



Manufacturing of lightweight insulation foam from renewable henequen fibres

Qasim Shoaib, Vaidurya Mukherjee, Vasiliki Marmaridou, Aleksei Sereda, Peterasp Satarawala, Jose Gonzalo Carillo Baeza, Eral Bele





Table of contents

- 1. Introduction
 - i. Design process Principles
 - ii. Intent of Study
 - iii. Market Gap & Innovation
- 2. Methodology
 - i. Fibrillation
 - ii. Foaming Process
 - iii. Optimization
 - iv. Structure

- 3. Assessment of performance
 - i. Mechanical Properties
 - i. Flame resistance
 - iii. Comparison with Porous Materials
 - iv. Flame resistance
 - V. Recyclability
 - Vi. Biodegradability
 - Vii. Life cycle performance

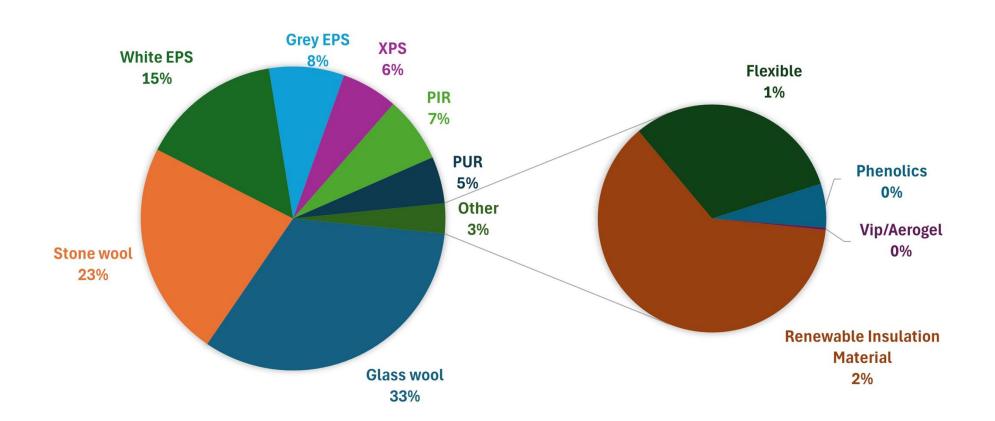
Introduction

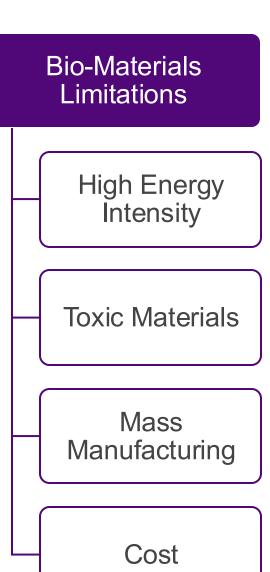


• The building sector accounts for 37% of global CO₂ emissions.

Problem

- Current materials have high environmental costs and are difficult to recycle.
- Bio-based options have limitations





Design process - Principles





Principles

- Abundant & Underutilized Resources
- Eco-Friendly Processing
- Flame Resistant
- Scalable & Low-Energy



Process

- Develop sustainable, high-performance insulation
- Optimize thermal, mechanical & fire resistance properties
- Compare environmental impact with traditional materials

Intent of Study



Intent

 Develop a simple and scalable manufacturing process for lightweight fibrous networks form lignocellulosic henequen fibre.

Why henequen?

- Natural long fibre which is underutilised and in abundance
- Sustainability benefits during growth and harvesting

Research objectives

- Assess performance of designed foams in areas of:
 - Density & Porosity
 - Thermal Conductivity
 - o Flame Resistance
 - Recyclability & Biodegradability
 - Life-Cycle Impact
- Assess viability in comparison to current solutions

Market Gap & Innovation



Gap in the market

Ease of Manufacturing

Easy Fibrillation

Flame Retardant

Sustainable

100% non-toxic foam

Investigation

Mechanical Fibrillation via blending

Chemical cross-linking

Flame Retardant using Borax

Xanthan Gum

Use of abundant raw material





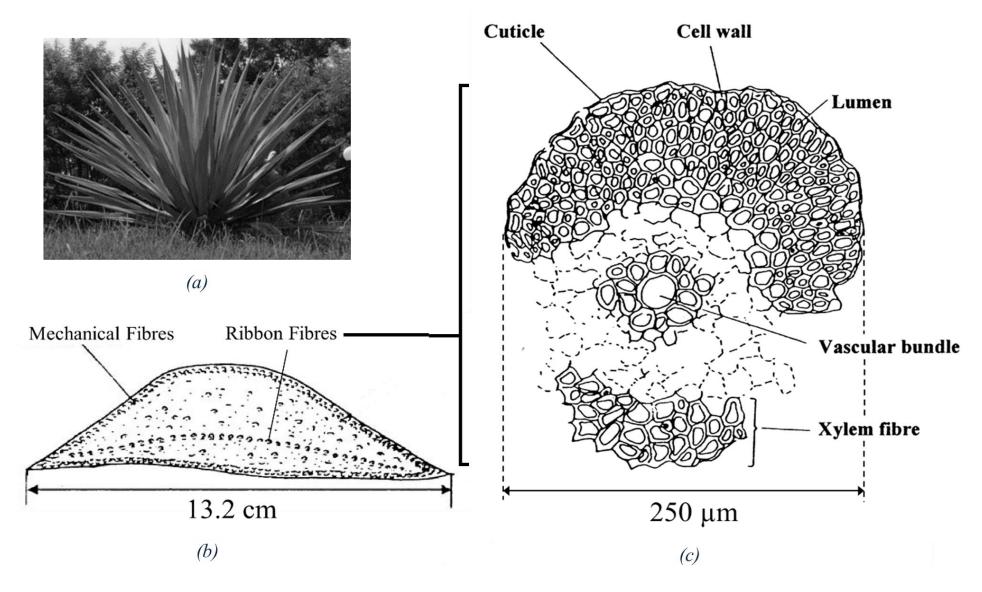
Table of contents

- 1. Introduction
 - i. Design process –Principles
 - ii. Intent of Study
 - iii. Market Gap & Innovation
- 2. Methodology
 - i. Fibrillation
 - ii. Foaming Process
 - iii. Optimization
 - iv. Structure

- 3. Assessment of performance
 - i. Mechanical Properties
 - ii. Flame resistance
 - iii. Comparison with Porous Materials
 - iv. Recyclability
 - V. Biodegradability
 - Vi. Life cycle performance

Henequen Fibre Structure

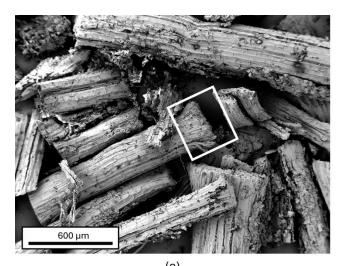


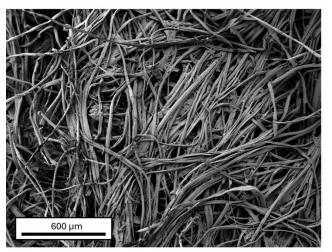


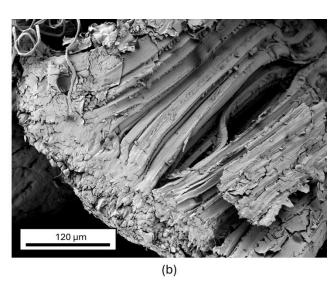
Fibrillation

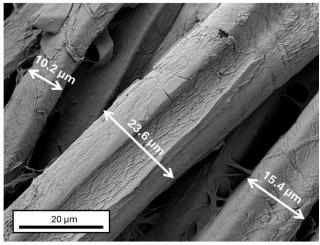


- Breaks fibre bundles into smaller 'fibrils' that branch out
- Allows for greater physical fibre-fibre interaction – fibres mechanically 'lock together'
- Effective strategy to deliver stability of fibre network -> Interlocking prevents relative displacement of fibres



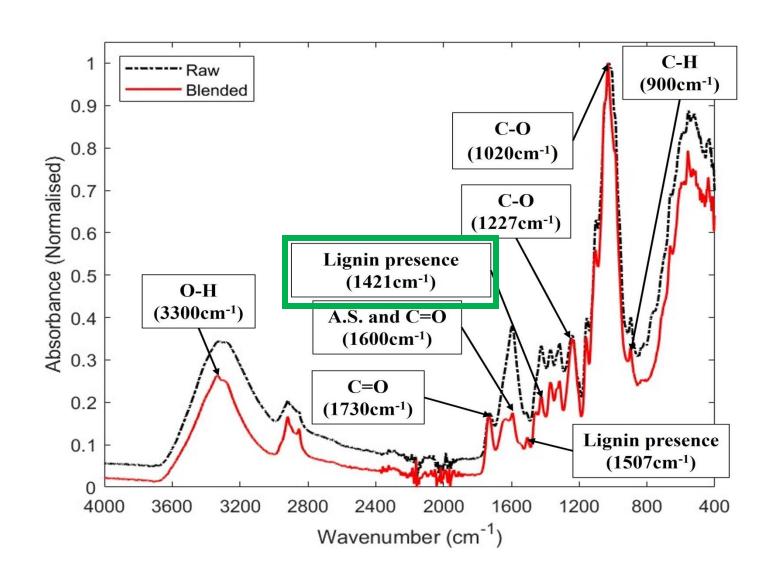






Simple Fibrillation Strategy using Blender





FTIR shows:

- Reduction (loss) of lignin
- Preservation of Hemicellulose

Blender observed to repeatably alter fibre geometry:

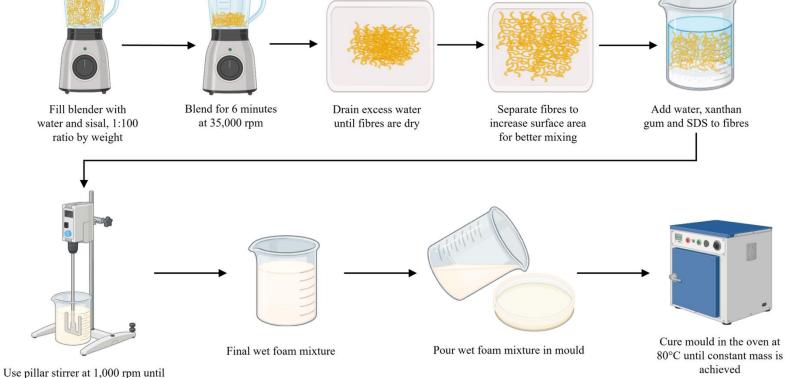
- Diameter Reduction: 381 µm
 → 21 µm (94.5%)
- Length Reduction: 5.1 cm
 → 2.9 cm (45%)
- Aspect Ratio Increase: 130 → 1400 (1076%)

Foam 1 – Xanthan Gum Stabilised



Xanthan gum (a common food additive) used as binder for cellulose fibre foam)

volume reaches steady state



This foam is:

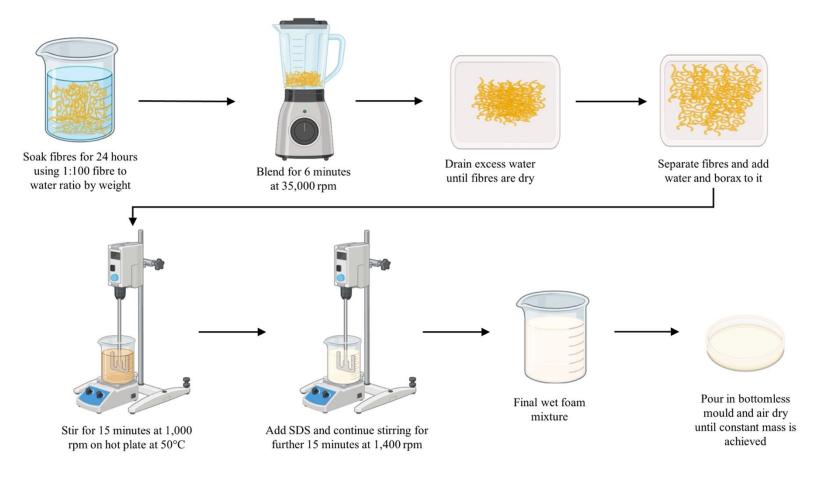
- Non-Toxic
- Easily Accessible
- Has not been explored for such uses
- Effective at very low concentrations of xanthan gum (0.5wt%)



Foam 2 – Borate Stabilised Foam



Addition of borate can help to stabilise cellulose fibre network (chemical interaction and cross-linking).



This foam is:

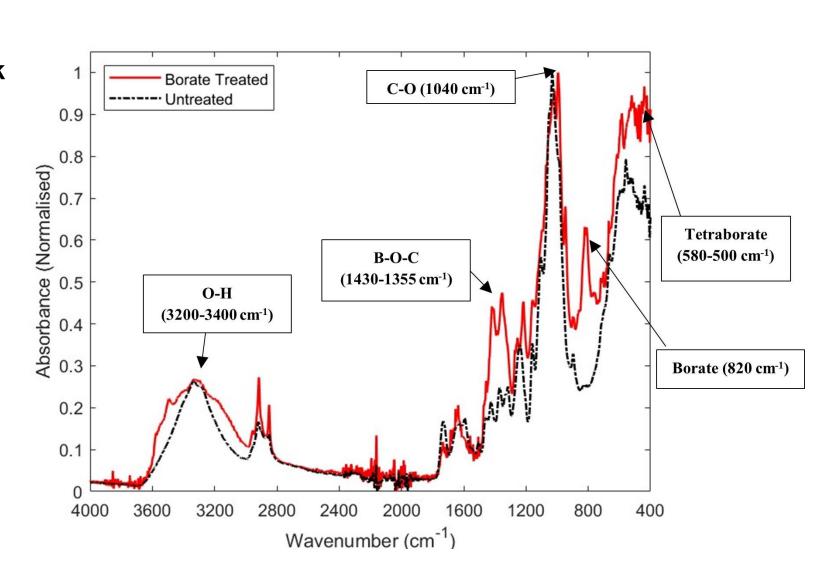
- Relatively safe/cheap
- Provides benefit at very low borax concentrations (1.5 – 2 wt%)
- Added to fibre solution prior to foaming process with minimal heating (~50°C)



Borate Cross-Linking & Chemical Validation

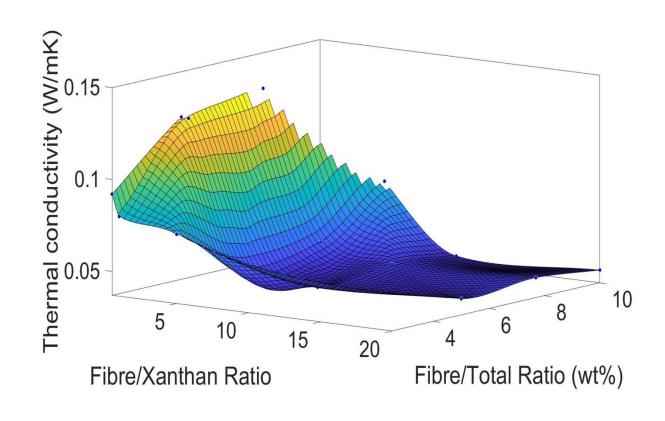


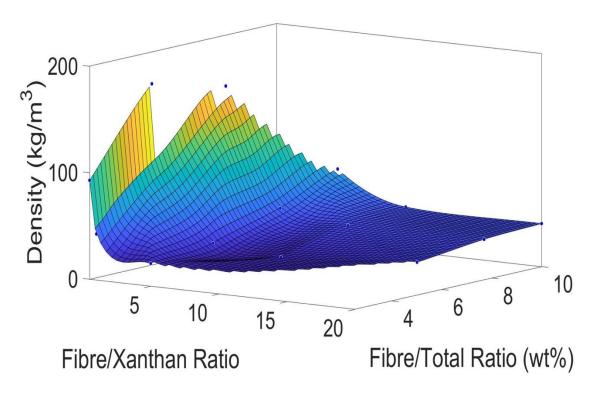
- **Chemical Cross-Linking in** Borate Foam: New B-O-C peak (1355-1430 cm⁻¹) indicates Formation of borate linkage
- Residual borate (820 cm⁻¹) & tetraborate (580-500 cm⁻¹) observed
- Synergistic with delivery of flame resistant properties from borate.



Optimising Thermal Resistance I

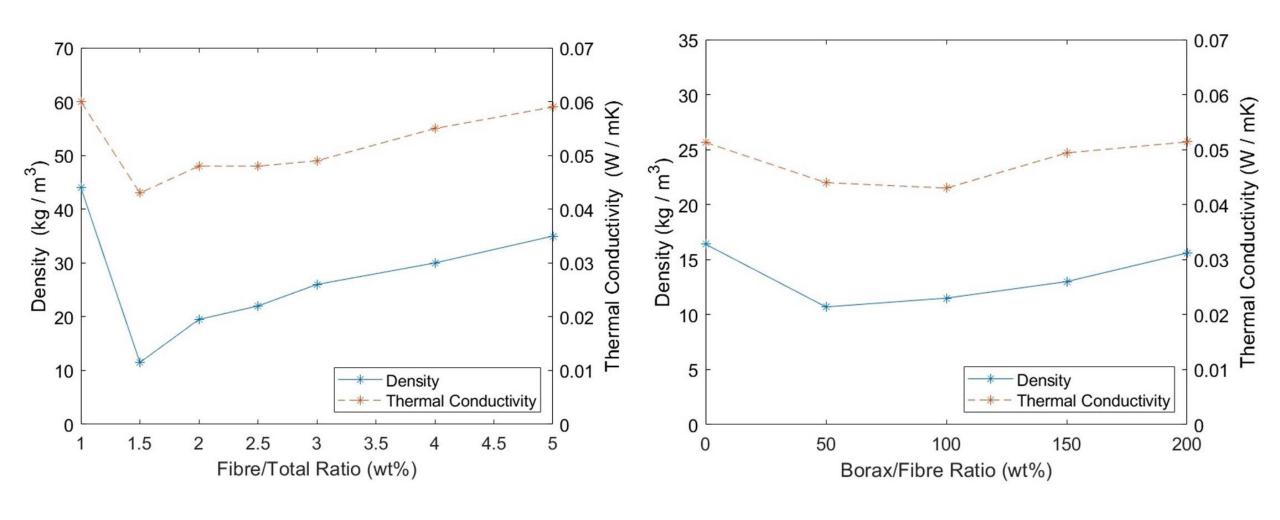






Optimising Thermal Resistance II





Structure – Density & Porosity

Foam Morphology & Porosity:

- CT scans confirm a randomly oriented fibrous network
- 94.4% porosity

 Fibres are coated and physically adhered by a gum layer

 Thin film at intersections stabilizes the structure

• 94.3% porosity

- Fibres physically linked forming branched junctions
- No binding agent required

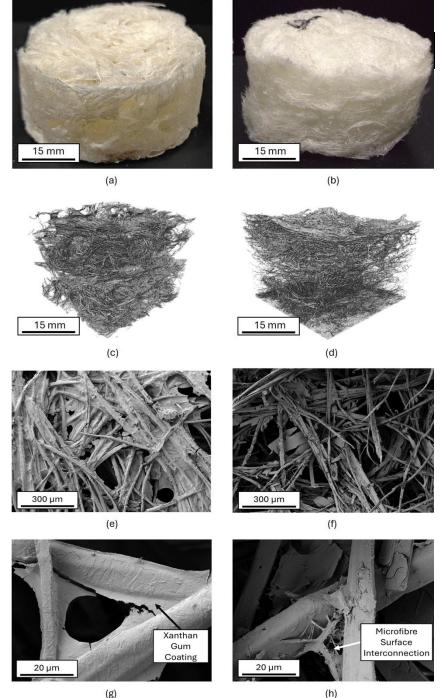






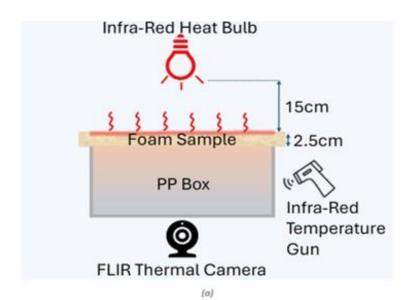
Table of contents

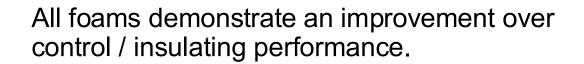
- 1. Introduction
 - i. Design process –Principles
 - ii. Intent of Study
 - iii. Market Gap & Innovation
- 2. Methodology
 - i. Fibrillation
 - ii. Foaming Process
 - iii. Optimization
 - iv. Structure

- 3. Assessment of performance
 - i. Mechanical Properties
 - ii. Flame resistance
 - iii. Comparison with porous materials
 - V. Recyclability
 - V. Biodegradability
 - Vi. Life cycle performance

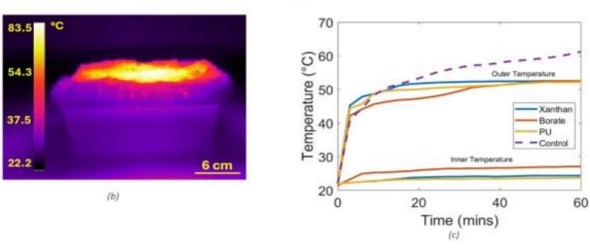
Thermal Imaging







New design foams are comparable in performance to current PU solutions



Material	Outer Temperature (°C) Temperature in box (°C)		
Xanthan	83.5	27.3	
Borate	52.5	27.0	
PU Foam	54.1	24.9	
Control	-	61.2	

Mechanical Performance

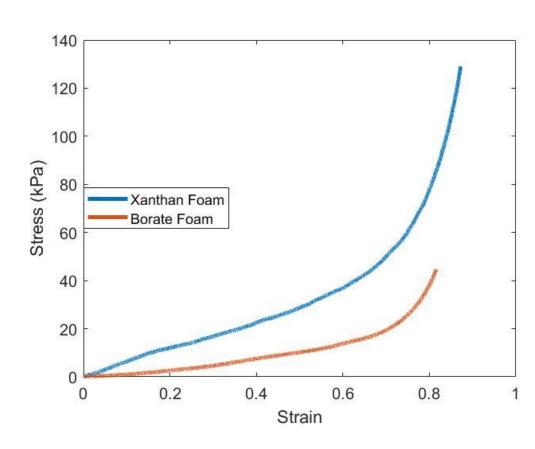


Two Distinct Deformation Phases:

- Elastic Regime: Initial fibre rearrangement and bending.
- Strain Stiffening: Increased resistance due to fibre buckling.
 - Densification Effect: New fibre-fibre contacts form under large strains, increasing resistance.

Key Observations:

- Xanthan Foam: Higher stiffness (56.4 kPa) due to mechanical bonding.
- Borate Foam: Lower stiffness (19.4 kPa) with chemical cross-linking.
- Both foams exceed BS ISO 21844 (7 kPa at 10% strain).

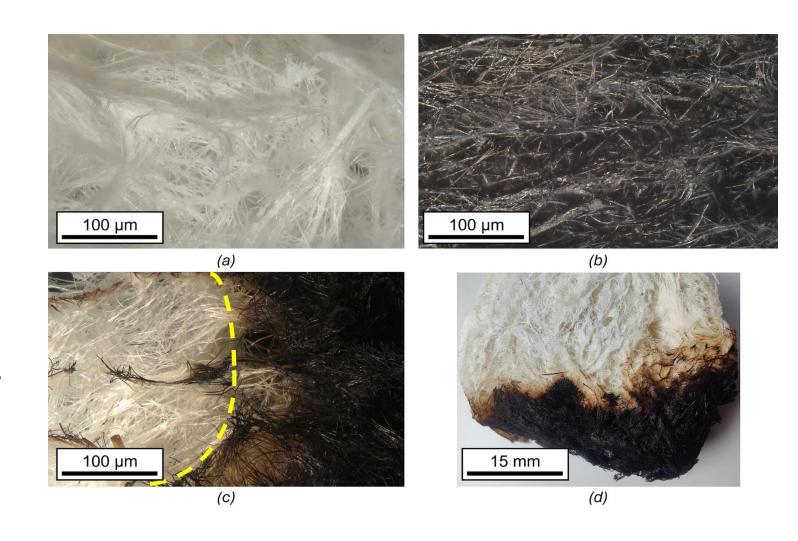


Flame Resistance - Borates



Borate:

- Addition serves dual purpose can be used to deliver both
 - flame resistance
 - stabilisation for foam design
- Natural sodium salt
- Effective at small concentrations (1.5-2 wt%)
- Homogenizes well

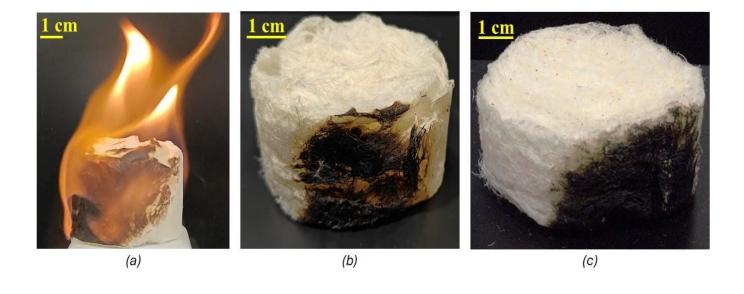


Flame Resistance - LOI

≜UCL

Results:

- Limiting Oxygen Index (LOI) Test (ASTM D2863)
- LOI Value: 29.5% → Classified as "Self-Extinguishing" (LOI >28%).





Comparison with Other Porous Materials



Density

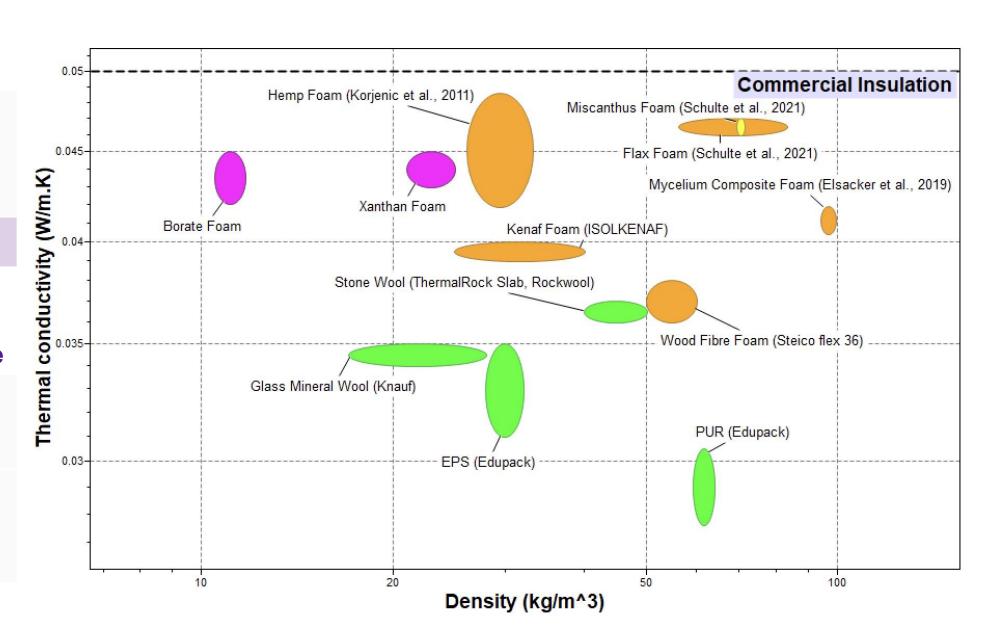
Lower than fossil-fuel and lignocellulosic insulation

→ Resource-efficient

Thermal performance

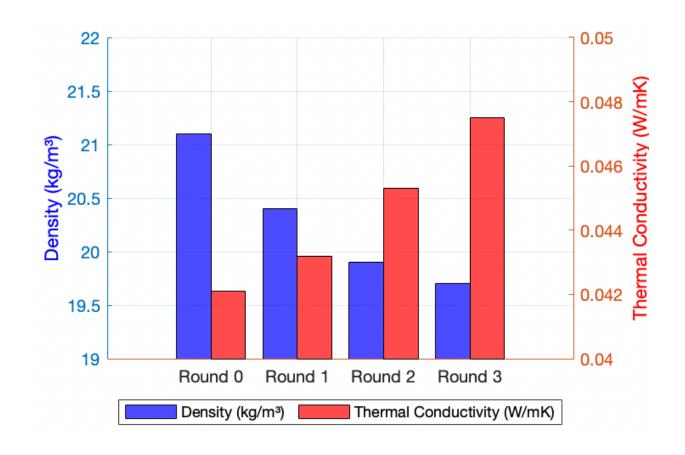
Slightly poorer than **mineral wool & EPS**

Comparable to biobased insulation



Recyclability





Xanthan foam can be recycled successfully: density and thermal conductivity were retained in 3 cycles.

Recycling achieves

- Minimising waste and energy consumption
- Minimising carbon footprint
- Maintaining similar results to the original foam

	Initial	Round 1	Round 2	Round 3
Density (kg/m3)	21.1	20.4	19.9	19.7
Thermal Conductivity (W/mK)	0.0421	0.0432	0.0453	0.0475

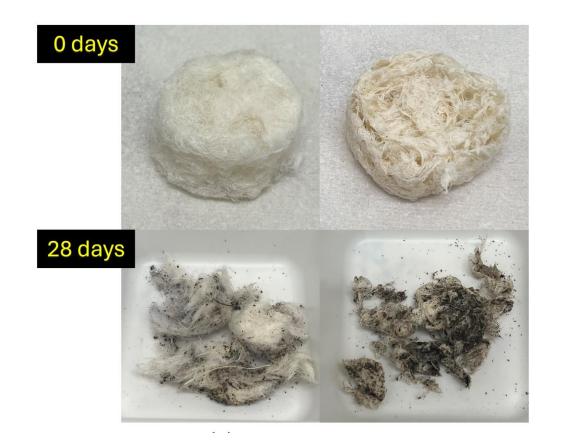
Biodegradability

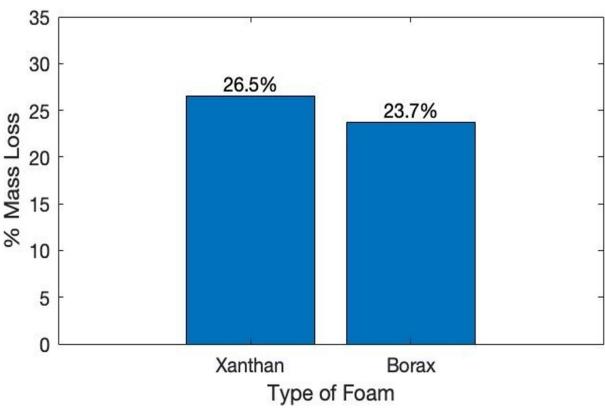


Biodegradability shows promising results:

Xanthan Foam loses 26.5% of its mass

Borate Foam loses 23.7% of its mass



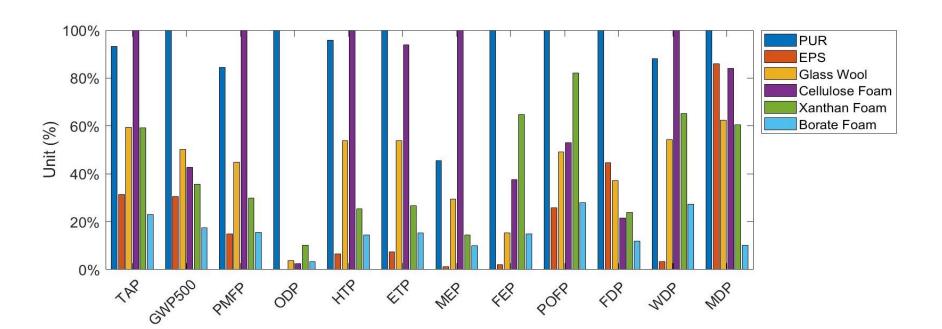


Life Cycle Assessment

≜UCL

- Borate Foam: 3.81 kg CO₂ / functional unit
- Xanthan Foam: 7.62 kg CO₂ / functional unit
 - Main contributors: Diesel (30-55%)
 Electricity (30-50%)
- Glass Wool Comparison: lower impact overall but higher freshwater eutrophication & water use

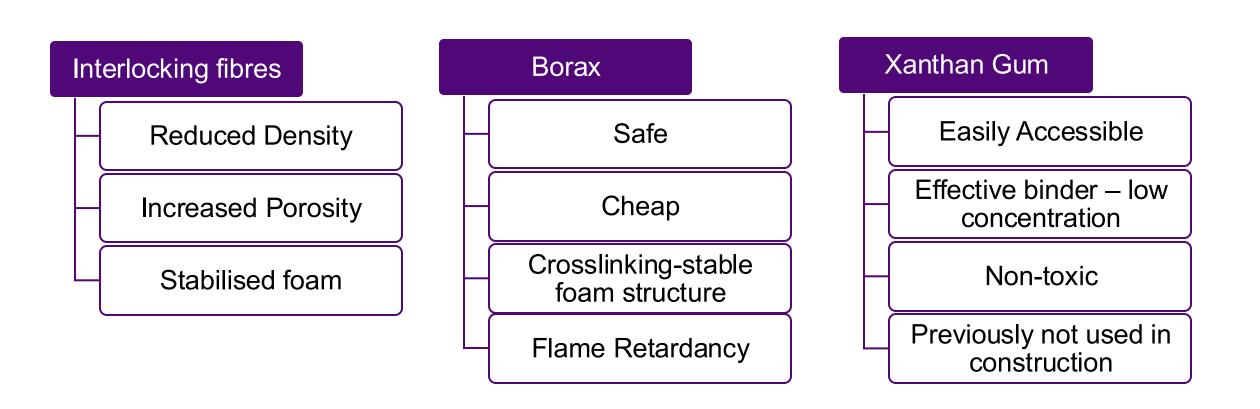
- PUR Foam Comparison
 - Lower toxicity (HTP, ETP) (up to 90%)
 - Lower global warming potential (80% Borate)
 - Lower fossil fuel depletion (90% Borate, 80% Xanthan)
 - Lower water use (90% Borate, 40% Xanthan)



Key Takeaways



Combination of materials and methods as potential building blocks of high-performing foams for thermal insulation applications



Thank you!





UCL Mechanical Engineering

Supported by:

- Royal Society: International Exchanges Scheme 2020 (Grant No.507 IES/R1/201026)
- CONAHCYT: Ciencia Básica 2017–2018 (Grant No.509 A1-S-8864)

