

3D-printed vs thermoformed orthosis for trigger finger and hyperextension of the proximal interphalangeal joint

Submission date 08/11/2025	Recruitment status Recruiting	☒ Prospectively registered ☐ Protocol
Registration date 11/11/2025	Overall study status Ongoing	☐ Statistical analysis plan ☐ Results
Last Edited 29/07/2026	Condition category Musculoskeletal Diseases	☐ Individual participant data ☒ Record updated in last year

Plain English summary of protocol

Background and study aims

This study is exploring a new way to make orthoses (hand supports) using 3D printing technology. These supports are used to help people with a condition called trigger finger, which causes pain and stiffness in the fingers. Although 3D printing has many advantages over traditional methods, it's not yet commonly used in clinics. The aim of this study is to see how acceptable and useful 3D-printed orthoses are for patients and healthcare professionals.

Who can participate?

Adults aged 18 years and over who have trigger finger can take part in the study. Both men and women will be included, and participants will be recruited from different healthcare centres.

What does the study involve?

Participants will join online meetings using Microsoft Teams or Zoom. They will receive a custom-made 3D-printed orthosis to wear. They will also be asked to fill out questionnaires about their pain and hand function at five different times: before wearing the orthosis, and then 24 hours, 1 week, 1 month, and 2 months after starting to wear it. All questionnaires can be completed from home using a phone or computer. The medication used by the participants will also be evaluated in the first questionnaire using the Cumulative Analgesic Consumption Score.

What are the possible benefits and risks of participating?

Participants may benefit from reduced pain and improved finger movement thanks to the orthosis. However, wearing it might be uncomfortable and could cause skin irritation or pain due to limited finger movement. The study also requires some time commitment—about 30 minutes online and around 15 minutes to complete questionnaires at each of the five time points.

Where is the study run from?

The study is being run by Université du Québec à Trois-Rivières (UQTR) in Canada.

When is the study starting and how long is it expected to run for?

June 2025 to December 2026

Who is funding the study?
Projet 2024-2025 de la Planification stratégique institutionnelle de l'UQTR (Canada)

Who is the main contact?
The main contact for the study is Tokiko Hamasaki, an occupational therapist and PhD researcher at UQTR.
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Principal investigator

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Additional identifiers

Study information

Scientific Title

Comparison of 3D-printed and thermoformed orthosis fabrication methods for trigger finger and hyperextension of the proximal interphalangeal joint: a pilot study

Study objectives

This project aims to evaluate whether 3D-printed orthosis is acceptable in a clinical setting, both for professionals and for patients with trigger finger and hyperextension of the proximal interphalangeal joint. This technology, which is still rarely used in this field, could offer practical and clinical advantages over traditional methods.

Ethics approval required

Ethics approval required

Ethics approval(s)

Approved 14/10/2025, Comité d'éthique de la recherche du CIUSSS de la Mauricie-et-du-Centre-du-Québec (2700, boulevard des Forges, bureau 302, Trois-Rivières, G8Z 1V2, Canada; +1 819 372-3133, poste 32303; 04ethiqueciusssmcq@ssss.gouv.qc.ca), ref: MP-29-2026-895, 1025

Primary study design

Interventional

Study design

Quasi-experimental crossover study with repeated measures

Study type(s)

Quality of life, Treatment

Health condition(s) or problem(s) studied

Trigger finger and hyperextension of the proximal interphalangeal joint

Interventions

The study design is semi-experimental; thus, there is no randomization, and there is only one arm for which the treatment is a 3D-printed orthosis. Every participant will wear the 3D orthosis and the thermoformed orthosis (control). They will wear each orthosis for a total of 2 months. The participants will answer a questionnaire assessing their symptoms before wearing the orthosis and questionnaires about the orthosis 24 hours, 1 week, 1 month and 2 months after beginning to wear the orthosis. They will not wear the 3D orthosis and the thermoformed orthosis at the same time.

Intervention Type

Procedure/Surgery

Primary outcome(s)

1. Pain and upper limb function measured using 0-10 rating scale for pain at the start of the study and after beginning to wear the orthosis at multiple intervals (24 hours, 1 week, 1 month, 2 months)
2. QuickDASH questionnaire for upper limb function at the start of the study and after beginning to wear the orthosis at multiple intervals (24 hours, 1 week, 1 month, 2 months)

Key secondary outcome(s)

User satisfaction measured using QUEST 2.0 questionnaire after beginning to wear the orthosis at multiple intervals (24 hours, 1 week, 1 month, 2 months)

Completion date

01/12/2026

Eligibility**Key inclusion criteria**

1. The patient must be 18 years or older
2. The patient must have a trigger finger and hyperextension of the proximal interphalangeal joint
3. The patient must be able to use Microsoft Teams or Zoom with their cell phone or computer
4. The patient must be able to read, write and answer the questionnaires

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Mixed

Lower age limit

18 Years

Upper age limit

100 Years

Sex

All

Total final enrolment

0

Key exclusion criteria

1. The patient is already wearing an orthosis
2. The patient received a cortisone injection in the last 4 months
3. The patient received surgery for this condition

Date of first enrolment

01/12/2025

Date of final enrolment

01/12/2026

Locations**Countries of recruitment**

Canada

Study participating centre

Clinique Multidisciplinaire en Santé de l'UQTR

Pavillon de la Santé, Local 2827

Trois-Rivières

Canada

G9A 5H7

Study participating centre

CBI Santé - Trois-Rivières

1785 Bd du Carmel local 103

Trois-Rivières

Canada

G8Z 3R8

Study participating centre

Centre hospitalier affilié universitaire régional de Trois-Rivières

1991 Bd du Carmel

Trois-Rivières
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G8Z 3R9

Sponsor information

Organisation

Université du Québec à Trois-Rivières

Funder(s)

Funder type

University/education

Funder Name

Université du Québec à Trois-Rivières

Alternative Name(s)

Universidad de Quebec en Trois-Rivières, UQTR

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

Canada

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated and/or analysed during the current study will be available upon request from Tokiko Hamasaki (tokiko.hamasaki@uqtr.ca)

IPD sharing plan summary

Available on request