



Cite this: *RSC Adv.*, 2018, 8, 24773

Received 20th April 2018
 Accepted 21st June 2018

DOI: 10.1039/c8ra03409e

rsc.li/rsc-advances

Electrorheological behavior of iron(II) oxalate micro-rods

E. Kutalkova, T. Plachy, J. Osicka, M. Cvek, M. Mrlik and M. Sedlacik *

Electrorheological (ER) fluids represent smart materials with extensive application potential due to their rheological properties which can be readily changed under an external electric field. In this study, the iron(II) oxalate particles with rod-like morphology were successfully synthesized by the co-precipitation method using sulphate heptahydrate and oxalic acid dihydrate. The characterization of particles was performed via X-ray diffractometry and scanning electron microscopy. Subsequently, the ER fluids were prepared by dispersing the synthesized particles in silicone oil. The optical microscopy demonstrated the formation of chain-like particle structures upon the application of an electric field. Rheological properties were determined by means of rotational rheometry including creep-recovery experiments. The viscoelastic behavior of systems under investigation in the presence of the electric field was confirmed by the presence of recoverable strain of the system.

Introduction

Electrorheological (ER) fluids are well-known intelligent systems whose rheological parameters can be controlled by application of an external electric field. They are typically two-phase systems consisting of electrically-polarizable particles dispersed in a non-conducting liquid medium. After application of the electric field, the particles start to join themselves into chain-like structures spanning the electrodes due to dipole-dipole interactions.¹ This behaviour is accompanied by a significant change in rheological parameters and is called the ER effect. The intensity of the ER effect depends on the electric field strength,² E , the volume fraction of the particles,^{3,4} ϕ , their morphology, size,⁵⁻⁷ and conductivity,^{8,9} σ . Moreover, dielectric properties such as dielectric permittivity of the particles, dielectric relaxation strength, $\Delta\epsilon$, and relaxation time, τ_{rel} , of the ER fluids,¹⁰ and temperature,^{11,12} T , play a significant role in achieving a high ER effect. Depending on these parameters the ER fluids thus undergo transitions from liquid to viscoelastic systems exhibiting a portion of elastic (recoverable) deformation.^{1,13-15}

One of the important parameters of the ER fluids is the yield stress, τ_y , representing the value of stress that the ER fluids can withstand before they start to flow. It has been shown that rod-like particle-based ER fluids exhibit higher ER effects (higher τ_y) than those containing the equivalent amount of spherical particles not only due to stronger friction forces but also stronger dipole moments developed in the built-up chains of polarized particles after switching on the electric field.^{6,7,16,17} On

the other hand, another important feature of the ER fluids is their efficiency, e . In the case of the ER fluids containing the rod-like particles their η_0 is often higher since the rod-like particles possess higher viscous drag forces.¹⁸ The higher viscosity of the systems containing the rod-like particles can then be explained as a higher energy required for the orientation of such particles in the direction of the flow when compared to spherical ones.¹⁸

The co-precipitation method is a synthetic route that supports crystal growth in a certain direction, thus a material with unique properties resulting from the shape anisotropy can be obtained. The molar ratio of the precursors and the conditions during the co-precipitation determine morphology and size of the particles which can be thus optimized in order to obtain material with preferable ER effects.¹⁹ Moreover, this method is facile, cost effective and enables large-scale production.¹⁹

Iron(II) oxalate dihydrate ($\text{FeC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$) is an inorganic compound often employed in solid-state chemistry as precursor for thermally-induced syntheses of various nanocrystalline metal oxides.²⁰ A wider potential of this material recently appeared when Plachy *et al.*²¹ successfully employed the iron oxalate as a stabilizing agent in magnetorheological applications. In this study, rod-like iron(II) oxalate dihydrate particles prepared *via* co-precipitation method were introduced to electrorheology as a novel dispersed phase to fabricate the ER fluids. A dependence of the ER effects on the particles concentration was investigated and viscoelastic properties of the ER fluids were further studied through creep-recovery test confirming their transition from a liquid state to a viscoelastic solid in the presence of the electric field.

Centre of Polymer Systems, University Institute, Tomas Bata University in Zlín, tr. T. Bati 5678, 760 01 Zlín, Czech Republic. E-mail: msedlacik@utb.cz



Experimental

Particle synthesis

Iron(II) oxalate particles were synthesized *via* the co-precipitation method using iron(II) sulphate heptahydrate (Sigma Aldrich; USA) and oxalic acid dihydrate (Lach-Ner; Czech Republic).^{21,22} Briefly, iron(II) sulphate heptahydrate ($\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$, $\geq 99\%$) and oxalic acid dihydrate ($\text{C}_2\text{H}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$, p.a.) were used in an equivalent amount (20 mmol) and were separately dissolved in a 28 mL mixture of ethyleneglycol (Penta; Czech Republic) and demineralized water 3 : 1 (v/v). The oxalic acid solution served as a precipitating agent, which was slowly added into the iron sulphate solution under stirring using a magnetic stirrer. While the solution was further treated in a pressurized reactor in the previous study,²² in this case the stirring proceeded until yellow precipitate was developed. Subsequently, the product was thoroughly filtered-off with a distilled water and dried overnight at the temperature of 70 °C.

Particle characterization

The prepared iron(II) oxalate particles were investigated using an X-ray diffractometry (XRD) to confirm their successful preparation. The XRD data was collected monitoring the diffraction pattern appearing in the 2θ angle range of 10–95° (scan speed of 1° min⁻¹) with Co K α 1 radiation using an X-ray diffractometer Miniflex 600 (Rigaku, Japan). The conductivity of the particles (1.5×10^{-8} S m⁻¹) was measured by the two-point method at ambient temperature using an electrometer (Keithley 6517B, USA). A scanning electron microscope (SEM; Nova NanoSEM 450, FEI, USA) was used to observe size, morphology, and uniformity of the prepared particles. Density of the prepared particles was determined with the help of a gas pycnometer (UltraFoil 1200e, Quantachrome Instruments, Germany) using a nitrogen as a gaseous medium. The density of synthesized particles was 2.35 ± 0.05 g cm⁻³. The obtained density is further expressed as a mean value and standard deviation of 10 independent measurements. The ER fluids were obtained by mixing the dried particles with silicone oil (Lukosiol M200, Chemical Works Kolín, Czech Republic, dynamic viscosity $\eta_c = 194$ mPa s at the temperature of 25 °C) at various volume concentrations of the particles (1, 5, and 10 vol%). Before each measurement, the prepared ER fluids were manually stirred for 5 min with a glass stick.

Optical microscopy

Optical microscopy was used in order to evaluate the formation of internal chain-like structures in the prepared ER fluid. The representative sample of concentration 0.5 vol% was placed on the glass plate between two copper electrodes while the external electric field strength of 0.5 kV mm⁻¹ was supplied by the DC high-voltage source (Keithley 2400, USA). The chain-like structures were observed using an optical microscope (Leica DVM2500; Leica Microsystems, UK).

Rheological measurements

Electrorheological measurements were performed using a rotational rheometer Bohlin Gemini (Malvern Instruments, UK) with a plate–plate geometry (diameter of 20 mm with a gap of 0.5 mm) at 25 °C. The external electric field strengths of 0.3–1.5 kV mm⁻¹ were induced by the high-voltage source DC TREK (TREK 668B, USA). In the case of the on-state characterization, the electric field was applied for 1 min before starting the shearing in order to provide time enough for the particles to create equilibrium chain-like structures. After each on-state test run, the constant shear rate of 20 s⁻¹ was applied for 1 min to disrupt the residual structures.

Creep and creep-recovery experiments were performed in order to investigate the viscoelastic properties of prepared ER fluids containing the iron(II) oxalate particles. In the presence of the external electric field, recoverable strains, γ_{REC} , of the ER fluids were observed, while during the measurements in its absence, zero recoverable strain was found. The recoverable strain can be calculated as (eqn 1):

$$\gamma_{\text{REC}} = (\gamma_i - \gamma_f) / \gamma_i \times 100 (\%) \quad (1)$$

where γ_i is the maximum strain before removing the applied shear stress and γ_f is the final strain.

Results and discussion

X-ray diffraction patterns of the prepared particles possess all the characteristic peaks confirming dominant presence of the FeC_2O_4 phase (Fig. 1).^{21,22} In the X-ray spectrum the diffraction peak is dominant at approximately 21° 2θ indexed for material as the (111) peak.

The prepared particles possess rod-like morphology with relatively high aspect ratio. The length of the particles was ranging from 15 to 30 μm and their thickness was ranging from 1 to 2 μm (Fig. 2).

As can be seen, in the absence of the electric field the investigated ER fluids exhibited random distribution of

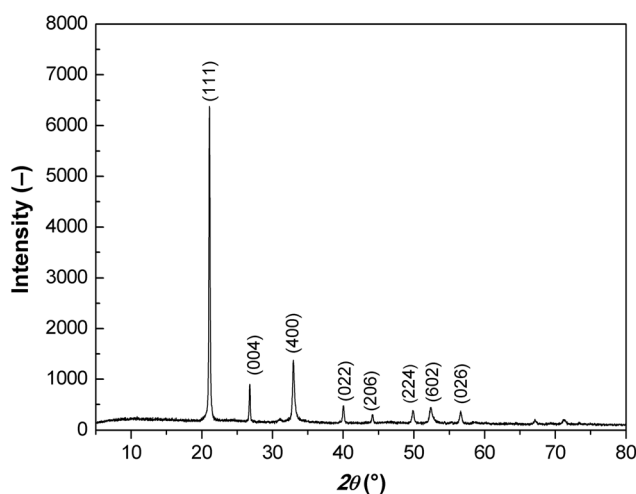


Fig. 1 XRD results of the prepared iron(II) oxalate particles.



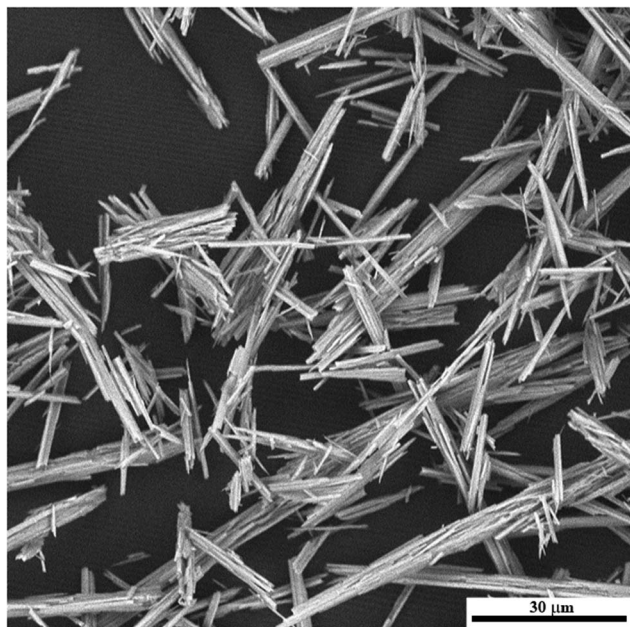


Fig. 2 SEM image of the prepared rod-like iron(II) oxalate particles.

dispersed particles (Fig. 3a). However, after its application the particles start to span the gap between the electrodes creating oriented chain-like structures (Fig. 3b). This particle alignment is associated with a significant increase in the viscosity of the system as will be described further in detail.

Fig. 4 demonstrates rheological properties of silicone-oil suspensions containing the iron(II) oxalate particles at the concentrations of 1, 5, and 10 vol% in the absence as well as in the presence of the external electric field strengths of 0.3; 0.6; 0.9; 1.2, and 1.5 kV mm^{-1} . A substantial increase of shear stress, τ , can be observed upon the application of the external electric field, when the particles in silicon-oil suspensions start to develop internal chain-like structures along the direction of electric field applied. This effect also leads to the increase in viscosity of the system. As can be seen in Fig. 4, in the absence of the electric field, the shear stress of prepared ER fluids is almost linearly dependent on the shear rate illustrating nearly newtonian behavior. In the presence of the electric field, however, the ER fluids start to exhibit τ_y as a rule the higher electric field the higher value of τ_y due to polarization forces between the particles. With increasing particles concentration in the system τ_y values further increase due to enhanced electrostatic interactions acting between dispersed particles. The ER fluid with particle loading of 10 vol.% possesses the shear stress values substantially higher (about more than 1.5 order of a magnitude) at low shear rate values than the ER fluids with lower particles loading (Fig. 4c). Upon the application of the external electric field strength of 1.5 kV mm^{-1} in the case concentration of the particles 10 vol%, the yield stress values are around 80 Pa. The observed ER performance is thus comparable with that of recently-developed ER fluids containing the particles prepared *via* multi-step syntheses.^{23,24} Hence, the presented material in this study can be characterized by two benefits, the sufficient ER

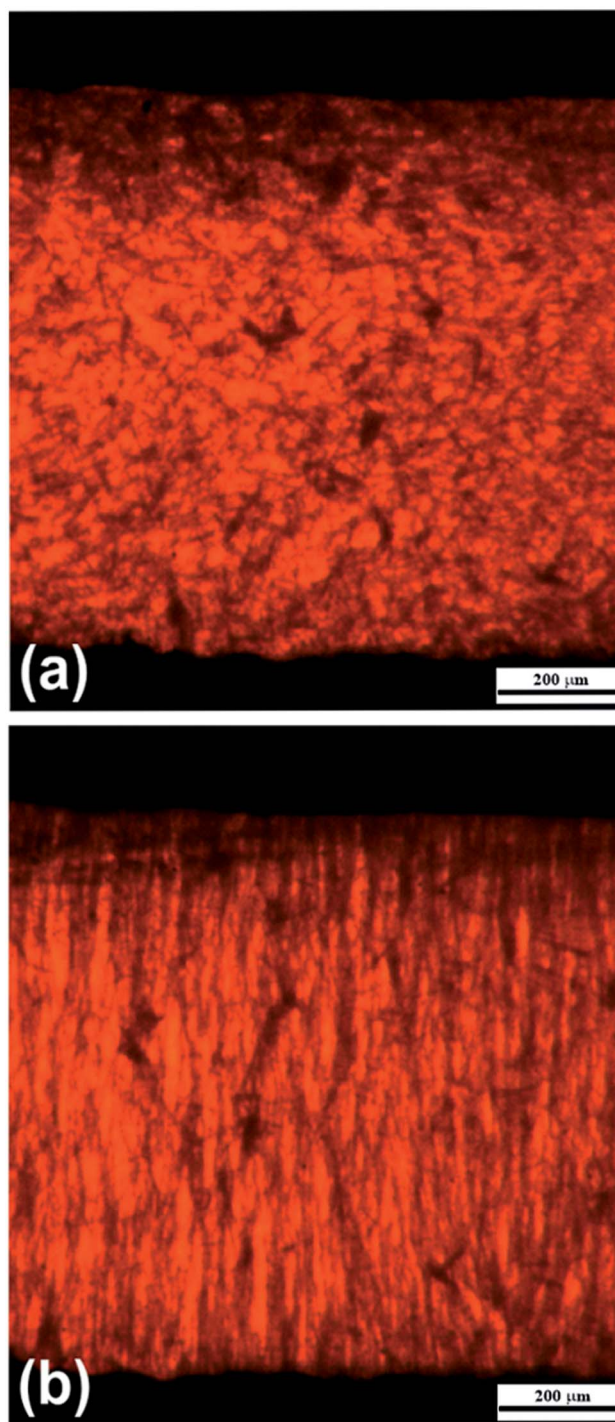


Fig. 3 Optical microscopy images of silicone-oil suspensions containing the iron(II) oxalate particles (0.5 vol%) at 0 kV mm^{-1} (a) and 1 kV mm^{-1} (b) of the external electric field.

performance and its facile synthesis not requiring any special or toxic chemicals.

In the case of ER fluids, Cho-Choi-Jhon (CCJ) model has been proposed very well for fitting the flow curves of ER fluids (eqn (2)):²⁵



$$\tau = \frac{\tau_y}{1 + (t_1 \dot{\gamma})^\alpha} + \eta_\infty \left(1 + \frac{1}{(t_2 \dot{\gamma})^\beta} \right) \dot{\gamma} \quad (2)$$

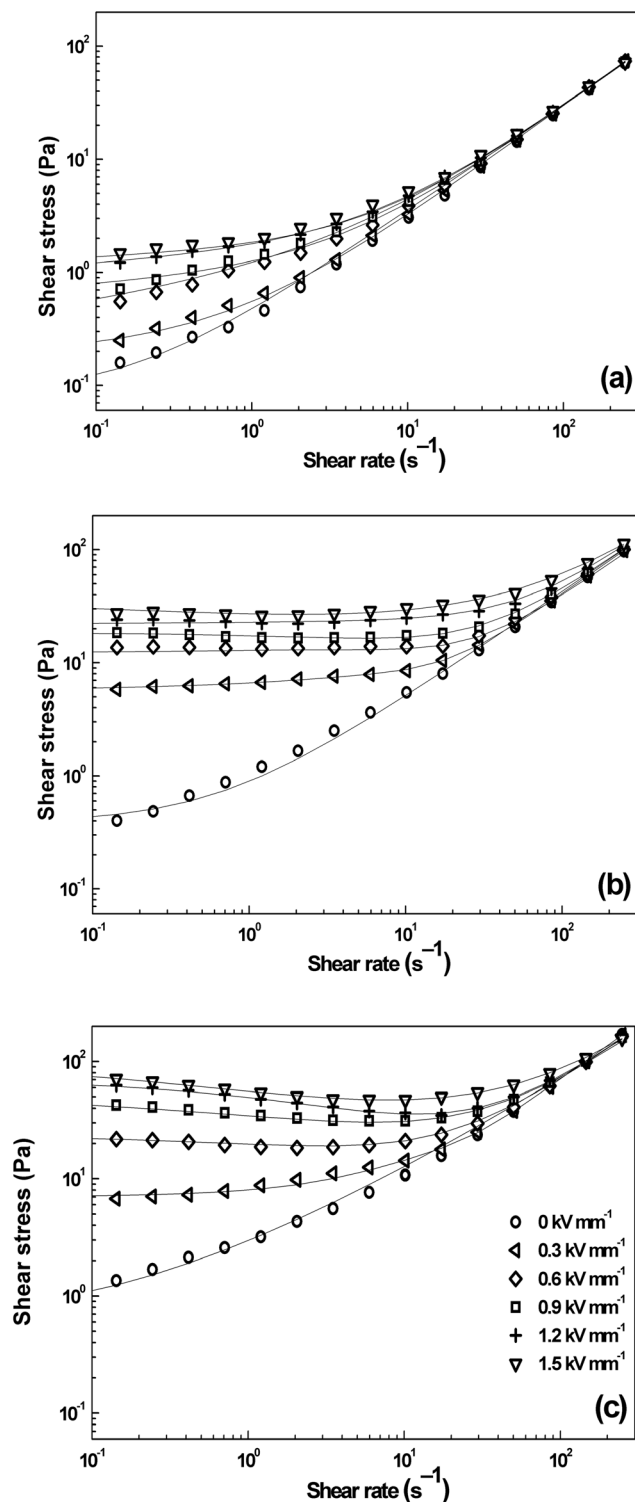


Fig. 4 Dependences of the shear stress on the shear rate for the ER fluids containing the iron(II) oxalate prepared at particle concentrations of 1 vol% (a), 5 vol% (b), and 10 vol% (c) investigated in the absence of the electric field 0 kV mm⁻¹ (circles) and in the presence of the electric field strengths of 0.3 kV mm⁻¹ (left triangles), 0.6 kV mm⁻¹ (diamonds), 0.9 kV mm⁻¹ (squares), 1.2 kV mm⁻¹ (crosses), and 1.5 kV mm⁻¹ (down triangles). The solid lines refer to the CCJ model predictions.

where τ_y is the dynamic yield stress defined as the extrapolated stress from low shear rate region, the t_1 and t_2 are time constants inverse of the shear rate representing the region where the shear stress exhibits a minimum at the low shear rate and inverse to the shear rate at which a pseudo-newtonian behavior starts. The exponent α is related to the decrease in the shear stress and the values of β are between 0–1, since $d\tau/d\dot{\gamma} > 0$, and the η_∞ represents shear viscosity at high shear rates. The yield stresses and other parameters are summarized for ER fluid with 5 vol% of the iron(II) oxalate particles in the Table 1. The CCJ data fit is represented in Fig. 4 by solid lines.

Another important rheological property for the description of the ER fluids behavior is ER efficiency, e (eqn (3)). This factor shows the behavior changes in the viscosity of the ER fluids in the absence and in the presence of external electric field. It can be calculated as:

$$e = (\eta_E - \eta_0)/\eta_0 \quad (3)$$

where η_E is viscosity in the presence of the external electric field and η_0 is field-off viscosity.

Fig. 5 shows the dependence of the ER efficiency on the shear rate. The ER efficiency of ER fluids under investigation is enhanced with increasing intensity of electric field at each concentration.

The ER efficiency is significantly higher at low shear rates compared to high shear rates as the electrostatic forces are dominating over hydrodynamic ones. However, the latter forces increase with increasing shear rate and the domination of electrostatic forces is disappearing since the rupturing of formed internal structures is starting to continuously predominate over the self-healing ability of the chain-like structure given just by the polarization of particles. The highest ER efficiency (Fig. 6) is determined at the concentration of 5 vol% and the electric field strength of 1.5 kV mm⁻¹. It is evident that with further increase in particle concentration the efficiency is not increasing anymore due to the effect of increased field-off viscosity. Similar phenomenon with optimum concentration of the particles on the ER performance of ER fluids has been already introduced.⁴ The effect of competition between electrostatic forces and hydrodynamic ones described above can be also observed in this case.

The τ_y values obtained applying eqn (2) on experimental data for all investigated ER fluids was further used to show the increasing rigidity of created internal structures within the ER fluids after the application of the external electric field in Fig. 7. The dependence of the τ_y of ER fluids on the electric field strength, E_c , applied obeys mostly the power law dependence (eqn (4)):²⁶

$$\tau_y = q \times E_c^\alpha \quad (4)$$

where parameter q describes a rigidity of the system and α is a slope of linear curve fitting the data. This power law dependence demonstrates the conduction is the predominant factor



Table 1 Calculated CCJ model parameters for the ER fluid containing 5 vol% iron(II) oxalate particles under various electric field strengths

Parameters	0 kV mm ⁻¹	0.3 kV mm ⁻¹	0.6 kV mm ⁻¹	0.9 kV mm ⁻¹	1.2 kV mm ⁻¹	1.5 kV mm ⁻¹
τ_y	0.4	5.1	12.1	18.1	22.2	26.5
t_1	0.05	0.08	0.06	0.03	0.01	3.40
α	0.41	1.75	1.20	0.68	0.93	0.46
η_∞	0.28	0.38	0.34	0.39	0.39	0.32
t_2	0.59	0.25	0.29	0.90	0.23	0.01
β	0.16	0.84	0.65	0.99	0.76	0.90

for developing internal structures of the particles when the exponent α is close 1.5, while the polarization mechanism is predicted by α exponent close to 2.²⁷ The slope of the dependence in Fig. 7 is very close to 1.5 irrespectively the concentration of iron(II) oxalate particles used indicating the conduction model is dominant factor for ER performance for these particles.

A creep test involves two distinct phases, the creep phase and the recovery phase. In the former one, a constant stress is applied to the sample for a fixed time period. After removal of the τ some deformation may be recoverable (the recovery phase). For a viscoelastic material, recoverable strain, γ_{REC} , is expressed as in eqn (1).

The strain of viscoelastic material increases with time under the application of constant τ to the samples for 30 s. After the removal of applied τ , the deformation in the case of viscoelastic materials is partly recoverable. Shear stress values were taken below the elastic stress, τ' , evolved from storage modulus, G' , of the ER fluid by the relation expression of $\tau' = G'\gamma$, where γ represents deformation strain, since τ' represents the maximum value to support complete recovery when τ is eliminated.²⁴ These τ values (0.35 Pa, 0.7 Pa, 1.4 Pa) were applied to the ER fluid (5 vol%) under investigation in the presence of the electric field strength of 0.9 kV mm⁻¹ (Fig. 8). After application of the

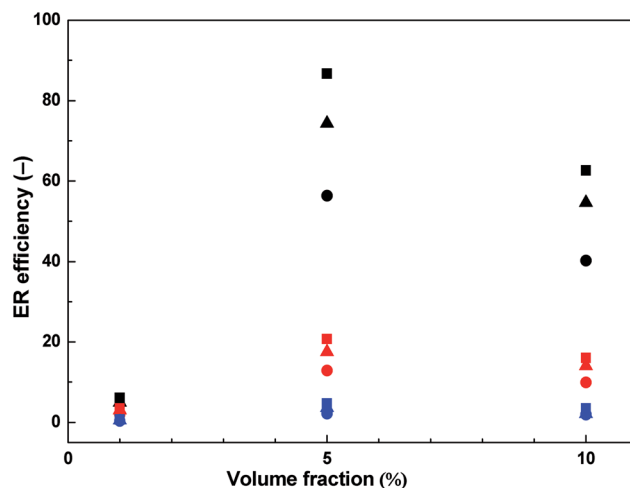


Fig. 6 Dependences of the ER efficiency on the particle volume fraction at shear rates of 0.1 s⁻¹ (black), 1 s⁻¹ (red) and 10 s⁻¹ (blue) in the presence of the electric fields 0.9 kV mm⁻¹ (circles), 1.2 kV mm⁻¹ (up triangles) and 1.5 kV mm⁻¹ (squares).

external electric field, the ER fluid changes its solely viscous behaviour to viscoelastic one and exhibits a certain level of the γ_{REC} depending on the intensity of the applied τ (Fig. 8). The γ_{REC} was evaluated according to eqn (1) (Table 2). These observations confirmed the transition of the investigated ER

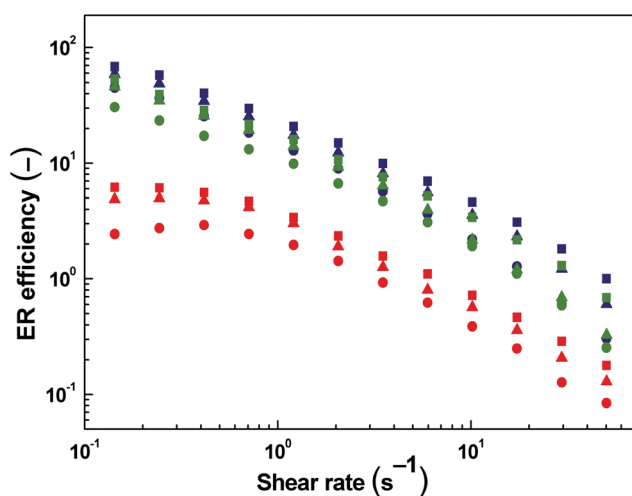


Fig. 5 Dependences of the ER efficiency on the shear rate for the ER fluids containing the iron(II) oxalate prepared at particle concentrations of 1 vol% (red), 5 vol% (blue) and 10 vol% (green) in the presence of the electric field strengths of 0.9 kV mm⁻¹ (circles), 1.2 kV mm⁻¹ (up triangles), and 1.5 kV mm⁻¹ (squares).

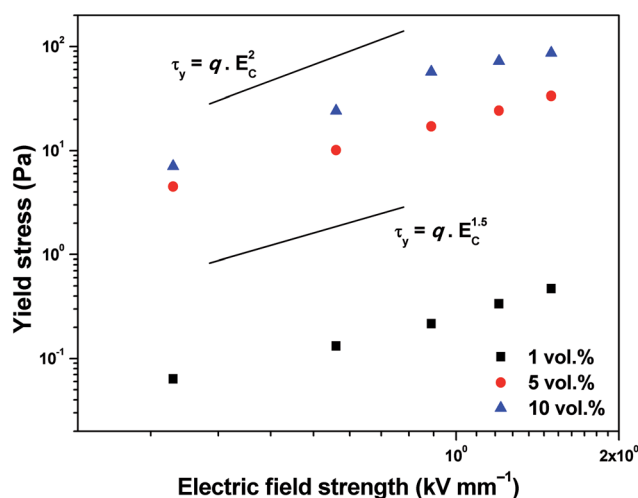


Fig. 7 Dependence of the dynamic yield stress on the applied electric field strength for ER fluids iron(II) oxalate particles of various concentrations.



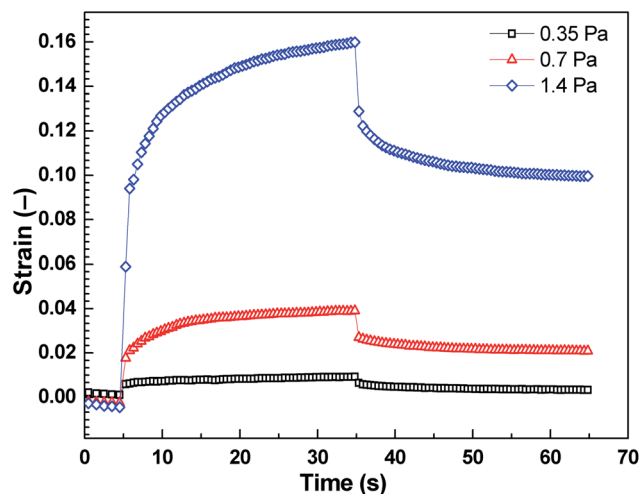


Fig. 8 Creep and recovery responses for the ER fluids (5 vol%).

systems from liquid to viscoelastic solids upon the application of the electric field. As can be seen in Fig. 9, the applied τ value above the τ' caused continuous flow resulting in almost no γ_{REC} .

Electrorheological reproducibility (Fig. 10) of the system after the switching on and off an external electric field has been confirmed by means of the time course of shear stress at the shear rate 1 s^{-1} . In the absence of field, polarization of the particles disappears and viscosity is relatively quickly returned

Table 2 Recoverable strains for the ER fluid containing 5 vol% of iron(II) oxalate particles in the presence 0.9 kV mm^{-1}

Shear stress, τ (Pa)	Recoverable strain, γ_{REC} (%)
0.35	~64
0.7	~46
1.4	~38
2.1	~0

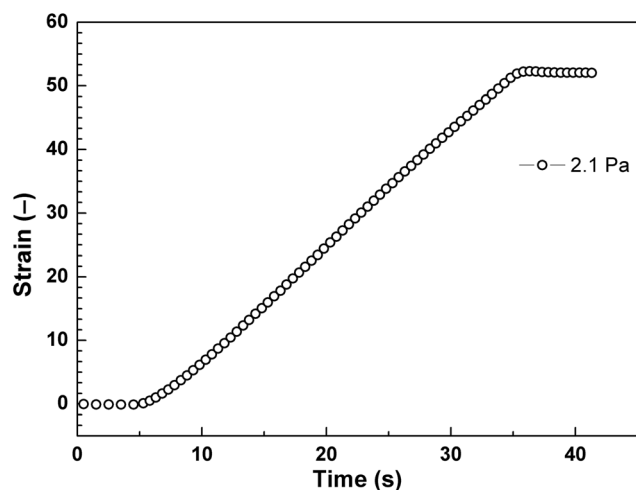


Fig. 9 Creep and recovery responses for the ER fluid (5 vol%) containing the iron(II) oxalate particles in the presence of the electric field strength of 0.9 kV mm^{-1} at the applied shear stress of 2.1 Pa.

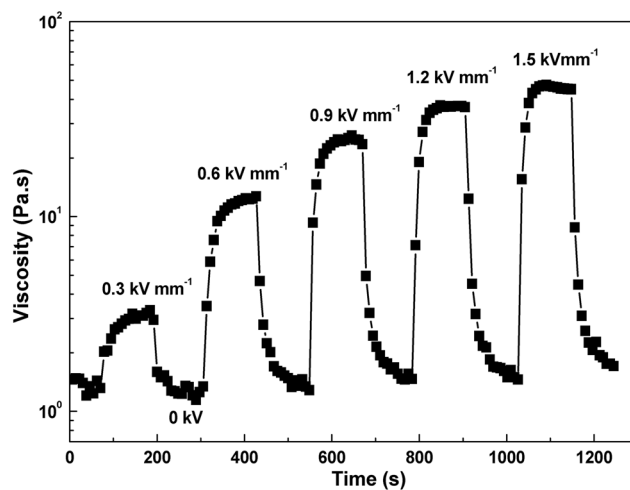


Fig. 10 Time dependence of the shear stress alternating between 0 and 1.5 kV mm^{-1} at the shear rate of 1 s^{-1} for 10 vol% ER fluid.

to its original value. Since the ER performance of ER fluids based on iron(II) oxalate particles is driven by conduction model and these particles have rod-like structure at the same time, their orientation along the electric field is a little bit slower when compared with globular nanoparticles ER systems.²⁸ However, when the rigid structure along the field direction is formed, this is stable within the wide shear rate region as illustrated in Fig. 4 and Table 1 just probably due to the conduction model and friction between oriented rod-like particles in the particulate chains.

The important issue of contemporary ER fluids is their sedimentation stability. The particles are generally dispersed in a liquid medium which possess lower density than the dispersed particles. This leads to undesirable effects and consequently possible problems in their potential applications. As can be seen in Fig. 11, the ER fluid based on 5 vol% of iron(II) oxalate particles keeps stable for a long time probably because of the particles' rod-like structure hindering their

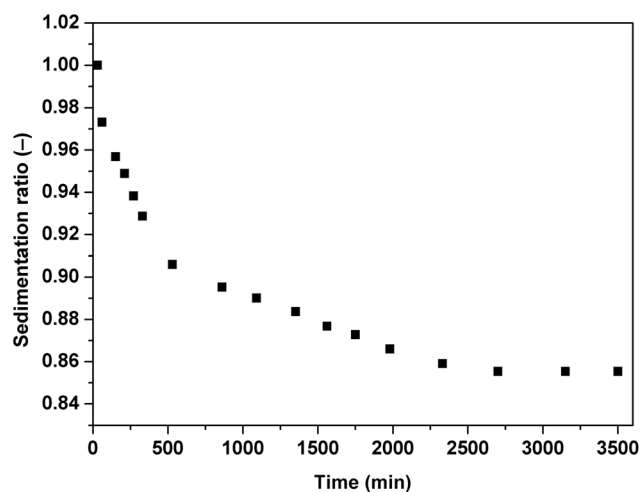


Fig. 11 The sedimentation stability of ER fluids containing 5 vol% of iron(II) oxalate particles.



sedimentation.²⁹ containing the iron(II) oxalate particles in the presence of the electric field strength of 0.9 kV mm^{-1} under the applied shear stresses of 0.35 Pa (black squares), 0.7 Pa (red triangles) and 1.4 Pa (blue diamonds).

Conclusions

Iron(II) oxalate particles were synthesized *via* co-precipitation method using iron(II) sulphate heptahydrate and oxalic acid dihydrate as precursors. These particles with rod-like morphology were subsequently mixed with silicone oil at concentrations of 1, 5, and 10 vol% in order to create novel ER fluids. Utilization of particles with rod-like morphology led to significant ER effects. Furthermore, the highest ER efficiency was found for the system containing 5 vol% of synthesized particles. The viscoelastic behavior investigation of the ER fluids was conducted through creep-recovery test. At levels of shear stress where electrostatic forces dominated over hydrodynamic ones the systems exhibited certain elastic portion of deformation. In the case of utilizing shear stress values of 0.35 Pa, 0.7 Pa, and 1.4 Pa the recoverable strain was 64%, 46%, and 38%, respectively. The creep-recovery test confirmed the transition of the ER fluids from a liquid state to viscoelastic solid.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors wish to thank the Internal Grant Agency of Tomas Bata University in Zlín (projects nos. IGA/CPS/2017/004 and IGA/CPS/2018/004) for the financial support. This work was supported by the Ministry of Education, Youth and Sports of the Czech Republic – Program NPU I (LO1504). This research was also carried out with support of the Operational Program Research and Development for Innovations co-funded by the European Regional Development Fund (ERDF) and national budget of the Czech Republic, within the framework of the Centre of Polymer Systems project (CZ.1.05/2.1.00/19.0409).

References

- 1 J. Stejskal, M. Mrlik, T. Plachy, M. Trchova, J. Kovarova and Y. Li, *React. Funct. Polym.*, 2017, **120**, 30–37.
- 2 M. Mrlik, M. Ilcikova, T. Plachy, R. Moucka, V. Pavlinek and J. Mosnacek, *J. Ind. Eng. Chem.*, 2018, **57**, 104–122.
- 3 Y. D. Liu and H. J. Choi, *Electrorheological fluids: smart soft matter and characteristics*, *Soft Matter*, 2012, **8**, 11961–11978.
- 4 A. Lengalova, V. Pavlinek, P. Saha, O. Quadrat, T. Kitano and J. Stejskal, *Eur. Polym. J.*, 2003, **39**, 641–645.
- 5 A. Lengalova, V. Pavlinek, P. Saha, O. Quadrat and J. Stejskal, *Colloids Surf., A*, 2003, **227**, 1–8.
- 6 M. Sedlacik, M. Mrlik, Z. Kozakova, V. Pavlinek and I. Kuritka, *Colloid Polym. Sci.*, 2013, **291**, 1105–1111.
- 7 J. B. Yin, X. X. Wang, R. T. Chang and X. P. Zhao, *Soft Matter*, 2012, **8**, 294–297.
- 8 M. Stenicka, V. Pavlinek, P. Saha, N. V. Blinova, J. Stejskal and O. Quadrat, *Colloid Polym. Sci.*, 2009, **287**, 403–412.
- 9 T. Plachy, M. Sedlacik, V. Pavlinek and J. Stejskal, *J. Mater. Chem. C*, 2015, **3**, 9973–9980.
- 10 L. C. Davis, *J. Appl. Phys.*, 1992, **72**, 1334–1340.
- 11 T. Plachy, M. Sedlacik, V. Pavlinek, J. Stejskal, M. P. Graca and L. C. Costa, *J. Intell. Mater. Syst. Struct.*, 2016, **27**, 880–886.
- 12 J. B. Yin, X. A. Xia, L. Q. Xiang and X. P. Zhao, *Smart Mater. Struct.*, 2011, **20**, 015002.
- 13 O. Erol, H. I. Unal and B. Sari, *Polym. Compos.*, 2010, **31**, 471–481.
- 14 K. Oz, M. Yavuz, H. Yilmaz, H. I. Unal and B. Sari, *J. Mater. Sci.*, 2008, **43**, 1451–1459.
- 15 B. Sari, N. Yavas, M. Makulogullari, O. Erol and H. I. Unal, *React. Funct. Polym.*, 2009, **69**, 808–815.
- 16 X. A. Xia, J. B. Yin, P. F. Qiang and X. P. Zhao, *Polymer*, 2011, **52**, 786–792.
- 17 J. B. Yin, X. P. Zhao, X. Xia, L. Q. Xiang and Y. P. Qiao, *Polymer*, 2008, **49**, 4413–4419.
- 18 D. Vincenzi, *J. Fluid Mech.*, 2013, **719**, 465–487.
- 19 Z. G. Jia, D. P. Ren, L. X. Xu and R. S. Zhu, *Mater. Sci. Semicond. Process.*, 2012, **15**, 270–276.
- 20 M. Hermanek, R. Zboril, M. Mashlan, L. Machala and O. Schneeweiss, *J. Mater. Chem.*, 2006, **16**, 1273–1280.
- 21 T. Plachy, M. Cvek, Z. Kozakova, M. Sedlacik and R. Moucka, *Smart Mater. Struct.*, 2017, **26**, 025026.
- 22 Z. Kozakova, I. Kuritka, P. Bazant, M. Pastorek and V. Babayan, *Mater. Lett.*, 2015, **138**, 116–119.
- 23 X. Q. Ji, W. L. Zhang, W. P. Jia, X. X. Wang, Y. Tian, L. Deng and J. Q. Liu, *J. Ind. Eng. Chem.*, 2017, **56**, 203–211.
- 24 C. S. Jun, S. H. Kwon, H. J. Choi and Y. Seo, *ACS Appl. Mater. Interfaces*, 2017, **9**, 44811–44819.
- 25 M. S. Cho, H. J. Choi and M. S. Jhon, *Polymer*, 2005, **46**, 11484–11488.
- 26 S. G. Kim, J. Y. Lim, J. H. Sung, H. J. Choi and Y. Seo, *Polymer*, 2007, **48**, 6622–6631.
- 27 K. Zhang, Y. D. Liu, M. S. Jhon and H. J. Choi, *J. Colloid Interface Sci.*, 2013, **409**, 259–263.
- 28 S. H. Piao, C. Y. Gao and H. J. Choi, *Polymer*, 2017, **127**, 174–181.
- 29 C. G. Niu, X. F. Dong, H. Zhao and M. Qi, *Smart Mater. Struct.*, 2014, **23**, 075018.

