





Introduction

- 3D printers capable of printing hydrogel materials are prohibitively expensive, slow, and not scalable for large-scale fabrication.
- Traditional 3D printing uses an extrusion-based approach in which materials are ejected from a nozzle tip, but this process is extremely slow.¹
- Masked SLA (mSLA) 3D printers use light to photo-polymerize materials layer-by-layer at a time, creating high resolution prints in less time.¹
- It may be possible to use mSLA 3D printers for printing photopolymerizable hydrogel materials.

Purpose

To develop bioinks for commercial mSLA printers to produce highquality prints and methods for assessing print quality.

Methods

- Bioink formulations consisted of 10% GelMa (gelatin methacrylate),² LAP (Lithium phenyl-2,4,6-trimethylbenzoylphosphinate), and a yellow food-grade dye (to minimize unwanted light leakage). A commercial mSLA (masked stereolithography) printer, the Photon Mono X (AnyCubic, Shenzhen), was retrofitted with a custom
- temperature and humidity control kit.
- Printing process was performed at 40°C and 90% humidity to ensure that the GelMa remained in a liquid state.
- A set of matrix cubes of varying sizes with holes was printed and used as a standard control for comparing different formulations. Images of the cubes were taken with a camera, top-down and sidereview, analyzed with the ImageJ software and compared with the original CAD designs to derive an overall print quality score.
- Two print variables, exposure time (5 s to 40 s) and yellow dye concentration (1 - 7%), were analyzed in this study.

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Results

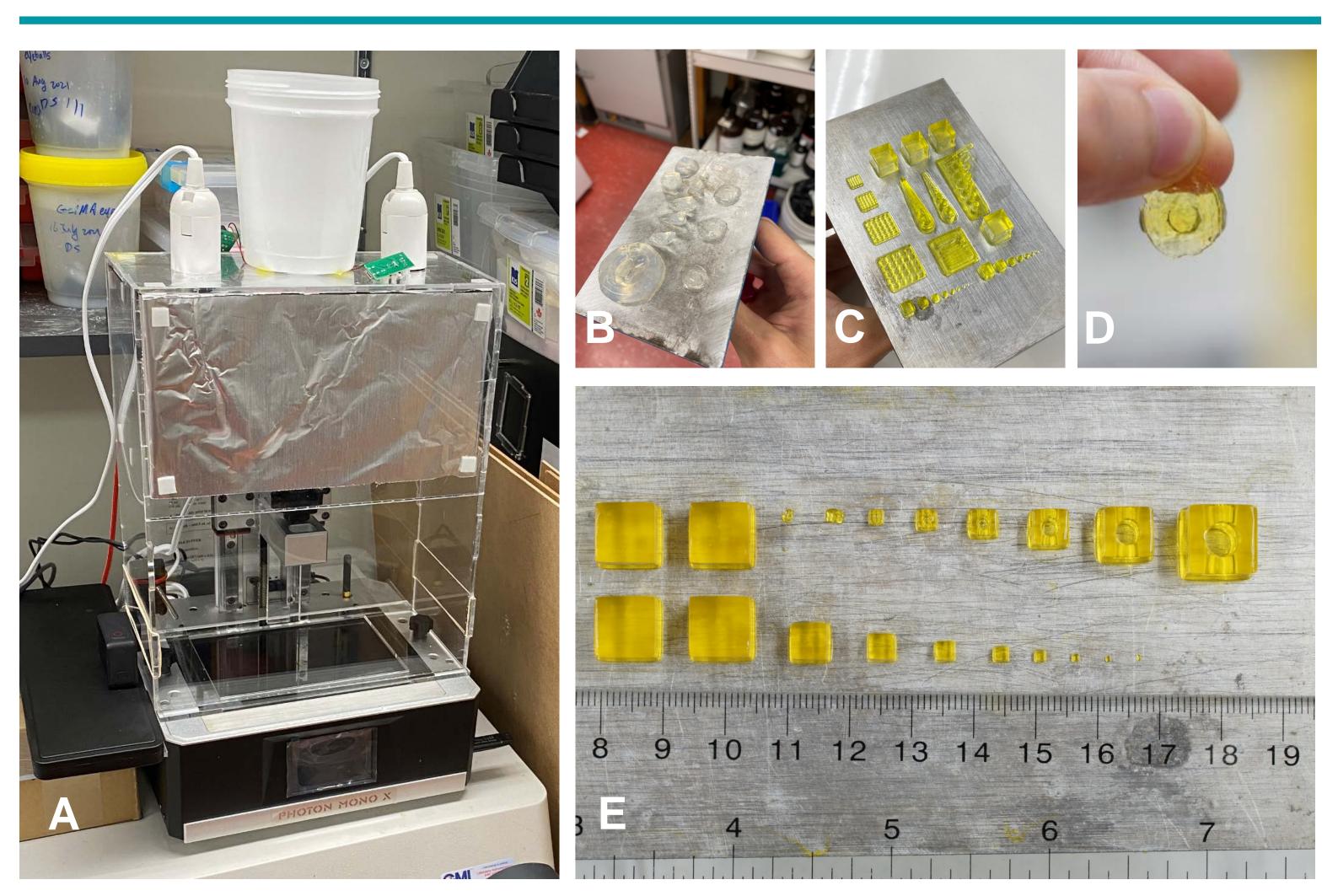


Figure 1 A Retro-fitted mSLA 3D printer with humidity and temperature control. B A sample 3D print without yellow dye and **C** with a yellow dye. **D** a model 3D printed contact lens. **E** A set of 3D-printed matrix cubes standards used to assess the quality of different bioinks.



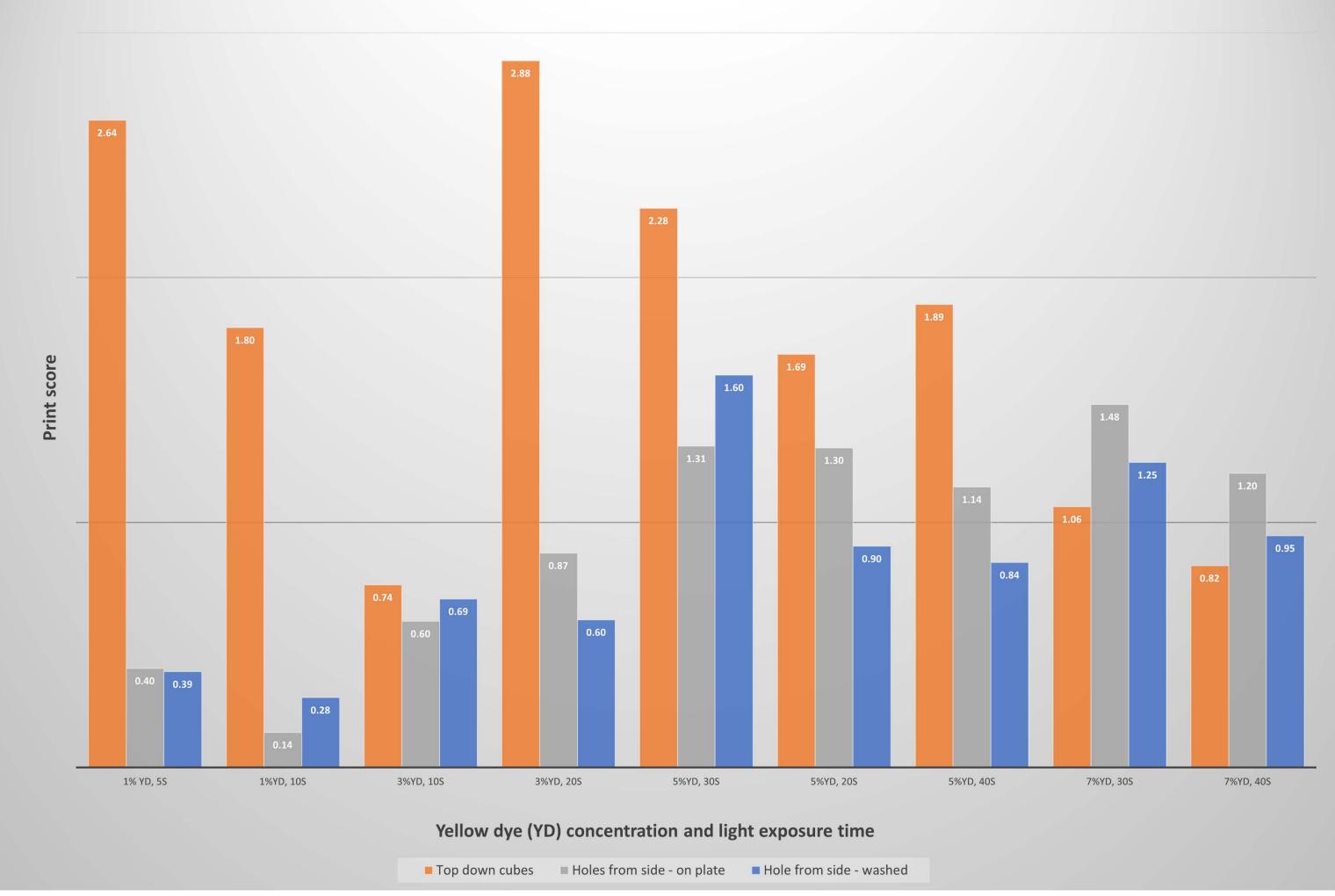


Figure 2 Print scores of the standard cubes for different yellow dye concentrations and exposure times

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Developing bioinks for commercial mSLA printers and a method for quantifying print quality Chau-Minh Phan,^{1,2} David Wulff,^{1,2} Lyndon Jones^{1,2}

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Conclusions

- reliability.
- polymers.

References

- 2020;15(6):065017.

Addition of the dye was critical to producing high-resolution prints. In the absence of dye, the printed cubes had edge deformations or did not properly resolve internal structures

Best resolution with the highest print scores were obtained at either 5% yellow dye concentration and 30 seconds exposure time, or 3% yellow dye concentration and 20 seconds exposure time.

A prototype contact lens with a 200 µm thickness was able to be 3D printed using the developed print methods and parameters, with a total print time of approximately 20 minutes.

• However, print designs with a very low surface area at the base (such as a contact lens) have a high rate of failure due to the prints falling off the print plate during the printing process.

The yellow dye can be removed post-printing by washing the print in phosphate buffered saline or by bleaching with UV light.

• The current study demonstrated that a low-cost commercial 3D mSLA printer could be used with bioinks for 3D bioprinting. • Further work is necessary to improve the print resolution and

• We hypothesize that each different bioink formulation will require different print parameters, so a standardized process to assess print quality, could immensely facilitate the optimization process.

• Future work will optimize this 3D printing method with other

. Arslan-Yildiz A, El Assal R, Chen P, Guven S, Inci F, Demirci U. Towards artificial tissue models: past, present, and future of 3D bioprinting. Biofabrication. 2016;8(1):014103.

2. Rizwan M, Chan SW, Comeau PA, Willett TL, Yim EKF. Effect of sterilization treatment on mechanical properties, biodegradation, bioactivity and printability of GeIMA hydrogels. Biomedical materials.

