From medical imaging data to 3D printed anatomical models

Thore M. Bücking, Emma R. Hill, James L. Robertson, Efthymios Maneas, Andrew A. Plumb, Daniil I. Nikitichev

1 Department of Medical Physics and Biomedical Engineering, University College London, London, United Kingdom, 2 Centre for Medical Imaging, University College London, London, United Kingdom

* Thore.Buecking.14@ucl.ac.uk

Abstract

Anatomical models are important training and teaching tools in the clinical environment and are routinely used in medical imaging research. Advances in segmentation algorithms and increased availability of three-dimensional (3D) printers have made it possible to create cost-efficient patient-specific models without expert knowledge. We introduce a general workflow that can be used to convert volumetric medical imaging data (as generated by Computer Tomography (CT)) to 3D printed physical models. This process is broken up into three steps: image segmentation, mesh refinement and 3D printing. To lower the barrier to entry and provide the best options when aiming to 3D print an anatomical model from medical images, we provide an overview of relevant free and open-source image segmentation tools as well as 3D printing technologies. We demonstrate the utility of this streamlined workflow by creating models of ribs, liver, and lung using a Fused Deposition Modelling 3D printer.

Introduction

Anatomical models have applications in clinical training and surgical planning as well as in medical imaging research. In the clinic, the physical interaction with models facilitates learning anatomy and how different structures interact spatially in the body. Simulation-based training with anatomical models reduces the risks of surgical interventions [1], which are directly linked to patient experience and healthcare costs. For example, improvement of central venous catheter insertions has been achieved by the use of anatomically and ultrasonically accurate teaching phantoms [2]. In addition, the phantoms can be used for pre-operative surgical planning, which has been shown to be beneficial in craniofacial surgery [3] and is being explored in a number of other surgical fields [4, 5]. Lastly, anatomical phantoms can be designed to mimic tissue when imaged with the modality of interest; most commonly ultrasound, Computed Tomography (CT), or Magnetic Resonance Imaging (MRI). Imaging phantoms are also important for the development of novel imaging modalities such as photoacoustics [6], or for validation of image-based biomarkers such as pore size estimation using nuclear magnetic resonance [7], where they provide controlled experimental environments.

Anatomically accurate models can be computer-generated from medical image data. CT and MRI are widely used to image biological features, ranging from whole-body imaging to
particular areas of interest such as tumours or specific parts of the brain. Depending on the imaging modality, different features can be observed and different image segmentation algorithms will be appropriate. CT pixel intensities directly correlate to tissue density. The modality thus lends itself well to segmenting structures such as bones (high density) or lungs (low density). MRI offers excellent soft tissue contrast, which, for example, enables differentiation between white and grey matter in the brain [8].

Recent advances in segmentation software have made it increasingly easy to automatically or semi-automatically extract the surface of structures of interest from three-dimensional (3D) medical imaging data. This has made it possible to generate anatomical models using a standard personal computer with little prior anatomical knowledge. At the same time 3D printers, traditionally used in industrial applications, are now available for home use thanks to low-cost desktop alternatives. This technology enables fast creation of 3D models without the need for classical manufacturing expertise.

Accessibility to 3D printers and advanced segmentation algorithms have led to an increase in use of 3D printing in medicine, which has received interest due to a multitude of potential medical applications [4, 9]. Models can be made patient-specific, and rapidly redesigned and prototyped, providing an inexpensive alternative to generic commercially available anatomical models. 3D printing thus found applications in teaching the structure of kidney [5], heart [10], and liver [11]. A number of studies have also investigated the potential of using 3D printing techniques to produce tissue-mimicking phantoms for research and teaching, with example applications in producing models of vessels [2, 12], parts of the skull [13], optic nerves [14], and renal system [15]. The process of going from medical imaging data to 3D printed models has been described for the brain [16,17], the human sinus [18], as well as from a general point of view [19], but challenges remain to make the process widely available to novice users.

In this work, we present a practical guide to creating a broad range of anatomical models from medical imaging data. In the next section, we provide an overview of the general workflow and include a table listing the relevant 3D printing technologies. We have implemented a streamlined processing pipeline on different regions of interest (ribs, liver, and lung), which will be described in the sections below. While the pipeline is illustrated using CT images, it is also applicable to other volumetric medical imaging modalities, like MRI. The workflow is broken down into three steps (Fig 1):

Image segmentation. After acquiring a medical image, structures of interest need to be segmented. Image segmentation is the process of partitioning an image into multiple labelled regions locating objects and boundaries in images. It can be used to create patient-specific, highly accurate computer models of organs and tissue. There are a number of image segmentation techniques, which each have advantages and disadvantages, but there is no single segmentation technique which is suitable for all images and applications. Basic segmentation
approaches rely on the principle that each tissue type has a characteristic range of pixel intensities. Hence, it is possible to distinguish between tissues and identify boundaries.

There is a wide range of software that is capable of performing image segmentation, ranging from multi-purpose commercial platforms with integrated physics simulations (e.g. Mimics [20], or Simpleware [21]), to open-source tools targeted to specific organs (e.g. FreeSurfer [22] for the brain). In this work, we have used the freeware software packages called Seg3D [23] and 3D Slicer [24], as they are capable of processing a range of medical imaging data. Furthermore, we provide a summary of comparable freeware software available at the time of writing in the discussion.

**Mesh refinement.** Following image segmentation, the 3D model can be further refined into a printable 3D mesh. There are a number of computer-aided design tools that can be used for this purpose and allow almost limitless mesh manipulation and refinement. However, the main reasons for such mesh post-processing of the segmentation are as follows:

Repairing: Errors and discontinuities that sometimes arise in the image segmentation and exporting process need to be repaired before printing.

---

**Fig 1. From medical image to 3D print workflow.** After the anatomical structure has been segmented (a), the resulting surface needs to be refined (b) to remove image artefacts, after which it can be 3D printed (c).

https://doi.org/10.1371/journal.pone.0178540.g001
Smoothing: Staircasing errors resulting from the resolution of the original medical image can be mitigated by smoothing the surface of the mesh model.

Appending: The segmentation will often only be one component of a final model. To convert the model into a useable form, it is often necessary to combine it with other structures or remove unneeded parts from the segmentation.

**3D printing.** There are many different 3D printing technologies available, each with their own characteristics. Here we provide an overview of the 3D printing methods, which are suitable for the creation of anatomical models and highlight their respective advantages. The relevant 3D printing technologies can be classified into three groups: extrusion printing, photopolymerisation and powder-based printing. The most common example of extrusion printing is known as Fused Deposition Modelling (FDM), which is based on melting and depositing a material via a nozzle, building the desired shape layer by layer. In photopolymerisation, liquid polymers are selectively cured, typically using UV light. Important examples are Stereolithography (SLA) and Digital Light Processing (DLP), which selectively cure a plastic in a bath. Moreover, the photopolymer can be sprayed onto the print in thin layers, where it is subsequently cured. This technique is known as Material Jetting (MJ). Lastly, in powder-based techniques, a powdered material is bound together. This can either be done using a liquid binding agent (Binder Jetting, BJ), or by fusing the particles together using heat (Selective Laser Sintering, SLS). The characteristics of these techniques are summarised in Table 1.

**Preparation of example anatomical models**

The models of the ribs and liver were segmented using Seg3D (v.2.2.1) while the lung was segmented using 3D Slicer (v.4.6). We smoothed the models using MeshMixer [27] (v.3.0) and printed all of them using Filament Deposition Modelling (FDM). All processing was done using Windows 10 as operating system.

**Image segmentation.** Seg3D v.2.2.1 was used to generate the rib model from the CT MECANIX dataset (Siemens Sensation 64, 3 mm slice thickness, 0.56 mm by 0.56 mm pixel size, 120 kV peak kilo-voltage, 100 mAs exposure) available on the OSIRIX website [28]. The ribs were segmented using thresholding, manual modification (cropping), a connected-component-filter as well as a fill-holes-filter. The segmentation was exported as a stereolithography (STL) file using the “Export Isosurface” command.

**Table 1. Overview of the most important 3D printing technologies with medical applications: Extrusion printing, photopolymerisation, and powder binding.**

| Printing techniques | Advantages | Disadvantages | Examples of medical application |
|---------------------|------------|---------------|--------------------------------|
| Extrusion printing: Filament Deposition Modelling (FDM) | Low material costs | Rippled and porous surface | Kidney [15], Liver [11], Sinus [13,18] |
| | Low cost printers available | Fragile along z-axis | |
| | Simple to use | | |
| Photopolymerisation: Stereolithography (SLA) & Digital Light Processing (DLP) | Moderate cost | Prints are prone to slight distortions | Prosthetics [9] |
| | Good surface finish / high resolution | Curing resins need to be handled with care | |
| Photopolymerisation: Material Jetting (MJ) | Very good surface finish / high resolution | High material cost | Vessels [12,25], Spine [26] |
| | Ability to gradually combine different polymers | Curing resins need to be handled with care | |
| Powder Binding: Binder Jetting (BJ) | Can include colour | | Brain [16], Heart [10] |
| | Quick | Printers are expensive | |
| | Low material costs | Rough surface finish | |
| | Many materials available | | |
| Powder Binding: Selective Laser Sintering (SLS) | Prints are strong | Printers are expensive | |
| | Many materials available | Rough surface finish | |

https://doi.org/10.1371/journal.pone.0178540.t001
3D Slicer v.4.6 was used to create a model of the liver and the right lung from the CT ARTIFIX dataset (Siemens Sensation 64, 1.5 mm slice thickness, 0.59 mm by 0.59 mm pixel size, 120 kV peak kilo-voltage, 300 mAs exposure) from the OSIRIX website [28]. This was done using the level tracing algorithm as well as manual modification. After segmentation, the “make model” tool was used to export the volume as a STL file.

Mesh refinement. The ribs model was refined using Meshmixer to improve its topology. This was done by adjusting the mesh density of the surface and by applying a global smoothing filter that removes step artefacts due to the finite voxel size. Furthermore, we have used FreeCAD (v.0.16) [29] to design a holder for a tissue phantom. As FreeCAD has limitations working with large mesh files, the holder was attached to the ribs model structure in another software package called Blender (v.2.6) [30].

The lung model was also smoothed using MeshMixer. The different segmentation method demanded a local smoothing approach utilising the “RobustSmooth” brush provided by the software. Furthermore, the “Flatten” and “Inflate” brushes were used to remove unphysiological holes in the model.

The smoothing of the liver was also done utilizing both a global smoothing filter as well as the “RobustSmooth” brush tool.

3D printing. We have used an Ultimaker 2 (Ultimaker, Chorley, England) FDM printer to create our models. They were prepared for the printer using the open-source slicing software Cura (v.15.04.6), which is provided for free by Ultimaker. All models were printed with a layer height of 0.12 mm and a shell, bottom, and top thickness of 0.8 mm with a nozzle size of 0.4 mm. The prints were created at 20% infill, except for the rib model, which was printed with 100% infill to be functionally similar to bone when viewed under a medical ultrasound scanner (Siemens Acuson S1000 with a 16 MHz ultrasound probe). The material used for printing was “enhanced Polymax” polylactic acid (PLA) (PolyMax; Polymakr, Changshu, China). To estimate the print accuracy, the dimensions of the models were quantified at different sites in silico using Meshmixer and compared to the dimensions of the 3D prints, which were measured using calipers and a micrometer.

Results and discussion
The 3D printed models of the ribs, liver and lung can be seen in Fig 2. On the ribs phantom the holder can be seen, which can be used to place tissue mimicking phantoms underneath the ribs in order to perform realistic ultrasound imaging experiments and training. The liver
phantom was printed in coloured PLA, while the lung was painted using acrylic colour to be used as a teaching model.

The print duration for lung, ribs, and liver were 43h, 87.5h, and 27.5h respectively. The cost of the PLA for each of the models was approximately £16, £25, and £10 respectively. The print accuracy of the models is summarised in Table 2.

The ribs phantom was used as an ultrasound imaging phantom for clinical training. By embedding the ribs into a mineral-oil based material (Mindsets, Waltham Cross, United Kingdom) to simulate surrounding musculature and soft tissues, and combining these with a chicken breast and an 18 gauge puncture needle, it was possible to perform a low-cost mock kidney fine needle aspiration (FNA) procedure (Fig 3) [31]. In Fig 3B, the reflection of part of the ribs phantom can be seen in the top right corner. The artificial ribs are creating a shadowing effect which can also be observed in real ultrasound imaging procedures.

| Phantom | Name of Feature | Measured in silico (mm) | Measured on 3D print (mm) | Percentage Error |
|---------|----------------|-------------------------|--------------------------|------------------|
| Ribs    | Thickness of superior rib | 14.4                   | 14.1                     | 2.1%             |
|         | Distance between spinous processes | 110.1                 | 110.4                    | 0.3%             |
|         | Depth of the spine | 75.8                   | 76.2                     | 0.5%             |
|         | Length of middle rib | 187.2                  | 187.5                    | 0.2%             |
|         | Mean Error        |                         |                          | 0.78%            |
| Liver   | Total height      | 99.5                   | 99.1                     | 0.4%             |
|         | Total width       | 201.1                  | 198                      | 1.6%             |
|         | Total depth       | 135.1                  | 132.5                    | 1.9%             |
|         | Mean Error        |                         |                          | 1.3%             |
| Lung    | Length of bronchus | 75.2                   | 74                       | 1.6%             |
|         | Thickness of bronchus | 10.8                  | 10.5                     | 2.9%             |
|         | AP of base of lung | 114.5                  | 116                      | 1.3%             |
|         | AP of pericardium | 57.5                   | 60                       | 4.3%             |
|         | Mean Error        |                         |                          | 2.53%            |

https://doi.org/10.1371/journal.pone.0178540.t002
When going from medical imaging data to 3D printed anatomical models, the choice of an appropriate image segmentation algorithm is arguably the most important step. An overview of relevant freeware segmentation tools can be seen in Table 3. We have illustrated the use of two different open-source tools, which offer a multitude of ways to achieve accurate segmentation.

Seg3D has both manual and automatic segmentation tools and provides a library of add-ons with additional algorithms and applications optimised for particular segmentation applications. A key feature is that the interface makes it possible to visualise images in 3D with multiple volumes managed as layers. This facilitates the manipulation of several segmentations, which is particularly useful when it is necessary to use a combination of segmentation techniques to obtain the final surface. For example, the images may be cropped prior to more advanced segmentation processes in order to isolate the volume of interest. Furthermore, it provides Boolean transforms to combine multiple segmentations into a single surface. Seg3D also provides the option of exporting the final segmentation to the STL file format.

3D Slicer has a multitude of other image manipulation options and can be used to register different scans to each other. Because the large range of tools provided by the software, the interface is more difficult to master. However, it provides a range of powerful segmentation algorithms and has a unique selection of extensions available, which can be utilised for more specific tasks.

The segmentations formed the basis of the 3D printed models, which we created using FDM, allowing easy, low-cost creation of anatomical models. It was possible to create the segmented structures with high detail, allowing them to be used as teaching models. Furthermore, the printed model of the ribs was found to be functionally close to a real rib cage when imaged by an ultrasound scanner [31].

Using FDM for the creation of anatomical models has limitations inherent to the printing technique: The surface of the models was rough and rigid, which is not a realistic representation.
of the real tissue, and support material has to be carefully removed without damaging the finished print. However, there is flexible PLA available and the model surface can be smoothed using 3D print coatings (e.g. XTC-3D by Smooth-On, Macungie, USA). An alternative approach is to use a material jetting technique, which can combine different polymers seamlessly in one print, offering the possibility of creating a gradient of flexibility.

Conclusions and future work
We have introduced a general workflow that can be used to generate 3D printed anatomical models from medical imaging data. This streamlined pipeline is applicable for volumetric medical imaging data and works for a wide variety of organs and other anatomical regions of interest. We have demonstrated its use in the creation of models of ribs, a liver and a lung from CT datasets.

Recent developments in image segmentation algorithms have enabled the use of a multitude of tools and strategies in delineating anatomical structures of interest. We have provided an overview of the most relevant open-source tools that can be used for anatomical structure segmentation by end-users who are not medical or image processing specialists.

Future work will focus on creating flexible phantoms and exploring different materials with regard to their tissue mimicking characteristics in US and MRI systems.

Acknowledgments
The authors wish to thank Eve Hatten and George Dwyer for technical support and wish to thank Elliott Magee, Nishat Ahmed, Shehzaad Chikte, Ferran Prados, and Tom Vercauteren for insightful discussions.

Author Contributions
Conceptualization: DIN.
Data curation: TMB.
Funding acquisition: DIN TMB ERH JLR EM.
Investigation: TMB JLR AAP.
Methodology: JLR ERH AAP.
Project administration: DIN TMB ERH JLR.
Resources: AAP.
Supervision: DIN.
Validation: AAP.
Visualization: TMB ERH JLR.
Writing – original draft: TMB.
Writing – review & editing: DIN JLR AAP ERH EM.

References
1. McGaghie WC, Issenberg SB, Petrusa ER, Scalese RJ. A critical review of simulation-based medical education research: 2003–2009. Med Educ. Wiley Online Library; 2010; 44: 50–63. https://doi.org/10.1111/j.1365-2933.2009.03547.x PMID: 20078756
2. Evans LV, Dodge KL, Shah TD, Kaplan LJ, Siegel MD, Moore CL, et al. Simulation training in central venous catheter insertion: Improved performance in clinical practice. Acad Med. 2010; 85: 1462–1469. https://doi.org/10.1097/ACM.0b013e3181ea9a3 PMID: 20736674

3. Müller A, Krishnan KG, Uhl E, Mast G. The application of rapid prototyping techniques in cranial reconstruction and preoperative planning in neurosurgery. J Craniofac Surg. 2003; 14: 899–914. PMID: 14600634

4. Rengier F, Mehdiratta A, Von Tengg-Kobligk H, Zechmann CM, Unterhinninghofen R, Kauczor HU, et al. 3D printing based on imaging data: Review of medical applications. Int J Comput Assist Radiol Surg. 2010; 5: 335–341. https://doi.org/10.1007/s11548-010-0476-x PMID: 20467825

5. Kusaka M, Sugimoto M, Fukami N, Sasaki H, Takenaka M, Anraku T, et al. Initial experience with a tailor-made simulation and navigation program using a 3-D printer model of kidney transplantation surgery. Transplant Proc. Elsevier Inc.; 2015; 47: 596–599.

6. Zhang E, Lauber J, Beard P. Backward-mode multiwavelength photoacoustic scanner using a planar Fabry-Perot polymer film ultrasound sensor for high-resolution three-dimensional imaging of biological tissues. Appl Opt. 2008; 47: 561–577. PMID: 18239717

7. Siow B, Drobnjak I, Chatterjee A, Lythgoe MF, Alexander DC. Estimation of pore size in a microstructure phantom using the optimised gradient waveform diffusion weighted NMR sequence. J Magn Reson. Elsevier Inc.; 2012; 214: 51–60. https://doi.org/10.1016/j.jmr.2011.10.004 PMID: 22116034

8. Wells WM, Crimson WEL, Kikinis R, Jolesz FA. Adaptive segmentation of MRI data. Int J Comput Assist Radiol Surg. 2010; 5: 335–341. https://doi.org/10.1007/s11548-010-0476-x PMID: 20467825

9. Trace AP, Ortiz D, Deal A, Retrouvey M, Elzie C, Goodmurphy C, et al. Radiology’s Emerging Role in 3-D Printing Applications in Health Care. J Am Coll Radiol. Elsevier Inc; 2016; 13: 856–862.e4. https://doi.org/10.1016/j.jacr.2016.03.025 PMID: 27236288

10. Biglino G, Capelli C, Wray J, Schievano S, Leaver L-K, Khambdakone S, et al. 3D-manufactured patient-specific models of congenital heart defects for communication in clinical practice: feasibility and acceptability. BMJ Open. 2015; 5: 7165.

11. Kong X, Nie L, Zhang H, Wang Z, Ye Q, Tang L, et al. Do 3D Printing Models Improve Anatomical Teaching About Hepatic Segments to Medical Students? A Randomized Controlled Study. J Ultrasound Med. 2016; 35: 1333–1339. https://doi.org/10.7863/ultra.15.06012 PMID: 27162278

12. Ceh J, Youd T, Mastrovich Z, Peterson C, Khan S, Sasser T, et al. Bismuth Infusion of ABS Enables Additive Manufacturing of Complex Radiological Phantoms and Shielding Equipment. Sensors. 2017; 17: 459.

13. Zeiler FA, Unger B, Kramer AH, Kirpatrick AW, Gillman LM. A Unique Model for Ultrasound Assessment of Optic Nerve Sheath Diameter. Can J Neurourol Sci. 2013; 40: 225–229. PMID: 23419572

14. Jarvis D, Griffiths PD, Majewski C. Demonstration of Normal and Abnormal Fetal Brains Using 3D Printing from In Utero MR Imaging Data. AJNR Am J Neuroradiol. 2016; 1–5.

15. Naftulin JS, Kimchi EY, Cash SS. Streamlined, inexpensive 3D printing of the brain and skull. PLoS One. 2015; 10: e0136198. https://doi.org/10.1371/journal.pone.0136198 PMID: 26295459

16. Sander IM, McGoldrick MT, Helms MN, Betts A, van Avermaete A, Owens E, et al. Three-dimensional printing of X-ray computed tomography datasets with multiple materials using open-source data processing. Anat Sci Educ. 2017; 0.

17. Fedorov A, Beichel R, Kalpathy-Cramer J, Finet J, Fillion-Robin J-C, Pujol S, et al. 3D Slicer as an Image Computing Platform for the Quantitative Imaging Network. Magn Reson Imaging. Elsevier; 2012; 30: 1323–1341. https://doi.org/10.1016/j.mri.2012.05.001 PMID: 22770690
25. Cloonan AJ, Shahmirzadi D, Li RX, Doyle BJ, Konofagou EE, Mcgloughlin TM. 3D-printed tissue-mimicking phantoms for medical imaging and computational validation applications. 3D Print Addit Manuf. 2014; 1: 14–13.

26. West SJ, Mari J-M, Khan A, Wan JHY, Zhu W, Koutsakos IG, et al. Development of an ultrasound phantom for spinal injections with 3-dimensional printing. Reg Anesth Pain Med. 2014; 39: 429–33. https://doi.org/10.1097/AAP.0000000000000136 PMID: 25105983

27. Autodesk Meshmixer [Internet]. [cited 1 Jan 2016]. Available: http://www.meshmixer.com/

28. Rosset Antoine and Spadola Luca and Ratib O. OsiriX: an open-source software for navigating in multi-dimensional DICOM images. J Digit Imaging. 2004; 17: 205–216. https://doi.org/10.1007/s10278-004-1014-6 PMID: 15534753

29. FreeCAD [Internet]. [cited 1 Jan 2016]. Available: http://www.freecadweb.org/

30. Blender [Internet]. [cited 1 Jan 2016]. Available: https://www.blender.org/

31. Robertson J, Hill E, Plumb A, Choong S, West S, Nikitichev D. 3D printed ultrasound phantoms for clinical training. IS&T's 32nd International Conference on Printing for Fabrication (NIP32). Manchester; 2016.

32. InVesalius [Internet]. [cited 1 Jan 2016]. Available: http://www.cti.gov.br/invesalius/

33. itk-SNAP [Internet]. [cited 1 Jan 2016]. Available: http://www.itksnap.org/

34. Osirix MD [Internet]. [cited 21 Mar 2017]. Available: http://www.osirix-viewer.com/

35. Horos [Internet]. [cited 25 Apr 2017]. Available: https://www.horosproject.org/

36. ImageJ [Internet]. [cited 27 Mar 2017]. Available: https://imagej.nih.gov/ij/