

Article

New simple and robust method for determination of polarity of deep eutectic solvents (DESs) by means of contact angle measurement

Łukasz Cichocki ¹, Dorota Warmińska ², Justyna Łuczak^{1,5}, Andrzej Przyjazny³, and Grzegorz Boczkaj ^{4,5*}

¹ Gdańsk University of Technology, Faculty of Chemistry, Department of Process Engineering and Chemical Technology, 80 – 233 Gdańsk, G. Narutowicza St. 11/12, Poland;

² Gdańsk University of Technology, Faculty of Chemistry, Department of Physical Chemistry, 80 – 233 Gdańsk, G. Narutowicza St. 11/12, Poland

³ Kettering University, 1700 University Avenue, Flint, MI 48504, USA

⁴ Gdańsk University of Technology, Faculty of Civil and Environmental Engineering, Department of Sanitary Engineering, 80 – 233 Gdańsk, G. Narutowicza St. 11/12, Poland

⁵ Advanced Materials Center, Gdańsk University of Technology, 80 – 233 Gdańsk, G. Narutowicza St. 11/12, Poland

* Correspondence: grzegorz.boczkaj@pg.edu.pl; +48 697970303

Abstract: The paper presents a new procedure for assessing the polarity and hydrophobicity of deep eutectic solvents (DESs) based on the measurement of the DES contact angle on glass. DESs consisting of benzoic acid derivatives and quaternary ammonium chlorides – tetrabutylammonium chloride (TBAC) and benzylidimethylhexadecylammonium chloride (16-BAC) in selected mole ratios were chosen for the study. To investigate the DESs polarity, an optical goniometer and an Er(30) solvatochromic scale based on the Reichardt's dye were used. The research demonstrated the high accuracy and precision of the developed procedure. The simplicity of the examination and the availability of basic equipment allow the implementation of the developed procedure in routine investigations of DESs.

Keywords: DESs, polarity, hydrophobicity, contact angle, solvatochromism, materials characterization

1. Introduction

Increasingly more research on DESs is carried out throughout the world and their properties are being improved for effective application in science and industry [1-5]. The first step to discover the nature of DESs is to study their physicochemical properties [6-7]. Some physicochemical properties can be easily measured. However, there are also those physicochemical properties that are difficult to measure and, sometimes, almost impossible to determine for DESs. Such properties include polarity and hydrophobicity of DESs. For example, the study of the octanol-water partition coefficient [8] requires contact of DESs with water, which, as a strong solvating agent, displaces DES intramolecular hydrogen bonds with bonds between DES substrates and water molecules [9]. The aqueous degradation of DESs to the starting materials means that it is not DESs that are actually investigated but an aqueous mixture or solution of the reactants forming DESs. Reversed-phase high-performance liquid chromatography can also be used for polarity measurement, but it also requires converting DESs into solution in a given solvent [10]. Although solvents other than water can be used but also in this case there is a high interference of the solvent in the DESs structure. Even nonpolar solvents can negatively affect the accuracy of the measurement due to the fact that practically the properties of two-component DESs in a solution can be treated as three-component DESs in terms of the physicochemical properties investigated. Consequently, in this study, a

method of determining DESs polarity and hydrophobicity was sought without the need to prepare solutions, *i.e.* without interfering with the internal structure of DESs. The structure of DESs, as a mixture of at least two components having hydrogen bond donors (HBD) and hydrogen bond acceptors (HBA) is determined by the hydrogen bonds formed between HBD and HBA. The system of hydrogen bonds in the DES molecule is sensitive to changes related to the presence of an additional component, such as water or another solvent. One of the methods of determination of DESs hydrophobicity without the need to prepare their solutions is the contact angle test which was one of the aspects of this work. The contact angle is measured for pure DESs by the direct method of depositing a drop of DES on the reference surface. The method does not require the use of any solvents, only an optical goniometer, and a reference surface. The method of measurement of contact angle is simple and provides direct results in a short time. In the present study, for comparative purposes, the polarity measurement was also carried out using the second solvent-free method based on solvatochromism based on the Reichardt's dye [11]. The solvatochromic responses of UV-vis absorption probes have been so far applied by Florindo et al. [12] for DESs based on cholinium chloride, DL-menthol and TBAC and by Pandey et al. [13] for DESs containing ChCl and glycerol, urea, malonic acid, and ethylene glycol, respectively. Pandey et al. identified that the high polarity of the studied DESs was significantly influenced by HBD nature. Among the above four combinations, ChCl:Gly exhibited the highest $E_T(30)$ value. This observation was consistent with the results obtained by Florindo et al. who found that the polarity of DESs changes in the order: choline chloride:malonic acid > choline chloride:glycolic acid > choline chloride: levulinic acid. Solvatochromic indices to examine the polarities of DESs were used also by Abbott et al. who determined the polarities of choline chloride-glycerol DESs of different molar ratios, revealing a linear polarity increase with increasing ChCl concentration. In summary, the higher the number of carboxyls, hydroxyls, or carbonyls groups is in HBD, the higher polarity of the DESs composed of such HBD. This relationship results from the structure of the carboxyl group, which as a combination of the $-C=O$ (carbonyl group) and the $-OH$ (hydroxyl group) is more polar than sole hydroxyl group or the carbonyl group.

2. Results and discussion

The measurement of polarity of DESs using the contact angle technique is an alternative to tests requiring the preparation of DES solutions in solvents. The high viscosity and ease of crystallization of DESs make working with pure DESs difficult and may require optimization of test conditions in order to obtain correct results. Polarity studies with the use of solvents eliminate the problem of crystallization and high viscosity of DESs, but they completely disrupt the internal structure of DESs, which significantly changes physicochemical properties of DESs. The contact angle measurement carried out in an optical goniometer allows the determination of contact angles of pure DESs. Measurement of the contact angle while meeting the requirements of the procedure provides the contact angles of DESs relative to the tested surface - glass. The contact angle, therefore, indirectly defines the affinity of DES molecules to the surface of the reference material. Two reference substances with extreme properties were also used in the study – water as a representative of a highly hydrophilic liquid and rapeseed oil as a representative of a highly hydrophobic liquid. The contact angle test itself requires obtaining a DES drop of the smallest possible diameter. A drop diameter as small as possible is very important in preventing the DES drop from spreading out. The spreading out of DESs drops and other factors leading to the asymmetry of the drops distort the result of the test. The contact angle should be identical, measured along the entire circumference of the drop's contact with the surface. In the case of a goniometric measurement, the drop is projected onto a plane, and therefore only two contact angles on the opposite sides of the drop can be measured. The final results of the contact angle determination were analyzed in terms of the similarity of the behavior of a given DES on a given material to water and rapeseed

oil as reference substances. The results of contact angle measurements are summarized in Table 1.

Table 1. Contact angle (CA) values of DESs on glass.

Sample	HBA	HBD	Mole ratio	CA glass
1		rapeseed oil		39.5
2	TBAC	salicylic acid	3:2	46.9
3	TBAC	3,5-dinitrobenzoic acid	1:1	59.6
4	TBAC	3,5-dinitrosalicylic acid	1:1	50.6
5	TBAC	4-chlorosalicylic acid	1:1	41.8
6	TBAC	acetylsalicylic acid	3:2	45.8
7	TBAC	4-tert-butylbenzoic acid	1:1	46.8
8	16-BAC	salicylic acid	3:2	40.6
9	16-BAC	acetylsalicylic acid	3:2	40.2
10		water		74.5

DESs for which the contact angle values on glass were closest to water were classified as DESs with predominantly hydrophilic properties (DES 3) while DESs for which the contact angle values on glass were close to oil on glass were classified as hydrophobic (DESs 5, 8 and 9). DESs with contact angles intermediate between water and oil were classified as intermediate between hydrophilic and hydrophobic (DESs 2, 4, 6, and 7). The results are presented in Figure 1.

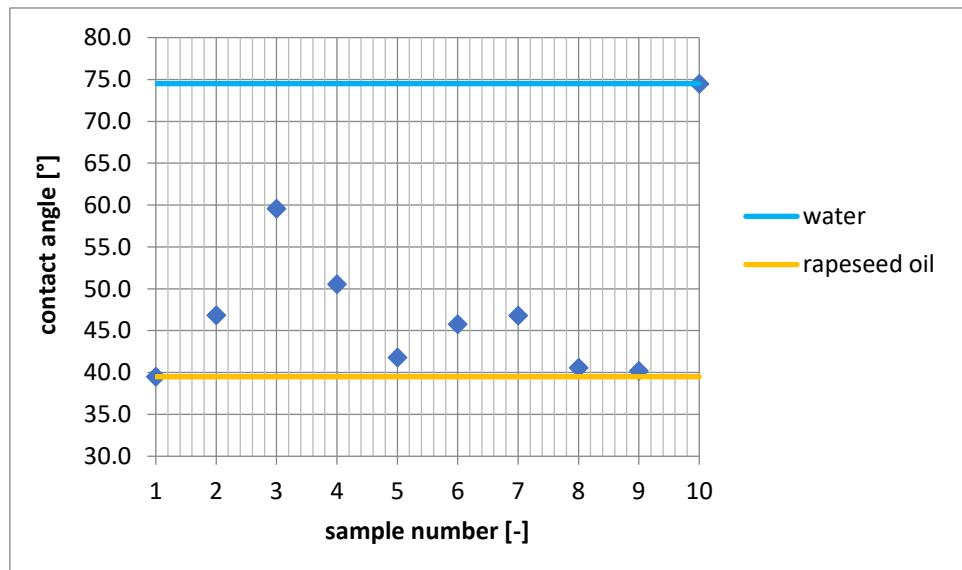


Figure 1. Contact angle: 1 rapeseed oil, 2-9 DESs, 10 water - on glass.

Inspection of the above graph reveals that DESs have different contact angle values depending on the molecular structure, which is reflected in the polarity and hydrophobicity of DESs. DES no. 3 consisting of TBAC and 3,5-dinitrobenzoic acid was found to be the most hydrophilic. TBAC as a quaternary ammonium salt has four lone pairs of electrons on the chloride anion, which is a potential HBA, and the 3,5-dinitrobenzoic acid bonded to TBAC has two nitro groups that strongly deactivate the benzene ring, increasing the effect of the HBA – HBD interaction. The study also revealed the two most hydrophobic DESs – nos. 8 and 9. The highly hydrophobic nature of these DESs is mainly due to the n-hexadecyl groups from 16-BAC, showing much stronger hydrophobicity than the n-butyl groups of TBAC.

For comparative purposes, the polarity measurement was carried out using the second solvent-free method based on solvatochromism, i.e. changes in absorption of electromagnetic radiation by dyes depending on the solvent in which are dissolved. One of the largest solvatochromic effects (highest shift of absorption bands) is observed for Reichardt's dye. Solvatochromism is another method of measurement of DES polarity without the need to prepare DES solutions, thus without affecting the internal structure of DESs. The structure of DESs and the DES-dye interactions affect the structural changes and charge distribution in the Reichardt's dye molecule. As a result, the dye has different absorption of radiation at a particular wavelength depending on the environment in which it is present. Such a relationship is caused by shifts in charge transfer (CT) bands in the dye. The shifts of the CT bands result from the effect of DESs on the position of conjugated double bonds in the dye molecule. The excited state of the dye is stabilized by polar solvents through the phenomenon of solvation of the dye molecule by solvent molecules, hence the predominance of this form of betaine in the polar environment (Figure 2).

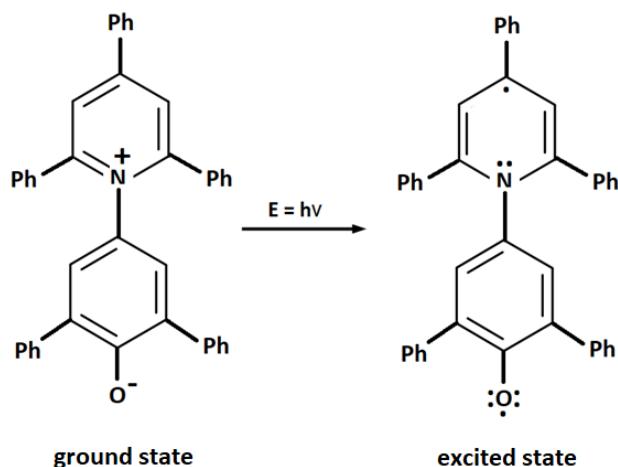


Figure 2. Reichardt's betaine (own drawing based on [14]).

The wavelengths corresponding to the maximum absorbance of the Reichardt's dye in DESs relative to the reference substances – water and rapeseed oil reveal the shift in the absorption band of the dye. The comparison of the shifts in absorption maxima allows to determine whether a given DES affects the dye in the same way as water (polar substance) or rapeseed oil (nonpolar substance) – Figure 3.

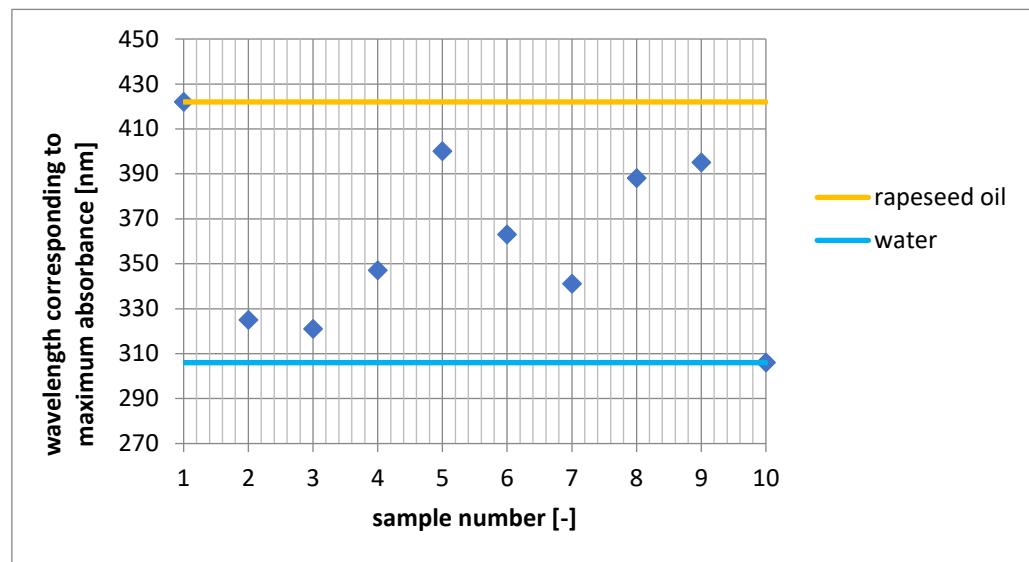


Figure 3. Wavelength corresponding to maximum absorbance for Reichardt's dye: 1 - water, 2-9 - DESs, 10 – rapeseed oil.

Measurement of the CT band shifts in the Reichardt's dye became the basis for the creation of a polarity scale called $E_T(30)$. $E_T(30)$ values are based on measurements of the wavelength corresponding to the highest absorbance (equation 1). In addition to $E_T(30)$, the so-called normalized value $E_T^N(30)$ has been introduced, in which the least polar substance was tetramethylsilane (TMS): $E_T^N(30) = 0$ and the most polar substance was water: $E_T^N(30) = 1$. Hence, the normalized scale ranges from 0.000 for TMS to 1.000 for water. In this work, for the calculation of the normalized polarity scale, rapeseed oil was used as the least polar substance: $E_T^N(30) = 0$ and water was used as the most polar substance: $E_T^N(30) = 1$.

$$E_T(30)[\text{kcal/mol}] = hcN_A \nu_{max}(\text{cm}^{-1}) = \frac{28591}{\lambda_{max}[\text{nm}]} \quad (1)$$

$$E_T^N(30) = \frac{E_T(30) - E_T(\text{rapeseed oil})}{E_T(\text{H}_2\text{O}) - E_T(\text{rapeseed oil})} \quad (2)$$

On the basis of the wavelength corresponding to maximum absorbance and equations (1) and (2) [8], the normalized values of the $E_T^N(30)$ polarity scale were calculated (Table 2).

Table 2. Calculated normalized values for $E_T^N(30)$ polarity scale.

Sample no. *	λ_{\max} [nm]	$E_T(30)$ [kcal/mol]	$E_T^N(30)$ [-]
1 (water)	306	93.4	1.00
2	325	88.0	0.79
3	321	89.1	0.83
4	347	82.4	0.57
5	400	71.5	0.15
6	363	78.8	0.43
7	341	83.8	0.63
8	388	73.7	0.23
9	395	72.4	0.18
10 (rapeseed oil)	422	67.8	0.00

*Sample no. 1 – water, 2-9 DESs and 10 – rapeseed oil.

Based on Table 2, it is possible to assess the similarity of a given DES to water, which demonstrates the hydrophilic nature of DES, or to rapeseed oil, which reveals the hydrophobic nature of DES. Values of $E_T^N(30)$ ranging from 0 to 0.25 were obtained for DESs number 5, 8 and 9, which proves the hydrophobic nature of these DESs. Intermediate values of the polarity coefficient, ranging from 0.25 to 0.8, were obtained for DESs no. 2, 4, 6 and 7 while the highest similarity to water and the lowest to rapeseed oil was obtained for DES no. 3: $E_T^N(30)$ was in the 0.8 to 1 range.

Table 3. Classification of DESs samples in terms of polarity.

scale	polarity/hydrophilicity		
	high	intermediate	low
	DESs sample no.		
goniometric	3	2, 4, 6, 7	5, 8, 9
solvatochromic	3	2, 4, 6, 7	5, 8, 9

Inspection of Table 3 reveals that both methods of DESs classification in terms of polarity are compatible. The two test methods (goniometric and solvatochromic) are independent and based on different DESs parameters, but ultimately they indicate the same property of DESs - polarity. In the case of goniometric measurement of contact angle, the results obtained were based on the interactions between DESs and the reference surface. The contact angle indicates the affinity of DESs for a particular structural group constituting the reference material (glass – oxygen groups of silicon oxide). For the solvatochromic measurement, shifts in charge transfer (CT) bands in the Reichardt's dye due to the presence of DES are used. Both parameters indirectly indicate DESs polarity and related to its hydrophilicity, but are based on completely different phenomena. Obtaining similar results using such different methods demonstrates the correctness of the determination. It is also apparent that both methods of polarity assessment are universal and particularly useful for testing complex (multicomponent) mixtures, in which the prepa-

ration of aqueous solutions is either impossible due to the low solubility of DESs or due to hydrolysis of DESs substrates in aqueous solutions.

The methods of determination of polarity described in this paper, especially the contact angle technique, can be widely used in the study of very complex mixtures. The goniometric method also has a great advantage over the solvatochromic procedure due to the simplicity of the test, the possibility of investigation of opaque samples as well as the low cost of the test and its short duration. In many cases, the measurement can be simplified by taking a picture with a camera and image processing with an appropriate graphics software. The only limitations of the goniometric method are substances whose very high viscosity and density prevent the free formation and detachment of drops from the goniometer needle.

3. Materials and methods

3.1. Materials

The following reagents were used for the synthesis of DESs: as HBA: TBAC and 16-BAC with purity over 97% - (Sigma Aldrich) and as HBD acids: salicylic, 3,5-dinitrobenzoic, 3,5-dinitrosalicylic, 4-chlorosalicylic, acetylsalicylic, 4-tert-butylbenzoic, 4-methylbenzoic and 5-sulfosalicylic with purity over 98% - (Sigma Aldrich). Glass was used as the reference material for contact angle measurement.

3.2. Methods

3.2.1. Synthesis of DESs

The synthesis of DESs was carried out in previously selected mole ratios, depending upon DES: HBA:HBD 1:1 and 3:2. In order to prepare DES, proper amounts of the reactants were weighed on a model AS.310.R2 analytical balance (RADWAG) with an accuracy of 0.1 mg, so that the total mass of HBA and HBD ranged from 2 to 3 g. Next, the weighed amounts of the reactants were placed in a 5-mL screw cap vial along with a magnetic stir bar. The vials were placed in a heating mantle using water as a heating medium and equipped with a magnetic stirrer (set model: 06 MSH PRO T, CHEMLAND). Reactant mixtures were left for 120 minutes at 340-350 K with stirring at 1400-1500 rpm. After 120 min all of the synthesized DESs were clear liquids. The molar compositions of the examined DESs are listed in Table 4.

Table 4. Mole ratios of investigated DESs.

HBA	HBD	Molar ratio
TBAC	salicylic acid	3:2
TBAC	3,5-dinitrobenzoic acid	1:1
TBAC	3,5-dinitrosalicylic acid	1:1
TBAC	4-chlorosalicylic acid	1:1
TBAC	acetylsalicylic acid	3:2
TBAC	4-tert-butylbenzoic acid	1:1
16-BAC	salicylic acid	3:2
16-BAC	acetylsalicylic acid	3:2

3.2.2. Polarity measurement

Contact angle

The contact angle was measured with a model OCA 25 optical goniometer (DataPhysics) using the sessile drop technique (Figure 4). Silicate glass, which is considered a material with a predominance of hydrophilic properties (oxygen groups of silicon(IV) oxide in the glass structure), was used as the reference surface. The glass surface was prepared by thoroughly cleaning it with methanol and leveling. Prior to sampling, DESs were heated and then brought to room temperature in order to ensure the representativeness of the sample. The samples were collected using a 1-mL plastic syringe. Based on the publication [15], a needle diameter of 0.55 mm was selected as the optimum to generate DES drops on the tested surface. The same needle diameter was used for all samples to facilitate the comparison of the obtained results. DESs drops were produced by the syringe needle placed perpendicularly to the tested surface. The tip of the needle was 30 mm above the test surface. The movement of the syringe plunger was controlled by the stepper motor, which ensured a slow increase of the droplet at the tip of the needle until the drop was detached by gravity.

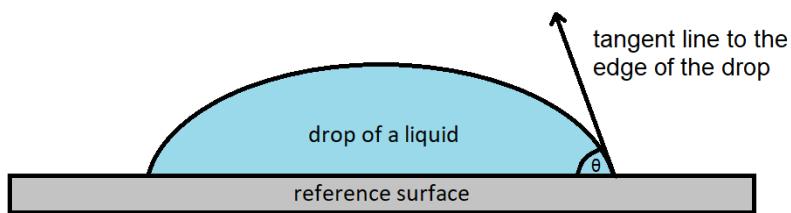


Figure 4. Graphical representation of contact angle determination using the sessile drop technique (θ – the contact angle). Own drawing based on [16].

The results from the goniometer software were recorded after obtaining identical values of both contact angles on both sides of the drop. Each measurement was carried out on glass at room temperature. Contact angle values were computed by goniometer software using the Young-Laplace fit (equation 1). This model uses the DES drop shape through the radius of the drop projection onto the plane. The knowledge of the Laplace pressure (ΔP) by the software allows the calculation of the contact angle on surface.

$$\Delta P = \gamma \left(\frac{1}{R_1} - \frac{1}{R_2} \right) \quad (3)$$

γ - surface tension [N/m], ΔP – Laplace pressure [Pa], R - radius [m] [16]

Solvatochromism

This method requires a solution of Reichardt's dye in DES at a concentration of approximately 0.3 mg/mL of sample. The dye concentration in DESs was optimized so that the absorbance as a function of dye concentration ranged from 0 to 1 (linear range of the Lambert-Beer law). A model DR 5000 spectrophotometer (Hach Lange) was used in the experiments. Pure DESs were used as a reference in spectrophotometric analysis in order to eliminate the effect of background on UV-VIS absorbance measurements. Next, the absorption spectra of the dye solutions in DESs were taken and the wavelength of maximum absorbance determined.

4. Conclusions

The results obtained in this study revealed that the goniometric (contact angle measurement) method of testing the polarity of DESs on glass can be used to determine the hydrophilicity/hydrophobicity of DESs with respect to each other or with respect to reference substances. This method has a number of advantages over the methods using solvents. Firstly, it does not require the use of solvents, most of which are toxic and require proper disposal after use. Secondly, the goniometric method is consistent with the principles of green chemistry and it is completely emission-free. The study also demonstrated the compatibility of the goniometric method with the solvatochromic method, which requires expensive and difficult to synthesize dyes. A very small amount of sample is needed for a goniometric test and the entire sample can be recovered from the reference surface. In the case of other polarity testing methods, for example the solvatochromic method, the preparation of dye solutions in DESs is required, which contaminates the DES samples and precludes their subsequent use. Thus, the goniometric method is currently the most ecological and least expensive methods of polarity testing, and there are also opportunities for its further development and optimization.

References

1. Cichowska-Kopczyńska, I., Warmińska, D., Nowosielski, B. Solubility of carbon dioxide in deep eutectic solvents based on 3-amino-1-propanol and tetraalkylammonium salts at low pressure. *Materials* **2021**, *14*(3), 1-14.
doi.org/10.3390/ma14030594
2. Momotko, M., Łuczak, J., Przyjazny, A., Boczkaj, G. First deep eutectic solvent-based (DES) stationary phase for gas chromatography and future perspectives for DES application in separation techniques. *J. Chromatogr. A* **2021**, *1635*.
doi.org/10.1016/j.chroma.2020.461701
3. Tang, B., Zhang, H., Row, K. H. Application of deep eutectic solvents in the extraction and separation of target compounds from various samples. *J. Sep. Sci.* **2015**, *38*(6), 1053-1064.
doi.org/10.1002/jssc.201401347
4. Marcus Y. *Deep Eutectic Solvents: Synthesis, Properties, and Applications*; Springer International Publishing, 2019
5. Hansen, B. B., Spittle, S., Chen, B., Poe, D., Zhang, Y., et al., Deep Eutectic Solvents: A Review of Fundamentals and Applications. *Chem. Rev.* **2020**.
doi:10.1021/acs.chemrev.0c00385
6. Zhu, J., Yu, K., Zhu, Y., Zhu, R., Ye, F., Song, N., Xu, Y. Physicochemical properties of deep eutectic solvents formed by choline chloride and phenolic compounds at $T = (293.15 \text{ to } 333.15) \text{ K}$: The influence of electronic effect of substitution group. *J. Mol. Liq.* **2017**, *232*, 182-187.
doi.org/10.1016/j.molliq.2017.02.071
7. Jafari, K., Fatemi, M. H., & Estellé, P., A short overview of the thermophysical properties and current use as base fluid for heat transfer nanofluids. *J. Mol. Liq.* **2020**, *114752*.
doi:10.1016/j.molliq.2020.114752
8. Farias, F. O., Passos, H., Lima, Á. S., Mafra, M. R., Coutinho, J. A. P. Is It Possible to Create Ternary-like Aqueous Biphasic Systems with Deep Eutectic Solvents? *ACS Sustain. Chem. Eng.* **2017**, *5*(10), 9402-9411.
doi.org/10.1021/acssuschemeng.7b02514
9. Ma, C., Laaksonen, A., Liu, C., Lu, X., Ji, X. The peculiar effect of water on ionic liquids and deep eutectic solvents. *Chem. Soc. Rev.* **2018**, *47*(23), 8685-8720.
doi.org/10.1039/c8cs00325d
10. Du, C. M., Valko, K., Bevan, C., Reynolds, D., & Abraham, M. H. Rapid method for estimating octanol-water partition coefficient (LOG POCT) from isocratic rp-hplc and a hydrogen bond acidity term (A). *J. Liq. Chromatogr. Relat.* **2001**, *24*(5), 635-649.
doi:10.1081/jlc-100103400
11. Abbott, A. P., Harris, R. C., Ryder, K. S., D'Agostino, C., Gladden, L. F., Mantle, M. D. Glycerol eutectics as sustainable solvent systems. *Green Chem.*, **2011**, *13*(1), 82-90.
doi: 10.1039/c0gc00395f
12. Florindo, A. J. S. McIntosh, T. Welton, L. C. et al., A closer look into deep eutectic solvents: Exploring intermolecular interactions using solvatochromic probes, *Phys. Chem.*, **2017**, *20*(1), 206-213
doi: 10.1039/c7cp06471c.
13. Pandey, A., Pandey, S. Solvatochromic probe behavior within choline chloride-based deep eutectic solvents: Effect of temperature and water. *J. Phys. Chem. B*, **2014**, *118*(50), 14652-14661,
doi: 10.1021/jp510420h.
14. Reichardt, C. Solvatochromic Dyes as Solvent Polarity Indicators. *Chem. Rev.* **1994**, *94*(8), 2319-2358.
doi:10.1021/cr00032a005
15. Vuckovac, M., Latikka, M., Liu, K., Huhtamäki, T., Ras, R. H. A. Uncertainties in contact angle goniometry. *Soft Matter* **2019**, *15*(35), 7089-7096.
doi.org/10.1039/c9sm01221d
16. Song, B., Ju, J., Springer, J. Determination of Interfacial Tension from the Profile of a Pendant Drop Using Computer-Aided Image Processing 2. Experimental. *J. Colloid Interface Sci.* **1996**, *184*(1), 77-91.