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A 3D reconstruction method of the body envelope from biplanar X-rays: Evaluation of its accuracy and reliability

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A B S T R A C T

The aim of this study was to propose a novel method for reconstructing the external body envelope from the low dose biplanar X-rays of a person. The 3D body envelope was obtained by deforming a template to match the surface profiles in two X-rays images in three successive steps: global morphing to adopt the position of a person and scale the template's body segments, followed by a gross deformation and a fine deformation using two sets of pre-defined control points. To evaluate the method, a biplanar X-ray acquisition was obtained from head to foot for 12 volunteers in a standing posture. Up to 172 radio-opaque skin markers were attached to the body surface and used as reference positions. Each envelope was reconstructed three times by three operators. Results showed a bias lower than 7 mm and a confidence interval (95%) of reproducibility lower than 6 mm for all body parts, comparable to other existing methods matching a template onto stereographic photographs. The proposed method offers the possibility of reconstructing body shape in addition to the skeleton using a low dose biplanar X-rays system.

Keywords:

Body envelope
Geometric modeling
Biplanar X-rays
Reproductibility
Joint center

1. Introduction

3D body scanning technologies are becoming accessible making it possible to easily obtain 3D body shape (Daanen and Haar, 2013; Liu et al., 2015). For instance, Park et al. (2015) recently reported that full child shape could be measured with a good accuracy using low-cost depth cameras and a statistical body shape model. To create personalized human model with an articulated linkage for re-posturing or movement analysis, an accurate estimation of internal joint centers is required. Anatomical landmarks (ALs) generally need to be manually identified on the body mesh so as to estimate joint centers. Not only the palpation of ALs requires specific training, but this is also a very time consuming process. Recently, methods were proposed to automatically locate ALs on body shape scans by non-rigid model fitting (Yamazaki et al., 2013) or using a statistical shape model (Reed et al., 2015). But relationships between ALs and joint centers are missing in general due to lack of in vivo data from human

beings (Reed et al., 1999). Therefore there is a need to collect data which contains both external body surface and internal skeleton geometries.

Recent development of low dose bi-planar X-rays (BPXr) (Dubousset et al., 2010) allows the 3D reconstruction of subjects' skeleton in an upright posture (Mitton et al., 2006; Humbert et al., 2009; Chaibi et al. 2012; Quijano et al., 2013; Aubert et al., 2014) while the skin contour is also visible on the radiographs. Using this technology, Sandoz et al. (2010) proposed a method for reconstructing the external body envelope. Eleven template meshes representing the 11 different body segments were morphed on skin contour and then combined to obtain an entire reconstruction of the body. However the method was only validated on a few male subjects and the separate reconstruction of different body segments was a time-consuming process requiring a work day per subject. Global morphing of a template mesh of the whole body envelope would allow reduction of the reconstruction time.

The aim of the present study was to propose a new method for reconstructing the 3D external body envelope from BPXr as well as a procedure for its validation. A 3D surface mesh template was morphed to match body surface profiles in two orthogonal BPXr. The method was validated by comparing the reconstructed surface

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to the locations of up to 172 radiopaque markers uniformly attached to the body skin surface on 12 participants.

2. Material and methods

2.1. Data collecting

After approval of the ethical committee (CPP 06036) and written informed consent, 12 volunteers (6 males, 6 females, average age: 26.8 ± 3.1 , average BMI: $23.6 \pm 2.9 \text{ kg/m}^2$) participated in data collecting. 172 radio-opaque markers (150 of $\varnothing 2 \text{ mm}$ and 22 of $\varnothing 5 \text{ mm}$) were uniformly placed on the participants' skin from head to foot excluding the arms, including a hair cap with 10 markers. Participants were asked to adopt a standard position (Steffen et al., 2010). Two calibrated orthogonal BPXr (EOS Imaging, Paris, France) were simultaneously acquired in 18 s (exposure parameters: 0.5 mGy, face radiograph: 90 kV, 190 mA, profile: 105 kV, 250 mA).

2.2. Envelope reconstruction method

The 3D body envelope was obtained by deforming a template to match the surface profiles in two BPXr. The template matching procedure included three successive steps: global morphing to adopt the participant's position and scale the template's body segments, followed by a gross deformation, and a fine deformation to fit the template surface profiles to the X-rays. As the arms exceeded the acquisition area, only an armless mesh was reconstructed.

Two reference templates were created corresponding to an average height male (1.75 m, 70 kg) and female (1.68 m, 60 kg) in the EOS standing position whose BPXr were available from a previous study. A male (12,035 nodes, 24,068 triangles) and female (12,243 nodes, 24,484 triangles) artist representation from the MakeHuman™ Open Source application 1.0.1 (The MakeHuman team, <https://www.makehuman.org>) were initially used and manually deformed to match the two reference subjects with help of the Blender software v.2.72b (Blender Foundation, <https://www.blender.org>). Three sets of control points were pre-defined on the two templates (Fig. 1a) and successively used for the body envelope reconstruction for each participant from his/her BPXr.

The first set of control points contained the location of the crotch and 10 joint centers in order to scale the body envelope template onto the participants' radiographs. The corresponding points of a target participant had to be manually identified on the radiographs by an operator (Fig. 1b). Then, they were used to globally morph the template to the position of the participant by a moving least square algorithm (MLS) (Cuno et al., 2007) providing an "as-rigid-as-possible" deformation of the template model (called 'global morphing'). The deformed template was automatically projected on the two radiographs (Fig. 1c) together with all control points.

The second set included 49 skin points, used to locally deform the envelope by using MLS in the second step (called 'first deformation'). They were disposed on the body parts of high shape variation between individuals such as the back of the calf and the waist which depend on subjects' weight and muscle structures. By

manually moving each of these skin control points, the template could be deformed to grossly match the contour from the front and profile radiographs. A pre-customized surface model called "initial solution" could thus be achieved (Fig. 1d).

Finally an additional set of 71 points were proposed for a fine deformation of the template (called 'fine deformation'). They were once again located depending on human shape variations which need a refinement such as the buttocks curvature and the abdomen bulge. Each of these 71 points could be dragged with help of the mouse to control the local deformation of a body region using a kriging algorithm (Trochu, 1993) (Fig. 1e).

2.3. Assessment of the reconstruction method

2.3.1. Reference points

Among 172 radio-opaque spherical markers stuck to the body surface, on average 123 (± 13) markers per participant was automatically detected on the radiographs using a pixel-wise comparison of the image with a reference pattern (i.e. cross-correlation). They were used as reference points to evaluate the accuracy and reproducibility of the reconstructed surface. Accuracy of the markers detection method was assessed with an average error of 0.045 mm (2SD < 0.06 mm). Because of the markers' size, the marker skin contact point was defined as the marker center translated by the marker's radius in the direction normal to the external surface of the reconstructed envelope. As the hair cap with markers worn by the participants was not directly in contact with the skull, an offset of 5 mm (representing the average distance between skull and hair cap) was used.

2.3.2. Corrected envelope

To evaluate the error map, a reference envelope was obtained using the radio-opaque markers as new control points by deforming the reconstructed envelope, resulting in a corrected envelope passing through these reference points.

2.3.3. Bias and reproducibility

An inter-intra operator variability study was performed to evaluate the error due to manual adjustments of control points. Three operators performed the reconstruction of 12 participants three times. The instruction given to the operators was to match the reference template to the two radiographic profiles as closely as possible. All operators were trained with the help of an experienced person using a sample set of six subjects' radiographs of different anthropometric dimensions which were not used for the method's evaluation.

Bias and reproducibility variance of the method were assessed as recommended by the International Organization for Standardization (ISO 5725-1-2).

- For each reference marker that was successfully detected, the bias was defined as the average signed point-to-surface distance between the reference radio-opaque markers and the reconstructed surfaces obtained from all operators.
- For each participant i , the reproducibility variance (S_{Ri}^2) expressed the variability in the mesh nodes distribution of reconstructed envelopes that is due to the variability among operators. It involved the interoperator (S_{Ri}^2) and intraoperator

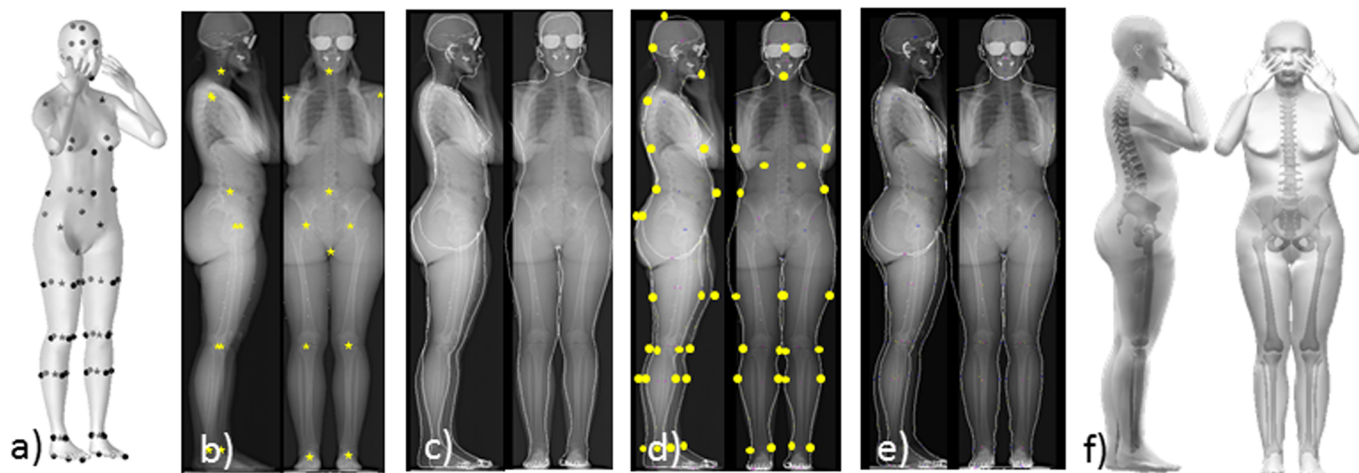


Fig. 1. Reconstruction steps: (a) Prerequisite: allocation of control points on the template. Control point used for the global morphing (11 asterisks) and first deformation (49 circles) steps are represented on the female template. (b) Manual identification of the crotch and 10 joints centers for the global morphing step. (c) Projection and global morphing of the model on the front and profile radiographs: the template adopts the position of the person and body segments are scaled onto the participant's radiographs. (d) Projection and manual positioning of 49 surface points on the skin profiles for the first gross deformation step. (e) Adjusted model after the fine adjustment step. (f) 3D reconstruction of subject-specific body envelope. Bones reconstructions were added as an illustrative purpose of the possibility to obtain the external and internal geometry of the body.

variances (S_{ri}^2):

$$S_{Ri}^2 = S_{Li}^2 + S_{ri}^2$$

where S_{ri}^2 was the mean of the variances of three reconstructions performed by each operator (intraoperator variance); S_{Li}^2 quantified the variance of the mean reconstructions obtained between operators (interoperator variance)

Finally the global reproducibility variance (S_R) over the n participants ($n=12$) was calculated as follow:

$$S_R = \sqrt{\frac{\sum_{i=1}^n S_{Ri}^2}{n}}$$

A confidence interval at the 95% confidence level (95% CI) was considered as $2 * S_R$ as an indicator of the global uncertainty.

3. Results

The bias was assessed for each subject using detected markers and was comprised between -6.7 and 2.3 mm (Table 1). Mean (respectively max) marker-to-surface absolute distance was 4.9 (9.0) mm. 95% CI of reproducibility was smaller than 5.8 mm (Table 1). Higher accuracy errors were distributed along the lateral parts of the thorax and the abdomen compared to the middle area (Fig. 2a). Regarding reproducibility, local maxima did not exceed 20 mm (Fig. 2b).

4. Discussion

A method allowing the reconstruction of the body envelope from BPXr by matching a 3D body shape template was proposed in this study. Reconstruction errors were comparable to other previous studies (Lin and Wang, 2012; Lee, 2000; Zhu et al., 2013), with mean absolute values ranging from 8 to 13 mm. Moreover with BPXr, both the external envelope and internal skeleton of the whole body become accessible from a same set of images (Fig. 1f) while other medical imaging techniques such as MRI (Bauer et al., 2007; Cheng et al., 2000; Mungiole and Martin, 1990) or CT-Scan (Huang and Suarez, 1983; Pearsall et al., 1994) can only generate partial reconstructions due to high cost, long post-processing and/or exposure to radiation. In addition this method allows straight-forward identification of ALs for joint centers estimation by projecting subcutaneous bony landmarks on the envelope surface. This approach was implemented in a preliminary study (Nérot et al., 2015) for hip joint center (HJC) prediction proposing new

predictors from the body surface in addition to commonly used ALs. Parameters such as distances between anatomical landmarks (ALs), barycenter of the body segments or elliptic shape descriptors at the section of the anterior superior iliac spine, were used as predictors. Results were comparable or even better than other existing methods which require the accurate location of some ALs based on manual palpation.

The proposed method allows for template scaling which makes it not sensitive to subject's height and weight. A very different body shape from the template may require more time to get a good fitting. In our test sample of 6 males and 6 females with different height and weight (Table 1), high errors in reconstructed envelopes were not observed for extreme height and weight participants. Work is in progress to evaluate the method for an elderly population whose overall posture and surface contour may be very different from that of the template used in the present study. If necessary, additional templates and/or control points could be added for a better matching with the target subject's contour, requiring more time for the morphing process.

However, it should be mentioned that the EOS bi-planar X-ray system is limited to 34 cm in width for the radiographic acquisition. The front view of the medial and lateral contours of the torso was frequently missing even for fairly lean participants. Missing contours were approximately interpolated using visible contours yielding higher reconstruction errors. Improvement of the method could consider estimating the missing information based on statistical shape models.

Another limitation is that template matching is operator dependent and still a time consuming process. Even though the reconstruction time was drastically reduced to 20 min compared to the one proposed by Sandoz et al. (2010), matching the control points on the radiographic profiles remained mostly manual. Therefore operator training (approximately 5 h) is necessary to use the software, especially for better handling of available control points. Work is in progress for automatic contour detection from the images for a less operator dependent process.

As conclusion, the proposed method allows for simultaneously collecting the external envelope and internal skeleton on a broad population. It opens the way to a large exploration for locating joint centers from the characteristics of external body shape in addition to the traditional ALs for a more accurate prediction of internal skeleton location.

Table 1
Bias and total uncertainty (95%CI repro) per subject and body region (mm): bias was considered as the average signed point-to-surface distance (\pm standard deviation) between the reference radio-opaque markers and the reconstructed surfaces obtained from all operators; 95% CI of reproducibility (repro) (corresponding to $2 * S_R$) provides an indicator of the global uncertainty on mesh nodes distribution due to the variability among subjects (in mm).

	Subject	Height	Weight	Thorax	Abdomen	Head	Thighs	Lower legs	Feet
Men	1	1.88	84.4	1.7 ± 9.1	0.9 ± 7.1	-3.6 ± 2.3	-1.0 ± 6.0	-1.7 ± 3.9	-3.0 ± 2.6
	2	1.75	69.1	-1.8 ± 8.6	2.3 ± 11.1	-1.3 ± 3.2	-1.5 ± 5.6	-3.4 ± 4.0	-3.7 ± 2.9
	3	1.82	76.5	1.0 ± 11.3	0.8 ± 6.9	-0.8 ± 3.5	-2.5 ± 5.6	-2.6 ± 3.3	-5.3 ± 3.5
	4	1.87	88.1	0.5 ± 11.8	1.6 ± 10.3	-1.5 ± 6.7	-1.5 ± 6.1	-3.3 ± 3.6	-3.5 ± 3.3
	5	1.87	103.3	1.5 ± 10.7	0.1 ± 8.6	-2.2 ± 7.1	-3.2 ± 6.5	-4.8 ± 3.6	-6.5 ± 4.5
	6	1.75	63.2	1.1 ± 7.1	-0.8 ± 6.5	-0.8 ± 3.5	-2.7 ± 5.4	-6.7 ± 3.6	-1.5 ± 7.7
			Mean bias	0.70	0.8	-1.7	-2.1	-3.8	-3.9
		95% CI repro	5.8	4.9	4.0	2.9	3.0	3.7	
Women	7	1.74	64.3	-0.5 ± 5.7	0.2 ± 5.8	1.7 ± 6.4	-1.2 ± 3.9	-3.2 ± 2.5	-2.2 ± 3.9
	8	1.66	55.1	0.1 ± 8.5	-0.6 ± 4.4	-2.8 ± 3.8	-1.4 ± 4.0	-3.2 ± 2.4	-2.6 ± 2.8
	9	1.63	60.7	-1.9 ± 8.5	-2.2 ± 6.0	-1.1 ± 2.5	-2.4 ± 4.5	-4.2 ± 3.2	-1.3 ± 5.7
	10	1.62	74.5	-5.7 ± 5.6	-3.8 ± 6.7	-1.9 ± 2.4	-3.9 ± 4.3	-3.6 ± 2.1	-3.9 ± 2.6
	11	1.65	59.1	1.1 ± 9.9	-1.5 ± 7.8	-2.4 ± 3.2	-2.2 ± 4.6	-4.3 ± 3.1	-4.0 ± 3.8
	12	1.63	65	-1.5 ± 7.1	-1.1 ± 6.1	-2.8 ± 4.1	-2.9 ± 4.7	-3.4 ± 2.3	-3.4 ± 3.6
			Mean bias	-1.4	-1.5	-1.6	-2.4	-3.7	-2.9
		95% CI repro	1.2	5.9	4.1	3.7	3.5	4.6	
All		Mean bias	-0.4	-0.4	-1.6	-2.2	-3.7	-4.0	
		95% CI repro	0.9	5.4	4.0	3.30	3.3	4.1	

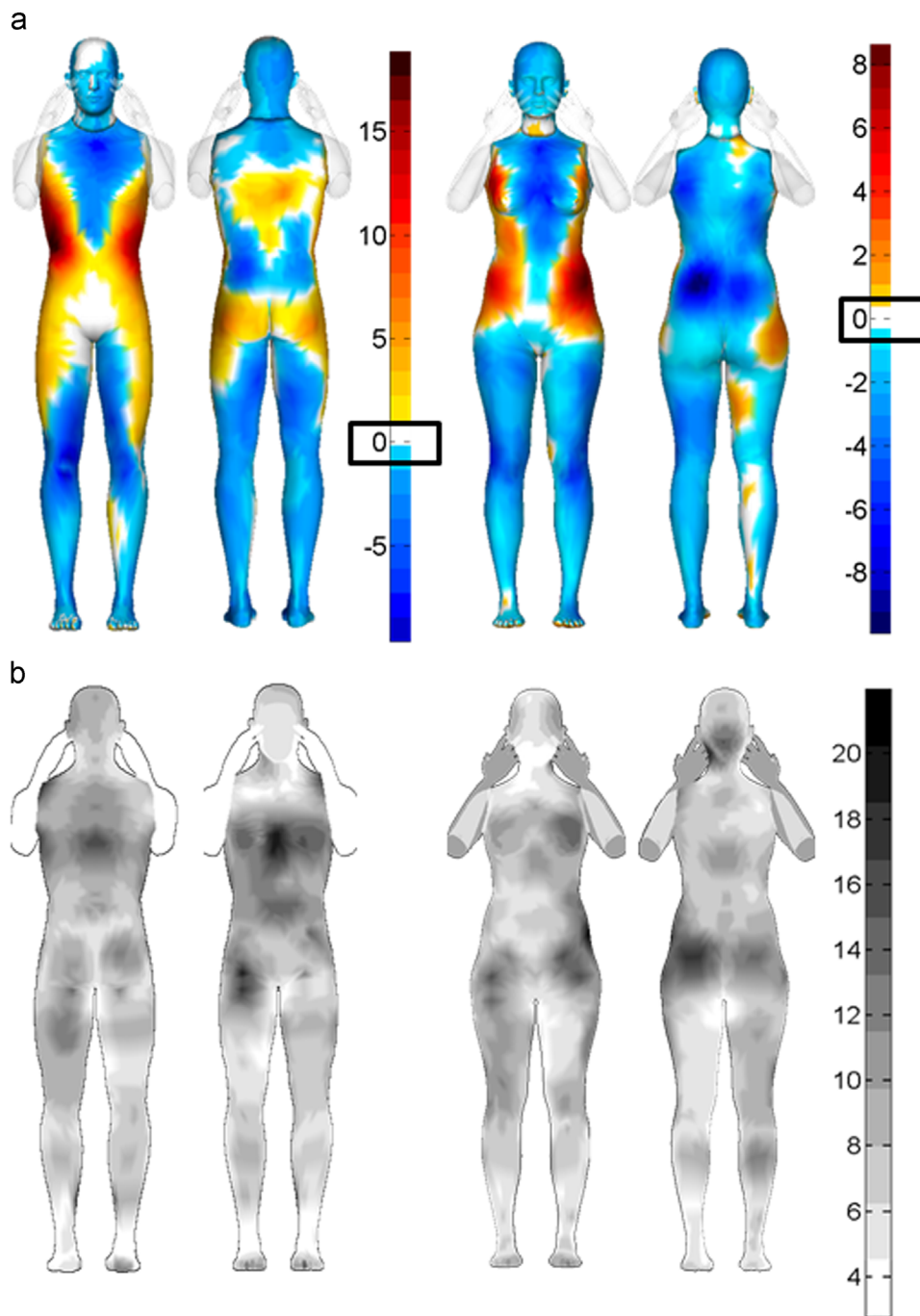


Fig. 2. (a) Average error map displayed on the templates showing the bias (signed distance between the reconstructed envelopes and the corrected envelopes used as reference) in mm for male (a) and female (b). (b) Error map showing the 95% CI of reproducibility ($2*S_R$) as an indicator of the global uncertainty on mesh nodes distribution due to the variability among subjects (in mm). Only an armless mesh was reconstructed. Generic arm meshes were afterwards merged to the reconstructed surface for visual purpose.

Conflict of interest statement

The authors have no conflicts of interest to declare.

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