# **Examples of Failure Analysis of Multiphase Materials**

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Al2O3/Zr02

Problem statement  $\mathbf{KT} + \mathbf{CT} = \mathbf{F}$ 

Thermal equation of equilibrium, fulfills boundary and initial conditions

T,  $\dot{T}$  are the temperature vector and temperature rate vector

 ${\bf K}$  is the conductivity matrix,  ${\bf C}$  is the heat capacity matrix (diagonal),  ${\bf F}$  is the vector of thermal source and fluxes

 $M\ddot{u} = f - p$  M is the diagonal mass matrix, u is the displacement vector,  $\ddot{u}$  is the nodal acceleration vetor, f is the internal force vetor and p is the nodal load vector

$$\ddot{\mathbf{u}}_n = (\mathbf{M})^{-1} (\mathbf{f}_n - \mathbf{p}_n)$$
 where n denotes the current time step and  $\Delta t$  is the time interval

 $\dot{\mathbf{u}}_{n+1/2} = \mathbf{u}_{n-1/2} + \ddot{\mathbf{u}}_n \Delta t$  Velocity;  $\mathbf{u}_{n+1} = \mathbf{u}_n + \dot{\mathbf{u}}_{n+1/2} \Delta t$  displacements

 $\mathbf{T}_{n+1} = (\mathbf{C})^{-1} (\mathbf{F}_n - \mathbf{K}_n \mathbf{T}_n)$  Nodal temperature vector



f is the heat flux due to plastic strain;  $\sigma$  is the Cauchy stress tensor, epl is the plastic strain deformation rate

$$\sigma^{o} = \left(A + B\left(\overline{\varepsilon}^{pl}\right)^{n}\right) \left(1 + C\ln\dot{\varepsilon}^{*}\right) \left(1 - \zeta^{m}\right)$$

 $f = \chi \mathbf{\sigma} : \mathbf{e}^{pl}$ 

A is the yield stress, B is the hardening coefficient, C is the strain rate coefficient,  $\overline{\varepsilon}^{pl}$  is the equivalence plastic strain,  $\dot{\varepsilon}^*$  is the rate of the equivalent plastic strain

$$\zeta = \begin{cases} 0 & for \quad T < T_{trans} \\ \left(T - T_{trans}\right) / \left(T_{melt} - T_{trans}\right) & for \quad T_{trans} \le T \le T_{melt} \\ 1 & for \quad T > T_{melt} \end{cases}$$

where *T* is the current temperature, *T*<sub>trans</sub> is the temperature around which the yield stress becomes temperaturedependent, and *T*<sub>melt</sub> is the melting temperature.

### **Taylor bar**

Parameters of the Johnson-Cook material model.

Parameters	Values and units
Young's modulus	$2.1 \times 10^5 \text{ MPa}$
Poisson's coefficient	0.296
Yield stress (A)	455.0 MPa
Hardening coefficient (B)	2475.0 MPa
Hardening exponent (n)	0.9
Strain rate coefficient C	0.0235
Thermal softening exponent (m)	1.0
Melt temperature T <sub>melt</sub>	1728 K
Transition temperature T <sub>trans</sub>	293 K
Specific heat $C_H$	440 J/kg K
Thermal conductivity K	150 W/m K
Thermal expansion $\alpha_t$	$5.0 \times 10^{-5}  1/K$
Mass density	9130 kg/m <sup>3</sup>



#### Maximum displacement, equivalent plastic strain, temperature in analyzed meshing cases, and a comparison between the adiabatic and coupled solutions.

Case		Number of nodes and elements	$\begin{array}{l} \text{Displacement} \\ \times \ 10^{-2} \ \text{m} \end{array}$	Equivalent plastic strain (Element <i>E</i> o)	Temperature, [K] (Element <i>E</i> o)	Equivalent plastic strain (Node Po)	Temperature, [K] (Node <i>P</i> o)
B1	Adiabatic	Nds 21,917	1.068	1.206	759.4	1.156	759.3
	Coupled	Els 19,200	1.072	1.203	707.6	1.154	707.6
B2	Adiabatic	Nds 32,767	1.068	1.195	780.4	1.195	780.3
	Coupled	Els 28,800	1.070	1.193	720.3	1.193	720.3
B3	Adiabatic	Nds 43,617	1.067	1.206	786.2	1.206	786.2
	Coupled	Els 38,400	1.071	1.203	724.3	1.203	724.3
B4	Adiabatic	Nds 54,467	1.069	1.210	788.0	1.210	788.0
	Coupled	Els 48,000	1.071	1.206	725.7	1.206	725.6



Impact velocity 350 m/s

#### Eq plastic strain

#### Temperature

Impact velocity 550 m/s

#### Adiabatic solution 4.1236 3.6655 3.2075 Z 2.7494 X 2.2913 **X** 1.8332 1.3752 0.91709 0.45901 0.00094 (c) 1666.3 1513.7 1361.1 Z Z 1208.6 1056 X V 903.41 750.83 Temperature 598.26 445.68 293.11 (e)

### Coupled solution

3.0904 2.7471 2.4038 2.0606 1.7173 1.374 1.0307 0.68748 0.34421 0.00093 (d)

1178.3

1079.9

981.59

883.24

784.89

686.54

588.19

489.84 391.49

**293.14** (f)

#### Eq plastic strain

	Displacement $\times 10^{-2}$ m	Equivalent plastic strain (element Eo)
Adiabatic	1.742	4.124
Coupled	1.740	3.090
	Temperature [K] (Element Eo)	Equivalent plastic strain (Node Po)
	1666.3 1178.3	4.134 3.092



Relationship between the maximum equivalent plastic strain (a) and the maximum temperature (b) and the impact velocity in the adiabatic and coupled solutions.



Time variation of the maximum equivalent plastic strain for different impact velocities; (a) adiabatic solution, (b) coupled solution



Time variation of the maximum temperature for different impact velocities; (a) adiabatic solution, (b) coupled solution.





(d)

(c)

### Mesh dependency check

Mesh dependency check. Maximum displacement, equivalent plastic strain, temperature for different meshes, and a comparison between the adiabatic and coupled solutions.

Case	ł	Number of nodes and elements	Displacement x 10 <sup>-6</sup> m	Equivalent plastic strain	Temperature [K]
P1	Adiabatic Coupled	Nds 40,726/ Els 15,516/	1.7353 1.3320	3.6238 2.0727	1646.7 386.7
P2	Adiabatic	18,666 Nds 75,634/	1.6199	3.5469	1637.9
	Coupled	Els 31,032/ 37,332	1.2579	2.0431	385.7
P3	Adiabatic	Nds 101,486/	1.7254	3.5429	1637.3
	Coupled	Els 24,252/ 61,554	1.3419	2.1045	389.6
P4	Adiabatic	Nds 130,482	1.6259	3.5189	1634.4
	Coupled	Els 32,336/ 82,072	1.2776	2.0772	389.0



(a) region of the maximumequivalent plastic strain andtemperature, (b) detail of themesh with Element Mo



Comparison of the adiabatic and coupled solutions depending on the impact velocity at Element Mo; (a) equivalent plastic strain, (b) temperature



	Equivalent pl	astic strain	
	30 m/s		
	Adiabatic	Coupled	%
	0.331	0.309	7.2
	0.293	0.285	2.7
	0.283	0.268	5.8
	0.011	0.011	0.0
60	m/s		
Ad	liabatic	Coupled	%
1.1	151	0.897	22.1
0.8	805	0.742	7.8
0.8	897	0.773	13.8
0.0	042	0.042	0.0
75	m/s		
Adi	iabatic	Coupled	%
1.7	62	1.205	31.6
1.1	17	0.995	10.9
1.2	92	1.033	20.0
0.0	60	0.070	1.4
0.0		0.070	1.4

Velocity

Junction

A B

С

D

10.0

x10<sup>-8</sup>

10.0

x10-8

Equivalent plastic strain in the adiabatic an coupled solutions; (a) Junction A, (b) Junction B, (c) Junction C, (d) Junction D. Impact velocity versus equivalent plastic strain at Junctions A, B, C and D towards the end of the process.



1116

770

868

305

338

245

354

302

230.2

214.3

145.2

1.0

Temperature in the adiabatic and coupled solutions; (a) Junction A, (b) Junction B, (c) Junction C, (d) Junction D.



Temperature in the adiabatic and coupled solutions; (a) Junction A, (b) Junction B, (c) Junction C, (d) Junction D.

1	'em	pe	rat	tur	e
		-			

Velocity	30 m/s			60 m/s			75 m/s		
Junction	Adiabatic	Coupled	%	Adiabatic	Coupled	%	Adiabatic	Coupled	%
A B C D	382 367 364 294	300 302 301 294	21.5 17.2 17.3 0.0	793 600 650 300	322 328 331 298	59.4 45.3 49.1 0.7	1116 770 868 305	338 245 354 302	230.2 214.3 145.2 1.0

#### Temperature at the end of the process

HMH stress in the adiabatic and coupled solutions; (a) Junction A, (b) Junction B, (c) Junction C, (d) Junction D.



HMH stress versus impact velocity at Junctions A, B, C and D, time instant: 6.0 imes 10<sup>-8</sup> s.

	HMH stress										
Velocity	30 m/s			60 m/s			75 m/s				
Junction	Adiabatic $\times 10^{+9}$	Coupled $\times$ $10^{+9}$	%	Adiabatic $\times 10^{+9}$	Coupled $\times$ $10^{+9}$	%	Adiabatic $\times 10^{+9}$	Coupled $\times$ $10^{+9}$	%		
Α	1.465	1.537	4.9	2.341	3.294	40.7	2.087	3.891	86.4		
В	1.333	1.291	3.2	2.256	2.827	25.3	2.413	3.407	47.2		
С	1.271	1.264	0.6	2.138	2.684	25.5	2.283	3.292	44.2		
D	0.102	0.116	13.7	0.488	0.487	0.20	0.638	0.635	0.5		



Temperature at Elements Aa, Ba, Ca and Da in the grains adjacent to the binders, coupled solution; (a) Junction A, (b) Junction B, (c) Junction C, (d) Junction D.

	Temperature					
Velocity	30 m/s	60 m/s	75 m/s			
Element adjacent to Junction	[K]	[K]	[K]			
Α	301	326	344			
В	301	325	344			
C	299	324	343			
D	294	297	300			

Impact velocity versus temperature in the grains at Elements Aa, Ba, Ca and Da adjacent to Junctions A, B, C and D, time instant: 6.0 × 10–8 s.

14.0 x10+9 12.0 x10+9 adiabatic, 30 m/s ---adiabatic, 30 m/s ---coupled, 30 m/s -----adiabatic, 60 m/s ----coupled, 30 m/s -----12.0 10.0 adiabatic, 60 m/s ·· ·· coupled, 60 m/s .---coupled, 60 m/s -----10.0 adiabatic, 75 m/s ..... adiabatic, 75 m/s ..... HMH stress (Pa) HMH stress (Pa) 8.0 coupled, 75 m/s coupled, 75 m/s 8.0 6.0 6.0 4.0 4.0 2.0 2.0 n 6.0 8.0 10.0 8.0 10.0 0 2.0 4.0 2.0 4.0 6.0 0 x10-8 x10<sup>-8</sup> Time (s) Time (s) (a) (b) 8.0 x10+5 6.0 x10+9 adiabatic, 30 m/s -----coupled, 30 m/s adiabatic, 60 m/s ---adiabatic, 30 m/s ----coupled, 30 m/s ---adiabatic, 60 m/s ··· ·· coupled, 60 m/s ----coupled, 60 m/s -----6.0 HMH stress (Pa) HMH stress (Pa) adiabatic, 75 m/s ..... adiabatic, 75 m/s ..... 4.0 coupled, 75 m/s ---coupled, 75 m/s ----4.0 2.0 2.0 0 4.0 10.0 2.0 6.0 8.0 0 0 2.0 4.0 6.0 8.0 10.0 x10<sup>-8</sup> x10<sup>-8</sup> Time (s) Time (s) (c) (d)

HMH stress at Elements Aa, Ba, Ca and Da in the grains adjacent to the binders in the adiabatic and coupled solutions; (a) Junction A, (b) Junction B, (c)

Junction C, (d) Junction D.

H	IMH	stress
		10.04 20000

Velocity	y 30 m/s			60 m/s			75 m/s		
Junction	Adiabatic $\times 10^{+9}$	Coupled $\times$ 10 $^{+9}$	%	Adiabatic $\times 10^{+9}$	Coupled $\times$ $10^{+9}$	%	Adiabatic $\times 10^{+9}$	Coupled $\times$ $10^{+9}$	%
Α	2.803	2.837	1.2	5.371	5.779	32.2	6.526	7.525	15.3
B	2.076	2.631	27	5.260	5.772	9.7	6.752	7.329	7.2
С	1.429	1.416	1.0	3.378	3.430	1.5	4.286	4.443	3.7
D	0.279	0.309	11.0	0.691	0.664	3.9	0.840	0.828	1.4



Displacement (m) fields (magnified 5 times); (a) adiabatic solution; (b) coupled solution.

HMH stress (Pa), time instant:  $2.5 \times 10-9$  s; (a) adiabatic solution, (b) coupled solution.





HMH stress (Pa) in the binders, time instant:  $10.0 \times 10-9$  s; (a) adiabatic solution, (b) coupled solution.





HMH stress (Pa), time instant: 75.0 × 10–9 s; (a) adiabatic solution, (b) coupled solution.



HMH stress (Pa), time instant: 100 × 10–9 s; (a) adiabatic solution; (b) coupled solution.



Equivalent plastic strain variations at the junctions; (a) adiabatic solution; (b) coupled solution; impact velocity 75 m/s





Temperature distribution (K); (a) adiabatic solution; (b) coupled solution`, End of the process, Impact velocity 75 m/s





• The displacement fields obtained for the two solutions are qualitatively similar;

• Although the HMH stress fields in both cases are similar with respect to quality, the maximum stress in the binders is lower than that in the grains;

• The equivalent plastic strains in the binders are lower in the coupled solution;

- The maximum temperature is significantly lower in the coupled solution;
- The polycrystalline material grains are affected by temperature increase;

• Given the significant differences in the solutions, the adiabatic solution should not be used for the analysis of polycrystalline materials.

# **Peridynamics method**

 ${f Q}$  and  ${f x}$  are the points inside an undeformed body. The bond definition

 $\boldsymbol{\xi} = \boldsymbol{Q} - \boldsymbol{x}$ 

The reference state  $\mathbf{X}$  is a function that is valid on the bond  $\mathbf{X}(\boldsymbol{\xi})$ . The deformation state is dependent on the new position of the coordinate  $\mathbf{x}$  in the deformed body  $\mathbf{y}(\mathbf{x})$  in the following way

$$\mathbf{Y}(\mathbf{x},\boldsymbol{\xi}) = \mathbf{y}(\mathbf{x}+\boldsymbol{\xi}) - \mathbf{y}(\mathbf{x}),$$

$$\mathbf{Y}(\mathbf{x},\boldsymbol{\xi}) = \mathbf{y}(\mathbf{Q}) - \mathbf{y}(\mathbf{x}).$$

The state of displacements is:

$$\mathbf{U}(\mathbf{x},\boldsymbol{\xi}) = \mathbf{u}(\mathbf{x}+\boldsymbol{\xi}) - \mathbf{u}(\boldsymbol{\xi}),$$
  
$$\mathbf{U}(x,\boldsymbol{\xi}) = \mathbf{u}(Q) - \mathbf{u}(x).$$

The scalar extension state  $e(\mathbf{Y})$  is of the form:

 $e(\mathbf{Y}) = \big|\mathbf{Y}\big| - \big|\mathbf{X}\big|.$ 

The decomposition of the scalar extension state into spherical and deviatoric components





The force state  $t(\mathbf{Y})$  is shown in a form similar to the standard stress-strain relation that is the sum of its spherical and deviatoric parts

$$t(\mathbf{Y}) = \left(\frac{3k\theta}{m}\right)\omega x + \alpha \omega e^d.$$

k is the bulk modulus,  $\theta$  is the dilatation, m is the weighted volume,  $\omega$  is the influence function,  $x = |\xi|$  is the basic scalar state,  $\alpha = (15\mu)/m$  is the coefficient being in relation to the shear modulus  $\mu$ .



Integration scheme

Constitutive law

A special case of the state-based model is the bond-based model

$$f = ce_{\zeta}(\mathbf{x}, t, \xi), \qquad c = (18k)/(\pi h^4)$$

c depends on the bulk modulus k and the horizon h

The failure is predominantly assumed on cracks appearance, the  $e_{cr}$  evaluation is based on fracture energy evaluation,

$$e_{cr} = \sqrt{\frac{5G_{ci}}{9kh}}$$

where  $G_{ci}$  is the fracture energy depending on the mode of failure

# **Peridynamics method**

The material failure model is based on the following assumptions: the bonds fail when their elongation overcomes the critical elongation; the total damage is due to the accumulation of broken bonds; and the bond breaking is an irreversible process.

The damage definion *d* reads:

$$d(x,t) = 1 - \frac{\int_{\Omega} \zeta(x,t,\xi) d\Omega}{\int_{\Omega} d\Omega}.$$

The damage variable varies between 0 and 1. The value 0 means that the material is pristine while 1 means that the material is completely damaged.

$$G_I = \frac{\left(1 - \nu^2\right)K_I}{2E}$$

## Contact



The contact is checked between the triangular facets of the bodies that are contact candidates. The sphere is transformed into an icosahedron

Transformation of a sphere into an inscribed icosahedron: (**a**) sphere; (**b**) icosahedron.

$$f_{ct} = \begin{cases} C_t (r_r - d) V_1 V_2 & \text{for } d \leq r_r \\ 0 & d > r_r \end{cases}$$

where *d* is the distance between the bodies,  $V_1$  and  $V_2$  are the volumes associated with the corresponding calculation points and  $C_t=18K_s/\pi h^5$ . The constant *Ks* is the artificial spring stiffness assumed as a high **penalty number** of the range  $1.0E+12\div1.0E+20$ .





#### Al2O3/ZrO2



Schemes of the analyzed samples (dimension x1.0E-07 m),  $Al_2O_3 - grey$ ,  $ZrO_2$  - blue: (**a**) Low content of  $Al_2O_3$ , Case A (41%); (**b**) High content of  $Al_2O_3$ , Case B (75%).



Damage advancement (dimension x 1.0E-5 m), Case A, impact velocity 60 m/s: (a) time 1.125E-08 s; (b) time 2.375E-08 s; (c) time 3.000E-07 s; (d) time 4.000E-08 s.



Damage advancement (dimension x 1.0E-5 m), Case B, impact velocity 60 m/s: (a) time 1.125E-08 s; (b) time 2.375E-08 s; (c) time 3.000E-07 s; (d) time 4.000E-08 s.



Damage advancement, the highest distance of a point of d>0.8 from the hitting edge: (a) Case A; (b) Case B.



Damage advancement, the highest distance of a point of d>0.8 from the hitting edge, microstructure comparison: (a) impact velocity 10 m/s; (b) impact velocity 60 m/s.



Damage advancement (dimension x 1.0E-5 m), the highest distance of a point of d>0.8 from the hitting edge at the impact velocity 10 m/s: (**a**) time 2.375e-8 s ; (**b**) time 3.000e-8 s.



Comparison of damage variable distributions in the cross-section of the plate along x-axis at the mid-span of the plate, from the top: Case A, impact velocity 30 m/s, Case A, impact velocity 60 m/s, Case B, impact velocity 30 m/s and Case B, impact velocity 60 m/s.



Comparison of variations of damage variable in time for Case A and B at different points: (a) Point A; (b) Point B; (c) Point C; (d) Point D.

Damage in Case B starts later



Variation of damage variable in time for different impact velocities, Case A: (a) Point A; (b), Point B; (c) Point C; (d) Point D.



Variation of percentage of total damage in time for different impact velocities: (a) Case A; (b), Case B.



Dependence of percentage of total damage at particular time instants on impact velocity: (a) Case A; (b), Case B.

Following the considerations, we summarize and arrive at the conclusions as follows:

•Damage analysis of the impacting plates should be performed in threedimensions,

•Damage is concentrated close to the mid-span of the depth of the cross-section and is the lowest close to the surfaces of the plates,

•Damage growth during impact that is characterized by percentage of total damage of the plate is strongly nonlinear,

•Damage appears early at sparse distributed points well before damage front where the damage becomes massive,

•At least in the investigated impact velocity range, the traces of damage appears practically in the entire plate independently of the impact velocity what means that almost the entire plate appears to be damaged to the different extent,

•When taking into account total damage variation dependence on impact velocity, the damage variation is almost linear at particular time instants. The latter allows for the damage prediction for different impact velocities using linear interpolation.

•At the beginning of the process, damage forms lines that are parallel to the hitting edge. We concern the phenomenon as a characteristic feature of the damage process of thin ceramic platelets.

### **Material parameters:**

- Skeleton:Young's modulus(SiC)Poison's ratioMass densityFracture toughness
- 430.0 GPa 0.75 3200 kg/m3 4.1 MPa.(m)^(1/2)

#### **Base and**

piston: Young's modulus (steel) Poison's ratio Mass density 210.0E+09 Pa 0.3 7800.0 kg/m3





internal structure of the SiC foam

R 19,7 Y X 0 11,3 P 1,1 4,0 21,8 Base

18,6



**Contact model** – general contact, penalty formulation, friction coefficient 0.3

**Discretization**:

total 461,496 calculation points Skeleton - 261,496 Base - 100,000 Piston - 100,000

## **Impact velocities:**

40 m/s and 365 m/s

Critical stretch: SiC – 9.5716E-05

Horizon; skeleton: 30.0E-04 m, base and piston 6.0E-04 m Process time: 3.06E-05, time increment 2.0e-08 s (below critical time), 1500 increments, solution time 3800 s, 1920 CPUs

Piston

SCF

Steel base

Positions of the points in the details: region P, region Q, region R



Total damage variation

Local damage (at points) variation



#### Points where d>0.95



#### V=40 m/s, end of the process



# Numerical example



### V=385 m/s Points where d>0.95

#### V=385 m/s, end of the process

- When the sample is subjected to the high-velocity impact, the structure undergoes self-contact in the pores.
- The destruction of the SCF into fragments of the structure appears in the course of a high-velocity impact.
- During high velocity process the self-contact appears.
- In the low-velocities impact, the damage of the structure is mainly due to local microcracks in the material.
- During high velocity impact appears out of plane failure mode.



### **5. Numerical model**





The **SCF** sample is discretized with 261,496 volumes, while the steel base counts 100,000 volumes.

In the calculations, the **horizon** *h* value for the:

- foam is assumed of 30.0E-04 m,
- base is 6.5E-04 m.



Percent of damage of the sample for low velocities, and for high velocities.



Damage at the observed points: low velocities, high velocities.



Impact velocity **15 m/s**, damage at time 2.4E-06 s, 11.4E-06 s , 19.8E-06 s



Impact velocity 15 m/s, points of damage d < 0.8 at time 2.4E-06 s, 11.4E-06 s, 19.8E-06 s



Impact velocity 800 m/s, damage distribution at time 2.4E-06 s, 11.4E-06 s, 19.8E-06 s.



Impact velocity 800 m/s, points of damage d < 0.8 at time 2.4E-06 s, 11.4E-06 s, 19.8E-06 s

A qualitative comparison of the high and low-velocity impact of a SiC foam has been presented in the paper. The analysis is performed using the peridynamics method. It has been found that the behaviour of the foam is different when the impact velocity is low in comparison to the high velocity response. The distinction of the low- and highvelocity process is based on evaluating total damage accumulation in the structure.

- In the high-velocities impact range, the total damage of the foam sample grows almost linearly with slight deviations from the linearity.
- When the foam sample is subjected to the high-velocity impact, the structure undergoes self-contact in the pores.
- The destruction of the SCF into fragments of the foam structure appears in the course of a high-velocity impact.
- While the impact is performer with the low-velocity, the fragmentation and selfcontact does not appear.
- In the low-velocities impact, the damage of the foam structure is mainly due to local microcracks in the material.

#### **Remarks on HPC**

Number of processes	Adiabatic so	lution	Coupled sol	Coupled solution		
	Time (s)	Efficacy	Time (s)	Efficacy		
4	5396	1	6362	1.00		
8	2746	1.97	3248	1.96		
16	1548	3.49	1879	3.39		
24	1126	4.79	1335	4.77		
32	1005	5.37	1199	5.31		
48	777	6.94	935.0	6.80		
96	733	7.36	837.0	7.60		

Number of processes and solution time for the adiabatic and coupled solutions.







We used 960, 1920 and 3840 processes with the wall-clock times 930 s, 530 s and 290 s, respectively

The calculations were done using PL-GRID national computational resources at the Interdisciplinary Centre for Mathematical and Computational Modeling, University of Warsaw, the CYFRONET, Krakow, and Academic Computer Centre in Gdańsk, Poland.

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# Prerequisites for modelling

- grain size distribution,
- interface thickness distribution
- pore size distribution
- pore placement inside the material
- mechanical properties: Young modulus, Poisson coefficient, yield stress, material hardening, material viscosity