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Introduction of an adhesion factor to *cube in cube* models and its effect on calculated moduli of particulate composites

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The cube in cube approach was used by Paul and Ishai-Cohen to model and derive formulas for filler content dependent Young's moduli of particle filled composites assuming perfect filler matrix adhesion. Their formulas were chosen because of their simplicity, and recalculated using an elementary volume approach which transforms spherical inclusions to cubic inclusions. The EV approach led to expression of the composites moduli that allows introducing an adhesion factor k_{adh} ranging from 0 and 1 to take into account reduced filler matrix adhesion. This adhesion factor scales the edge length of the cubic inclusions, thus reducing the stress transfer area between matrix and filler. Fitting the experimental data with the modified Paul model provides reasonable k_{adh} for PA66, PBT, PP, PE-LD and BR which are in line with their surface energies. Further analysis showed that stiffening only occurs if k_{adh} exceeds $\sqrt{E_M/E_F}$ and depends on the ratio of matrix modulus and filler modulus. The modified model allows for a quick calculation of any particle filled composites for known matrix modulus E_M , filler modulus E_F , filler volume content v_F and adhesion factor k_{adh} . Thus, finite element analysis (FEA) simulations of any particle filled polymer parts as well as materials selection are significantly eased. FEA of cubic and hexagonal EV arrangements show that stress distributions within the EV exhibit more shear stresses if one deviates from the cubic arrangement. At high filler contents the assumption that the property of the EV is representative for the whole composite, holds only for filler volume contents up to 15 or 20% (corresponding to 30 to 40 weight %). Thus, for vast majority of commercially available particulate composites, the modified model can be applied. Furthermore, this indicates that the *cube in cube* approach reaches two limits: (i) the occurrence of increasing shear stresses at filler contents above 20% due to deviations of EV arrangements or spatial filler distribution from cubic arrangements (singular), and (ii) increasing interaction between particles with the formation of particle network within the matrix violating the EV assumption of their homogeneous dispersion.

Sustainability demands on performance of polymer parts are steadily increasing. Not only polymeric systems from renewable sources are currently developed, but contemporary established materials commodities are recycled. To meet these requirements, polymers are often modified by blending, copolymerization and reinforcement with particulate fillers and fibers. The access of these materials to emerging applications brings necessity of fast availability of relevant physical properties as stiffness, strength, and thermal properties e.g. thermal length expansion or heat conductivity.

In this perspective, use of reinforced composites will increase substantially¹, and their performance has to be determined with respect to varying or even undefined mechanical properties of raw materials. This can be achieved by an elaborate thermo-mechanical analysis of a particular composite system (which is expensive and time consuming) or via accurate predictions of composite properties using analytical approaches.

Performance of composites is determined by properties of a matrix, dispersed phase and interface between filler and matrix². A filler-matrix adhesion can be modified by coupling agents. He and Jiang³ showed that glass

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Models	Young's modulus of particle filled composite
Voigt ¹⁵	$E_C = E_M \left((1 - v_F) + \frac{E_F}{E_M} v_F \right) = E_M (1 - v_F) + E_F v_F$ (1)
Reuss ¹⁶	$E_C = E_M \frac{\frac{E_F}{E_M}}{v_F + \frac{E_F}{E_M} (1 - v_F)} = \frac{E_F E_M}{E_F (1 - v_F) + E_M v_F}$ (2)
Guth ¹⁷	$E_C = E_M \left(1 + K_E v_F + 14.1 v_F^2 \right)$ (3)
Paul ²⁰	$E_C = E_M \left(\frac{1 + \left(\frac{E_F}{E_M} - 1 \right) v_F^{\frac{2}{3}}}{1 + \left(\frac{E_F}{E_M} - 1 \right) \left(v_F^{\frac{2}{3}} - v_F \right)} \right)$ (4)
Hirsch ²²	$E_C = E_M \left(\chi \left((1 - v_F) + \frac{E_F}{E_M} v_F \right) + \frac{(1 - \chi) \frac{E_F}{E_M}}{v_F + \frac{E_F}{E_M} (1 - v_F)} \right)$ (5)
Counto ²³	$E_C = E_M \left(\frac{\left((1 - v_F^{\frac{1}{2}}) + v_F^{\frac{1}{2}} \frac{E_F}{E_M} \right)}{\left((1 - v_F^{\frac{1}{2}}) \right)^2 + \left((1 - v_F^{\frac{1}{2}}) \right) v_F^{\frac{1}{2}} \frac{E_F}{E_M} + v_F^{\frac{1}{2}}} \right)$ (6)
Takayanagi et al. ¹⁹	$E_C = E_M / \left(\frac{5v_F}{\left(1 - \frac{(2+3v_F)}{5} \right) + \frac{(2+3v_F)}{5} \frac{E_F}{E_M}} + \frac{1 - 5v_F}{(2+3v_F)} \frac{E_F}{E_M} \right)$ (7)
Ishai-Cohen ²¹	$E_C = E_M \left(\frac{\frac{E_F}{E_M} + \left(\frac{E_F}{E_M} - 1 \right) \left(v_F - v_F^{\frac{1}{3}} \right)}{\frac{E_F}{E_M} - \left(\frac{E_F}{E_M} - 1 \right) v_F^{\frac{1}{3}}} \right)$ (8)
Halpin-Tsai ²⁴	$E_C = E_M \left(\frac{1 + \left(\frac{E_F}{E_M} - 1 \right) \xi v_F}{\left(\frac{E_F}{E_M} + \xi \right)} \right) \left(\frac{\left(\frac{E_F}{E_M} - 1 \right)}{1 - \left(\frac{E_F}{E_M} + \xi \right) v_F} \right)$ (9)
Nielsen ^{25,27}	$E_C = E_M \left(\frac{1 + (K_E - 1) \left(\frac{E_F}{E_M} - 1 \right) v_F}{1 - \left(\frac{E_F}{E_M} - 1 \right) \left(1 + \frac{(1 - v_{F,max}) v_F}{v_{F,max}^2} \right)} \right)$ (10)

Table 1. Summary of two-phase models to predict Young's moduli of particulate composites. Moduli of matrix E_M and filler E_F , filler volume content v_F , parameter χ determining stress transfer between fiber and matrix, geometry factor ξ , Einstein coefficient KE and maximum volume fraction $v_{F,max}$.

bead reinforced polyepichlorhydrin pretreated with a coupling agent yielded in an enhanced stiffness behavior over the whole filler volume range (5–35%). Demir et al.⁴ investigated the effect of different coupling agents on the performance of luffa fiber/polypropylene composites and found out that the moduli of the composites containing 15 wt% fiber increased by 52 to 98% after modification. They concluded that the addition of the coupling agents was accompanied by the decrease in water absorption due to a better adhesion between fibers and matrix. Similar conclusion was presented by Jacob et al.⁵, which studied the effect of various silane coupling agents on viscoelastic properties of sisal/oil palm hybrid fiber reinforced natural rubber composites, and reported that the chemical modification of fibers with respect to the used coupling agents led to improved wettability and consequently to an increase in storage and loss moduli of the composites.

Ku et al.⁶ reviewed an effect of various pretreatments and coupling agents on tensile properties of natural fiber reinforced polymer composites. They summarized the studies, which confirmed the increase in Young's modulus due to the enhanced interfacial adhesion caused by the pretreatment of the fillers and the coupling agents. On the other hand, Dekkers et al.⁷ and Dibenedetto et al.⁸ used glass beads as fillers for polystyrene and epoxy resin matrices and found that the Young's moduli were not affected by the surface treatment. Wang et al.⁹ observed increasing moduli of polypropylene/barium sulfate composites using stearic acid, silane and maleic anhydride as surface modifications, but they attributed it not to the improved interfacial adhesion, but rather to higher crystallinity and the formation of the crystal lattices in the matrix.

Numerous models have been developed to calculate Young's moduli of particulate polymer composites considering elastic properties of fillers and matrices, volume content and aspect ratio, Table 1. For modelling purposes, a two-phase model in terms of the representative volume element is subjected to unidirectional stresses and strains for calculating expressions of elastic constants^{10,11}. Three-phase models are not considered in this work because their higher complexity requires additional material parameters^{12–14}.

The models of Voigt¹⁵ and Reuss¹⁶ represent the upper bound for the uniform stress distribution and the lower bound for the uniform strain distribution, respectively, and differ significantly. In most cases the measured Young's moduli lie in between. This indicates that they can only serve as a rough estimation, and that real loading states are more complex than uniform stress or uniform strain. Guth¹⁷ proposed a two-phase model based on Einstein's approach¹⁸ to determine the viscosity of a suspension with spherical inclusions.

The *cube in cube* models (4–10) take into account a dispersed structure of particle filled composites. Takayanagi et al.¹⁹ combined the models of Voigt and Reuss in a way that allows for introducing a dependency on filler volume content. Paul²⁰ assumed uniform stress states with a perfect filler-matrix adhesion, whereas Ishai and Cohen²¹ assumed uniform strain states. Their models represent more precise upper and lower bounds for

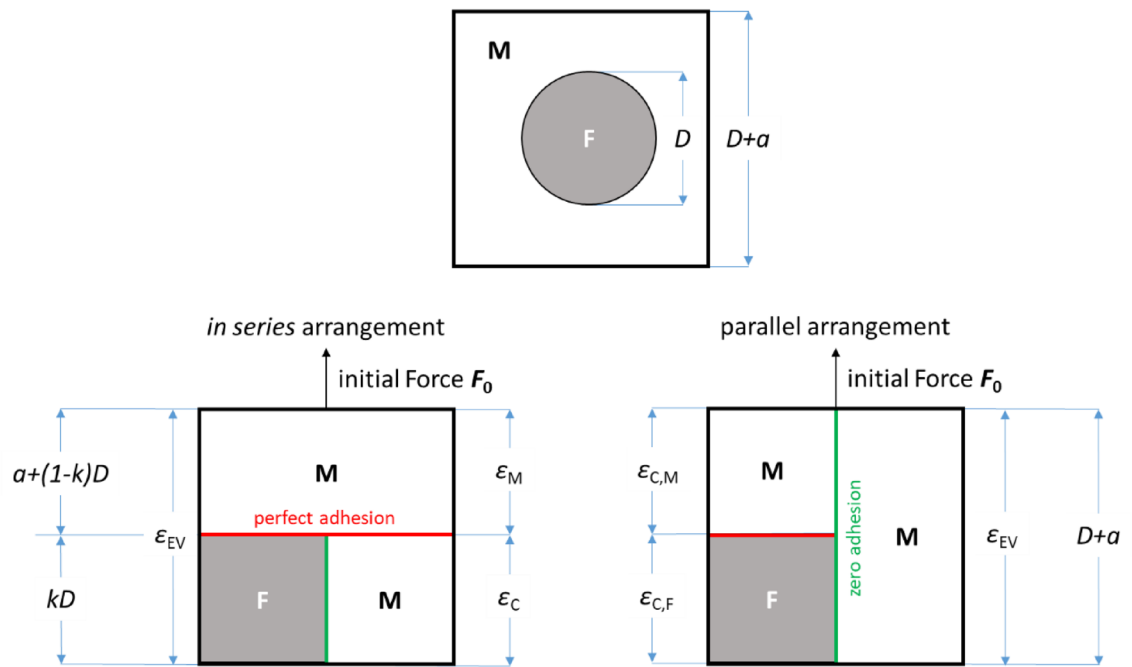


Figure 1. Cubic elementary volume (EV) containing a spherical inclusion with diameter D and distance a between inclusions (top), and its *cube in cube* consideration in *series* and *parallel* arrangements with adhesion boundary conditions.

composites moduli than those of Voigt and Reuss. More complicated models require additional parameters, which are determined by a fitting procedure:

- Hirsch²² combined the models of Voigt and Reuss balancing their contributions with a factor χ .
- Counto²³ developed a simple model for concrete systems assuming perfect filler matrix adhesion; it coincides with the Hirsch model for $\chi=0.5$.
- Halpin and Tsai²⁴ developed a model to describe anisotropic properties of fiber and filler reinforced composites with one equation in which an efficiency factor ξ takes into account fiber or filler geometry and spatial orientation of fibers or platelets; due to its simplicity it became popular regardless of limited accuracy.
- Nielsen²⁵, Lewis and Nielsen²⁶ and Nielsen²⁷ proposed an alternative model based on Halpin–Tsai²⁴ and Kerner²⁸ in which mechanical properties depend additionally on fiber or filler geometry and load direction, maximum filler content $v_{F,max}$; moreover, Nielsen assumed a Poisson ratio-dependent Einstein coefficient K_E .

Moduli of matrix E_M and filler E_F , filler volume content v_F , parameter χ determining stress transfer between fiber and matrix, geometry factor ξ , Einstein coefficient K_E and maximum volume fraction $v_{F,max}$.

All models depicted in Table 1 assume a perfect filler–matrix adhesion. Reduced adhesion between a filler and a matrix has not been considered yet in *cube in cube* models. The aim of this study is to introduce an adhesion factor having values between “0” and “1” to the EV models of Paul and Ishai–Cohen to express adhesion quantitatively, and to investigate how the adhesion factor and spatial arrangement of EV affect the filler content dependent composites moduli.

Theoretical considerations

If filler particles are homogeneously dispersed in a matrix, one can define an elementary volume (EV) containing a single particle that is to be representative for composite properties. A cube of length $D+a$ is shown with a spherical inclusion of diameter D consisting of matrix M and filler, Fig. 1 top. The “cube in cube” approach requires that a particle of any geometry is transformed to a cube of length kD with the efficiency factor k . It takes into account that less than the maximum cross-section contributes to the stress transfer, and thus depends on particle shapes. For spheres it is determined by the condition

$$V_{sphere} = \frac{\pi}{6}D^3 = k^3D^3 = V_{cube} \Rightarrow k = k_{sphere} = \sqrt[3]{\frac{\pi}{6}} \cong 0.81 \quad (11)$$

with volume of sphere V_{sphere} and volume of cube V_{cube} . In order to calculate the filler volume content dependent Young’s modulus of the composite, the cube has to be divided into a matrix part and a composite part done either *in series* or *parallel* to account for microscopic mechanical properties, Fig. 1 bottom left and right.

This allows calculation how the modulus of the EV depends on the arrangements of matrix part and composites part. Furthermore, the analysis of the EV separation shows that certain assumptions were made with respect to the adhesions acting between matrix and filler:

- For the *in series* arrangement “perfect” adhesion is assumed over the entire cross-section $(D+a)^2$ perpendicular to the load direction between matrix part and composites part, and “zero” adhesion is required between filler and matrix of the composite part along the load direction. If the EV cube is strained to ε_{EV} by the load F_0 , the matrix part experiences the strain ε_M and the composites part the strain ε_C .
- For the *parallel* arrangement “perfect” adhesion is assumed only over the cross-section $(kD)^2$ of the composite part between the filler and the matrix perpendicular to the load direction, and “zero” adhesion is required between the composite part and the matrix part along the load direction. If the EV cube is strained to ε_{EV} by the load F_0 , the matrix of the composite part experiences the strain $\varepsilon_{C,M}$ and the filler of the composite part the strain $\varepsilon_{C,F}$.

If the aspect ratio is not close to “1”, one has to distinguish the extremes “along the long axis” and “perpendicular to the long axis” e.g. with a square column consideration. In this consideration a reduction of adhesion can only apply to “perfect” adhesion, whereas “zero” adhesion leads to neglecting of shear stresses.

Case 1: in series EV arrangement and Paul model. The first step is to define the elastic stress–strain–relation of the EV to represent the macroscopic mechanical behavior:

$$\sigma = \sigma_{EV} = E_{EV} \frac{\Delta(D+a)}{D+a} = E_{EV} \varepsilon_{EV} \Rightarrow \varepsilon_{EV} = \frac{\sigma}{E_{EV}} \quad (12)$$

with external stress σ , stress acting on EV σ_{EV} , distance a between particles, modulus of EV E_{EV} and strain of EV ε_{EV} . The second step is to define the stress–strain–relation of the matrix part with respect to the microscopic strain ε_M of the matrix part:

$$\sigma = \sigma_M = E_M \frac{\Delta a}{a} = E_M \varepsilon_M \Rightarrow \varepsilon_M = \frac{\sigma}{E_M} \quad (13)$$

with stress acting on matrix part σ_M , matrix modulus E_M , and matrix strain ε_M , and the stress–strain–relation of the composites part:

$$\sigma = \sigma_C = \sigma_{C,M} + \sigma_{C,F} \Rightarrow \varepsilon_C = \frac{\sigma}{E_C} \quad (14)$$

with stress acting on composite σ_C , stress acting on matrix of composites part $\sigma_{C,M}$, stress acting on filler of composites part $\sigma_{C,F}$, strain of composites part ε_C , and the composites modulus E_C . The stress acting on the matrix of the composites part is given by:

$$\sigma_{C,M} = \left(1 - \frac{(kD)^2}{(D+a)^2} \right) E_M \varepsilon_C, \quad (15)$$

and the stress acting on the filler of the composites part is given by:

$$\sigma_{C,F} = \frac{(kD)^2}{(D+a)^2} E_F \varepsilon_C \quad (16)$$

with filler modulus E_F . Introduction of (15) and (16) in (14) links the external stress to the microscopic strain ε_C yielding

$$\sigma = \sigma_C = \left(\left(\underbrace{1 - \frac{(KD)^2}{(D+a)^2}}_{\text{relative matrix cross-section}} \right) E_M + \underbrace{\frac{(KD)^2}{(D+a)^2}}_{\text{relative filler cross-section}} \right) \varepsilon_C. \quad (17)$$

If the interfacial adhesion is not perfect, the stress transfer to the inclusion is reduced. The structure of (17) allows to identify where non-perfect reduced filler matrix adhesion may come into play as it has to decrease the stiffening capability of the filler. Thus, the second term in the brackets on the right side of (17) describing the contribution of filler modulus E_F to the stress–strain–relation of the composites does not contribute to full extent. In this context the EV representation of stress–strain–relation (17) provides a hint, where to introduce the dimensionless adhesion factor k_{adh} that ranges from “0” (no adhesion) to “1” (perfect adhesion). In Eq. (17) one deals with the coefficients attached to E_M and E_F which represent relative cross-sections of matrix and filler, respectively. Therefore, the adhesion factor is introduced quadratically as it is to reduce the edge length of the filler cube. In that respect the physical meaning of the adhesion factor is that it scales the edge length of the filler cube, and thus the available contact area of stress transfer between filler and matrix to account for the reduced

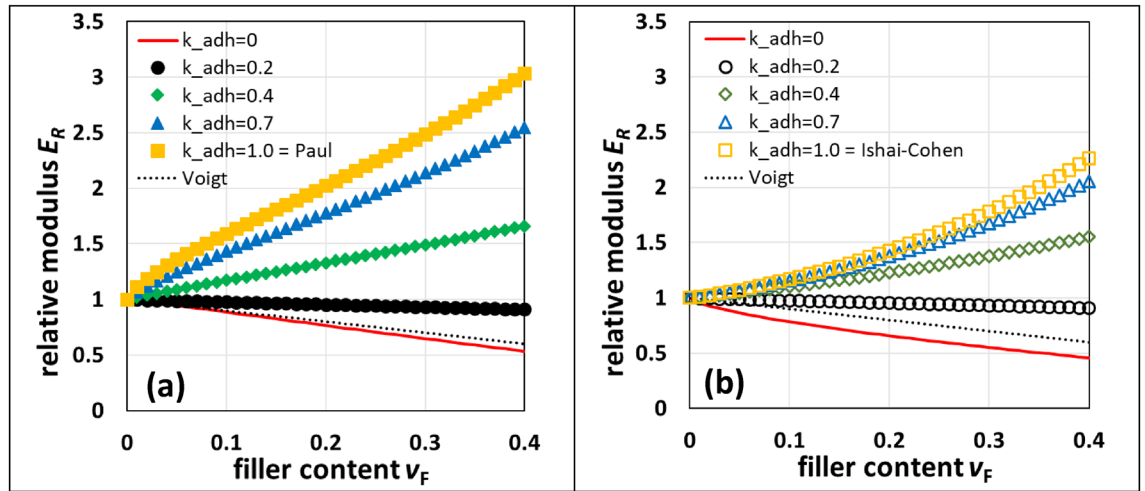


Figure 2. Calculated filler volume content dependent relative moduli E_R for different adhesion factors k_{adh} with $E_M = 3200$ MPa and $E_F = 63,000$ MPa for (a) *in series* EV arrangement and (b) *parallel* EV arrangement. The dotted line represents the decreasing E_R for a foam with $E_F = 0$ MPa according to Voigt.

adhesion due to the surface energies of filler and matrix, respectively. After elimination of the particle diameter D , one gets:

$$\sigma \underset{d=\frac{a}{b}}{\approx} \left(\left(1 - \frac{k^2}{(1+d)^2} \right) E_M + k_{adh}^2 \frac{k^2}{(1+d)^2} E_F \right) \varepsilon_C \tag{18}$$

Solving (18) for ε_C yields the strain of the composite part.

$$\varepsilon_C = \frac{\sigma}{E_M \left(1 + \frac{k^2}{(1+d)^2} \frac{k_{adh}^2 E_F - E_M}{E_M} \right)} \tag{19}$$

The strain of the EV ε_{EV} depends on the strains of both the matrix part ε_M and the composites part ε_C (see Appendix A):

$$\varepsilon_{EV}(\varepsilon_M, \varepsilon_C) = \left(1 - \frac{k}{(1+d)} \right) \varepsilon_M + \frac{k}{(1+d)} \varepsilon_C \tag{20}$$

Substituting the strains from (12), (13) and (19) in (20), and solving for E_{EV} yields:

$$E_{EV} = E_M \left(1 + \frac{\frac{(k)^3}{(1+d)^3} \frac{k_{adh}^2 E_F - E_M}{E_M}}{1 + \frac{(k)^2}{(1+d)^2} \frac{k_{adh}^2 E_F - E_M}{E_M} - \frac{(k)^3}{(1+d)^3} \frac{k_{adh}^2 E_F - E_M}{E_M}} \right) \tag{21}$$

The efficiency factor k is related to the filler volume content v_F by

$$v_F = \frac{(k)^3}{(1+d)^3}; v_F^{2/3} = \frac{(k)^2}{(1+d)^2}; v_F^{1/3} = \frac{k}{(1+d)}, \tag{22}$$

and (21) becomes:

$$E_{EV} = E_M \left(1 + \frac{v_F \frac{k_{adh}^2 E_F - E_M}{E_M}}{1 + v_F^{2/3} \frac{k_{adh}^2 E_F - E_M}{E_M} - v_F \frac{k_{adh}^2 E_F - E_M}{E_M}} \right) \tag{23}$$

For perfect adhesion with $k_{adh} = 1$, Eq. (23) becomes identical to Paul's relation in Table 1.

From Eq. (23) follows that a stiffening effect only occurs if $k_{adh} > \sqrt{E_M/E_F}$. Calculating the filler volume content dependent relative modulus $E_R = E_C/E_M$ with $E_M = 3200$ MPa and $E_F = 63,000$ MPa, respectively, and varying k_{adh} between 0 and 1 shows that for $k_{adh} > 0.23$ one gets stiffening, and for $k_{adh} < 0.23$ softening, Fig. 2a.

Case 2: parallel EV arrangement and Ishai-Cohen model. The external stress σ acts macroscopically on the EV according to (12). The stress-strain-relation of the matrix part is given by

Matrix	GB volume content	GB diameter r_F	Matrix density ρ_M	Matrix modulus E_M	Poisson ratio of matrix μ_M	Surface energy ²⁹⁻³¹ σ_{surface}
	v_F					
PA66	0.16/0.23	25	1.14	3100 to 3200	0.42	38 to 55
PBT	0.12/0.19	25	1.30	2600 to 2800	0.41	44 to 49
BR	0.15/0.30/0.45	58	0.95	7.6	0.48	26 to 27
PE-LD	0.08/0.19/0.35	40 to 60	0.93	98	0.48	33 to 35
iPP	0.10/0.20/0.30	Not given	0.91	1070	0.45	31 to 42

Table 2. Properties of matrices and dispersed phase (GB).

$$\sigma = \sigma_M = E_M \varepsilon_{EV} \Rightarrow \varepsilon_{EV} = \frac{\sigma_M}{E_M}, \quad (24)$$

and the stress-strain-relations of the composites part are given by

$$\sigma = \sigma_C = E_C \varepsilon_C = E_C \varepsilon_{EV} \quad (25a)$$

$$\text{with } \sigma = \sigma_{C,M} = E_M \varepsilon_{C,M} \quad (25b)$$

$$\sigma = \sigma_{C,F} = k_{adh}^2 E_F \varepsilon_{C,F} \quad (25c)$$

In (25c) the adhesion factor k_{adh} is already introduced to account for possibly reduced load transfer from matrix to filler. The external stress σ is distributed on the matrix part by σ_M and the composites part by σ_C due to the corresponding cross-sections yielding

$$\sigma = \sigma_M \frac{A_M}{A_{EV}} + \sigma_C \frac{A_C}{A_{EV}} = \sigma_M \left(1 - \frac{k^2}{(1+d)^2} \right) + \sigma_C \frac{k^2}{(1+d)^2} \quad (26)$$

With cross-section of matrix A_M , cross-section of composites part A_C , and cross-section A_{EV} of EV. The strains of ε_{EV} , $\varepsilon_{C,M}$ and $\varepsilon_{C,F}$ of the composites part are connected by

$$\varepsilon_{EV} = \left(1 - \frac{k}{(1+d)} \right) \varepsilon_{C,M} + \frac{k}{(1+d)} \varepsilon_{C,F} \quad (27)$$

Insertion of (25a), (25b) and (25c) in (27) and solving for E_C yields

$$E_{EV} = E_M \left(1 + \frac{\frac{k^3}{(1+d)^3} (k_{adh}^2 E_F - E_M)}{k_{adh}^2 E_F - \frac{k}{1+d} (k_{adh}^2 E_F - E_M)} \right) \quad (28)$$

or in terms of filler volume content v_F using (22)

$$E_{EV} = E_M \left(1 + \frac{v_F \frac{k_{adh}^2 E_F - E_M}{E_M}}{\frac{k_{adh}^2 E_F}{E_M} - v_F^{1/3} \frac{k_{adh}^2 E_F - E_M}{E_M}} \right). \quad (29)$$

For perfect interfacial adhesion with $k_{adh} = 1$, the Eq. (28) coincides to Ishai-Cohen's relation, Table 1.

From Eq. (28) also follows that a stiffening effect only occurs if $k_{adh} > \sqrt{E_M/E_F}$. The filler volume content dependent relative modulus E_R is calculated with $E_M = 3200$ MPa and $E_F = 63,000$ MPa, respectively, and varying k_{adh} between 0 and 1, Fig. 2b.

From Fig. 2 it is obvious that Eq. (23) produces larger filler volume content dependent Young's moduli than Eq. (29). Introducing the adhesion factor k_{adh} causes a decrease of Young's moduli. Therefore, only Eq. (23) representing the upper bound is used to derive adhesion factors from measured Young's moduli.

Materials and methods

Materials. Five kinds of polymer matrices filled with various contents of glass beads (GB) as spherical inclusions were used in this study (Table 2):

- polyamide 66 (PA66): RADIPOL A45 (0 wt% GB), AKROMID A3 GK 30 1 natur (30 wt% GB), AKROMID A3 GK 40 1 natur (40 wt% GB)
- polybutyleneterephthalate (PBT): Ultradur B 2550 (0 wt% GB), Ultradur B 4300 K4 (20 wt% GB), Ultradur B 4300 K6 (30 wt% GB)
- butadiene rubber (BR): isomer ratio cis:trans:vinyl = 20:60:20, $M_N = 2$ to 3×10^5 , cross-linked with dicumyl peroxide, 0, 15, 30 and 45 vol% GB unsized and sized
- polyethylene (PE-LD): Nova Chemicals LDPE LA-0219-A, Canada with 0, 20, 40, and 60 wt% GB

- isotactic polypropylene (iPP): Petolen MH418 with 0, 10, 20 and 30 vol% GB

The properties of polar PA66 (Technical data sheet, Radici Group, RADIPOL[®] A45, 2021) and PBT (Technical data sheet; BASF, Ultradur[®] B 2550–PBT, 2021) composites were determined experimentally, whereas the properties of nonpolar BR³², PE-LD³³ and iPP³⁴ were taken from literature. As in Lohrmann³² two compound series—one with unsized GB and one with sized GB—were investigated giving the chance to check how surface treatment of GB affects the stiffness. Glass beads filler has density $\rho_F = 2.5 \text{ g/cm}^3$, Young's modulus $E_F = 63 \text{ GPa}$ and Poisson ratio $\mu_F = 0.22$.

Methods. *Preparation of tensile test bars.* Injection molded test bars of PA66 and PBT compounds (type 1A according to ISO 527-2) were tested after annealing for 4 h at 180 °C to minimize effects of a thermal history.

Tensile tests. Universal testing machine (Zwick Z100, Zwick/Roell, Ulm, Germany) was used to perform tensile tests according to DIN EN ISO 527-1 with $n = 5$, equipped with a 10 kN load cell (resolution: 0.12%) and a multiXtens extensometer (resolution: 0.1 μm), an initial sample length of 50 mm, a preload of 0.1 MPa. Differing to ISO 527, the tensile tests of PA66 and PBT compounds were performed with a strain rate of 10%/min. The stress–strain-curves were evaluated using the viscoelastic stress strain function (VSSF)³⁵:

$$\sigma(\varepsilon) \underset{\varepsilon < \varepsilon_R}{\cong} E\varepsilon_R \left(1 - e^{-\frac{\varepsilon}{\varepsilon_R}}\right) \quad (30)$$

with strain rate dependent Young's modulus E and relaxation strain ε_R . Equation (27) is derived from the time dependent solution of the Maxwell model if one substitutes the time by the strain assuming constant strain rates in the tensile test. The fitting procedure was as follows:

- export of raw data of a force F and a length change ΔL
- error analysis of F and ΔL
- consideration of an error propagation of a conversion to stress and the strain
- fitting of measured σ - ε -curves using VSSF
- determination of E and ε_R as well as corresponding standard deviations
- determination of adhesion factors k_{adh} of filled compounds using Eq. (18) with E_M and E_F as input parameters from Table 2.

The stress–strain curves of BR compounds were taken from³², digitized and evaluated correspondingly to include an example of a rubber matrix composite.

Determination of adhesion factors. If a filler matrix adhesion is reduced, a composite modulus decreases. As the modulus E_{EV} according to the Eq. (18) represents the upper bound for $k_{\text{adh}} = 1$, it is used to determine the adhesion factors of the composites depicted in Table 2. The fitting was done for each composite modulus E_C using the Excel solver tool. In cases of several measured E_C this allows for providing standard deviations (STD).

Scanning electron microscopy (SEM). Tensile test bars of PA66 and PBT compounds (type 1A according to ISO 527-2) were fractured under cryogenic conditions in a liquid nitrogen after approximately 5 min of storage. The fracture surfaces were sputtered by gold layer for 90 s at 20 mA, 0.1 mbar in an argon atmosphere. SEM (JEOL JSM-IT100, Japan) equipped with a tungsten cathode was used to investigate the structure of the fracture surfaces at 5.0 kV under vacuum conditions.

Finite element analysis (FEA). FEA using the software SIMULIA/ABAQUS2020 was performed in 2D for the middle plane of the EV, where maximum stresses occur. A 3D simulation would provide the same stress distribution for the middle plane. A mixed mesh of triangle (S3) and rectangle (S4) shell elements with linear shape functions were used. The total number of elements for each arrangement was approximately 53,000. The nodes at the top and bottom were coupled in y - and z -directions with a reference node using a kinematic coupling constraint. For the reference node at the bottom all translational and rotational degrees of freedom were fixed to $u_1 = u_2 = \dots = u_6 = 0$. For the reference node at the top, all degrees of freedom except for the translation in y -direction were fixed as well. The displacement in the positive y -direction (u_2) was defined at the top node to apply a uniaxial tensile load. Static linear analysis was carried out by implementing materials properties as a filler modulus E_F and a matrix modulus E_M , a filler Poisson ratio μ_F and a matrix Poisson ratio μ_M , and a filler volume content v_F . Filler content dependent EV arrangements in a cubic and a hexagonal lattice were investigated, Fig. 3. Maximum achievable filler contents are $v_{F,\text{max}} = \pi/6 \approx 0.52$ for cubic and $v_{F,\text{max}} = \pi/\sqrt{27} \approx 0.60$ for hexagonal arrangements. For given technically relevant filler volume contents below 0.4 both arrangements are identical with respect to a space filling but not to a force flow under uniaxial loads.

Results and discussion

The tensile properties of PA 66 and PBT composites were determined according to ISO 527 and evaluated using the VSSF, Table 3. Both PA66 and PBT compounds showed a stiffening effect with increasing GB contents, whereas reinforcement is only found for the PA66 compounds indicating a higher filler matrix adhesion. Moduli

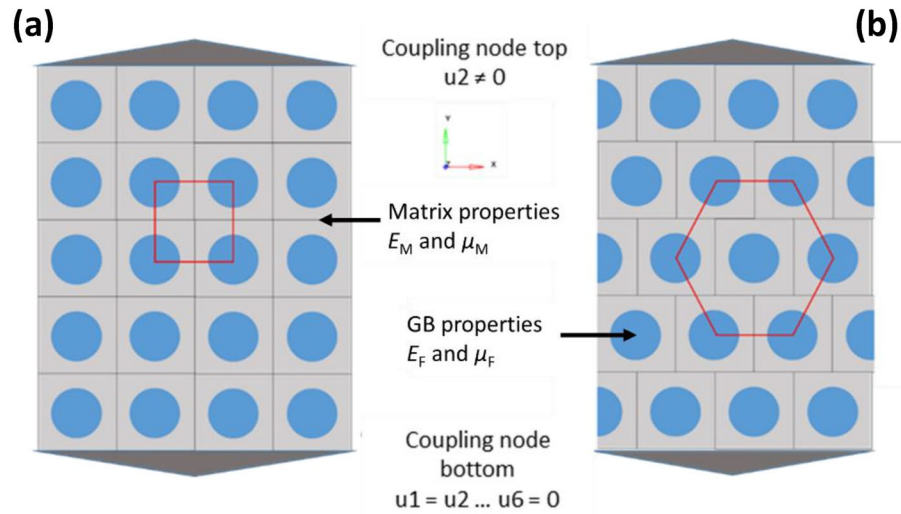


Figure 3. Cubic elementary volume model for FEA simulations: (a) cubic and (b) hexagonal lattice arrangements. Note that this EV representation holds also for unidirectional fiber systems if the load acts only perpendicular to the fiber axis.

Matrix	Filler content v_f	Modulus $E_{ISO\ 527}$ MPa	Tensile strength σ_{max} MPa	Tensile elongation ϵ_{max} %	Modulus E_{VSSF} MPa	Relaxation strain ϵ_R %
PA66	0	3143 ± 14	81 ± 0.3	3.9 ± 0.5	3197 ± 28	9.4 ± 0.2
PA66	0.16	4811 ± 22	91 ± 1.1	3.1 ± 0.8	4955 ± 44	7.8 ± 0.1
PA66	0.23	5299 ± 45	86 ± 0.2	3.7 ± 0.5	5452 ± 48	7.2 ± 0.1
PBT	0	2704 ± 18	59 ± 0.1	10.8 ± 1.4	2754 ± 25	9.4 ± 0.1
PBT	0.12	3581 ± 32	57 ± 0.1	3.2 ± 0.2	3691 ± 33	5.9 ± 0.1
PBT	0.19	4254 ± 30	55 ± 0.1	2.0 ± 0.6	4408 ± 39	5.4 ± 0.2
BR _{unsized}	0	–	–	–	7.0	6.2
	0.15	–	–	–	11.0	6.3
	0.30	–	–	–	16.8	8.0
	0.45	–	–	–	48.0	0.7
BR _{sized}	0	–	–	–	6.9	5.0
	0.15	–	–	–	9.9	4.6
	0.30	–	–	–	19.2	5.5
	0.45	–	–	–	52.2	1.1
PE-LD	0	–	8.4 ± 0.2	–	98 ± 5	–
	0.15	–	7.8 ± 0.5	–	124 ± 6	–
	0.30	–	6.2 ± 0.3	–	175 ± 27	–
	0.45	–	3.3 ± 0.1	–	219 ± 76	–
iPP	0	–	31.1 ± 0.5	14.7 ± 0.4	1070 ± 20	–
	0.10	–	26.0 ± 1.7	10.6 ± 0.8	1230 ± 10	–
	0.20	–	18.8 ± 4.9	6.1 ± 1.6	1280 ± 20	–
	0.30	–	15.9 ± 1.3	45 ± 1.2	1310 ± 20	–

Table 3. Tensile properties of composites. For BR compounds the standard deviation could not be determined because there was only one stress strain curve for each compound available in Lohrmann³². For PE-LD and iPP the data were taken directly from Chimeni et al.³³ and Balkan and Demirer.³⁴

determined by ISO 527 are smaller than those determined by the VSSF because the strain rate of 10%/min is not reached yet at strains $\epsilon_1 = 0.05\%$ and $\epsilon_2 = 0.25\%$.

The adhesion factors k_{adh} were determined by a fitting procedure that brings calculated moduli E_{EV} according to the Eq. (18) to coincidence to measured moduli E_{VSSF} . The comparison of E_{EV} to calculated moduli E_C according to Ishai-Cohen, Paul and Halpin-Tsai with $\xi = 2$ for the transversal case shows that fitting the composites moduli according to Eq. (18) provides reasonable values of the adhesion factors, Table 4. It confirms that Paul

Matrix	Filler content v_F	Calculated Young's modulus E_C				Adhesion factor k_{adh}	Stiffening limit $k_{adh,min}^{stiffening}$
		Ishai-Cohen	Paul	Halpin-Tsai	EV ($E_{EV} \equiv E_{VSSr}$)		
	–	MPa					–
PA66	0.16	4229 ± 35	5963 ± 44	4770 ± 32	4955 ± 44	0.604 ± 0.017	0.225
	0.23	4899 ± 40	6940 ± 51	5408 ± 36	5452 ± 48	0.570 ± 0.013	
PBT	0.12	3370 ± 30	4744 ± 37	3754 ± 26	3691 ± 33	0.478 ± 0.015	0.209
	0.19	3895 ± 34	5608 ± 43	4449 ± 30	4408 ± 39	0.530 ± 0.013	
BR ^{unsized}	0.15	9.2	14.9	10.7	11.0	0.031	0.011
	0.30	13.3	21.1	16.0	16.8	0.042	
	0.45	20.4	29.9	24.1	29.9 ^a	n.f.	
BR ^{sized}	0.15	9.2	14.8	10.6	9.9	0.025	0.011
	0.30	13.2	21.0	15.8	19.2	0.073	
	0.45	20.3	29.6	23.9	29.7 ^a	n.f.	
iPP	0.10	1263	1876	1407	1230	0.220	0.130
	0.20	1565	2414	1824	1280	0.190	
	0.30	1993	3010	2351	1310	0.180	
PE-LD	0.08	112	173	125	124	0.100	0.039
	0.19	143	232	167	175	0.130	
	0.35	215	331	257	219	0.110	

Table 4. Comparison of calculated GB content dependent Young's moduli with evaluated adhesion factors k_{adh} and its minimal stiffening limit $k_{adh,min}^{stiffening} = \sqrt{E_M/E_F}$. ^aTo calculate E_{EV} the adhesion factor was set to $k_{adh} = 1$. n.f. means "no fit available".

and Ishai-Cohen models represent the upper and lower bounds, respectively. The calculated moduli according to Halpin-Tsai provide reasonable E_C values, but the efficiency factor ξ is hardly related to filler matrix adhesion as the calculated E_C can be larger or smaller compared to the measured ones.

For the polar polymer matrices PA66 and PBT, k_{adh} is found to be around 0.6 and 0.5, respectively, whereas it is smaller than 0.25 for the nonpolar polymer matrices iPP, PE-LD and BR, which is in accordance with the surface energies shown in Table 1. In spite of rather low k_{adh} all analyzed composites exhibit stiffening due to GB introduction as they always exceed the stiffening limit.

The comparison of measured Young's moduli of GB filled PA66 and PBT composites to calculated ones, Fig. 4, confirms that the models of Ishai-Cohen and Paul represent the lower and the upper bounds. The measured Young's moduli lie between these bounds as well as the calculated moduli according to Halpin-Tsai with $\xi = 2$ and EV with $k_{adh} = 0.6$ (PA66) and 0.5 (PBT). Both reproduce well the measured Young's moduli, but only the EV provides the quantitative information concerning the filler-matrix adhesion.

SEM images of PA66 and PBT composites show a fracture surface with embedded GB, which are partly covered by the matrix indicating rather poor filler-matrix adhesion, Fig. 5.

Better adhesion seen for the PA66 composite correlates with its higher k_{adh} . Small fibrils partly visible on the fractured surfaces of both the PA66 and PBT composites correspond to ductile failure of the matrices. It seems that the fracture behavior becomes more ductile, the higher the GB content is.

Similar increases of filler content dependent Young's moduli of elastomer (BR) composites were found for all models, Table 4, showing a stiffening effect although the adhesion factors become small. The introduction of a coupling agent to one BR composite has hardly an effect. The Halpin-Tsai model produces similar Young's moduli as the EV with k_{adh} between 0.02 and 0.05 for $v_F = 0.15$ and 0.30. For $v_F = 0.45$ all models generate too small Young's moduli. This can be attributed to the fact that at high filler contents, particle interactions become stronger going along with the formation of a particle network with significantly higher modulus³⁶. On first sight this issue could be eliminated by allowing adhesion factors exceeding 1. However, it turned out that the models of Ishai-Cohen and Paul limit the filler volume content dependent stiffening factors to:

$$\text{Ishai-Cohen } s(v_F) = \frac{E_{EV}}{E_M} = \frac{v_F}{1 - v_F^{\frac{1}{3}}} \xrightarrow{v_F=0.45} 2.9 \quad (31)$$

$$\text{Paul } s(v_F) = \frac{E_{EV}}{E_M} = \frac{1}{1 - v_F^{\frac{1}{3}}} \xrightarrow{v_F=0.45} 4.3 \quad (32)$$

From Fig. 3 it is obvious that the cubic arrangement of EV leads to less shear stresses compared to the hexagonal arrangement although the only difference is that each 2nd layer of EV is shifted by one half of the edge length. This is confirmed by purely elastic FEA simulations showing that the stress distributions differ significantly for the cubic and hexagonal EV arrangement, Fig. 6. The stresses have to be interpreted with respect to tensile strengths of matrix and GB, whereas $\sigma_{y,GB} > \sigma_{y,M}$. For all cubic arrangements the maximum stresses occur between the

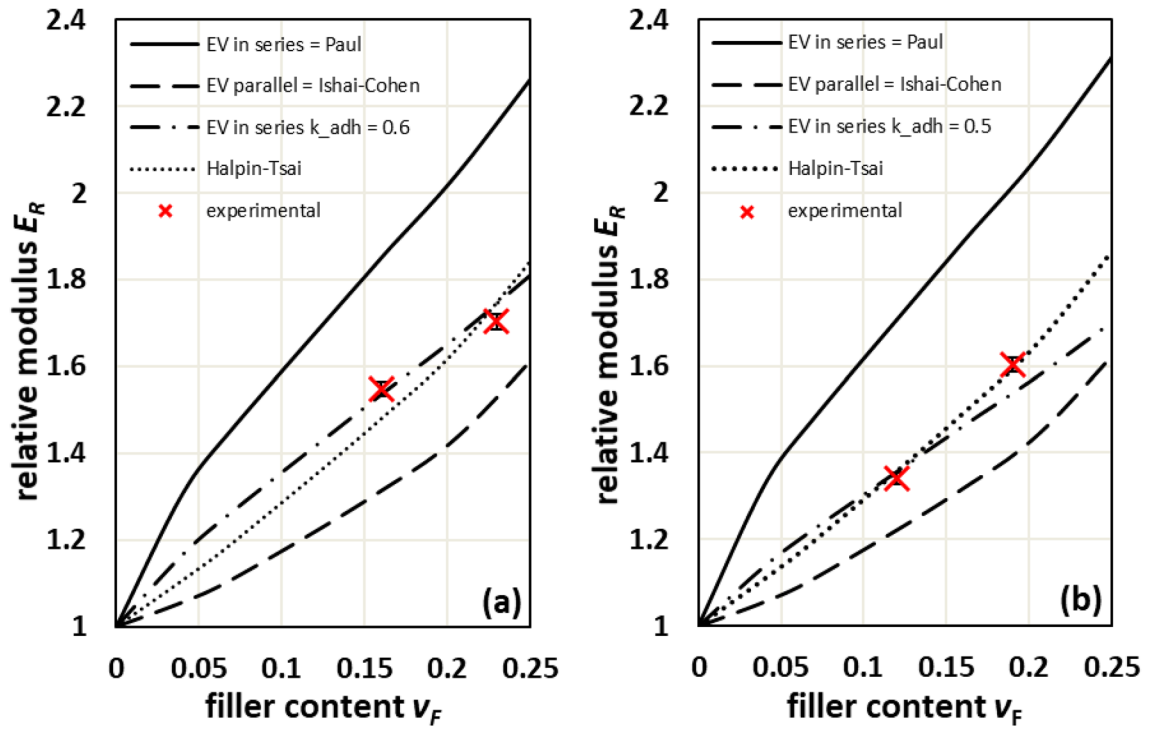


Figure 4. Comparison of relative Young's moduli calculated according to Ishai-Cohen, Paul, Halpin-Tsai with $\xi=2$, and EV for $k_{adh}=0.5$ and 0.6 (lines) with measured ones (symbols) for (a) PA66 and (b) PBT.

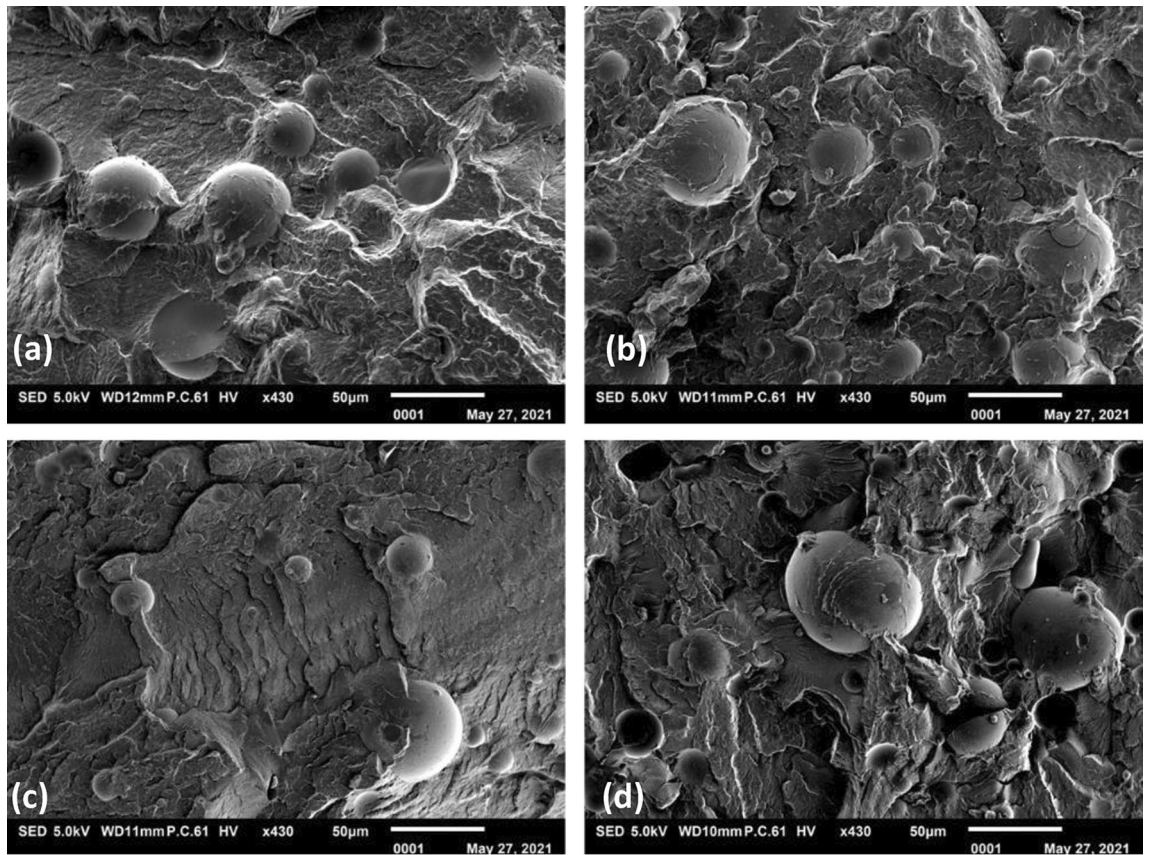


Figure 5. Fracture surface of composites: (a) PA66/GB16, (b) PA66/GB23, (c) PBT/GB12 and (d) PBT/GB19.

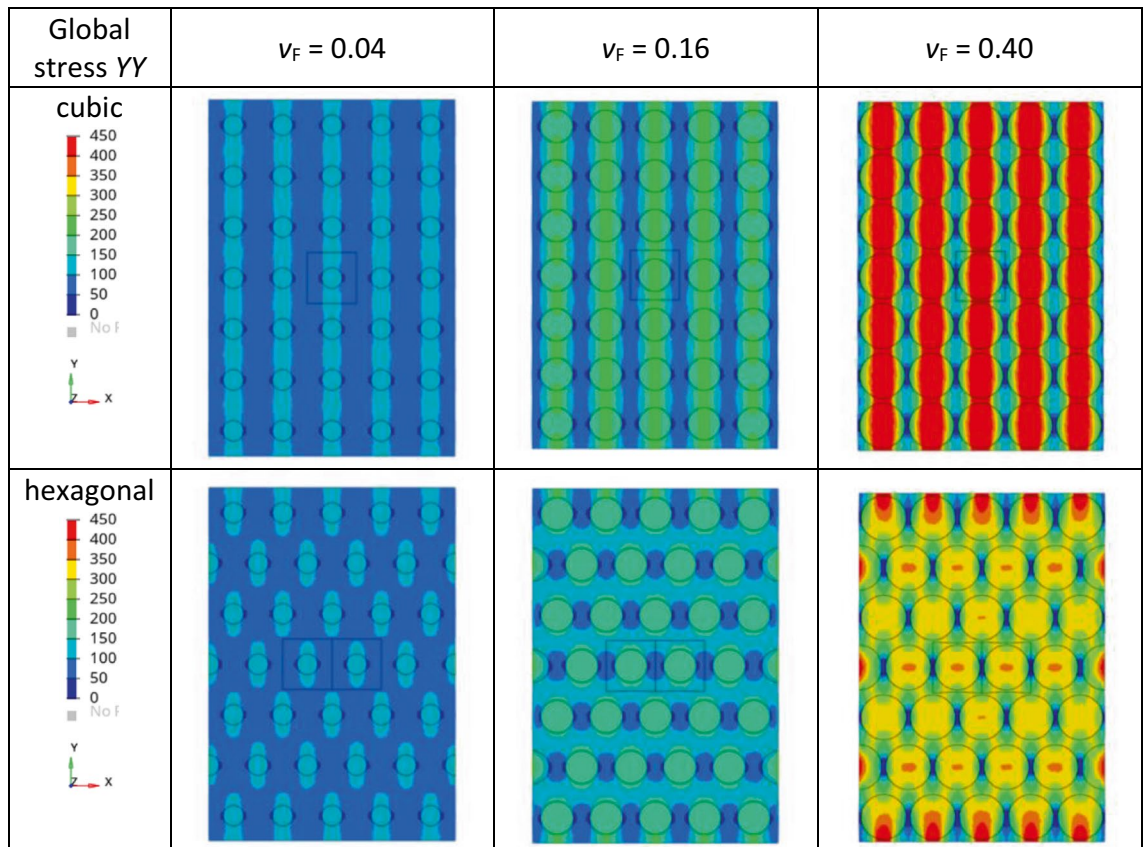


Figure 6. Filler content dependent stress distribution in tensile direction (YY) of cubic and hexagonal lattice arrangements of EV under tensile load in y -direction. Note that the displacements in y -direction are represented in a magnified manner.

Filler content, v_F	E_C (cubic lattice)	E_C (hexagonal lattice)	E_C ratio cubic:hexagonal
	MPa	MPa	
0.04	3950	3914	1.01
0.07	4389	4308	1.02
0.16	5988	5656	1.06
0.22	7461	6827	1.09
0.40	14,534	11,632	1.25

Table 5. FEA calculated filler volume content dependent Young's moduli of cubic and hexagonal lattice arrangements of EV with $E_M = 3200$ MPa, $E_F = 63,000$ MPa, $\mu_M = 0.42$ and $\mu_F = 0.22$.

GB in the matrix forming the parallel stress lines in the composite part along the load direction, whereas in the interjacent matrix parts the stresses remain smaller. With the increasing filler content, the stresses in tensile direction increase leading to a failure of the interface, and cracks propagation perpendicular to the load. Nevertheless, the stresses in both parts are always oriented in the parallel and uniaxial manner in tensile direction.

The hexagonal arrangement of EV shows stress pattern with in-load direction oriented oval "stress islands" around the GB particles. The minimum stresses occur perpendicular to the load direction in the matrix between the GB. The matrix stresses in the load direction between the GB seem to be more homogeneous compared to the cubic arrangement providing more deformability. With increasing the GB content, the stress islands unify and form shear stress band under an angle of 45° seen in Fig. 6—hexagonal/ $v_F = 0.40$. Both reduces moduli of the hexagonal arrangement in the load direction, Table 5, but the difference remains less than 10% for volume filler contents not exceeding 20%.

From Table 5 one can conclude that the properties of the EV are representative of composites properties for filler volume contents not exceeding 15 to 20%. In this filler content range, the EV arrangement effects remain in the order of the experimental error. For higher filler contents, the arrangement of EV starts to affect the composites moduli due to increasing shear stresses. The two considered EV arrangements are boundary cases. The real EV arrangements lie rather in between, and thus the measured moduli should be smaller than those of the

cubic arrangement, but larger than those of the hexagonal arrangement. It means that the adhesion factor k_{adh} is under-estimated if the Eq. (18) is used for fitting.

Increasing of filler content ease crystallization for semi-crystalline polymers due to more nucleation sites. Thus, matrix moduli are increased due to higher crystallinities⁹. However, higher matrix moduli have the consequence that the adhesion factors k_{adh} are determined slightly smaller. This shows that adhesion factors are also affected by the choice of the matrix polymer.

Conclusion

The *cube in cube* models of Paul (upper bound) and Ishai-Cohen (lower bound) determine the moduli of particle filled composites assuming perfect filler-matrix adhesion. To take into account reduced adhesion between filler particles and matrix polymer their models were recalculated using an EV approach, and extended by an adhesion factor k_{adh} that scales the edge length of the cubic inclusion. It leads to the reduction of the filler content dependent composites moduli $E_C(v_F)$. The modified Paul model in EV version as the upper bound of $E_C(v_F)$ was used to fit experimental moduli of glass bead filled polymers and provided reasonable adhesion factors: 0.6 for polyamide (PA66), 0.5 for polybutylene terephthalate (PBT), 0.2 for polypropylene (iPP), 0.13 for low density polyethylene (PE-LD) and 0.05 for butadiene rubber (BR), which are in line with the corresponding surface energies. The modified model allows for design engineers to calculate realistic modulus of any particulate composite using only matrix and filler moduli, filler content and adhesion factor without performing tests. Further analysis of the modified model elucidated that stiffening only occurs if k_{adh} exceeds $\sqrt{E_M/E_F}$ and relates stiffening to the ratio of matrix modulus E_M and filler modulus E_F . This means almost all technical relevant filler contents lead to stiffening of the polymer. Furthermore, the EV approach shows directly that the moduli of particulate composites depend on particle shape—quantified by the efficiency factor k —and the dimensions within the EV—expressed by the normalized filler distance d which depends on a filler volume content.

Additionally, finite element analyses of cubic and hexagonal EV arrangements show that the spatial EV arrangement leads to differences of the filler content dependent composites moduli especially for high volume contents. Thus, the assumption “The properties of the EV are representative for the whole composites” only holds for filler volume contents up to 15 or 20%. The increasing difference between composites moduli calculated for cubic and hexagonal EV arrangement can be attributed to two limits of the *cube in cube* model: first, the occurrence of increasing shear stresses if EV arrangements deviate from cubic arrangements, and second, increasing interaction among filler particles with the formation of particle networks within the matrix disturbing the EV assumption of homogeneously dispersed particles. The modulus increase of rubber composite by factor 2 or 3 if the filler content changes from 30 to 45%, respectively, cannot be explained within the *cube in cube* model anymore. From a practical point of view, where commercially available particle filled polymers have filler volume contents less than 20%, it is positive that the EV arrangement affects calculated composites moduli within the experimental error of 3 to 5%.

Data availability

The datasets generated during and/or analysed during the current study are available in the Zenodo open repository maintained by <https://zenodo.org/record/6460262>.

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Author contributions

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Competing interests

The authors declare no competing interests.

Additional information

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