Current challenges in thermodynamic aspects of rubber foam

Supitta Suethao¹, Worachai Ponloa¹, Saree Phongphanphanee¹, Jirasak Wong-Ekkabut¹,² & Wirasak Smithipong¹,³,⁴*

Natural rubber (NR) foam can be prepared by the Dunlop method using concentrated natural latex with chemical agents. Most previous studies have focused on the thermodynamic parameters of solid rubber in extension. The main objective of this study is to investigate the effect of the NR matrix concentration on the static and dynamic properties of NR foams, especially the new approach of considering the thermodynamic aspects of NR foam in compression. We found that the density and compression strength of NR foams increased with increasing NR matrix concentration. The mechanical properties of NR foam were in agreement with computational modelling. Moreover, thermodynamic aspects showed that the ratio of internal energy force to the compression force, $F_u/F$, and the entropy, $S$, increased with increasing matrix concentration. The activation enthalpy, $\Delta H_a$, also increased with increasing matrix concentration in the NR foam, indicating the greater relaxation time of the backbone of the rubber molecules. New scientific concepts of thermodynamic parameters of the crosslinked NR foam in compression mode are proposed and discussed. Our results will improve both the knowledge and the development of rubber foams based on the structure–properties relationship, especially the new scientific concept of the thermodynamical parameters under compression.

Polymer foams are formed by adding gas bubbles to the polymer matrix, leading to porosity and polymer continuous phases. The porosity, otherwise known as the cells, can be classified from its structure as either open–cell or closed–cell. The open–cell foam drives the flexibility, whereas the closed–cell foam drives the rigidity. Therefore, product applications depend on the structural properties of the foam¹–⁴.

Natural rubber (NR) foam is an interesting natural polymer foam which can be made into lightweight products and is suitable for comfort applications such as pillows and mattresses. Generally, rubber foams are porous, elastic and have a ventilated surface⁵,⁶. Rubber foam has good elasticity⁷, and its mechanical properties can be tuned with the choice of the type of latex, foam structure, and filler loading, for example⁷–¹⁰.

The most important characteristic of a rubber material is that it consists of long flexible molecules¹¹. Rubber molecules have a backbone of many covalent bonds which can rotate rapidly because of thermal agitation. Such long molecules convert their form easily at specific temperatures due to Brownian motion¹²,¹³. When no force is applied, they make random conformations but may adopt specific conformations if an external force is loaded. Changes in rubber elasticity are associated with changes in the configurational entropy and the system's internal energy during the deformation process⁷. For any elastic mechanism in extension mode, the stretching force is proportional to the temperature at a given state of strain. The thermodynamic study of the elastic deformation in rubber has shown that the stress depends significantly on the temperature in the extended state. It is possible to derive the separate contributions of the internal energy and entropy to the deformation process. The early experiments of Meyer and Ferri¹⁴ showed this statement to be substantially correct over a wide range of temperatures, provided that the extension was sufficiently large. The elastic behaviour seems to be irregular at lower strains because the force increases more slowly than would be expected from the theory. The deviations from the theoretical form of the force–extension curve for mechanical extension have been extensively studied by previous workers¹⁵–¹⁷; however, Mooney and Rivlin proposed a semi–empirical formula consistent with the experiment, referred to as the Mooney–Rivlin equation¹⁸,¹⁹.

¹Specialized Center of Rubber and Polymer Materials in Agriculture and Industry (RPM), Department of Materials Science, Faculty of Science, Kasetsart University, Bangkok 10900, Thailand. ²Department of Physics, Faculty of Science, Kasetsart University, Bangkok 10900, Thailand. ³Office of Research Integration on Target-Based Natural Rubber, National Research Council of Thailand (NRCT), Bangkok 10900, Thailand. ⁴Office of Natural Rubber Research Program, Thailand Science Research and Innovation (TSRI), Bangkok 10900, Thailand. ⁵Email: fsciwssm@ku.ac.th
When non-Gaussian theory has been applied to the experimental data at high strain process, the possible effects of strain-induced crystallisation on the mechanical properties of solid rubber have not been taken into account. The non-Gaussian theory of the strain state is significant within the crystallisation region, and it has been suggested that the rise of the force–deformation curve should be attributed primarily to this factor. The crystallisation that occurs would certainly be expected to produce some stiffening of the rubber, but it is not easy to predict the magnitude of the effect. However, it has been noted by Wang and Guth that the characteristic form of the force–extension curve for natural rubber is only slightly affected by increasing the temperature to 100 °C, though this will substantially reduce the crystallinity. Most existing studies investigated the solid rubber in extension mode; fewer studies have focused on the thermodynamics of polymer foams, especially rubber foam. In the present work, a new scientific concept of the thermodynamic parameters of the natural rubber (NR) foam is investigated. The mechanical properties are modelled. In particular, we consider some features of the compression of crosslinked rubber foams which are still not fully understood.

Materials and methods
Materials. The concentrated natural latex (60% dry rubber content and 1.7% non-rubber content), from Hevea brasiliensis in the southern part of Thailand, was supplied from Num Rubber & Latex Co., Ltd., Trang, Thailand. The chemical agents consist of 10% potassium oleate solution, 50% sulphur dispersion, 50% zinc diethylthiocarbamate (ZDEC) dispersion, 50% zinc-2-mercaptopbenzothiazole (ZMBT) dispersion, 50% Wingstay L dispersion, 50% zinc oxide (ZnO) dispersion, 33% diphenylguanidine (DPG) dispersion, and 12.5% sodium silicofluoride (SSF) dispersion. All the chemical agents were supplied from Thanodom Technology Co., Ltd., Thailand. The chemical agents used for the rubber foam preparation are summarised in Table 1.

Rubber foam preparation. The rubber foams were prepared in the following way: first, concentrated natural latex was stirred 80 rpm in a blender for 1 min to remove the ammonia. Second, the potassium oleate solution was added as the stirring speed was increased to 160 rpm for 10 min. Then, dispersions of sulphur, ZDEC, ZMBT and Wingstay L were added to the rubber compound with the stirring speed decreased to 80 rpm for 1 min. Next, ZnO and DPG dispersions were added into the rubber compound at the same mixing speed for 1 min. After that, the SSF dispersion was added into the rubber compound and mixing continued until the rubber foam had nearly reached the gel point. Finally, the rubber foam was transferred to a mould and allowed to set for 45 min. The vulcanisation of the rubber foam was performed by a hot air oven at 90 °C for 2 h. The rubber foam was then removed from the mould, washed, and dried in the hot air oven at 70 °C for 4 h.

Rubber foam characterisation. The density (kg/m³) of the foam was evaluated by the relationship between the weight (kg) and volume (m³) of the foam as described elsewhere. The chemical functional groups present in the foam sample were measured by Attenuated Total Reflection-Fourier Transform Infrared (ATR-FTIR) spectroscopy with a Ge crystal probe (VERTEX 70, Bruker, Billerica, MA, USA).

The morphology of foam samples was examined by a scanning electron microscope (SEM, FEI, Quanta 450, Eindhoven, Netherlands). The foam sample was coated with gold, and three replicants of each foam formula were tested. ImageJ software was used to evaluate the average pore size and porosity of the foam samples. The cell density (d) of the foam sample was calculated as in a previous study for comparison with the solid phase density of natural rubber (NR 0.93 g/cm³).

The computational modelling of the mechanical properties of the foam sample used the hyperfoam-polynomial strain energy function from 1st to 6th order. The finite element method (FEM) and curve-fitting analysis of foam sample data were performed using ABAQUS for uniaxial compression.

The relationship between the stress, σ, and the compression limit, λ, of each foam sample was plotted on the Mooney–Rivlin Eq. where C₁ and C₂ were the constant values:

| Chemical agents | Control + 10% NR or control added 10% NR (g) | Control (g) | Control − 10% NR or control reduced 10% NR (g) |
|-----------------|--------------------------------------------|------------|--------------------------------------------|
| 60% concentrated natural latex | 183.33 | 166.67 | 150.00 |
| 10% potassium oleate solution | 16.50 | 16.50 | 16.50 |
| 50% sulphur dispersion | 4.00 | 4.00 | 4.00 |
| 50% ZDEC dispersion | 2.00 | 2.00 | 2.00 |
| 50% ZMBT dispersion | 2.00 | 2.00 | 2.00 |
| 50% Wingstay L dispersion | 2.00 | 2.00 | 2.00 |
| 50% ZnO dispersion | 10.00 | 10.00 | 10.00 |
| 33% DPG dispersion | 2.00 | 2.00 | 2.00 |
| 12.5% SSF dispersion | 8.00 | 8.00 | 8.00 |

Table 1. Chemical agents used for the preparation of various foam samples.
The crosslinking density of the foam samples was evaluated by the swelling method according to the Flory–Rehner equation. We also used the Flory–Huggins equation to calculate the change in Gibbs free energy, $\Delta G$, and entropy, $\Delta S$, as follows:

$$ \frac{\sigma}{\left(\lambda - \frac{1}{\lambda^2}\right)} = 2C_1 + 2C_2 \frac{1}{\lambda} $$

where $V_r$ is the volume fraction of foam sample in the rubber network, $\chi$ is the parameter between the foam sample and the solvent interaction (defined as 0.43 + 0.05 $V_r$), $R$ is the ideal gas constant (8.3145 J/mol·K), and $T$ is the test temperature (298.15 K).

The thermodynamic parameters of the foam sample compression were measured by the texture analyser (TA.XT Plus, Stable Micro Systems, Godalming, Surrey, UK). The foam sample was also incubated at different temperatures (298.15, 308.15, 318.15, 328.15, and 338.15 K) during the compression process from 10 strain to 70% strain. Next, the relationship between force and temperature was plotted to obtain the ratio of internal energy to the compression force, $F_u/F$.

The activation enthalpy of the transition process, $\Delta H_a$, of the foam sample was evaluated by dynamic mechanical analysis (DMA1, Mettler Toledo, Columbus, OH, USA) from −193.15 to 353.15 K. From this, the $\Delta H_a$ value of the foam sample can be calculated as described elsewhere.

Results and discussion
Here we investigate the underlying thermodynamic relations of these rubber foam phenomena in more detail and explore how they may be applied experimentally to obtain quantitative information about the deformation process mechanism.

Physical and morphological properties. First, we discuss the chemical function of the rubber foams. Figure 1 presents an ATR–FTIR spectrum of the control sample from 500 to 4000 cm$^{-1}$, which confirmed the chemical functional group of crosslinked NR foam. However, there are no significant differences between the spectra of the foam samples, even at different rubber concentrations. To investigate the density of the foam foams (Table 2), rubber foam samples of each type with the same volume ($4.86 \times 10^{-5} m^3$) were prepared. Each rubber foam sample was weighed in kilograms. The results showed that the amount of matrix has a significant effect on the density of the foam sample. The density of the foam sample decreased by around 10% when the matrix was reduced by 10%, while the addition of 10% matrix content increased the density of the foam sample of around 10%.
In terms of the morphological properties, the images obtained from the SEM are presented in Fig. 2. We found that all the foam samples exhibited the open-cell structure with heterogeneous sizes of bubble (heterogeneous pore size), a known effect of the Dunlop process\textsuperscript{34,35}. However, the amount of NR matrix present affects the morphological properties of the foam samples. The foam of samples with a higher NR concentration is more interconnected (Fig. 2a–c), which agrees with the density measurements. The ImageJ software\textsuperscript{23} was used to evaluate the morphological parameters of the foam samples (Fig. 2d–f and Table 2). In Fig. 2d–f, the white areas are related to the interconnected foam, whereas the black areas relate to the pore or cell.

| Sample            | Foam density (± 3 kg/m$^3$) | Average pore size (± 150 µm) | Porosity (± 1.00%) | Cell density (± 500 cm$^{-3}$) |
|-------------------|-----------------------------|------------------------------|--------------------|--------------------------------|
| Control + 10% NR  | 123                         | 352                          | 45.97              | 38,054                         |
| Control           | 110                         | 522                          | 48.73              | 11,844                         |
| Control − 10% NR  | 99                          | 555                          | 55.37              | 9,966                          |

Table 2. Foam density, average pore size, porosity, and cell density of various foam samples.

Mechanical properties. Figure 3 shows the stress–strain curves from experiments and hyperfoam material modelling of rubber foam samples. For the reduced polynomial model used in this study, the 6th order is in good agreement with the experimental result. Table 3 presents the parameters from the fit to the 6th order,
and the computer modelling of the foam samples in compression is in good agreement with the experimental results (Fig. 3). Concerning the compression strength, which is the maximum compression stress at 75% from the rubber foam surface, we found that the compression strength increases with increasing NR matrix content. This result is in agreement with the density of the foam samples. Generally, NR exhibits good mechanical properties because it has a high molecular weight. Interestingly, the mechanical property, compression strength, is more sensitive to the matrix content than the density. The compression strength of rubber foam is decreased around 37% by a 10% reduction in matrix content while 10% addition to the matrix content results in an increase of compression strength of around 23%. In general, the compressive axial tests of foams show three main distinct regions. The deformation process starts with an initial linear elastic response on cell edges or cell walls. Deformations are increasing and lead to the cells to collapse while stresses remain roughly unchanged, known as the plateau region. This effect results in the ability to absorb impact and vibrating loading. This collapse progresses until opposing walls meet and touch. After the opposing walls touch, the deformation stops with increasing stresses (densification or locking) i.e. cellular solids exhibit deformation until the densification is reached. Alzoubi et al. showed that the natural latex is highly elastic and low viscous characteristics than any other foam samples such as Polyurethane. Therefore, the latex material has no clear distinct three regions as the case for other samples. However, at high strains, the cell walls of NR foam are completely collapsed: higher compression strength is represented by a higher concentration of the NR matrix.

The mechanical properties of rubber foam can also be investigated in terms of the stress, $\sigma$, and the compression limit, $\lambda$, based on the Mooney–Rivlin equation. We found that the slope of the straight line above 1.5 at x-axis for all samples is almost identical. The y–intercept for all samples is related to the crosslinking density of the foam samples; higher crosslinking density is represented by a higher absolute value of the y–intercept. The resulting pattern of curves is in good agreement with the Mooney–Rivlin experiment, although our curves were in the opposite direction compared to the Mooney–Rivlin result. However, the Mooney–Rivlin result was obtained in extension mode of solid rubber while our study was in compression mode of rubber foam.

**Thermodynamic aspects.** Based on the crosslinking density of the foam samples, a foam with higher NR matrix concentration possesses a higher volume fraction of rubber, $V_r$, and greater crosslinking density. We can calculate the change in the Gibbs free energy, $\Delta G$, and entropy, $\Delta S$, from the swelling test based on the Flory–Huggins equation, results are given in Table 4. A negative $\Delta G$ was found for all the foam samples, and the $\Delta G$ decreases with increasing NR matrix concentration. Moreover, $\Delta S$ increases with increasing matrix concentration, which indicates favourable thermodynamics. This is because of the NR matrix concentration on the foam sample: rubber with good mechanical properties (high compression strength and modulus) and high relaxation stress result in a more thermodynamically favourable system.
It is extremely interesting to investigate the thermodynamic aspects of rubber foam related to the mechanical properties in more detail. We focused on the compression of the foam samples. The compression force came exclusively from an entropic mechanism, for example, from the tendency of the rubber molecule to transform to random conformations. The compression force was then directly proportional to the absolute temperature. The model of the unloaded foam sample, which corresponds to a high degree of freedom for the rubber molecules, and the model of the loaded foam sample from compression, which corresponds to a lower degree of freedom for the rubber molecules. When the compression force is unloaded, the foam sample returns to its original shape, which is more favourable in terms of the entropy, \( S \), of the entangled molecules.

Consider the consequence of an external force causing compression on a rubber foam. The first law of thermodynamics can be written as:

\[
\dot{U} = \dot{Q} - \dot{W}
\]

Where \( \dot{U} \) is the rate of change of internal energy, \( \dot{Q} \) is the rate of heat transfer, and \( \dot{W} \) is the rate of work done. The relationship of stress, \( \sigma \), and compression limit, \( \lambda \), of foam samples based on the Mooney–Rivlin Eq. 26. The relative scatter on the results is estimated equal to about 5%.

**Figure 4.** The relationship of stress, \( \sigma \), and compression limit, \( \lambda \), of foam samples based on the Mooney–Rivlin Eq. 26.

**Table 4.** Thermodynamic parameters determined from the crosslinking density of various foam samples based on the Flory–Huggins equation.

| Sample          | Volume fraction of rubber, \( V_r \) (± 0.001% a.u.) | \( \Delta G \) (J/mol) | \( \Delta S \) (J/mol K) |
|-----------------|------------------------------------------------------|------------------------|--------------------------|
| Control + 10% NR| 0.2605                                               | −36.7143               | 0.1231                   |
| Control         | 0.2595                                               | −36.3674               | 0.1220                   |
| Control − 10% NR| 0.2575                                               | −35.6386               | 0.1195                   |

**Figure 5.** Unloaded (left) and loaded at z–direction (right) models of rubber foam samples. Unloaded rubber molecules (pink lines) have freedom to move, whereas loaded rubber molecules cannot move and rubber foam is expanded at x– and y–directions due to the compression. The model was drawn using Adobe illustrator software version 24.2.3.
where \( dU \) is the change in the rubber foam’s internal energy resulting from the absorption of heat, \( dQ \), and the distribution of work, \( dW \), on it by the external force. If we assume that the compression process is reversible based on the porous structure of rubber foam, the heat flow can be expressed as:\(^4\):

\[
dQ = T dS
\]

where \( T \) is the temperature and \( dS \) is the change in entropy, and thus:

\[
dU = T dS + dW
\]

This equation concerns the reversible compression of a foam sample. The work is done by applying a force, \( F \), to the foam, resulting in the change of length, \( dL \), from its original length. From the perspective of porous structure of rubber foam, when a foam is compressed at the \( z \)-direction, it can be expanded at the \( x \)- and \( y \)-directions. So, the uniaxial work done on the foam based on the assumption of constant volume is thus:

\[
dW = F dL
\]

Then, we combine (4), (5), (6), and (7) and produce:

\[
dU = T dS + dW
\]

We can take the partial differential of (8) with respect to \( L \) as follows\(^1\):

\[
\frac{dU}{dL} = T \frac{dS}{dL} + F
\]

\[
F = \frac{dU}{dL} - T \frac{dS}{dL} = F_u + F_s
\]

where \( F_u = dU/dL \) and \( F_s = -T(dS/dL) \). These two thermodynamic parameters relate to the internal energy and entropy changes on compression of the rubber foam sample.

Equation (9) is of basic significance in rubber elasticity since it provides a direct measurement of the changes of the internal energy and the entropy during a deforming. Its application is illustrated by Figs. 6, 7, and 8, in which the linear curve represents the variation with temperature of the force at a constant compressive strain. From (9), the slope of this curve gives the entropy change per unit compression, \( dS/dL \), for isothermal compression at the temperature \( T \). Correspondingly, the \( y \)-intercept at \( T = 0 \) is \( dU/dL \), the change of internal energy per unit compression.

Thus, the internal energy and entropy contributions to the force at any given compression strain can be obtained by the experimental force–temperature curve of the foam samples with different compression strains and temperatures (Figs. 6, 7, 8). The internal energy and entropy terms are independent of the temperature when the force–temperature curve is linear. However, at high compression strains, the effect of crystallisation could become significant. Strain–induced crystallisation of solid uncrosslinked rubber has previously been reported\(^4\), while stress–induced crystallisation of crosslinked rubber foam has been reported\(^7\). Figures 6, 7 and 8 show the increase of compression force with increasing compression strain (from 10 to 70% strain) in all types.

**Figure 6.** Force–temperature curves at a given strain for the control + 10% NR sample with \( R^2 = 0.9 \) minimum at each strain. The scatter on the results is on the order of the size of the figures.
of foam sample studied. At a certain compression strain, the compression force seems to be stable, indicating the high–density foam samples which are in good agreement with a previous study on high–density rubber foams with silica filler7. Furthermore, the slope decreases slightly at higher strains, indicating the decrease degree of freedom of the rubber molecules.

Concerning the elasticity of the foam sample in compression mode (Figs. 6, 7, 8), samples under high length or low compression strain, L, exhibits high entropy, S, values. Thus, the entropy of a foam sample is proportional to the length at a constant temperature, as shown in Eq. (10).

\[ S \propto L \quad (10) \]

Table 5 shows the values of the \( F_u/F \) ratio of various foam samples at different compression strains and two temperatures (298.15 and 338.15 K). The values of \( F_u \) and \( F \) grow as the compression strain increases for all types of foam sample. Moreover, the \( F_u/F \) value also increases with increasing compression strain, indicating the stability of the entropy during the deformation process. This result agrees with previous work on rubber foam with high silica loading, which showed high density and better mechanical properties. While the NR matrix concentration affects the mechanical properties of the foam samples, the ratio of \( F_u/F \) relates to the thermodynamic aspects of the mechanical properties of the foam samples. This result is in good agreement with the calculations of \( \Delta G \)
and $\Delta S$. Moreover, the ratio of $F_u/F$ increases as the temperature decreases from 338.15 to 298.15 K. This can be explained by the effect of the flow property of natural rubber at higher temperatures. The $F_u/F$ values obtained from this study (0.7–0.9) are higher than those from previous studies: 0.1–0.2 values for uncrosslinked rubber in extension and 0.6–0.8 values for lower density crosslinked foam samples in compression. The difference in $F_u/F$ values could be due to the differences in the rubber structure and the test method used.

Table 5. Compression strain, $F_u$, $F$, and $F_u/F$ values of foam samples at 298.15 K and 338.15 K.

| Sample          | Compression strain (%) | $F_u$ (± 5% N) | $F$ (± 5% N) | $F_u/F$ | $F$ (± 5% N) | $F_u/F$ |
|-----------------|------------------------|---------------|--------------|---------|--------------|---------|
| Control + 10% NR| 10                     | 5.42          | 7.48         | 0.7248  | 7.75         | 0.6990  |
|                 | 20                     | 8.21          | 10.86        | 0.7556  | 11.22        | 0.7317  |
|                 | 30                     | 10.76         | 13.83        | 0.7780  | 14.25        | 0.7555  |
|                 | 40                     | 13.79         | 17.13        | 0.8050  | 17.57        | 0.7845  |
|                 | 50                     | 17.77         | 21.14        | 0.8406  | 21.59        | 0.8230  |
|                 | 60                     | 25.52         | 29.24        | 0.8726  | 29.74        | 0.8579  |
|                 | 70                     | 39.75         | 41.96        | 0.9474  | 42.26        | 0.9408  |
| Control         | 10                     | 5.74          | 8.48         | 0.6766  | 8.85         | 0.6485  |
|                 | 20                     | 7.71          | 10.84        | 0.7112  | 11.26        | 0.6846  |
|                 | 30                     | 9.32          | 12.60        | 0.7398  | 13.04        | 0.7148  |
|                 | 40                     | 11.52         | 15.04        | 0.7660  | 15.51        | 0.7427  |
|                 | 50                     | 14.82         | 18.63        | 0.7952  | 19.15        | 0.7739  |
|                 | 60                     | 21.73         | 26.50        | 0.8200  | 27.14        | 0.8006  |
|                 | 70                     | 36.82         | 43.14        | 0.8535  | 43.99        | 0.8370  |
| Control − 10% NR| 10                     | 5.94          | 9.46         | 0.6281  | 9.93         | 0.5982  |
|                 | 20                     | 7.67          | 11.82        | 0.6494  | 12.38        | 0.6202  |
|                 | 30                     | 9.11          | 13.64        | 0.6677  | 14.25        | 0.6392  |
|                 | 40                     | 11.12         | 16.19        | 0.6869  | 16.87        | 0.6592  |
|                 | 50                     | 13.90         | 19.53        | 0.7115  | 20.29        | 0.6850  |
|                 | 60                     | 19.16         | 25.78        | 0.7433  | 26.67        | 0.7185  |
|                 | 70                     | 25.58         | 35.17        | 0.8126  | 36.05        | 0.7927  |

Figure 9. Internal energy contribution vs the compression force, $F_u/F$, of various foam samples: (a) 298.15 K, and (b) 338.15 K. The scatter on the results is on the order of the size of the figures.

The thermodynamic parameters of foam samples were also calculated from the perspective of dynamic mechanical analysis. The storage modulus, $E'$, and $\tan \delta$ of the foam samples as a function of temperature are presented in Fig. 10. In general, the storage modulus relates to the dynamic mechanical properties whereas $\tan \delta$ relates to the dissipation energy of a material. Rubber chains are freezing at the glassy plateau below the glass transition temperature, the foam sample with high NR content (control + 10% NR) represents a high...
storage modulus at the glassy plateau, indicating the lower free volume for high density samples. However, the storage modulus of other two samples is quite similar. The NR matrix concentration also affects the dissipation energy or hysteresis (maximum tan δ) of a foam sample. Samples with a high NR concentration possess more network structure, generating the low hysteresis. This result is in good agreement with the existing literature, and the hysteresis of the rubber foam could be due to either the molecular friction of short molecules or the reduced network structure.

Table 6 presents the dynamic mechanical parameters extracted from Fig. 10, from which the activation enthalpy, ∆H_a, of various foam samples can be calculated. This activation enthalpy focuses on the transition process between the glassy and rubber states of rubber molecules. This average value also corresponds to the relaxation of the backbone motion of rubber molecules. Interestingly, the NR matrix concentration again affects the activation enthalpy of the foam sample, where a higher NR concentration represents a higher average activation enthalpy, ∆H_a, due to the greater relaxation time of the rubber chains. These average activation enthalpy values are in good agreement with the previous literature.

Conclusions
In this study, we applied the Dunlop process to prepare rubber foam samples with different NR matrix concentrations. Then we investigated the thermodynamic relations of the system in more detail and how they may be applied experimentally to obtain quantitative information during the compression process. We found that the NR matrix content has a significant effect on the density and compression strength of the foam sample; nevertheless, the compression strength is more sensitive to the matrix concentration than the density. The foam samples have an open–cell structure with heterogeneous cell sizes, samples with higher NR matrix concentration exhibit higher interconnectivity and cell density.

The computational modelling using the hyperfoam model of the 6th order is in good agreement with the experimental result of the foam samples in the stress–strain curve. Moreover, the mechanical properties of the foam samples with different matrix concentrations are in good agreement with that of the Mooney–Rivlin experiment. However, our work was in compression mode while Mooney–Rivlin’s was in extension mode.

Based on the crosslinking density of the foam sample, the higher NR matrix concentration results in a higher volume fraction of rubber, V_r, and crosslinking density. When the Flory–Huggins equation is applied, the change in Gibbs free energy, ∆G, decreases, and the change in entropy, ∆S, increases with the increasing NR matrix concentration, which is thermodynamically favourable. The force and temperature relationship corresponded to the internal energy and entropy, which were experimentally determined by compression of foam samples. Interestingly, the ratio of the internal energy force to the compression force, F_u/F, increases with the NR matrix concentration, and the F_u/F ratio is in good agreement with the literature reviews. Thus, the change of foam length, ∆L, or compression strain, is directly influenced by the entropy change.

Table 6. Parameters obtained from the dynamic mechanical analysis of various foam samples.

| Sample               | E'_{glassy} at 203.15 K (± 5% MPa) | E'_{rubber} at 273.15 K (± 5% MPa) | T_g dynamic (K) | Tan δ max | t_A (area under tan δ peak) | ∆H_a (kJ/mol) |
|----------------------|------------------------------------|-----------------------------------|-----------------|-----------|---------------------------|----------------|
| Control + 10% NR     | 337.28                             | 0.37                              | 232.07          | 1.68      | 33.44                     | 143.36         |
| Control              | 293.53                             | 0.31                              | 229.98          | 1.80      | 34.50                     | 137.30         |
| Control – 10% NR     | 324.73                             | 0.49                              | 226.65          | 2.02      | 33.22                     | 131.28         |

Figure 10. Dynamic mechanical analysis of various foam samples: (a) storage modulus, E’, as a function of temperature, and (b) tan δ as a function of temperature. The relative scatter on the results is estimated equal to about 5%.
Dynamic mechanical analysis was applied to evaluate the activation enthalpy of the transition process, \( \Delta H_\text{a} \), of the foam samples. The results showed that a higher NR matrix concentration has the effect of increasing the average activation enthalpy related to the relaxation time of rubber molecules. Therefore, the NR matrix concentration affects the static and dynamic parameters resulting from the relationship between the rubber foam structure and the material properties. New approaches in the thermodynamic aspects of foam samples related to the matrix concentration effect were investigated and proposed.

**Data availability**

Correspondence and requests for materials should be addressed to W.S.

Received: 5 January 2021; Accepted: 1 March 2021
Published online: 17 March 2021

**References**

1. Mohabbi, A., Mighri, F., Aji, A. & Rodrigue, D. Current issues and challenges in polypropylene foaming: a review. *Cell. Polym.* 34, 299–338 (2015).
2. Andrieux, S., Quell, A., Stubenrauch, C. & Drenckhan, W. Liquid foam templating: a route to tailor-made polymer foams. *Adv. Colloid Interface Sci.* 256, 276–290 (2018).
3. Trofa, M., Di Maio, E. & Maffettone, P. L. Multi-graded foams upon time-dependent exposition to blowing agent. *Chem. Eng. J.* 362, 812–817 (2019).
4. Suetsao, S., Shah, D. U. & Smithhipong, W. Recent progress in processing functionally graded polymer foams. *Materials* 13, 4060. [https://doi.org/10.3390/ma13184060](https://doi.org/10.3390/ma13184060) (2020).
5. Najib, N. N., Ariiz, Z. M., Bakar, A. A. & Sipaut, C. S. Correlation between the acoustic and dynamic mechanical properties of natural rubber foam: effect of foaming temperature. *Mater. Des.* 32, 505–511 (2011).
6. Yuan, Q. et al. Comparison of fire behaviors of thermally thin and thick rubber latex foam under bottom ventilation. *Polymers* 11, 88. [https://doi.org/10.3390/polym11010088](https://doi.org/10.3390/polym11010088) (2019).
7. Prasopdee, T. & Smithhipong, W. Effect of fillers on the recovery of rubber foam: from theory to applications. *Polymers* 12, 2745. [https://doi.org/10.3390/polym12112745](https://doi.org/10.3390/polym12112745) (2020).
8. Chollakup, R., Suwanruji, P. & Smithhipong, W. New approach on structure-property relationships of stabilized natural rubbers. *J. Polym. Res.* 26, 37. [https://doi.org/10.1007/s10965-018-1685-1](https://doi.org/10.1007/s10965-018-1685-1) (2019).
9. Sukup, R., Sun, Y., Sukatta, U. & Smithhipong, W. Foam rubber from centrifuged and creamed latex. *J. Polym. Eng.* [https://doi.org/10.org/10.1515/polymeng-2018-0219](https://doi.org/10.org/10.1515/polymeng-2018-0219) (2019).
10. Khuntawee, W., Suphopphatrongtong, T., Phongghanphanee, S., Karttunen, M. & Wong-ekkabut, J. Molecular dynamics study of natural rubber–fullerene composites: connecting microscopic properties to macroscopic behavior. *Phys. Chem. Chem. Phys.* 21, 19403–19413. [https://doi.org/10.1039/c9cp03155c](https://doi.org/10.1039/c9cp03155c) (2019).
11. James, H. M. & Guth, E. Theory of the elasticity of rubber. *J. Appl. Phys.* 15, 294–303 (1944).
12. Einstein, A. *Investigations on the Theory of the Brownian Movement* (Dover Publications, 1956).
13. Lampo, A., March, M. A. G. & Lewenstein, M. Quantum Brownian Motion Revisited: Extensions and Applications (Springer International Publishing, 2019).
14. Meyer, K. H. & Ferri, C. The elasticity of rubber. *Rubber Chem. Technol.* 8, 319–334 (1935).
15. Roe, R. J. & Krigbaum, W. R. The contribution of internal energy to the elastic force of natural rubber. *J. Polym. Sci.* 61, 167–183 (1962).
16. Treloar, L. R. G. *The Physics of Rubber Elasticity* (Oxford University Press, 1975).
17. Pakornpadungsit, P., Smithhipong, W. & Chworos, A. Self-assembly nucleic acid-based biopolymers: learn from the nature. *J. Polym. Res.* 25, 45. [https://doi.org/10.1007/s10965-018-1441-6](https://doi.org/10.1007/s10965-018-1441-6) (2018).
18. Smith, T. L. Large deformation tensile properties of elastomers: I: temperature dependence of c1 and c2 in the Mooney-Rivlin equation. *J. Polym. Sci. Part C Polym. Symp.* 16, 841–858 (1967).
19. Spalath, G. D. Polyurethane elastomers studied by the Mooney-Rivlin equation for rubbers. *J. Appl. Polym. Sci.* 43, 613–620 (1991).
20. Flory, P. J. Thermodynamics of crystallization in high polymers: I. Crystallization induced by stretching. *J. Chem. Phys.* 15, 397–408. [https://doi.org/10.1063/1.1746537](https://doi.org/10.1063/1.1746537) (1947).
21. Wang, M. C. & Guth, E. Statistical theory of networks of non-Gaussian flexible chains. *J. Chem. Phys.* 20, 1144–1157 (1952).
22. Hasany, S. M., Saeed, M. M. & Ahmed, M. Adsorption isotherms and thermodynamic profile of Co(II)—SCN complex uptake on polyurethane foam. *Sep. Sci. Technol.* 35, 379–394 (2000).
23. Schneider, C. A., Rasband, W. S. & Eliceiri, K. W. NIH Image to ImageJ: 25 years of image analysis. *Nat. Methods* 9, 671–675. [https://doi.org/10.1038/nmeth.2089](https://doi.org/10.1038/nmeth.2089) (2012).
24. Mase, G. T., Smelser, R. E. & Rossmann, J. S. *Continuum Mechanics for Engineers* (CRC Press, 2020).
25. Smith, M. *ABAQUS/Standard User's Manual, Version 6.9* (Dassault Systèmes Simulia Corp, 2009).
26. Fukahori, T. & Seki, W. Molecular behaviour of elastomeric materials under large deformation: 1: re-evaluation of the Mooney-Rivlin plot. *Polymer* 33, 502–508 (1992).
27. Flory, P. J. & Rehner, J. Statistical mechanics of cross-linked polymer networks II: swelling. *J. Chem. Phys.* 11, 521–526 (1943).
28. Smithhipong, W., Nardin, M., Schultz, I. & Suchika, K. Adhesion and self-adsorption of rubbers, crosslinked by electron beam irradiation. *Int. J. Adhes. Adhes.* 27, 352–357 (2007).
29. Crook, S. G. Application of the Flory-Rehner equation and the Griffith fracture criterion to paint stripping. *J. Coat. Technol. Res.* 7, 397–408. [https://doi.org/10.1007/s11894-009-9166-1](https://doi.org/10.1007/s11894-009-9166-1) (2009).
30. Tangboriboon, N., Rortchanakarn, S., Petchaoen, K. & Sirivat, A. Effects of foaming agents and eggshell calcium carbonate (CaCO3) filler on natural rubber foam physical-thermal: mechanical properties. *J. Rubb. Res.* 19, 71–96 (2016).
31. Phomrak, S., Nimpaiboon, A., Newby, B.-M. & Phisalaphong, M. Natural rubber latex foam reinforced with micro- and nano-bridged cellulose via Dunlop method. *Polymers* 12, 1599. [https://doi.org/10.3390/polym12091599](https://doi.org/10.3390/polym12091599) (2020).
32. Poyavanaraphan, T. & Magarapahan, R. Pre Vulcanized natural rubber latex/clay aerogel nanocomposites. *Eur. Polym. J.* 44, 1968–1977 (2008).
33. Sadeghi Ghari, H. & Jalali-Arani, A. Nanocomposites based on natural rubber, organoclay and nano-calci carbonate: study on the structure, cure behavior, static and dynamic-mechanical properties. *Appl. Clay Sci.* 119, 348–357 (2016).
34. Rathnayake, L., Ismail, H., Azahari, B., Darsanaisiri, N. D. & Rajapaks, S. Synthesis and characterization of nano-silver incorporated natural rubber latex foam. *Polym. Plast. Technol. Eng.* 51, 603–611 (2012).
35. Panploo, K., Chalermsinsuwan, B. & Poompredub, S. Natural rubber latex foam with particulate fillers for carbon dioxide adsorption and regeneration. *RSC Adv.* 9, 28916–28923 (2019).
36. Forest, C., Chaumont, P., Casagnau, P., Swoboda, B. & Sonntag, P. Polymer nano-foams for insulating applications prepared from CO2 foaming. *Prog. Polym. Sci.* 41, 122–145 (2015).
37. Smithipong, W., Nardin, M., Schultz, J., Nipithakul, T. & Suchiva, K. Study of tack properties of uncrosslinked natural rubber. *J. Adhes. Sci. Technol.* **18**, 1449–1463 (2004).
38. Messinger, R. J., Marks, T. G., Gleimam, F. S., Milstein, F. & Chmelka, B. F. Molecular origins of macroscopic mechanical properties of elastomeric organosiloxane foams. *Macromolecules* **48**, 4835–4849. https://doi.org/10.1021/acs.macromol.5b00532 (2015).
39. Alzoubi, M., Al-Waked, R. & Tanbour, E. Compression and hysteresis curves of nonlinear polyurethane foams under different densities, strain rates and different environmental conditions. *J. Mech. Eng.* **9**, 101–109. https://doi.org/10.1115/IMECE2011-62290 (2011).
40. Adobe Inc. Adobe Illustrator. Available from: https://adobe.com/products/illustrator (2019).
41. Klein, S. & Nellis, G. *Thermodynamics* (Cambridge University Press, 2011).
42. Huneau, B. Strain-induced crystallization of natural rubber: a review of x-ray diffraction investigations. *Rubber Chem. Technol.* **84**, 425–452 (2011).
43. Brüning, K., Schneider, K., Roth, S. V. & Heinrich, G. Kinetics of strain-induced crystallization in natural rubber studied by WAXD: Dynamic and impact tensile experiments. *Macromolecules* **45**, 7914–7919 (2012).
44. Boel, M. & Eirich, F. Thermodynamics, thermal effects and dilatation of natural. https://doi.org/10.21236/ad0644112 (1966).
45. Stephen, R. *et al.* Flow properties of unvulcanised natural rubber/carboxylated styrene butadiene rubber latices and their blends. *J. Appl. Polym. Sci.* **104**, 2528–2535. https://doi.org/10.1002/app.25855 (2007).
46. Phuhiangpa, N., Ponloa, W., Phongphanphanee, S. & Smithipong, W. Performance of nano- and microcalcium carbonate in uncrosslinked natural rubber composites: new results of structure-properties relationship. *Polymers*. https://doi.org/10.3390/polym12092002 (2020).
47. Promhuad, K. & Smithipong, W. Effect of stabilizer states (solid Vs liquid) on properties of stabilized natural rubbers. *Polymers*. https://doi.org/10.3390/polym12040741 (2020).
48. Suethao, S. & Smithipong, W. Adhesion and heat build-up of rubber for energy-saved tyre. *IOP Conf. Ser. Mater. Sci. Eng.* **526**, 012011. https://doi.org/10.1088/1757-899X/526/1/012011 (2019).

**Acknowledgements**

This research was financially supported by the graduate school of Kasetsart University, Bangkok, Thailand. This research was also supported by the Specialized center of Rubber and Polymer Materials in agriculture and industry (RPM), Faculty of Science, Kasetsart University, Bangkok, Thailand.

**Author contributions**

S.S. carried out the preparation and characterisation of foam samples and wrote the manuscript; W.P. carried out the finite element analysis; S.P. contributed to the discussion of the manuscript; J.W. contributed to the discussion and review of the manuscript; W.S. conceived and designed the study, supervised, and revised the manuscript. All authors approved the manuscript before submission.

**Competing interests**

The authors declare no competing interests.

**Additional information**

**Correspondence** and requests for materials should be addressed to W.S.

**Reprints and permissions information** is available at www.nature.com/reprints.

**Publisher’s note** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

---

**Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2021