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DOI
10.1016/j.burns.2022.12.008

Publication date
2023

Document Version
Final published version

Published in
Burns

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Citation for published version (APA):
A clinimetric assessment of the validity and reliability of 3D technology for scar surface area measurement

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ARTICLE INFO

Article history:
Accepted 19 December 2022

Keywords:
Scars
Surface area
Measurement instrument
(3D) depth sensor
Clinimetric evaluation

ABSTRACT

Introduction: The quality of scars has become an important outcome of burn care. Objective scar assessment through scar surface area measurement enables quantification of scar formation and evaluation of treatment efficacy. 3D technology has proven valid and reliable but often remains cumbersome, expensive, and time-consuming. 3D technology with depth sensors on mobile devices has become available and might surpass these limitations. This study provides a clinimetric assessment of the validity and reliability of a 3D system with a depth sensor for scar surface area measurement.

Methods: A technology involving a depth sensor mounted on a mobile device was used. Images and analyses were made with a custom-made software application. A standardized one-keyframe image capturing procedure was followed. To assess validity, stickers with predefined dimensions (8.01 cm² – 77.70 cm²) were imaged in a single observer setting on various body parts of healthy volunteers. To assess reliability, hypertrophic scars, keloids, and normotrophic scars were imaged and rated by two observers independently. Data are
expressed as mean (+/- SD), Coefficient of Variation (CV), Intraclass Correlation Coefficients (ICC), and Limits of Agreement (LoA).

Results: Eighty stickers placed on 20 healthy volunteers showed validity with CV between 0.62%– 1.67% for observer A and 0.75%– 1.19% for observer B. For the reliability study, 69 scars on 36 patients were included. Mean scar surface area ranged from 0.83 cm² to 155.59 cm². Mean scar surface area measurement was 13.83 cm² (SD 23.06) for observer A and 13.59 cm² (SD 23.31) for observer B. Adjusted interobserver CV for trained observers is estimated as 5.59%, with corresponding LoA = 0 ± 0.15 x mean surface area. Interobserver ICCs were 0.99–1.00.

Conclusion: This 3D technology with a depth sensor for measuring scar surface area provides valid and reliable data and thereby surpasses expensive and time-consuming 3D cameras.

1. Introduction

Burn injuries are considered to be one of the most common types of traumatic injuries [1,2]. Throughout the last decades, the survival rate among these burn patients has increased significantly due to major improvements in burn care [3]. Because more people nowadays survive a severe burn injury, there are more survivors with extensive scars. Therefore, scar formation has become an important outcome of burn care [4]. Scar formation may lead to aesthetic, functional, social, and psychological problems, which consequently may influence patients’ health-related Quality of Life [5,6].

Besides subjective scar assessment tools, such as Vancouver Scar Scale (VSS) [7,8] and ‘Patient and Observer Scar Assessment Scale’ (POSAS) [9–11], objective scar measuring instruments are of interest in daily practice [4,12]. Objective scar quantity assessment, through scar surface area or scar volume measurement, enables quantification of scar formation over time, comparison of outcome measures, and evaluation of scar treatment efficacy [13]. The surface area measurement could potentially be used to assess hypopigmented, hyper-pigmented, or scar contraction. Scar volume measurement could potentially be used to evaluate pathologic scarring, such as hypertrophic or atrophic scars and keloids.

The most basic method of surface measurement is calculating length times width while using a ruler. However, digital innovation has led to scar surface area measurement by three-dimensional (3D) imaging. The advantage of 3D imaging is that it also takes body curvature into account. Therefore, 3D scar surface area measurement is more valid and reliable than measurements with 2D techniques [14–17]. Devices range from stereo photogrammetry such as Vectra H1 (Canfield Scientific, Fairfield, NJ, USA) [18,19] and 3D Lifeviz™ (Quantiﬁcare S.A., Sophia Antipolis, France) [15] to (laser-assisted) three-dimensional imaging systems such as Artec Eva (Artec Group, Luxembourg, Luxembourg) [14]. These systems are the most advanced products in their product range. However, very advanced devices often go along with high costs ranging from € 8.000 – 40.000 [20]. Moreover, these devices are time-consuming since the analysis of the 3D image is not performed directly on the cameras themselves. The images are mostly analyzed afterward on a computer with additional software [14,19]. These high costs and time-consuming procedures make it less feasible in a clinical setting.

To overcome these limitations, 3D imaging techniques develop towards other depth sensors. These depth sensors are low-cost (range € 500–800) and low definition or high definition image quality devices, which are attached to or integrated into mobile devices. They are therefore easily applicable in a clinical setting [21–23]. Analysis of 3D images is performed through real-time mobile applications and can be performed within a one-minute procedure [24,25]. In this clinimetric assessment, we evaluate the validity and reliability of a low definition depth sensor system on scar surface area measurement, to evaluate its potential as an objective scar measurement instrument [26].

2. Patients and Methods

2.1. Patients and volunteers

The Medical Ethics Review Committee (MERC) of Amsterdam University Medical Centers (Amsterdam UMC), location VUmc classified this study as not subject to the Medical Research Involving Human Subjects Act (WMO). Therefore, no further assessment by the MERC was needed. The principles outlined in the Declaration of Helsinki were followed. Oral informed consent was obtained from 20 volunteers and 36 patients which were 18 years of age or older. Staff members volunteered for the validity measurements. For reliability assessment, patients were included from the outpatient burn scar clinic of the Red Cross Hospital (RKZ), Beverwijk, and the scar outpatient clinic of the Department of Plastic, Reconstructive, and Hand surgery of the Amsterdam UMC, location VUMc in Amsterdam. All types of scars were eligible. Scars were classified into three categories: hypertrophic scars, keloids, or normotrophic scars. Hypertrophic scars were defined as scars with a prolonged period of collagen accumulation up to 6–12 months, resulting in a scar that is elevated and thickened, and is very stiff. Erythema will be prolonged and the scar is often intensely pruritic or even painful [27]. Keloids are defined as scars, which unlike hypertrophic scars behave more like tumors in that growth can occur even years after the original injury and extend far...
beyond the confines of the original scar [27]. Patient- and scar characteristics are listed in Table 1.

### 2.2. Three-dimensional image capturing

3D image capturing was performed with the Occipital© (San Francisco, CA, USA) Structure Sensor, a depth sensor (Fig. 1). The device was attached to an Apple© (Cupertino, CA, USA) iPad Air 2. The sensor casts thousands of invisible infrared dots onto an object and calculates the distance between the sensor and the infrared dot on the object to create a 3D image irrespective of curvature. The depth sensor is aligned with the build-in iOS camera thereby creating a colored live 3D image in the software application (3DU HealthCare). Thereafter, the software uses this 3D image to calculate the requested surface area of the object. The Structure Sensor has a VGA resolution of 640 x 480 pixels, its precision ranges from 0.5 mm to 3.0 mm on respectively 40–350 cm distance and it has a field of view of 58 x 45 degrees (horizontal x vertical) [26]. For the fact sheet of the Occipital© (San Francisco, CA, USA) Structure Sensor see appendix 1.

### 2.3. Real-time mobile application measurement

Real-time image capturing and calculations on the 3D image was performed using the 3DUniversum (Amsterdam, The Netherlands) (3DU) Healthcare iOS application [28]. The application is suitable on both iOS and Android platforms and analyzes images from Structure Sensor, Google Tango, and Intel RealSense depth sensors. The technology allows users to capture images, view and analyze surface area by the circumference and depth of the object. Additional software measurement data are color and volume measurement. Borders were drawn manually using the iPad touchscreen. To improve manual alignment object enlargement was performed on the device screen. Image capturing and analyzing of objects are performed in real-time and within a one-minute standardized procedure [28] (Fig. 1).

### 2.4. Image capturing procedure

To achieve standardized three-dimensional images, the image capturing procedure was performed perpendicular to the sticker or scar. The device is slowly pulled back until it meets the minimum threshold distance of 60 cm, thereby automatically starting the 3D image capturing process. Each object was captured in one keyframe. One keyframe is defined as an image capturing procedure in which the entire object is visible and captured within one 3D image. Overlapping keyframes from different angles, particularly used in circular scarring patterns, were not taken into account to diminish artifacts. Despite of the keyframe setting, the depth sensor always takes body curvature into account. Image capture was performed in a pre-arranged room with consistent lighting conditions. Patients had to remain
motionless for a few seconds to diminish image-capturing artifacts. Observers received instructions from colleagues with experience in 3D imaging of scars. Moreover, observers had access to the online device manuals of the Occipital® (San Francisco, CA, USA) Structure Sensor and, had support of 3DUniversum (Amsterdam, The Netherlands). All users were familiar with the use of the Apple® (Cupertino, CA, USA) iPad Air 2.

2.5. Validity procedure

Stickers with predefined dimensions were used to act as a gold standard measurement instrument29. Twenty volunteers permitted to apply four stickers of various sizes on their ankle (8.01 cm²), hand (16.33 cm²), forearm (44.42 cm²), and chest (77.70 cm²) to represent increasing scar surface area. A sample of volunteers provides measurement variability through difference in body size and curvature. Stickers are imaged once within a single-observer setting (D.R.) resulting in twenty 3D images per sticker and a total of eighty images. These eighty images were analyzed by both observer A (D.R.) and observer B (M.D.) independently using the device touchscreen. The validity image capturing procedure and flowchart are shown in Figs. 2 and 3.

2.6. Reliability procedure

Thirty-six patients with hypertrophic scars, keloids and non-motrophic scars who visited the outpatient clinic of the RKZ and Amsterdam UMC, location VUMc were considered eligible and gave oral informed consent. Following the pre-defined image capturing procedure, scars were imaged by two observers (D.R. and M.D.) separate from each other. Both 3D images were analyzed by both observers independently. This provided four surface area measurements per scar. The reliability image capturing procedure is shown in Fig. 4 and the flowchart for the reliability study in Fig. 5.

2.7. Statistical analysis

Data were analyzed using SPSS 24.0 (SPSS Inc. Chicago, USA). Descriptive statistics were performed on patient- and scar...
characteristics. Validity is expressed as mean sticker surface area measurements (cm$^2$) and mean differences between absolute sticker size and the mean sticker surface area measurements (cm$^2$). Boxplots illustrate the variance of measurement error per sticker for each observer independently. In addition, coefficients of variation (CV) were calculated. CV displays the percentage of one standard deviation (SD) in correspondence to the mean. A lower coefficient of variation represents a more accurate measurement instrument.

$$CV_{stickert1} = \frac{SD_{stickert1}}{Mean_{stickert1}} \times 100\%$$

Reliability is displayed through mean scar surface area measurements (cm$^2$), mean measurement difference inter- and intra-observer scar surface area measurements (cm$^2$), intra-observer and inter-observer coefficient of variation (CV),

Fig. 3 – Validity - Flowchart of the scanning procedure. flowchart of the scanning procedure for validity assessment of the camera and 3D software.

Fig. 4 – Reliability – Scar surface area measurement output of 3D program A) Manual annotation of a burn scar on the right cheek. Red line defines the manual boundaries for surface area measurement. B) A cut-out of the annotated scar area and presented 3D software output. The software provides data on scar surface area, volume and perimeter based on annotation and curvature.
limits of agreement (LoA), and Intraclass Correlation Coefficients (ICC). To account for an increase in observer measurement difference (cm²) alongside the increase in scar surface area (cm²), surface area measurements were 10 log-transformed to approximate a normal distribution. Transformed data were used to calculate standard error of measurement (SEM). SEM was transformed back to calculate CVs and LoA. CV is formulated as SEM * 100%. A Bland-Altman plot was made in which LoA were calculated as a function of the mean scar area. The LoA give an indication of the absolute agreement between observers based on 95% percent of surface area measurement differences between observers. ICC is a ratio of variance components of a random-effects model. ICC expresses the correlation of scar measurements within (ICC_intra) and between (ICC_inter) observers and ranges from 0.00 to 1.00. A value of 0.00 implies no correlation, whereas an ICC value of 1.00 indicates an excellent correlation. All ICCs were calculated on 10 log-transformed data.

Results are presented as an analysis of complete data, accompanied by an optimistic analysis in which data are used which could be obtained after training of observers. In this analysis, outliers expected to be due to clinical interpretation differences in the image capturing, protocol are excluded. We examined outliers with more than 20% interobserver measurement differences.

3. Results

3.1. Patients and scar characteristics

A total of 69 scars on 36 patients were included. All patients from the Amsterdam UMC presented with keloids, which were caused by trauma or surgery. At the Red Cross Hospital, mainly normotrophic scars (18/26 patients) and hypertrophic scars (11/26 patients) due to burns were included. Patient and scar characteristics are shown in Table 1.

3.2. Validity

Sticker surface area measurement showed small measurement errors. The CV for observer A was 0.62%, 1.53%, 1.67%, and 1.06% for ankle, hand, forearm, and chest measurements respectively. CV for observer B was 0.75%, 0.92%, 1.19% and 1.02% in the same sticker order. Results are shown in Table 2.
The boxplots in Fig. 6A and B show the distribution of sticker measurements for observers A and B separately. These figures illustrate an increase in measurement difference according to the increase in object size. Observers A and B show similar patterns due to the use of a standard image capturing protocol.

### 3.3 Reliability

Mean scar surface area ranged from 0.83 cm\(^2\) to 155.59 cm\(^2\). Table 3 shows that the mean scar surface area measurement was 13.83 cm\(^2\) (SD 23.06) for observer A and 13.59 cm\(^2\) (SD 23.31) for observer B. Intra-observer CV was 6.23% for observer A and 2.06% for observer B, indicating a difference in measuring accuracy. Inter-observer CV on the same scar were 8.11% (image 1) and 8.66% (image 2), indicating a discrepancy in clinical interpretation between observers. In addition, inter-observer CV on different images was 8.77%. This indicates that no significant additional measurement error is due to the image capturing device and image capturing procedure.

Seven outliers differing more than 20% between inter-observer measurements (20.39 – 45.78%) were examined. Four of these outliers were caused by interobserver differences in clinical interpretation of scar borders. Three outliers were based on the unknown intra-observer differences on the same 3D image. Adjusted analyses, excluding the seven outliers, resulted in an intra-observer measurement error of 4.07% (observer A) and 1.88% (observer B). Corresponding with an improved adjusted interobserver CV improved from 8.77% to 5.59%.

Limits of agreements were calculated for the interobserver variation of measurement A2 versus B1: LoA = 0 ± 0.24 x mean surface area (Fig. 7a). This implies that 95% of all scar surface area measurements in general, image capturing error, and interobserver variability will range 24% above or below the actual scar surface area. Adjusted limits of agreement improved: LoA = 0 ± 0.15 x mean surface area (Fig. 7b). Since a significant part of the measurements concerned small scars, an additional Bland and Altman plot was performed on scar surface areas smaller than 5 cm\(^2\). LoA for

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**Table 2 – Results for validity measurement using stickers with predefined dimensions.** Mean surface area displays the mean measurement of twenty stickers per body part on twenty healthy volunteers. Mean difference area displays the difference between true sticker size and the mean surface area. Coefficient of Variation (CV) displays the percentage of one standard deviation (SD) in correspondence to the mean measurement per sticker. A = observer A. B = observer B. CV = (SDsticker / Meansticker) * 100%.

<table>
<thead>
<tr>
<th>Sticker size</th>
<th>Ankle (n = 20)</th>
<th>Hand (n = 20)</th>
<th>Forearm (n = 20)</th>
<th>Chest (n = 20)</th>
</tr>
</thead>
<tbody>
<tr>
<td>cm(^2)</td>
<td>8.01</td>
<td>16.33</td>
<td>44.42</td>
<td>77.70</td>
</tr>
<tr>
<td>Mean surface area (A)</td>
<td>cm(^2) (SD)</td>
<td>7.97 (0.05)</td>
<td>16.59 (0.25)</td>
<td>44.98 (0.74)</td>
</tr>
<tr>
<td>Mean difference area (A)</td>
<td>cm(^2) (SD)</td>
<td>0.04 (0.05)</td>
<td>-0.26 (0.25)</td>
<td>-0.56 (0.74)</td>
</tr>
<tr>
<td>Mean surface area (B)</td>
<td>cm(^2) (SD)</td>
<td>8.01 (0.06)</td>
<td>16.44 (0.15)</td>
<td>45.62 (0.53)</td>
</tr>
<tr>
<td>Mean difference area (B)</td>
<td>cm(^2) (SD)</td>
<td>-0.01 (0.06)</td>
<td>-0.11 (0.15)</td>
<td>-0.20 (0.53)</td>
</tr>
<tr>
<td>CV(_{observerA}) (%)</td>
<td>0.62</td>
<td>1.53</td>
<td>1.67</td>
<td>1.06</td>
</tr>
<tr>
<td>CV(_{observerB}) (%)</td>
<td>0.75</td>
<td>0.92</td>
<td>1.19</td>
<td>1.02</td>
</tr>
</tbody>
</table>

---

**Fig. 6 – Validity – Boxplot of measurement difference to absolute sticker surface area.** boxplots showing variance between measured sticker areas made by observer A and observer B compared to absolute sticker area (cm\(^2\)). Body location on the x-axis represent four sticker locations and corresponding sizes; ankle (8.01 cm\(^2\)), hand (16.33 cm\(^2\)), forearm (44.42 cm\(^2\)) and chest (77.70 cm\(^2\)). Y-axis displays the measurement difference to absolute sticker surface area (cm\(^2\)). Each location is measured once per observer on twenty volunteers. The boxplots illustrate an increase of measurement difference according the increase in object size. Observer A and B show similar patterns due to the use of a standard scanning protocol.
In this study, we showed that scar surface area measurement could be valid and reliably performed by using three-dimensional technology on mobile devices. The technology provides an affordable and feasible system that can be used quite easily in scar care worldwide. In absence of a clinical gold standard, validity was expressed as measurement error (cm$^2$) and coefficient of variation (%) around stickers with predefined dimensions. These sticker dimensions and locations represented increasing scar surface area and differences in body curvature. The latter could interfere with calculations on the 3D image. Overall mean measurement differences and corresponding CV were found to be very low and thereby illustrate the accuracy of this 3D depth sensor in combination with the used software on scar surface area measurement. In addition, the tablet enlarging ability of the 3D image might have led to the more precise delineation of sticker borders and therefore more accurate measurements.

Reliability was assessed in a clinical two-observer setting. Data was presented as mean surface area (SD) and mean difference (SD). Intra- and interobserver reliability was expressed as coefficient of variation, limits of agreement, and Intraclass Correlation Coefficients. For scars, the coefficient of variation between observers ranged from 8.11% to 8.77%, which was higher than found in the validity assessment of stickers with clear boundaries (0.62–1.67%). Border interpretation is more challenging in scars than with the use of white stickers. This is especially the case in normotrophic scars since these colors merge with non-traumatized skin tones, whereas keloids or hypertrophic scarring show a clearer transition. Software upgrades, device resolution, standardized lighting conditions, and the zoom function might decrease these measurement errors. Increased measurement error is to be expected in untrained personnel and the lack of a clear standard operating procedure.

In this study, data are presented both as an analysis of complete data and after the exclusion of seven outliers which could be avoided by appropriate training of the observers. The training should include a set of challenging scars, with a two-folded aim: to have some practice with image capturing and analysis of scars and to make clear agreements on clinical interpretation of scar borders. Before implementing it in daily practice, we recommend health care providers train personnel with such training to prevent these kinds of measurement errors.

Both intra-observer and interobserver ICC were 0.99 or higher, indicating excellent reliability. ICCs with a minimum of 0.7 are considered acceptable for research purposes and ICCs of at least 0.9 are required in a clinical setting [29]. The high ICC, however, can be explained by the wide variation of scar sizes in this study, ranging from 0.83 cm$^2$ – 155.59 cm$^2$.

### Table 3. Results for reliability. Reliability measurement on 69 scars within a multi-observer setting. Scar surface area displays the mean (SD) of measurement per observer or between observers. Standard error of Measurement (SEM) expresses the error margin around the measurement. Due to skewed distribution, data were first 10 log-transformed to calculate SEMs, then SEMs were transformed back and presented in this table.

<table>
<thead>
<tr>
<th>Observer</th>
<th>Scar surface area (mean, SD) cm$^2$</th>
<th>Measurement difference (mean, SD) cm$^2$</th>
<th>SEM cm$^2$</th>
<th>ICC</th>
<th>ICC cor.</th>
</tr>
</thead>
<tbody>
<tr>
<td>A1</td>
<td>13.83 (3.06)</td>
<td>0.11 (0.05)</td>
<td>0.06</td>
<td>0.99</td>
<td>0.99</td>
</tr>
<tr>
<td>A2</td>
<td>13.89 (3.31)</td>
<td>0.08 (0.07)</td>
<td>0.06</td>
<td>0.99</td>
<td>0.99</td>
</tr>
<tr>
<td>B1</td>
<td>13.71 (2.46)</td>
<td>0.02 (0.02)</td>
<td>0.06</td>
<td>0.99</td>
<td>0.99</td>
</tr>
<tr>
<td>B2</td>
<td>13.70 (2.81)</td>
<td>0.00 (0.00)</td>
<td>0.06</td>
<td>0.99</td>
<td>0.99</td>
</tr>
</tbody>
</table>

4. Discussion

Scars < 5 cm$^2$ slightly increased: LoA = 0 ± 0.27 x mean surface area indicating higher measurement error on smaller scars (Fig. 8a). Adjusted LoA were 0 ± 0.16 x mean surface area (Fig. 8b). ICCs were high and ranged from 0.99 to 1.00 for both intra-observer and interobserver reliability.
Fig. 7 – a. Reliability – Bland and Altman Plot Interobserver Agreement – Overall Limits of Agreement. a. A Bland and Altman plot presenting the interobserver agreement between two observers (measurement A2 versus B1). Due to expected skewed distribution of scar surface area measurements, the data was natural log-transformed to approximate a normal distribution and obtain a regression coefficient for limits of agreement. The limits of agreement indicate the absolute agreement between observers based on 95% percent of surface area measurement differences between observers. The interobserver agreement is based on the mean surface areas of scan 2 analyzed by observer A and scan 1 analyzed by observer B (see Fig. 5).

b. Reliability – Bland and Altman Plot Interobserver Agreement – Overall Adjusted Limits of Agreement. b. A Bland and Altman plot adjusted for outliers, presenting an interobserver agreement between two observers (measurement A2 versus B1). Due to expected skewed distribution of scar surface area measurements, the data was natural log-transformed to approximate a normal distribution and obtain a regression coefficient for limits of agreement. The limits of agreement indicate the absolute agreement between observers based on 95% percent of surface area measurement differences between observers. The interobserver agreement is based on the mean surface areas of scan 2 analyzed by observer A and scan 1 analyzed by observer B (see Fig. 5).
The influence of device, software, and observer measurement error is small compared to the variation in scar size and therefore has little influence on the correlation.

Limits of Agreement (LoA) project a more comprehensive measurement outcome since it takes the wide range of object sizes into account. Since the LoA showed that the

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Fig. 8 – a. Reliability – Bland and Altman Plot Interobserver Agreement – Limits of Agreement for Small Scars (< 5 cm²)

a. A Bland and Altman plot presenting an interobserver agreement between two observers (measurement A2 versus B1). The interobserver agreement is based on the mean surface areas of scan 2 analyzed by observer A and scan 1 analyzed by observer B (Fig. 5) on scar surface areas less than 5 cm².

b. Reliability – Bland and Altman Plot Interobserver Agreement – Adjusted Limits of Agreement for Small Scars (< 5 cm²)

b. A Bland and Altman plot adjusted for outliers, presenting an interobserver agreement between two observers (measurement A2 versus B1). The interobserver agreement is based on the mean surface areas of scan 2 analyzed by observer A and scan 1 analyzed by observer B (Fig. 5) on scar surface areas less than 5 cm².
measurement error increased with increasing scar size, data were log-transformed to approximate a normal distribution. LoA can be extrapolated toward a clinical setting and used in comparison to other instruments.

The adjusted 3D depth sensor LoA = 0 ± 0.15 x mean surface area. This implies that 95% of all scar surface area measurements in a clinical setting, adjusted for scar size, image capturing error and interobserver variability will range 15% above or below the scar surface area. LoA of scars < 5 cm² slightly increased to LoA = 0 ± 0.16 x mean surface area. These LoA found in this study are considered appropriate for clinical use compared with other 3D cameras [14,15]. A study by Rashaan et al., 2016 [14] assessed wound surface areas with the Artec 3D camera and Stekelenburg et al., 2013 [15] performed scar surface area measurements with stereo photogrammetry. The mean scar surface area of this study was 13.71 cm², whereas a mean surface area of 20.58 cm² and 59.65 cm² were found in the two compared studies respectively. Limits of Agreement of the Artec MHT™ 3D camera was LoA = 0 ± 0.17 x mean surface area [14] whereas stereo photogrammetry showed a LoA = 0 ± 0.19 x mean surface area [15]. In general, this mobile 3D technology with a depth sensor is not inferior to counterparts such as the Artec MHT™ 3D camera or stereo photogrammetry regarding scar surface area measurement. Since mobile devices and 3D analyzing software provide object zooming function it provides more accurate and reliable measurements of the 3D image. Alongside less time-consuming data analysis and lower costs, this demonstrates the strengths of these 3D depth sensors and its mobile software applications in comparison to 3D cameras mentioned above.

This mobile 3D technology does have limitations. The difficulty of capturing wounds or scars on highly curved body parts is a well-known problem within 3D imaging techniques [14–16,30–33]. Body curvature remains a bigger challenge than with 3D techniques mentioned above, due to a lower resolution and thus fewer data points. High curvature and circular objects require multiple keyframes, which, to date, show too much overlap and therefore higher measurement error. Especially, scars on noses, ears, and fingers could not be imaged. This is unfavorable, since a substantial part of the patients that visit the outpatient clinics of the Red Cross Hospital have scars on their face and hands.

Ongoing this 3D software innovation is expected to surpass these limitations as they have already shown its accuracy and reliability within one keyframe. Automatic annotation of object borders based on color segmentation is currently in development. It might account for more reliable scar assessment by surpassing clinical interpretation and increasing usability in a clinical setting. A High Definition depth sensor that is currently on the market [34] might be more capable to measure the volume of a scar, which is an important clinical outcome parameter. Moreover, it might measure the extent of circular scar patterns or pigmentation changes within a scar more accurately. Alongside the integration of 3D depth sensors in current smartphones, the next innovations in healthcare applications are on the horizon.

## 5. Conclusion

3D technology by using affordable and feasible 3D depth sensors mounted on mobile devices provides instant object data in a clinical setting and thereby surpasses expensive and time-consuming 3D cameras. With the ongoing software innovation and integration of depth sensors in mobile devices, the implementation of this 3D technology for scar surface area measurements can be considered worldwide.

### Funding

This research project was funded by the Dutch Burns Foundation, project number 16.02.

### Disclosure

Authors S. Karaoglu and T. Gevers are founders of 3DUuniversum (Amsterdam, The Netherlands) (3DU) and supervised the development of the 3DU Healthcare iOS application.

### Declaration of Competing Interest

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests, Authors S. Karaoglu and T. Gevers are founders of 3DUuniversum (Amsterdam, The Netherlands) (3DU) and supervised the development of 3DU Healthcare iOS application. 3DU had no interference with the study data.

### References

[10] van der Wal MB, et al. The modified patient and observer scar assessment scale: a novel approach to defining...


