesive material. However, this approach has many limitations, such as skin irritations due to the adhesive insulators, gel hydration over time and water leakage [78]. Although, biosignals can be observed as long as water does not penetrate the sealed electrodes, the quality of the acquired signal degrades over time. Further developments of electrodes for underwater use is required [76]. One method for facilitating underwater biopotential measurement without adhesive waterproofing was proposed by Ohtsu et al. [78]. This method employs a waterproof-designed electrode with a high-input impedance amplifier. The insulated electrodes are coated with a non-conductive, water resistant material which served as the electrode-skin interface. Electrode, insulator and skin created a capacitive coupling that permitted signal measurement on the body surface. Water resistance was ensured by using a five layers' electrode design covered with antistatic silicone. A driven-shield-technique was implemented to prevent gain loss in the high-frequency range when the signal was sent through an extended wire to the amplifier.

Another water-resistant electrode was designed by Reyes et al. [78]. This electrode consists of two active components with hydrophobic properties. The first component, Polydimethylsiloxane (PDMS), provides an elastomeric, hydrophobic matrix. The second component, Carbon Black (CB) was used as a conductive material uniformly distributed inside the PDMS polymer matrix to facilitate the transport of electrons [80].

TABLE I
OVERVIEW AND EVALUATION OF MEASUREMENT APPROACHES OF ELECTROCARDIOGRAPHY, BLOOD PRESSURE AND STROKE AND CARDIAC OUTPUT IN ORDER TO VALIDATE USABILITY FOR SCUBA DIVING.

	Method	Commercial Device / Study	Invasive / Non-invasive	Application in Scuba Diving
Electrocardiography	Smart Wearable Systems	Textile based [83], T-Shirt [84], Belt [85], Vest [86]	Non-invasive	Application in wet and dry suits
	Wireless Wearable ECG Sensors	ECG wireless sensor board [90]	Non-invasive	Successful application [88]
	Mobile based ECG	Variety of mobile phone implementation [98], [101]	Non-invasive	Successful application [101], (Fig. 3)
Blood pressure	Krotkoff Method	Brachial artery cuff [115]	Non-invasive	Only when operated manually
	Oscillometric Method	Oscillometric cuff based device [119]	Non-invasive	Successful trials [119]
	Finger cuff Method	Penaz et al [117]	Non-invasive	Doubtful
Cardiac Output & Stroke Volume	Thermodilution	Ganz et al. [122]	Invasive	No
	Thoracic Electrical Bioimpedance (TEB)	Newman [124], Marik [128] and Guijar [129]	Non-invasive	Possible with limitations
	Bioreactance (phase shift)	Kere [132] and Raval [133]	Non-invasive	Possible
	Photoplethysmography	Kuch [138]	Non-invasive	Successful trials [138]

ECG signal acquisition of the PDMS/CB electrodes were tested in three conditions (dry, wet, full immersion) and compared to the performance of sealed Ag/AgCl electrodes [75] – [77]. The signal quality of both electrodes was highly correlated during dry and wet conditions. In immersed conditions, the PDMS/CB electrodes, however, maintained or in some cases even amplified the ECG signal, whereas the Ag/AgCl electrodes failed to record the signal accurately. Noh et al. performed trials with varying pressure fixations of the PDMS/CB electrodes. They revealed that increasing pressure levels on the electrodes resulted in a decrease of input impedance.

To date, these are the only promising approaches for underwater bioimpedance measurement. The evaluation work performed by Reyes [79] and Noh [80] showed the potential for bioimpedance measurements during immersion. However, there is no evidence of accurate performance in scuba diving.

C. ELECTROCARDIOGRAM

The electrocardiogram (ECG) is the best known and most widely used bio-signal in medicine. Acquired information about cardiac response in divers may help determine if divers with pre-existing cardiovascular illness should be allowed to dive [8]. Real-time ECG monitoring with an alert system triggered by cardiac abnormalities may increase diving safety (Tab. I).

Wearable ECG monitoring systems have been described for domestic, fitness and diagnostic applications. The American Heart Association [81] published guidelines for ambulatory ECG measurement including information about recording techniques, risk assessment and efficiency advises for monitoring and therapy. Baig et al. [82] classified wearable ECG devices into three categories: smart wearables, wireless and mobile ECG systems.

Smart wearable monitoring systems are wearable textile and garment based technologies. These embedded wearable sys-

tems, collect and/or process acquired ECG data in textile and planar-fashionable systems, such as T-Shirts [83], [84], belts [85] or smart vests [86]. They consist of sensor boards for long-term monitoring in a non-invasive and unobstructed way. Wireless monitoring systems are based on body network areas [87]. These networks allow the integration of intelligent monitoring systems, minimized components and low-power sensors attached directly to a patients' body. Holter monitors are the state of the art device for wireless, long-term ECG monitoring and storage. Bosco et al [88] performed a study on divers using a 12-lead Holter monitor and concluded that the device produces accurate ECG tracing. An example of an ECG board with data processing is the ECG-Micro-Board (Cor-science, Erlangen, GER) [89]. In addition to acquisition and storage, the board includes heart rate variability evaluation. More advanced wireless technologies can include contactless or leadless ECG monitoring [90], [91].

Mobile ECG monitoring exploits the technology in mobile phones [92] – [100]. The set-up contains a mobile phone as a processing and display unit and an ECG sensor, attached directly to a patients' thorax. The mobile device and sensor, communicate wirelessly via Bluetooth or Wifi [94]. Gradl et al. [98] developed an Android based application, capable of QRS and arrhythmia detection. Acquired data was stored on a mobile phones internal memory and simultaneously displayed in real-time. Mobile tele-monitoring is considered one of the most promising means for daily vital sign monitoring [102]. Despite facing common delay issues, this technology provides a high level of accuracy regarding long term application and cardiac event detection [95], [98] – [100].

The application of mobile ECG monitoring in scuba diving has been successfully demonstrated by Cibis et al. [101], (Fig. 3). Heart rates were detected and an alert system was triggered when the heart rate exceeded or dropped below pre-defined critical limits. Any further application in scuba diving has yet to be evaluated.

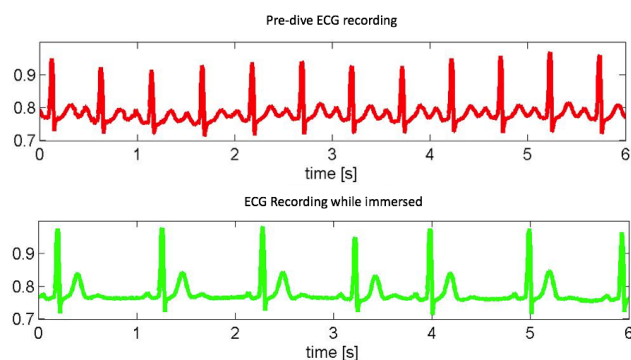


Fig. 3. ECG recordings of the same subject, taken prior and during a dive [101]. The immersed recording displays the onset of the diving reflex (bradycardia).

Von Tscherner et al. [102] recently proposed a novel current amplifier-based method to measure subcutaneous current flow, instead of surface potentials which become equipotential when immersed conditions. A current amplifier extracts current arising at the skin surface and thereby actively grounds the area beneath the electrodes enabling physiological measurements underwater without further electrode insulation [76]. In underwater trials, the information obtained in dry and wet environments was comparable [102].

For scuba diving, several of the listed ECG (Tab. I) measurement system seem applicable. Limitations are the ability to waterproof the system and the electrode-device connection [102]. The effects of body and water movement on the signal quality are unknown. A useful system has to operate accurately and be robust in the face of movement [103], [104].

D. BLOOD PRESSURE

Any changes in blood pressure (BP) over the course of a dive remains to be defined [11]. Accurate BP measurement is essential to understand hemodynamic effects and to classify cardio-vascular health risks [105] – [110]. Invasive methods are impractical for safety reasons [110] but have been used as the 'gold-standard' against which to validate non-invasive methods [111] – [114]. Non-invasive, automated methods include the oscillometric and the finger cuff based method (Tab. I).

The automated oscillometric method is derived from the manual Korotkoffs' BP method where the operator hears changes in arterial flow underneath a stethoscope during slow cuff deflation [115]. Oscillometry detects pressure oscillations under cuff rather than sound [105], [108]. The maximum oscillation corresponds to the mean arterial pressure [110], [112]. Because oscillations occur above the points of systolic and diastolic pressure, these values are only estimated indirectly by defined algorithms [116]. This method fails during physical activity because of movement artifacts, and this may make use during diving problematic [116]. Another method was first proposed by Penaz et al. in the 1970s [117], [118]. Penaz developed a finger cuff method, based on an 'unloaded arterial wall'

principle. Arterial pulsation in the finger is detected using a photo-plethysmograph, obtaining pressure oscillation from the measured pressure curve [105]. Trials demonstrated that the finger cuff method fails to measure absolute levels of blood pressure accurately, but does reflect changes over time accurately [118]. Because immersion at pressure tends to minimize blood flow in the peripheries, the use of this method is also problematic during diving.

Recently, a sphygmomanometer BP measurement device for underwater application was developed by Sieber et al. [119]. The aim was to obtain real-time BP measurements to evaluate BP response during breath-hold dives. The device was designed to fit in a waterproof housing, directly mounted to the cuff containing a microcontroller board, differential pressure sensor and display. The first trials were performed in a swimming pool at varying depths and in dry hyperbaric environments with good results.

The challenges for BP measurements in divers are to waterproof the control boards and the cuff operation (Tab. I). The impact of ambient pressure on the cuff must be considered [120]. Movement may compromise BP detection and have yet to be addressed in literature. The oscillometric method seems most likely to reach the exacting requirements [117].

E. STROKE VOLUME & CARDIAC OUTPUT

During scuba diving, increased ambient pressure causes a blood shift to the central compartment [5]. Measuring cardiac output (CO) and stroke volume (SV) will provide information about blood volume changes and cardiac work. This information can be used to understand blood shifts and may help to establish if divers with preexisting conditions can safely dive.

Technologies in SV and CO measurements predict the amount of fluid in the heart (Tab. I). The use of Fick's principle measuring the thermodilution of cold injective is the clinical standard to determine cardiac output [121], [122], but it is invasive and impractical for diving.

The thoracic electric bioimpedance (TEB) method was introduced by Kubicek to determine SV and CO non-invasively [123] – [127]. Kubicek proposed that pulsatile changes in resistance occur during ventricular systole and diastole. Four electrodes are applied on the neck and thorax and drive a current of known amplitude and frequency across the thorax. The pulsation is measured in voltage changes where the amplitudes of the output signal is compared to the input signal. The trans-thoracic impedance (Z_0) can be calculated and changes proportionally to the amount of fluid in the thorax. The stroke volume is proportional to the impedance change dZ/dt over the left ventricular ejection time, T_{LVE} . Early studies showed a poor agreement between TEB and thermodilution [128] and agreement was even worse with an increase of blood in the thorax (as is expected during scuba diving) [129]. TEB is also inaccurate in settings with electrical background noises, physical activity and environmental factors, while being sensitive to the electrodes placement [130], [131]. TEB seems of limited use for scuba diving.

Another method is bio-reactance, a phase shift in voltage

across the thorax [125] – [127], [132]. The human thorax was described as an electrical circuit containing a resistor R and a capacitor C , creating the thoracic impedance Z_0 . The values of R and C determine two components of the impedance; the amplitude and magnitude of the impedance measured in ohms; and the phase orientation measured in degrees. Pulsatile blood flow modifies R and C , leading to instant changes in amplitude and phase. The phase shift depends directly on the flow and constitutes the parameter of interest. The system operates on four dual electrodes. One electrode pair is driving an electric current across the thorax and the other pair measures voltage. The CO is determined by the relative phase shift, ϕ , between the input and output signals. The peak rate of the phase shift, $(\Delta\Phi)$, is proportional to the peak aortic flow. The SV is calculated using the phase shift peak rate and the ventricle ejection time T_{VET} . Unlike the TBE, the bioreactance method does not use a static impedance. Studies show a high correlation between bioreactance and thermodilution [133] – [135]. The good agreement, strong resistance against motion artifacts and low sensitivity to environmental and physical factors all suggest bioreactance may be usefully applied in scuba diving. An optical approach to measure peripheral volume changes is photoplethysmography (PPG) [136]. This method illuminates the skin and measures the amount of transmitted or reflected light to a photodiode. The amount of light detected at the photodiode is proportional to changes in the skin blood volume, however, these devices do not produce a quantitative measurement [137]. PPG can, however, be used to measure oxygen saturation and heart rate, and has been used in immersed conditions [138].

The challenges for scuba diving application are to ensure accuracy and safety underwater (Tab. I). In particular, the impact of immersion is unknown. Trials and studies are required to determine technical, physical and physiological factors for successful blood shift measurements.

V. OUTLOOK

This review has provided an overview of diving medicine and related monitoring technology. Each identified technological approach is considered to be applicable for advanced, wearable data acquisition underwater and may ultimately contribute to a detailed evaluation of diving physiology and diseases.

The long-term health consequences of compressed gas breathing are poorly defined and require further investigation as relatively few data are available and some are contradictory [139] – [142]. To date, some studies imply that cognitive and neurological impairments are the most serious long-term consequences of diving. While such long-term effects are assessed using questionnaires, coordination and reaction tests and imaging technologies (MRI, SPECT); wearable means to evaluated real-time impairment have also been described. A wearable critical flicker fusion frequency (CFFF) device has been used to determine alteration or arousal of cortical performance [143], [144]. Studies have shown that an increase of fusion frequency is equated with brain activation and a decrease equated in alteration of brain arousal with potential

application for real-time evaluation of nitrogen narcosis [145]. An other approach to measure brain activity is the application of a wearable electroencephalogram (EEG) [146]. EEG patterns can be used to investigate the impact of increased pressure [147] and nitrogen narcosis on the cortical performance [148]. Information obtained from a real-time measurement of cognitive functions could be used to prevent dangerous conditions and to investigate long-term impairments. Sensors for respiration and breathing gas analysis were not considered in this review. Gas analysis requires sensor attachment and modification of the breathing regulator and therefore was not considered in the scope of wearable technology. This review has concentrated on compressed-air diving and has not considered specific issues relating to the use of different breathing gas mixtures (such as those containing helium) [149]. These issues will need to be considered in relation to deep diving safety.

REFERENCES

- [1] M.D.J. Sayer, J. Barrington, "Trends in scientific diving: an analysis of scientific diving operation records, 1970-2004," *Underwater Technology*, 51-55, 2005.
- [2] M. Bennett, et al., "Recompression and adjunctive therapy for decompression illness," *Cochrane Database Syst Rev* 2, 2007.
- [3] M. Bennett, S. Mitchell & A. Dominguez, "Adjunctive treatment of decompression illness with a non-steroidal anti-inflammatory drug (tenoxicam) reduces compression requirement," *Undersea & hyperbaric medicine* 30.3, 195, 2003.
- [4] A. Spira, "Diving and marine medicine review part II: diving diseases," *Journal of travel medicine* 6.3, 180-198, 1999.
- [5] Y. Kawakami, B.H. Natelson, A.R. DuBois, "Cardiovascular effects of face immersion and factors affecting diving reflex in man," *Journal of Applied Physiology* 23.6, 964-970, 1967.
- [6] W. Hemelryck, et al., "Long term effects of recreational SCUBA diving on higher cognitive function," *Scandinavian journal of medicine & science in sports* 24.6 928-934, 2014.
- [7] R. Vann, M. Lang, "Recreational diving fatalities," *Undersea Hyperb Med.* 38(4):257-60, 2011.
- [8] A.O. Brubakk, T.S. Neuman, "Bennett and Elliott's physiology and medicine of diving," *Saunders Book Company*, 2003.
- [9] M.B. Strauss, "Diving medicine: contemporary topics and their controversies," *The American journal of emergency medicine* 19.3, 232-238, 2001.
- [10] S. Blogg, et al., "Proceedings of Validation of Dive Computers Workshop," 2012.
- [11] A. Sieber, et al., "Advanced instrumentation for research in diving and hyperbaric medicine," *Undersea and Hyperbaric Medical Society, Inc.*, 2010.
- [12] D.Y. Fei, et al., "A biomedical sensor system for real-time monitoring of astronauts? physiological parameters during extra-vehicular activities," *Computers in biology and medicine* 40.7 635-642, 2010.
- [13] A. Garbino, et al., "Physiological monitoring and analysis of a manned stratospheric balloon test program," *Aviation, space, and environmental medicine* 85.2, 177-182, 2014.
- [14] M. Pieri, et al., "Continuous real-time monitoring and recording of glycemia during scuba diving: pilot study," *Undersea & hyperbaric medicine, Inc* 43.3, 265, 2016.
- [15] B.H. Groh, et al., "IMU-based pose determination of scuba divers? bodies and shanks," *Wearable and Implantable Body Sensor Networks (BSN)*, IEEE, 2015.
- [16] T. Chowdhury, et al., "Trigemino-cardiac reflex: the current clinical and physiological knowledge," *Journal of Neurological Anesthesiology* 27(2): 136-147, 2015.
- [17] Diving Alert Network (DAN), "Annual Diving Report - 2012-2015 Edition", 2015.
- [18] T.J.R. Francis, S.J. Mitchell, "Bennett and Elliott's physiology and medicine of diving," *Saunders*, 530-556, 2003.
- [19] A. Boussuges, et al., "Hemodynamic changes induced by recreational scuba diving," *CHEST Journal* 129.5, 1337-1343, 2006.

- [20] S.J. Thornton, P.W. Hochachka, "Oxygen and the diving seal," *Undersea & Hyperbaric Medicine* 31.1, 81, 2004.
- [21] World Health Organization (WHO). Fact Sheet Drowning. http://www.who.int/violence_injury_prevention/publications/other_injury/en/drowning_factsheet.pdf, 2016.
- [22] E. Krug, "Injury, a leading cause of the global garden disease," *World Health Organization (WHO)*, 1999.
- [23] A.H. Idris, et al., "Recommended guidelines for uniform reporting of data from drowning: the 'Utstein style'," *Circulation* 108, 2565-74, 2003.
- [24] M.D. Weinstein, et al., "Near drowning: epidemiology, pathophysiology, and initial treatment," *J Emerg Med*, 14:461-7, 1996.
- [25] M. Bennett, "Cardiac problems and sudden death," *In Diving and Subaquatic Medicine*, 5th Edition, CRC Press, 449-457, 2015.
- [26] C. Marabotti, et al., "Cardiovascular changes during SCUBA diving: an underwater Doppler echocardiographic study," *Acta Physiologica* 209.1:62-68, 2013.
- [27] A. Tabib, et al., "Undetected cardiac lesions cause unexpected sudden cardiac death during occasional sport activity. A report of 80 cases," *European Heart Journal* 20.12:900-903, 1999.
- [28] D. Torbati, et al., "Free radical generation in the brain precedes hyperbaric oxygen induced convulsions," *Free Radical Biol Med*, 13:101-106, 1992.
- [29] A.B. Lumb, A.E. Taylor, "Hypoexia and Oxygen Toxicity," *Num?'s applied Respiratory Physiology*, 5th Ed. Butterworth Heinemann, 200, 491-510, 2003.
- [30] J.E. Clark, "Moving in extreme environments: inert gas narcosis and underwater activities," *Extreme physiology & medicine* 4.1, 2015.
- [31] D.F. Danzl, R.S. Pozos, "Accidental hypothermia," *New England Journal of Medicine* 331.26, 1756-1760, 1994.
- [32] D.J. Doolette, S.J. Mitchell, "The physiological kinetics of nitrogen and the prevention of decompression sickness," *Clin. Pharmacokinetics*, 40.1, 1-14, 2001.
- [33] A.E. Boycott, J.S. Haldane, "The prevention of compressed-air illness," *Journal of Hygiene* 8.03, 342-443, 1908.
- [34] S.J. Mitchell, D.F. Gorman, "The pathophysiology of cerebral arterial gas embolism," *J Extracorporeal Technology*, 34:18-23, 2002.
- [35] T.J.R. Francis, S.J. Mitchell, "Manifestations of decompression disorders," *Brubakk AO, Neuman TS (Eds.), Bennett and Elliott's Physiology and Medicine of Diving (5th ed)*, Harcourt Publisher, London, 530-599, 2003.
- [36] L.E. Howle, et al., "The probability and severity of decompression sickness," *PloS one* 12.3: e0172665, 2017.
- [37] B.R. Wienke, "Diving decompression models and bubble metrics: Modern computer syntheses," *Computers in biology and medicine* 39.4, 309-331, 2009.
- [38] K.A. Gault, et al., "US Navy Dive Computer Validation," 2012.
- [39] E.D. Thalmann, "Phase II Testing of Decompression Algorithms for Use in the US Navy Underwater Decompression Computer," *Navy Experimental Diving Unit Panama City Florida*, 1984.
- [40] A. Sieber, et al., "Dive computers: the need for validation and standards," *Rubicom Foundation*, 2012.
- [41] E. Azzopardi, et al., "A review of the technical specifications of 47 models of diving decompression computer," *Underwater Technology* 29.2, 63-72, 2010.
- [42] S.A. Angelini, "Validation of dive computer algorithms," *Rubicom Foundation*, 2012.
- [43] S.M. Egi, N.M. Gurmen, "Computation of decompression tables using continuous compartment half-lives," *Undersea & hyperbaric medicine* 27.3, 143, 2003.
- [44] D. Andri, et al., "Change of occurrence of type 1 and type 2 decompression sickness of divers treated at the Croatian Naval Medical Institute in the period from 1967 to 2000," *International maritime health* 54.1-4, 127-134, 2002.
- [45] B.R. Wienke, "Deep Stop Model Correlations," *J Bioengineer & Biomedical Sci* 5.155, 2015.
- [46] R.D. Vann, H.G. Clark, "Bubble growth and mechanical properties of tissue in decompression," *Undersea Biomed. Res.* 2, 185-194, 1975.
- [47] P.S. Epstein, M.S. Plesset, "On the stability of gas bubbles in liquid-gas solutions," *J. Chem. Phys.* 18, 1505-1509, 1950.
- [48] R.D. Vann, et al., "Decompression illness," *The Lancet* 377.9760, 153-164, 2011.
- [49] R.G. Buckles, "The physics of bubble formation and growth," *Aerospace Med.* 39, 1062-1069, 1968.
- [50] D.C. Pease, L.R. Blinks, "Cavitation from solid surfaces in the absence of gas nuclei," *J. Phys. Coll. Chem.* 51, 556-567, 1947.
- [51] B.R. Wienke, "Basic Decompression Theory and Application," *Best Publishing Company*, Flagstaff, 2001.
- [52] R.D. Workman, "Calculation of decompression schedules for nitrogen-oxygen and helium-oxygen dives," *No. NEDU-RR-6-65. Navy Experimental Diving Unit Panama City Florida*, 1965.
- [53] E.C. Baker, "Understanding M-values," *Immersed* 3, 23-27, 1998.
- [54] B.R. Wienke, "Computer validation and statistical correlations of a modern decompression diving algorithm," *Computers in biology and medicine* 40.3, 252-260, 2010.
- [55] B.R. Wienke, "Computational decompression models," *Int. J. Biomed. Comput.* 21, 205-221, 1987.
- [56] D.E. Yount, "On the evolution, generation, and regeneration of gas cavitation nuclei," *The Journal of the Acoustical Society of America* 71.6, 1473-1481, 1982.
- [57] B.R. Wienke, "Numerical phase algorithm for decompression computers and application," *Computers in biology and medicine* 22.6, 389-406, 2001.
- [58] C.R. Gutvik, et al., "Parameter estimation of the copernicus decompression model with venous gas emboli in human divers," *Medical & biological engineering & computing* 48.7, 625-636, 2010.
- [59] C.R. Gutvik, et al., "Use of heart rate monitoring for an individualized and time-variant decompression model," *European journal of applied physiology* 110.5, 885-892, 2010.
- [60] C.R. Gutvik, A.O. Brubakk, "A dynamic two-phase model for vascular bubble formation during decompression of divers," *IEEE Transactions on Biomedical Engineering* 56.3, 884-889, 2009.
- [61] J. Lautridou, et al., "Effect of simulated air dive and decompression sickness on the plasma proteome of rats," *PROTEOMICS-Clinical Applications* 10.5: 614-620, 2016.
- [62] I. Eftedal, et al., "Immune and inflammatory responses to freediving calculated from leukocyte gene expression profiles," *Physiological Genomics*, 2016.
- [63] D. Madden, et al., "Exercise before and after SCUBA diving and the role of cellular microparticles in decompression stress," *Medical hypotheses* 86, 80-84, 2016.
- [64] P.J. Butler, D.R. Jones, "Physiology of diving of birds and mammals," *Physiological reviews* 77.3, 837-899, 1997.
- [65] S. Atkinson, et al., "Stress physiology in marine mammals: how well do they fit the terrestrial model?" *Journal of Comparative Physiology B* 185.5, 463-486, 2015.
- [66] A. Fahlman, et al., "Physiological Monitoring in Diving Mammals," *Texas A&M Univ Corpus Christi TX*, 2013.
- [67] M.A. Lang, et al., "Diving physiology and decompression sickness: Considerations from humans and marine animals," *Smithsonian Contributions to Marine Sciences* 29, 23-38, 2013.
- [68] P. Ponganis, "After 73 years, still the foundation of diving physiology research," *Journal of Experimental Biology* 216.18, 3381-3383, 2013.
- [69] R.E. Moon, R.D. Vann, P.B. Bennett, "The physiology of decompression illness," *Scientific American* 273, 70-77, 1995.
- [70] S.R. Thom, et al., "Association of microparticles and neutrophil activation with decompression sickness," *Journal of Applied Physiology* 119.5, 427-434, 2015.
- [71] I. Eftedal, "Diving into the rat plasma proteome to get to the bottom of decompression sickness," *Proteomics-Clinical Applications*, 2016.
- [72] D. Garca-Praga, et al., "Decompression sickness ('the bends') in sea turtles," *Dis Aquat Organ*, 2014.
- [73] S.P. Evgenidis, et al., "Post-dive detection of bubbles in scuba divers employing electrical impedance spectroscopy measurements," *EUBS 2016, Geneva, Switzerland*, 13-16 Sept, 2016.
- [74] H.J. Seabrook, et al., "Medical applications: a database and characterization of apps in Apple iOS and Android platforms," *BMC research notes* 7.1, 573, 2014.
- [75] M. Laurino, et al., "Psycho-physiological tele-monitoring of human operators in commercial diving: the Life Support System in the SUONO project," *Engineering in Medicine and Biology Society (EMBC)*, 37th Annual International Conference of the IEEE, 2015.
- [76] J.W. Whitting, V. von Tscharnner, "Monopolar current amplifier electromyographic signals collected in air and underwater without insulation," *Journal of Electromyography and Kinesiology*, 24(6), 848-854, 2014.
- [77] M. Ohtsu, Y. Fukuka, A. Ueno, "Underwater electromyographic measurement using a waterproof insulated electrode," *Advanced Biomedical Engineering* 1.0, 81-88, 2012.
- [78] B. Reyes, et al., "Novel electrodes for underwater ECG monitoring," *IEEE Transactions on Biomedical Engineering* 61.6, 1863-1876, 2014.
- [79] B. Reyes, et al., "Performance evaluation of carbon black based electrodes for underwater ECG monitoring," *Engineering in Medicine and Biology Society (EMBC)*, 36th Annual International Conference of the IEEE, 2014.

- [80] Y. Noh, et al., "Novel Conductive Carbon Black and Polydimethylsiloxane ECG Electrode: A Comparison with Commercial Electrodes in Fresh, Chlorinated, and Salt Water," *Annals of Biomedical Engineering*, 1-16, 2016.
- [81] M.H. Crawford, et al., "ACC/AHA guidelines for ambulatory electrocardiography: executive summary and recommendations a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines developed in collaboration with the North American Society for Pacing and Electrophysiology," *Circulation* 100.8, 886-893, 1999.
- [82] M.M. Baig, et al., "A comprehensive survey of wearable and wireless ECG monitoring systems for older adults," *Medical & biological engineering & computing* 51.5 485-495, 2013.
- [83] P. Jourand, et al., "Textile integrated breathing and ECG monitoring system," *Procedia Chemistry* 1.1, 722-725, 2009.
- [84] J. Yoo, et al., "A wearable ECG acquisition system with compact planar-fashionable circuit board-based shirt," *IEEE Transactions on Information Technology in Biomedicine* 13.6, 897-902 (2009).
- [85] E. Sardini, M. Serpelloni, "Instrumented wearable belt for wireless health monitoring," *Procedia Engineering* 5, 580-583, 2010.
- [86] P.S. Pandian, et al., "Smart Vest: Wearable multi-parameter remote physiological monitoring system," *Medical engineering & physics* 30.4, 466-477, 2008.
- [87] J.Y. Khan, et al., "Wireless body area network (WBAN) design techniques and performance evaluation," *Journal of medical systems* 36.3, 1441-1457, 2012.
- [88] G. Bosco, et al., "12-lead Holter monitoring in diving and water sports: a preliminary investigation," *Diving and hyperbaric medicine* 44.4, 202-207, 2014.
- [89] Corscience, "ECG products", <http://www.corscience.de/en/>, 2016.
- [90] Y. Yoon, et al., "Non-constrained blood pressure monitoring using ECG and PPG for personal healthcare," *Journal of medical systems* 33.4, 261-266, 2009.
- [91] D. Scherr, et al., "Prospective comparison of the diagnostic utility of a standard event monitor versus a 'leadless' portable ECG monitor in the evaluation of patients with palpitations," *Journal of Interventional Cardiac Electrophysiology* 22.1, 39-44, 2000.
- [92] C.G. Scully, et al., "Physiological parameter monitoring from optical recordings with a mobile phone," *IEEE Transactions on Biomedical Engineering* 59.2, 303-306, 2003.
- [93] C. Chan, et al., "Energy efficient diagnostic grade mobile ECG monitoring," *IEEE 10th International New Circuits and Systems Conference (NEWCAS)*, 2012.
- [94] B. Yu, L. Xu, Y. Li, "Bluetooth low energy (BLE) based mobile electrocardiogram monitoring system," *Information and Automation (ICIA)*, IEEE, 2012.
- [95] C.D. Galloway, D.E. Albert, S. Freedman, "iPhone ECG application for community screening to detect silent atrial fibrillation: a novel technology to prevent stroke," *Int. J. Cardiol.* 165, 193-194, 2013.
- [96] Z. Sankari, H. Adeli, "HeartSaver: a mobile cardiac monitoring system for auto-detection of atrial fibrillation, myocardial infarction, and atrioventricular block," *Computers in biology and medicine* 41.4, 211-220, 2011.
- [97] J.J. Oresko, et al., "A wearable smartphone-based platform for real-time cardiovascular disease detection via electrocardiogram processing," *IEEE Transactions on Information Technology in Biomedicine* 14.3, 734-740, 2010.
- [98] S. Gradl, et al., "Real-time ECG monitoring and arrhythmia detection using Android-based mobile devices," *Engineering in Medicine and Biology Society (EMBC)*, IEEE, 2012.
- [99] M. Rosenberg, et al., "Feasibility and accuracy of using mobile phone images of electrocardiograms to initiate the cardiac catheterization process," *Journal of telemedicine and telecare*, 2015.
- [100] D.D. McManus, et al., "A novel application for the detection of an irregular pulse using an iPhone 4S in patients with atrial fibrillation," *Heart Rhythm* 10.3, 315-319, 2013.
- [101] T. Cibis, et al., "Wearable real-time ecg monitoring with emergency alert system for scuba diving," *Engineering in Medicine and Biology Society*, 6074-6077, 2015.
- [102] V. von Tschamer, et al., "Comparison of electromyographic signals from monopolar current and potential amplifiers derived from a penniform muscle, the gastrocnemius medialis," *J Electromyogr Kinesiol.* 2013.
- [103] G.D. Gargiulo, et al., "Unipolar ECG circuits: towards more precise cardiac event identification," *Circuits and Systems (ISCAS)*, IEEE, 2013.
- [104] T. Martin, E. Jovanov, D. Raskovic, "Issues in wearable computing for medical monitoring applications: a case study of a wearable ECG monitoring device," *The Fourth International Symposium on Wearable Computers*, IEEE, 2000.
- [105] T.G. Pickering, et al., "Recommendations for blood pressure measurement in humans and experimental animals part 1: blood pressure measurement in humans: a statement for professionals from the Subcommittee of Professional and Public Education of the American Heart Association Council on High Blood Pressure Research," *Hypertension* 45.1, 142-161, 2005.
- [106] L.A. Geddes, "The direct and indirect measurement of blood pressure," *Year Book Medical Publisher*, Chicago, 1970.
- [107] E. O'Brien, D. Fitzgerald, "The history of blood pressure measurement," *J. Hum. Hypertens* 8, 73-84, 1994.
- [108] H. Smulyan, M.E. Safar, "Blood pressure measurement: retrospective and prospective views," *American journal of hypertension* 24.6, 628-634, 2011.
- [109] P.A. Shaltis, A. Reisner, H.H. Asada, "Wearable, cuff-less PPG-based blood pressure monitor with novel height sensor," *Engineering in Medicine and Biology Society (EMBS)*, IEEE, 2006.
- [110] T. Weber, et al., "Validation of a brachial cuff-based method for estimating central systolic blood pressure," *Hypertension* 58.5, 825-832, 2011.
- [111] G. Parati, et al., "Comparison of finger and intra-arterial blood pressure monitoring at rest and during laboratory testing," *Hypertension* 13.6 Pt 1, 647-655, 1989.
- [112] SN. Hunyor, et al., "Comparison of performance of various sphygmomanometers with intra-arterial blood pressure readings," *Br Med J.* 2, 159-162, 1987.
- [113] J. van Egmond, et al., "Invasive v. non-invasive measurement of arterial pressure. Comparison of two automatic methods and simultaneously measured direct intra-arterial pressure," *Br J Anaesth.*, 57, 434-444, 1985.
- [114] E. Umana, "Comparison of oscillometric and intra-arterial systolic and diastolic blood pressures in lean, overweight and obese patients," *Angiology* 57, 41-45, 2016.
- [115] N.S. Korotkoff, "On the subject of methods of measuring blood pressure," *Bull Imp Military Med Acad* 11, 365-367, 1905.
- [116] E. Meaney, et al., "Formula and nomogram for the sphygmomanometric calculation of the mean arterial pressure," *Heart* 84.1, 64-64, 2000.
- [117] J. Penaz, "Photoelectric measurement of blood pressure, volume and flow in the finger," *Digest of the 10th international conference on medical and biological engineering*, Vol. 104. International Federation for Medical and Biological Engineering, Publishers New York, 1973.
- [118] B. Imholz, et al., "Non-invasive continuous finger blood pressure measurement during orthostatic stress compared to intra-arterial pressure," *Cardiovascular Research* 24.3, 214-221, 1990.
- [119] A. Sieber, et al., "Underwater study of arterial blood pressure in breath-hold divers," *Journal of Applied Physiology* 107.5, 1526-1531, 2009.
- [120] J. Fortin, et al., "Continuous non-invasive blood pressure monitoring using concentrically interlocking control loops," *Computers in biology and medicine* 36.9, 941-957, 2006.
- [121] MB. Visscher, JA. Johnson, "The Fick principle: analysis of potential errors in its conventional application," *Journal of applied physiology* 5.10, 635-638, 1953.
- [122] W. Ganz, H.J.C. Swan, "Measurement of blood flow by thermodilution," *The American journal of cardiology* 29.2, 241-246, 1972.
- [123] W.G. Kubicek, et al., "Development and evaluation of an impedance cardiac output system," *Aerospace medicine* 37.12, 1208-1212, 1966.
- [124] D.G. Newman, R. Callister, "The non-invasive assessment of stroke volume and cardiac output by impedance cardiography: a review," *Aviation, space, and environmental medicine* 70.8, 780-789, 1999.
- [125] S. Armstrong, R. Fernando, M. Columb, "Minimally-and noninvasive assessment of maternal cardiac output: go with the flow!" *Int j of obst anes* 20.4, 330-340, 2011.
- [126] J.C. Denniston, et al., "Measurement of cardiac output by electrical impedance at rest and during exercise," *Journal of Applied Physiology* 40.1, 91-95, 1976.
- [127] D.A. Hett, M.M. Jonas, "Non-invasive cardiac output monitoring," *Intensive and critical care nursing* 20.2, 103-108, 2004.
- [128] P.E. Marik, et al., "A comparison of hemodynamic parameters derived from transthoracic electrical bioimpedance with those parameters obtained by thermodilution and ventricular angiography," *Critical care medicine* 25.9, 1545-1550, 1997.
- [129] A.R. Gujjar, et al., "Non-invasive cardiac output by transthoracic electrical bioimpedance in post-cardiac surgery patients: comparison with thermodilution method," *Journal of clinical monitoring and computing* 22.3, 175-180, 2008.

- [130] B.D Spiess, et al., "Comparison of bioimpedance versus thermodilution cardiac output during cardiac surgery: evaluation of a second-generation bioimpedance device," *Journal of cardiothoracic and vascular anesthesia* 15.5, 567-573, 2001.
- [131] W.S. Sageman, et al., "Equivalence of bioimpedance and thermodilution in measuring cardiac index after cardiac surgery," *Journal of cardiothoracic and vascular anesthesia* 16.1, 8-14, 2002.
- [132] H. Keren, et al., "Evaluation of a noninvasive continuous cardiac output monitoring system based on thoracic bioimpedance," *American Journal of Physiology-Heart and Circulatory Physiology* 293.1, H583-H589, 2007.
- [133] N.Y. Raval, et al., "Multicenter evaluation of noninvasive cardiac output measurement by bioimpedance technique," *Journal of clinical monitoring and computing* 22.2, 113-119, 2008.
- [134] P. Squara, et al., "Comparison of monitoring performance of Bioimpedance vs. pulse contour during lung recruitment maneuvers," *Crit Care* 13.4, R125, 2009.
- [135] P.E. Marik, "Noninvasive cardiac output monitors: a state-of the-art review," *J Cardiothorac Vasc Anesth* 27.1, 121-134, 2013.
- [136] J. Allen, "Photoplethysmography and its application in clinical physiological measurement," *Physiological measurement* 28.3: R1, 2007.
- [137] A.A.R. Kamal, et al., "Skin photoplethysmography - a review," *Computer methods and programs in biomedicine* 28.4:257-269, 1989.
- [138] B. Kuch, et al., "A novel wearable apnea dive computer for continuous plethysmographic monitoring of oxygen saturation and heart rate," *Rubicom Foundation*, 2010.
- [139] K. Todnem, et al., "Neurological long term consequences of deep diving," *British journal of industrial medicine* 48.4, 258-266, 1991.
- [140] D. McQueen, G. Kent, A. Murrison, "Self-reported long-term effects of diving and decompression illness in recreational scuba divers," *British journal of sports medicine* 28.2, 101-104, 1994.
- [141] M. Knauth, "Long-Term Neurologic Damage and Brain Lesions in Recreational Divers," *Clinical Neuroradiology* 18.1, 54-59, 2008.
- [142] D.O. Slosman, et al., "Negative neurofunctional effects of frequency, depth and environment in recreational scuba diving: The Geneva 'memory dive' study," *British journal of sports medicine* 38.2, 108- 114, 2004.
- [143] G. Conte, et al., "A Wearable Critical Flicker Fusion Frequency Detector for SCUBA divers," *The 26th International Ocean and Polar Engineering Conference*. International Society of Offshore and Polar Engineers, 2016.
- [144] W. Hemelryck, et al., "Functional comparison between critical flicker fusion frequency and simple cognitive tests in subjects breathing air or oxygen in normobaria," *Diving Hyperb Med*, 43(3):138-142, 2013.
- [145] C. Balestra, et al., "Persistence of critical flicker fusion frequency impairment after 33 mfs scuba dive: evidence of prolonged nitrogen narcosis?" *Eur J Appl Physiol*, 112(12):4063-8, 2013.
- [146] S. Schneider, et al., "When neuroscience gets wet and hardcore: neurocognitive markers obtained during whole body water immersion," *Experimental brain research* 232.10: 3325-3331, 2014.
- [147] R. Dolmierski, et al., "EEG changes under hyperbaric conditions: spectral analysis during simulated diving," *Acta neurologica scandinavica* 77.6: 437-439, 1988.
- [148] L. Pastena, et al., "EEG patterns associated with nitrogen narcosis (breathing air at 9 ATA)," *Aviation, space, and environmental medicine* 76.11: 1031-1036, 2005.
- [149] D.J. Doolette, S.J. Mitchell, "Recreational technical diving part 2: decompression from deep technical dives," *Diving and hyperbaric medicine*, 43.2, 96-104, 2013.