

Fused filament fabrication (FFF) of biobased low-density polymer foams

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Zusammenfassung In einer Welt, in der der Leichtbau aufgrund der Notwendigkeit, Ressourcen zu schonen und die Treibhausgasemissionen zu senken, eine immer wichtigere Rolle spielt, ist es wichtig, neue Technologien zu entwickeln, die leichtere Produkte ermöglichen. Die additive Fertigung bietet aufgrund ihrer Gestaltungsfreiheit ein hohes Leichtbaupotenzial durch die Realisierung niedriger Bauteildichten, aber es ist noch nicht möglich, Polymerschäume mit niedriger Dichte zu drucken, um noch leichtere Teile zu erhalten. Daher wird in diesem Beitrag ein neuartiges, extrusionsbasiertes additives Fertigungsverfahren vorgestellt, bei dem ein biobasiertes Polymerfilament mit einem physikalischen Treibmittel zur Herstellung von Schaumteilen verwendet wird. Ein Extrusionsversuchsaufbau ermöglicht die Analyse der Randbedingungen der Schaumextrusion, d. h. der Beziehung zwischen Vorschubgeschwindigkeit, Heiztemperatur, Durchmesser des extrudierten Materials, Extrusionskraft und Dichte des extrudierten Materials. Niedrigste Dichten lassen sich durch niedrige Temperaturen und schnellstmögliche Einzugsgeschwindigkeiten bei diesen Temperaturen erreichen. Eine umfassende Parameterstudie zur additiven Fertigung eines einfachen Schaumstofftestteils zeigt durchgängig niedrige Dichten von nahezu 100 kg/m3 bei ausreichender Druckqualität. Die Analyse der Schaummechanismen von drei ausgewählten Szenarien zeigt, dass die verwendete Berechnungsmethode eine ausreichende Vorhersagekraft für die Druckergebnisse besitzt. Mit der verwendeten Charakterisierung lassen sich Korrelationen der Expansionsgeschwindigkeit und zeit des Polymers während des Drucks nach dem Druckprozess recht gut abbilden.

Abstract In a world in which lightweight construction is playing an increasingly important role due to the need to conserve resources and lower the greenhouse gas emissions, it is important to develop new technologies that enable lighter products. Additive manufacturing offers a high lightweight potential due to its freedom of design by realising low part densities, but it is not yet possible to print low-density polymer foams to achieve even lighter parts. Therefore, this paper presents a novel, extrusion-based additive manufacturing process that uses a bio-based polymer filament loaded with a physical blowing agent to manufacture foam parts. An extrusion test setup allows the analysis of foam extrusion boundary conditions, i.e. the relationship between feeding velocity, heater temperature, extruded material diameter, extrusion force and extruded material density. Lowest densities can be achieved by using low temperatures and fastest at this

temperature possible feeding velocities. A comprehensive parameter study on the additive manufacturing of a simple foam test part shows continuous low densities of nearly 100 kg/m³ with an adequate print quality. The analysis of the foam mechanisms of three selected scenarios demonstrates that the calculation method used has adequate predictive power for print results. With the characterisation used, correlations of the expansion velocity and time of the polymer while printing can be reasonably well mapped after the printing process.

Introduction and motivation

Conserving resources is an increasingly important task due to the increasing concentration of greenhouse gases in the earth's atmosphere. Fossil-based polymers in particular cause high greenhouse gas emissions, not only through their production and disposal, but also during their transport. The production part can be reduced by using recycled materials [1] or bio-based polymers [2], while the transport share can be achieved by reducing the mass of the transported products. Polymer foams can enable low masses due to their low density, but are currently often fossil-based [3] and are sometimes produced in simple moulds and then individualised by machining [4]. Additive manufacturing can enable individualisation and thus contribute to further resource conservation. This effect can be enhanced by using bio-based plastics.

Definitions

Additive manufacturing is a potentially well-suited manufacturing process for the production of polymer foam components due to its freedom of design. This is reflected in the number of foaming processes already described in the literature. A good overview is provided by Nofar et al. [5], who identified four main methods: 1) Construction of porous structures (lattice structures) [6, 7]; 2) Syntactic foaming [8–10]; 3) Post-foaming of premoulded solid foams [11, 12]; 4) In-situ foaming of filaments loaded with blowing agents [13–15]. The additive manufacturing of lattice structures is an established process that only enables macroscopically porous structures, i.e. no micropores and therefore no intrinsic density reduction. Syntactic foaming and post-foaming require a pre-process to insert the fillers and a post-process to remove the fillers respectively for subsequent foaming. All of the process variants mentioned enable a lower density than the bulk material down to a minimum density of approx. 300 kg/m³ [9] of the additively manufactured components. In conventional production, densities up to ten times lower can be achieved with a thermoplastic polylactide (PLA) [16]. In the following, the in-situ foaming of filaments loaded with blowing with the aim of achieving an adequately low density of produced parts will be discussed.

Method

This paper presents a fused filament fabrication (FFF) process that enables the processing of filaments loaded with a physical blowing agent. [Figure 1](#page-3-0) shows the process used in this approach schematically. After the thermoplastic filament has been produced, it is loaded with a gaseous blowing agent. After transport and storage, the loaded filament is processed using an FFF printer to produce a foam component.

The manufacturing process is simplified according to the principle of mass conservation.

$$
\sum_{i=1}^{n} m_i = \sum_{i=1}^{n} \rho_i \cdot \dot{V}_i = \sum_{i=1}^{n} \rho_i \cdot v_i \cdot A_i = 0
$$
\n(1)

The sum of the mass flows \dot{m}_i and the volume flows \dot{V}_i , which depend on the densities ρ_i ,

Figure 2: Schematic view of material deposition: (a) filament; (b) hotend; (c) deposited foam material; (d) print bed

is 0. The conditions in an FFF process are shown in [Figure 2.](#page-3-1)

By considering the hotend as a flow system, the following assumption can be made, neglecting thermal effects.

$$
\rho_F \cdot v_F \cdot A_F = \rho_E \cdot v_E \cdot A_E \tag{2}
$$

Here, ρ_F is the density, A_F the cross-sectional area of the filament and v_F the feeding velocity as well as ρ_E the density, A_E the cross-sectional area of the extruded material and v_E the velocity of the material at the nozzle outlet.

Comminal et al. [17] simulated possible cross-sectional areas of the deposited strand – in this case the cross-sectional area of the extruded material A_E – taking into account thermal effects for bulk material in the FFF and found that these depend on the print head height

Figure 1: Foam printing process: from filament extrusion and loading over storage to printing

or layer height h_{PH} . The lower the print head height, the more the geometry of the cross-

sectional area changes from a round circle to a cuboid with rounded corners. This must be taken into account in the calculation. As a low h_{PH} leads to compressive forces on the already formed foam, this could compact the deposited material in the still lower viscous, soft state and increase the density of the extruded material ρ_E undesirably. For a circular cross-section, the following relationship is assumed with d_F , the diameter of the filament and d_E , the diameter of the extruded material.

$$
\rho_F \cdot v_F \cdot \frac{a_F^2 \cdot \pi}{4} = \rho_E \cdot v_E \cdot \frac{a_E^2 \cdot \pi}{4} \tag{3}
$$

When printing with solid material, the print head velocity v_{PH} is often calculated on the assumption that the processing material is an incompressible fluid. As the extruded crosssectional area A_E does not always correspond to the geometry specified in the printer's control system calculation model and the material is not always

completely incompressible, a multiplier M is used to compensate for these effects, which cannot be predicted exactly. Depending on the cross-section under consideration, the continuity equation is used, whereby h_E is often also regarded as the height of the extruded material and w_{E} is the width of the extruded material.

Circle

e:
$$
v_F \cdot \frac{d_F^2 \cdot \pi}{4} = v_{PH} \frac{d_E^2 \cdot \pi}{4} \cdot M
$$

angle:
$$
v_F \cdot \frac{d_F^2 \cdot \pi}{4} = v_{PH} \cdot h_F \cdot w_F \cdot M
$$
 (4)

Rectangle: $v_F \cdot \frac{d_F^{-2} \cdot \pi}{4}$ $\frac{a}{4} = v_{PH} \cdot h_E \cdot w_E \cdot M$

In FFF with thermoplastic foam, a further factor, the internal expansion velocity of the material v_l , must be taken into account. Depending on when the expansion takes place and in which direction a free expansion of the material is possible, v_{PH} can have an increasing or decreasing effect on the extruded material density. It is assumed that the time of expansion of the material is primarily triggered by the temperature of the material and the surrounding pressure. As hotends are not currently designed to be gas-tight, gaseous blowing agents can leak. It can also be assumed that the foaming process already begins in the hotend in some cases. If the material expands immediately after exiting the nozzle, it has free space if $v_{PH} \ge v_I$. This should be the ideal case. If $v_{PH} < v_I$, only d_E or w_E may increase due to the lack of free volume in the print direction and in the direction of the print head. If the expansion takes place later, d_E or w_E and simultaneously h_E can become larger, as the upward-limiting print head has already moved away. The requirement for this is that the molten material at this point has a sufficiently high temperature and the viscosity is low enough to allow pore growth. Ideally, the temperature of the material will have dropped to such an extent by the time pore growth is complete that the increased viscosity of the material prevents the structure from shrinking. The material should therefore cool down as quickly as possible after leaving the nozzle and expansion in order to achieve a minimum density ρ_E in the extruded strand.

The tests were carried out in a two-step approach. (1) Free-flow extrusion tests were carried out as a function of the heater temperature T_H and the feeding velocity v_F in order to classify the extrudability of the material. At the same time, the extrusion force F_E was recorded. The diameter d_E and the density of the extruded material ρ_E were measured at room temperature. (2) Single-path tower samples with an edge length of 50 mm and a height of 16 mm were printed in an FFF printer using the suitable parameters found. The heating element temperature T_H , the feeding velocity v_F , print head velocity v_{PH} , multiplier M and additionally the print head height h_{PH} were varied.

Materials

The material used in this work is a blend of different PLA types from the "Luminy® PLA" portfolio of "TotalEnergies Corbion". Specifically, the types "L175" and "LX975" were used. These PLA types differ mainly in the L-isomer content and thus in the crystallinity. The "L175" has an L-isomer content of \geq 99 % and has the highest crystallinity of the materials used. The "LX975" has an L-isomer content of 88 % and is completely amorphous. The crystallinity plays an important role both for gas absorption and for the subsequent foaming process in the printer. The only additive used was 0.5 % "Talc L88" as a nucleating agent.

The filaments were extruded on a "HAAKE PolyLab" twin-screw extruder and wound onto a filament spool after cooling in a water bath. The filaments produced are physically loaded with a blowing agent in an autoclave process. The blowing agent used is the gas 1,3,3,3tetrafluoropropene, which is also known under the trade name "HFO 1234Ze". It was chosen because it has good solubility in PLA. During the loading process, the filaments are placed in the autoclave and exposed to the blowing agent at a selected temperaturepressure-time regime. The pressure results from the vapour pressure of the selected temperature. The amount of gas is selected so that a saturated atmosphere with a constant concentration is created and this remains constant when it is injected into the filament. With this method of loading, blowing agent contents of 10 wt % - 20 wt % can be achieved in the filament. Due to these high blowing agent contents in the polymer compared to foam extrusion [16, 18], a certain proportion of the blowing agent must be present in liquid form in the autoclave in order to fulfil the above-mentioned condition. The exact blowing agent content in the filament depends on the temperature and the saturation time. Maximum saturation is reached after approx. 18 hours at the selected loading temperature. After treatment, the loaded filament is apparently indistinguishable from unloaded filament. However, it is characterised by a greatly increased flexibility compared to the previous state. This flexibility can be explained by the fact that the blowing agent also acts as a plasticiser.

After treatment, it can be observed that the blowing agent content in the filament decreases again over time. This is to be expected, as storage in an air atmosphere corresponds to a reversal of the loading. In order to ensure the longest possible shelf life of the filament, various storage conditions were investigated. With regard to the highest possible blowing agent content over time, storage at $<$ -15 °C is the most favourable. This preserves the blowing agent in a thermodynamically and kinetically favourable way. This means that blowing agent contents of over 10 wt % can be guaranteed even after storage for several months. In this study, the above-mentioned storage conditions were maintained and a processing time of less than two weeks was always achieved, so that a low loss of blowing agent can be assumed.

Experimental setup

An extrusion force measuring setup according to Fischer el al. [19] was initially used to process the filament loaded with blowing agent. This allows the extrusion force to be measured with an annular force sensor inside the hotend. The 3D printer used for the tests is a modified "German RepRap X400" as shown in [Figure 3.](#page-6-0) The loaded filament (a) is fed to the print head via a tubing system and transported towards the hotend with the 0.8 mm nozzle (c) using the feeding unit (b). The printer has a water cooling system for strong thermal separation of the hotend and feeding unit. After exiting the nozzle, the processing material foams and is applied to a heated print bed (d). The printer has an insulated print chamber (e). The control display (f) can be used to start print jobs and monitor the current status of the print. The density measurement was carried out on a precision scale "Precisa LS 220A" according to Archimedes' principle in water. Samples with a mass of 0.1 g were

used throughout the material characterisation. The measurement was repeated six times and the mean value was calculated. The microscope images were taken using a "Leica DVM6" optical microscope.

Foam extrusion characterisation

Experimental design foam extrusion

Extrusion characterisation is an important tool for determining initial boundary conditions for the FFF process. Filaments with a length of 60 mm were extruded through a nozzle with a diameter of 0.8 mm. The heating element temperature T_H was varied in 10 °C steps from 160 – 230 °C. At the same time, the feeding velocity v_F was changed from $3 - 12$ mm/s in steps of 1.5 mm/s. The test plan can be found in [Table 1](#page-15-0) in the Appendix. The extrusion force F_E was averaged over the measuring length. The diameter of the extruded material d_E was measured at

Figure 4: Extruded material sample

Figure 3: Experimental set-up: (a) filament entrance; (b) filament feeder;(c) hotend; (d) heated print bed; (e) insulated print chamber; (f) control display

five points on the cooled strand and the average was calculated. The density of the extruded material ρ_E was also determined.

Results foam extrusion

Not all parameter combinations allowed the filament to be extruded. Particularly low T_H caused the extrusion to stop at higher v_F . But even high T_H at low v_F had the consequence that the filament buckled immediately after the feeding unit due to the flexibility and an

additional temperature-related loss of stiffness of the filament. An example of an extruded foam strand is shown in [Figure 4.](#page-6-1)

The results of the foam extrusion tests are shown in [Figure 5.](#page-7-0) With increasing v_F , d_F increases until it levels off at approx. $d_E = 1.8$ mm after a maximum of $d_E = 2.2$ mm. The density ρ_E shows a strong temperature dependence at $v_F = 3$ mm/s and is smaller at lower T_H. The higher v_F becomes, the higher ρ_E becomes. The lowest density of ρ_E = 99.8 kg/m³ was achieved at $v_F = 4.5$ mm/s and $T_H = 180$ °C. The extrusion force F_E shows a similar behaviour, when v_F is increased. A higher T_H shifts the curve in the direction of lower F_E

Discussion foam extrusion

This analysis shows, that low densities of the extruded material ρ_E of less than 100 kg/m³ can be achieved. With regard to the diameter d_E measured during extrusion characterisation, it is assumed that this is dependent on the temperature of the extruded material and consequently on its viscosity and the cooling time. The longer the material remains in a plasticised state after extrusion, the greater the chance that the material will collapse before d_E is measured and a lower d_E will be measured. This means that ρ_E is also influenced to higher values by a possible collapse of the extruded material. Compensation through faster ν_F and thus lower heat input into the material is only possible to a small extent, as the example of the density reduction of approx. 84 kg/m³ at $T_H = 230$ °C from $v_F = 9$ mm/s to $v_F = 12$ mm/s shows. The higher extrusion force F_E measured under these conditions is noticeable, which could be an indication for the assumption of pressuretriggered expansion. Consequently, a low *ρ^E* is unexpectedly only achieved with a low heater temperature T_H . As a result, only low v_F can be set in order to achieve the lowest ρ_E . Low T_H and high v_F mean, that the filament does not achieve a sufficiently low viscosity and therefore cannot be fed through the nozzle. This can confirm the result of the extrusion force measurement. Stable extrusion with only slightly increasing F_E always occurs at $v_F \leq 6$ mm/s (see [Figure 5,](#page-7-0) green area), which does not always result in a low density. The increasing ρ_E with increasing T_H at $v_F = 3$ mm/s for example, suggests that due to the low v_F the expansion already takes place primarily inside the hotend and therefore only compacted, molten material emerges from the nozzle.

Foam printing

The experimental design of foam production in FFF with a physical blowing agent-loaded filament must be prepared differently than in FFF with bulk material due to the expansion of the material that must be taken into account. A single-path, square geometry (see [Figure 6\)](#page-8-0) with a height of H_P was selected as the test specimen and up to four of the test geometries were loaded into the "Simplify3D" software as a slicer for each print job.

A skirt (a) around each test specimen should guarantee sufficient material output from the start of the test specimen print. The first layer (c) was printed at half the print head height (here: layer height) h_{PH} . The heater temperature T_{H_1} , the print head velocity v_{PH_1} , the extrusion width w_E , the print head height (here: layer height) h_{PH} and the multiplier M were identified as the most important variables to be set in the slicer and were specified individually for each test geometry. T_H was varied between 160 °C and 180 °C, starting at 170 \degree C, based on the findings for the lowest possible density from the extrusion tests. The reason for choosing $T_H = 170$ °C is that the lowest densities were achieved there for the velocities explained below. Velocities of 25 mm/s to 75 mm/s were selected for v_{PH} . It started with v_{PH} = 50 mm/s. In the path planning, the slicer calculates the required volume of polymer according to the principle of mass conservation described in Eq. [\(4\).](#page-4-0) However, a rectangular cross-sectional area of the extruded material A_E is assumed here. Since the formation of a circular A_E was assumed here, the extrusion width w_E and the print head height (layer height) h_{PH} were selected as the same in the settings and varied from w_E = 1.49 – 2.15 mm based on the limit diameter from the extrusion tests of $w_E \approx 1.9$ mm, initially in 0.25 mm steps and later in relative dimensions of 5 % of the initial value. Since the height of the test specimen H_P was specified as constant due to dimensional accuracy, the test specimen had a different number of layers n_L , which can be calculated as follows after subtracting the first layer printed with half the height.

$$
n_L \approx \frac{H_P - \frac{hp_H}{2}}{hp_H} \tag{5}
$$

The value was rounded to integer. The multiplier M was calculated by transforming Eq. [\(4\),](#page-4-0) whereby A_E was assumed to be quadratic and feeding velocities of $v_F \approx 4 - 11$ mm/s were specified. After conversion, this corresponds approx. to the values of $v_F \approx 3 - 9$ mm/s for the circular cross-section from the extrusion test, which resulted in a low density. The print bed temperature was set to 60 °C and a thin layer of "Magigoo Original" adhesive was applied to the print bed to improve adhesion. The complete test plan can be found in the Appendix in [Table 2.](#page-16-0) After printing, the actual extrusion width $w_{E,m}$ and the actual height of

the test specimen $H_{P,m}$ were measured at three points on each side of the cuboid and the respective mean value was determined. The measured, averaged layer height $h_{E,m}$ was determined after removal of the first layer in accordance with Eq. [\(5\)](#page-8-1).

Figure 6: Foam print test part: (a) skirt; (b) layers; (c) first layer

$$
h_{E,m} \approx \frac{H_{P,m} - \frac{h_{PH}}{2}}{n_L} = \frac{H_{P,m} - \frac{h_{PH}}{2}}{\frac{H_P - \frac{h_{PH}}{2}}{h_{PH}}}
$$

The density ρ_E was measured on each side of the cut samples and the average value was calculated.

Results foam printing

The overall results of the tests are summarised in [Table 3](#page-17-0) in the Appendix. The first printing tests showed strong fluctuations and there were results with either underextruded [\(Figure 7,](#page-9-0) left) or over-extruded [\(Figure 7,](#page-9-0) right) test samples. A total of ten of these results could not be utilised as the material did not adhere to the print bed. A further five test specimens collapsed due to under-extrusion and no clear height of the test specimen $H_{P,m}$ could be determined. In addition, there was a strong influence of the set parameters on each other so that suitable, smaller variation differences had to be found first. An example of an optically good component with a low density ρ_E = 113 kg/m³ is shown in the centre of [Figure 7.](#page-9-0) A microscopic image of the crosssection of this component can be seen in [Figure 8.](#page-9-1) The pores are larger at the edge of the cross-section than inside. Despite the low density, it can be concluded that the expansion inside the strand is not

sufficiently triggered by T_H .

Discussion foam printing

Firstly, a comparative test shows the reproducibility of the results P05, P72, P05, P72, P78 – P82 at $T_H = 170 \text{ °C}$, v_{PH} = 50 mm/s, w_E = 1.9 mm and $M = 0.1$ mm. The mean value of the density is $\rho_E \approx 118 \text{ kg/m}^3$ with a standard deviation of approx. 6.4 %. The average of the actual extrusion width is $w_{Em} \approx 1.86$ mm with a standard deviation of approx. 2.8 %, the measured average layer height $h_{E,m} \approx 1.82$ mm with a standard deviation of approx. 0.7 %. This means that the values of $w_{E,m}$ and $h_{E,m}$ are lower overall than the specified 1.9 mm, but are only subject to minor fluctuations. The density is exposed to somewhat greater fluctuations. This may be due to non-uniform blowing agent

(6)

Figure 8: Optical microscope image of the cross section of the extruded strand printed with P35 parameters

Figure 7: Printed foam test parts of poor (left and right) and good (middle) quality: (a) no print bed adhesion; (b) material carried over due to under-extrusion; (c) material lateral evasion due to over-extrusion

distribution in the filament and the different densities in different test part areas. The mean values mentioned are summarised below as the result for test P88. Three different scenarios for interpretation are explained with the aim of finding suitable parameter sets.

Scenario 1 analyses the influence of the print head velocity v_{PH} on the print result. The tests P01, P19, P20, P27 – P37, P88 are considered here, where $T_H = 170$ °C is constant. w_E is analysed in two configurations at 1.65 mm and 1.9 mm and M at 0.1 and 0.11. v_{PH} is varied between 50 mm/s and 75 mm/s in 5 mm/s steps. The density distribution over the print head velocity is shown in [Figure 9.](#page-10-0) No adhesion could be achieved at higher v_{PH} and therefore these cannot be analysed. At $w_E = 1.9$ mm, ρ_E increases steeply over v_{PH} and here also due to the constant M over v_F , which agrees with the results from the extrusion tests. The earlier print error at $w_E = 1.9$ mm can be explained by a smaller v_F compared to w_E = 1.65 mm. At w_E = 1.65 mm, ρ_E shows no clear tendency. This could be due to the measurement uncertainty of ρ_E . The result of the dimensional measurement via v_{PH} is shown in [Figure 10.](#page-10-1) At $w_E = 1.9$ mm, $w_{E,m}$ and $h_{E,m}$ drop sharply compared to the target value. This correlates with the increase in ρ_E and suggests that the blowing agent could not be fully activated at high v_F and expansion took place to a lesser extent. w_{Em} and h_{Em} show no clear trend at $w_E = 1.65$ mm, but are not the same at the respective print head velocity. The deposited cross-section is therefore not round. The change from $w_{Em} > h_{Em}$ to $w_{Em} < h_{Em}$ at $v_{PH} \approx 58$ mm/s can be interpreted as a change from a lateral over-extrusion to an under-extrusion. Considering the model from [Figure 2](#page-3-1) this could mean that a change from $v_{PH} < v_I$ to v_{PH} > v_I takes place here.

Scenario 2 considers experiments P82 – P87. There, $T_H = 170$ °C, $v_F = 7.5$ mm/s and $M = 0.1$ is constant. v_{PH} is changed from 50 mm/s to 75 mm/s in 5 mm/s steps and w_E is changed from 1.9 mm to 1.55 mm in accordance with the definition in Eq. [\(4\).](#page-4-0) The resulting ρ_E is plotted against w_E in [Figure 11.](#page-11-0) With increasing w_E , ρ_E tends to increase. This corresponds to the opposite of scenario 1, in which increasing v_{PH} also resulted in lower ρ_E . However, with a difference of $\Delta\rho_E \approx 14$ kg/m³ over the v_{PH} range,

the effect is much smaller than the difference $\Delta\rho_E\approx 38$ kg/m³ in scenario 1. The constant v_F results in a more constant ρ_E than a constant w_E. In other words, a good processing window is found here for the aim of a low density over different v_{PH} values. The comparison of w_{Em} and h_{Em} with the set extrusion width w_E shows that this was achieved in the height $h_{E,m}$, but initially not in the width $w_{E,m}$ [\(Figure 12\)](#page-11-1). This means that, as initially assumed, the extruded cross-section is more rectangular at low h_{PH} and thus w_E . With the simultaneously high v_{PH} and low v_F , it is possible that not enough material is extruded. However, since ρ_E is comparatively very low, it is assumed that $v_1 \approx v_{PH}$. This is because when v_{PH} decreases, i.e. $v_{PH} < v_I$, $w_{E,m}$ increases relative to $h_{E,m}$ until it exceeds $w_E = 1.9$ mm and over-extrusion occurs.

Figure 11: Scenario 2: extruded material density ρE as a function of extrusion width wE,m

Extrusion width, $w_{\vec{E}}$ in mm

÷

 $wE.m \longrightarrow hE.n$

Scenario 3 shows the selection of tests P01, P05, P35 – P42. v_{PH} = 50 mm/s constant, v_F at 6.2 mm/s and 7.5 mm/s, h_{PH} at 1.65 mm and 1.9 mm and M at 0.1 and 0.11 for two configurations, $T_H = 160 - 180$ °C in 5 °C steps are considered. The result for ρ_E is shown in [Figure 13.](#page-12-0) As with the extrusion tests, ρ_E is lowest at low T_H . The almost equal ρ_E at the respective T_H indicates a strong effect of the calculation of the set parameters with Eq. [\(4\)](#page-4-0) on ρ_E relatively independent of the temperature. A difference is only observed at T_H = 180 °C. [Figure 14](#page-12-1) could provide information on this. The measured, averaged layer height h_{Em} and the measured extrusion width w_{Em} decrease with increasing T_H . At h_{PH} = 1.65 mm, h_{Em} is usually lower than w_{Em} . All other parameters being equal, this could mean a collapse of the paths caused by the temperature, which in turn corresponds with the experience from the extrusion tests. At h_{PH} = 1.65 mm, the difference between $h_{E,m}$ and $w_{E,m}$ is greater than at h_{PH} = 1.9 mm. This indicates over-extrusion and therefore v_{PH} < v_I may always apply. As the set width w_E is reached at low $T_H(w_{E,m} \approx w_E)$, it is assumed that the low h_{PH} leads to a reduction in h_{Em} . At higher T_{H_I} the collapse of the material probably predominates.

Figure 14: Scenario 3: extruded material width $w_{E,m}$ and height h_{Em} as a function of heater temperature T_H at two different print heights h_{PH} (dashed lines) and Multipliers M

Conclusion and outlook

With the help of a new method of loading thermoplastic filaments with a physical blowing agent, it was shown that polymer foam components with a low density of approx. 105 kg/ $m³$ can be produced using FFF. Suitable parameters for use in the printer were found by studying the properties during extrusion. A parameter study with an FFF printer made it possible to analyse the relationships between printing temperature, feeding velocity, print head velocity, print head height, extrusion width and multiplier. This made it possible to identify suitable parameter combinations supported by a calculation model.

In the future, larger investigations, e.g. by means of Design of Experiments (DoE), will be necessary for an even more precise investigation of the processing window. In addition, the influence of the selected parameters on pore size and mechanical properties must be analysed. This will also allow the poor adhesion of the extruded material paths to each other, which is assumed due to the circular cross-section, to be verified.

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Appendix

Table 1: Test plan foam extrusion

Table 2: Test plan foam printing

Table 3: Overall results foam printing

