Overview of the three multicriteria approaches applied to a global assessment of analytical methods

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ABSTRACT

Critical and global evaluation of analytical methods should be one of the primary goals in analytical chemistry. A holistic approach, however, requires a look at the varied features: commonly discussed validation criteria, often underrated practical and economic aspects, and typically overlooked compliance with the principles of green analytical chemistry. Carrying out such an assessment in a critical and transparent way is extremely difficult without special tools. The purpose of this work is to discuss and compare the three different approaches that seem to be potential candidates: multi-criteria decision analysis methods (MCDA), HEXAGON, and RGB model. The basic principles of each methodology, individual possibilities offered, and the results of the assessment of selected model methods will be presented. Ultimately, the potential compatibility of assessing the same group of methods using different tools will be examined. This contribution can help to select optimal tool and conduct more thorough and insightful assessments.

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1. Introduction

Critical assessment of the potential of new analytical methods is crucial in analytical chemistry regardless of the specificity of the method. Currently, commonly formalized validation criteria are used for this purpose, including parameters such as precision, accuracy (trueness), sensitivity, recovery, etc., and the assessment of the potential of a method consists in separate consideration and comparison of individual parameters with commonly accepted standards, as well as between different alternative methods. This approach, despite many advantages, has the disadvantage that it is difficult to express the analytical potential of a method using one unified measure (performance indicator), which would cover all validation criteria and allow easy overall assessment.

An important trend observed in recent years is paying more attention to the assessment of the so-called “greenness” of an analytical method, i.e. its friendliness in terms of safety for user health and the environment. This entails the constant development of new algorithms for assessing the greenness [1,2], including qualitative and quantitative approaches, e.g. Eco-Scale developed by the Group of Professor Jacek Namieśnik [3]. However, a holistic and comprehensive assessment of methods in terms of both analytical efficiency (validation criteria) and greenness is impossible using these metrics. Finally, the global assessment of a method also requires a look at its other features, often underestimated or considered in a highly simplified and only intuitive sense. However, they are often as important in everyday life as analytical performance expressed in validation parameters. These features relate to the “productivity” of the method and its effectiveness, understood in purely practical/economic terms. Due to the above, conducting a comprehensive assessment of an analytical method covering all of the mentioned attributes is extremely difficult without special tools dedicated for this purpose.

Currently, there are three proven tools that can be used in the overall assessment of analytical methods taking into account the
various attributes mentioned above. These are: (i) multi-criteria decision analysis (MCDA) algorithms that result in an indication of the best procedure to solve a given specific analytical problem [4–7]; (ii) an algorithm that is the extension of the aforementioned Eco-scale to other features of a method, named HEXAGON due to its pictorial form [8]; and (iii) an RGB model [9], named so because of the analogy to the model commonly used in representation and coding of colors in electronic devices, enabling the expression of a method’s potential by means of its resultant color, depending on the participation of the three primary attributes (Red, Green, Blue). These algorithms are, as one can assume, still largely unknown to analytical chemists community. Furthermore, their comparison with each other has never been presented before. The purpose of this work is to present for the first time the principles of operation and possibilities offered by each of the three mentioned approaches, as well as their mutual comparison. It is intended to facilitate a more thorough and comprehensive evaluation of analytical procedures, both being in use and newly developed, as well as to simplify the selection of an optimal tool for a given reader.

2. Model methods

Six model methods of quantitative analysis of the popular colorant added to food products and beverage, Sunset Yellow FCF (E110, SY) [10], were selected for presenting the operation and comparing the considered algorithms. These methods include high performance liquid chromatography with diode array detection (HPLC-DAD) [11], high performance liquid chromatography with tandem mass spectrometry detection (HPLC-MS/MS) [11], capillary electrophoresis with UV detection (CE) [12], polyclonal antibody-based indirect enzyme-linked immunosorbent assay (ELISA) [13], photoacoustic spectroscopy (PAS) [14], and first derivative spectrophotometry (FDS) [14]. All methods were dedicated to the same analyte – SY, although some of them enabled multi-component analysis of other popular food dyes. The studies were performed on various food samples, including beverages and solid products, hence the sample preparation procedure depended on the type of sample material. In addition, CE and ELISA methods also involved the preparation of specific reagents needed for proper analysis. In the case of CE, they were diamino moiety-functionalized silica nanoparticles (NPs) [12], used in the extraction process and as an additive to the separation buffer, while in the case of ELISA they were synthetic polyclonal antibodies [13]. The antibody preparation procedure was multi-stage and involved: chemical modification of SY to obtain immunogen, immunization of rabbits, and production and acquisition of antibodies from animals. More details on the individual methods are available in the relevant references [11–14].

3. MCDA (TOPSIS) algorithm

3.1. Working principle

MCDA is a group of tools that are used for scoring and ranking of alternatives according to the given assessment criteria. MCDA techniques are widely used in solving of various analytical problems [15]. The most widely applied ones are Analytical Hierarchical Process (AHP) [16], Elimination and Choice Expressing Reality (ELECTRE) [17], Preference Ranking Organization Method for Enrichment Evaluations (PROMETHEE) [18], Technique for Order Preference by Similarity to Ideal Solution (TOPSIS) [19], Multi Attribute Utility Theory (MAUT) [20], Multi Attribute Value Theory (MAVT) and Simple Additive Weighting [21,22]. They were described in detail in several books [23–25]. The mathematical algorithm in each of the techniques is different but the general principles can be summarized by few simple steps:

1. Definition of the goal of analysis. The goal of the analysis is usually finding the best solution to the given problem. It might be finding the optimal process, material, situation, location or state. The goal might also be ranking of all or part of available solutions. In the context of this study the aim of the analysis is finding the best analytical procedure and ranking of all available ones.

2. Definition of alternatives. The alternatives are the possible ways to achieve the main goal of the analysis. They may be possible processes, materials situations, locations or states that fulfill the requirements of the main analysis goal. Alternatives must be fully characterized by the criteria, with no gaps in the dataset. Here, the alternatives are analytical procedures that are applied for the given analytical task.

3. Definition of criteria. Criteria are the characteristic features that describe the set of alternatives. They have to be relevant to the main goal of analysis, have to be measurable in reliable way and comprehensively. As an input to MCDA analysis, they have to be in form of numerical values, or they need to be transformable into numerical values. One of the main advantages and the reasons to apply MCDA is the possibility to deal with criteria that are contradictory to each other. The criteria applied in this study are metrological, economic and greenness criteria.

4. Application of weights. The assessment criteria should be relevant to the goal of analysis, but their relevance can be different.
To differentiate between the criteria importance weights are usually assigned, or not in case of equal importance.

5. Running the algorithm. One of algorithms is applied to rank the alternatives according to the criteria with appropriate weights. Because of its simplicity, TOPSIS algorithm is applied in this study and is presented below:

a) The first step is construction of normalised decision matrix

\[ r_{ij} = \frac{x_{ij}}{\sqrt{\sum x_{ij}^2}}, \quad i = 1, 2, \ldots, m \]  \( \text{and} \ j = 1, 2, \ldots, n \)  \[ (1) \]

where \( x_{ij} \) and \( r_{ij} \) are original and normalised scores in decision matrix.

b) The second step is construction of the weighted normalised decision matrix, it is done according to relative importance (reflected by the weights, subjectively assigned by decision maker) that is set in point 4 of MCDA analysis.

\[ v_{ij} = r_{ij} \times w_j, \quad i = 1, 2, \ldots, m \] \( \text{and} \ j = 1, 2, \ldots, n \) \[ (2) \]

where \( w_j \) is the weight of the criterion \( j \) and \( \sum_{j=1}^{n} w_j = 1 \)

c) The next step is determination of both positive ideal \( (A^+) \) and negative ideal \( (A^-) \) solutions

\[ A^+ = \{ (\max v_{ij} | j \in C_{b}), (\min v_{ij} | j \in C_{c}) \} = \{ v^*_i | j = 1, 2, \ldots, m \} \] \[ (3) \]

\[ A^- = \{ (\min v_{ij} | j \in C_{b}), (\max v_{ij} | j \in C_{c}) \} = \{ v^*_i | j = 1, 2, \ldots, m \} \] \[ (4) \]

d) Then calculation of the separation measures for each alternative is performed

\[ S_j^+ = \sqrt{\sum_{j=1}^{m} (v_{ij} - v^*_i)^2}, \quad j = 1, 2, \ldots, m \] \[ (5) \]

\[ S_j^- = \sqrt{\sum_{j=1}^{m} (v_{ij} - v^*_i)^2}, \quad j = 1, 2, \ldots, m \] \[ (6) \]

e) Calculation of the relative closeness to the ideal solution is done

\[ C_j^+ = \frac{S_j^-}{S_j^+ + S_j^-}, \quad i = 1, 2, \ldots, m \] \( \text{and} \ 0 < C_j^+ < 1 \) \[ (7) \]

f) In the last step the scenarios are ranked according to similarity to ideal solution – from closest to furthest. The full ranking is created. For each alternative the value of similarity to ideal solution is calculated that is crucial in interpreting the final ranking as the differences between two consecutive alternatives may be varied.

6. Final decision making. The last step is the interpretation of the result, either the most appropriate alternative or investigation of created ranking. In this study, it is important to find the optimal alternative and rank the remaining ones.

Some MCDA applications include the opinions of multiple experts integration \[ [26], \] and often sensitivity analysis is performed to understand the impact of changes to criteria weights and explore the robustness of the indicated preferred solution \[ [27]. \]
DAD with the value of similarity to ideal solution equal to 0.34. It is characterized by relatively poor analytical performance, high consumption of organic solvents and the time of analysis is rather long. However, its performance is good in terms of other analytes that can be determined, and the amount of sample needed for analysis.

Thus, TOPSIS is a very simple and effective tool as it works on raw data, therefore no transformations and no reference thresholds are required to carry our assessment. By the selection of criteria and the application of adequate weights it can be applied as a fit-for-purpose tool. The ranking is easy to be interpreted, however, it does not carry any information about weak or strong points of the assessed procedures, thus, a thorough analysis of a method’s characteristics is difficult.

4. HEXAGON algorithm

4.1. Working principle

The hexagon quantitative tool comprises the rating of five variables of a method through the assignment of penalty points (PPs). The variables are divided into five groups: analytical features or figures of merit, associated chemical and health risks, environmental friendliness, sustainability and economic cost [8]. Specifically, the figures of merit include the analytical performance of the method under evaluation and they are organized into different blocks as follows: figures of merit 1 (FM-1) involve the sample treatment, characteristics of the method and calibration procedure while figures of merit 2 (FM-2) account for the quality control and accuracy. Chemical toxicity, hazard and safety considerations are evaluated by the globally harmonized system (SGA) [29,30]. The residues derived from the analytical method and the possibility to recycle them are taken into account to assess the sustainability offered by the analytical procedure. Additionally, the environmental impact is quantified by the carbon footprint metrics [31], which considers the energy consumption of the equipment employed and the time to perform the analysis. The related annual cost of the analytical determination is estimated according to the cost of the equipment needed in addition to its electricity consumption cost, the cost of the reagents and materials used, and the salary assigned to skilled personnel. Carbon footprint and annual cost are quantified in absolute terms whereas penalty points are ascribed to the other variables. Finally, the sum of the PPs and estimated carbon footprint and cost values are ranked in an overall quantification for each variable using a 0–4 scale and organized in a hexagon as resulting pictogram [8]. The higher the score (that is, getting closer to 4), the following statements are accomplished: the worst the adaptation of the figures of merit for providing a reliable analytical result, the worst the contribution to health and safety, the worst the environmental impact, sustainability and cost-benefit relation. At the final stage, the arithmetic mean of the 0–4 score ($S_{ar}$) is calculated for ranking the analytical procedures and eventually compare the evaluation results when applying the other proposed algorithms in the present article. The scale is related with excellent, good, suitable, weak and fail performance of the tested analytical method for the scores: 0, 1, 2, 3 and 4, respectively. The hexagon algorithm has been applied to a wide variety of methods that employ different analytical techniques. Among them, atomic absorption spectroscopy (AAS), inductively coupled plasma mass spectrometry (ICP-MS) and inductively coupled plasma optical emission spectrometry (ICP-OES), liquid and gas chromatography as well as radioactivity have already been evaluated when analyzing water industry [8]. UV–Vis spectrophotometry, fluorescence, quimioluminescence and ISE potentiometry methods for ammonium analysis in water samples have also been recently compared [32].

4.2. Exemplary analysis

4.2.1. Algorithm specification

The assessment of the SY analysis in animal feed and meat by means of the high performance liquid chromatography with diode array detection (HPLC-DAD) versus tandem mass spectrometry detection (HPLC-MS/MS) techniques [11] have been taken as a model to show the evaluation procedure established by the hexagon tool. Initially, the adequacy of the analytical parameters relative to sample/method and quality control is assessed. The aspects parameters considered and the PPs assigned for figures of merit 1 (FM-1) are listed in Tables 2–4 as shown in Ref. [8]. With the aim of only comparing the intrinsic characteristics of each procedure, the number of samples per week in both cases is fixed to 50, that is, the greenest alternative. The HPLC-DAD analytical method implies the need of 5 times preconcentration during the sample treatment (extraction process) in comparison to HPLC-MS/MS. As shown in Fig. 1a, the sum of PPs is 11 for the HPLC-DAD whereas only 9 PPs are assigned to the HPLC-MS/MS. This can be understood by the fact that triple quadrupole tandem mass spectrometer offers better sensitivity (2.18 ng/mL) than detection via diode-array detector (74.81 ng/mL). In addition to this, HPLC-DAD presents worse adaptation to the figures of merit regarding the calibration procedure than the HPLC-MS/MS due to the limit of detection and working range of concentrations [11]. On the other hand, both techniques present similar penalization concerning quality control and accuracy criteria (FM-2), as it can be seen in Fig. 1a. Overall penalization of figures of merit FM-1 and FM-2 is indicated in Fig. 1b.

The global penalization assigned to health and safety variables is presented in Fig. 1b. The high penalization score of both methods is due to the use of hazardous chemical reagents such as methanol, ethanol, formic acid and acetonitrile organic solvent. Toxicity PPs correspond to the sum of the penalties attributed to the pictograms of the SGA system each reagent has. Concerning safety, the HPLC-DAD based analytical procedure implies the evaporation to dryness in water bath under nitrogen beam during the sample extraction process. This makes the HPLC-MS/MS based method more suitable related to safety considerations. In order to evaluate sustainability, Table 9 in Ref. [8] is employed. Taking into account the amount of waste generated and the principles of green chemistry [33,34], it is concluded that both methods lead to similar contribution in terms of sustainability, that is, 13 PPs as shown in Fig. 1b.

Although similar environmental friendliness is found for the evaluated methods regarding residues generation, the electricity consumed by the instrumental equipment provides a remarkable difference in the environmental impact. Mainly, the difference between the two methods relies on the fact that a 16-min HPLC-DAD run is needed whereas HPLC-MS/MS requires the equipment to be switched on the whole working day (8 h). This leads to a much higher carbon footprint when using HPLC-MS/MS, as depicted in Fig. 1c. The carbon footprint estimation [27] is computed considering the time analysis and the instrument power set to 0.44 KW.

| Procedure        | Arithmetic mean ($S_{ar}$) |
|------------------|----------------------------|
| HPLC-MS/MS       | 2.57                       |
| HPLC-DAD         | 2.14                       |
| ELISA            | 2.29                       |
| CE               | 2.29                       |
| PAS              | 1.43                       |
| FDS              | 1.43                       |
and 3.69 kW for HPLC-DAD and HPLC-MS/MS, respectively. The reference constant value emission factor equals to 0.247 kg CO₂/ kWh [35]. In conclusion, the HPLC-DAD method involves a greener procedure from the environmental point of view.

Last variable analysed is the annual economic cost (in €). To compute this value, the sum of the criteria listed in Ref. [8] is assumed. Both HPLC-DAD and HPLC-MS/MS methods are supposed to analyse 50 samples weekly, giving rise to the same annual average of samples. The salary assigned to skilled personnel taking into account an 8-h working day and the reagents and consumable material costs are similar for both methods. However, noticeable differences are found when defining electricity costs (0.15 €/kWh) according to the time of analysis for the annual average number of samples, and more remarkably, the equipment cost. It is well known that HPLC-MS/MS equipment with a triple quadrupole mass spectrometer is much more expensive than HPLC-DAD. Therefore, the global estimation of the economic cost indicates that HPLC-MS/MS method is more cost-effective than the HPLC-DAD, as represented in Fig. 1d.

4.2.2. Evaluation results

The results obtained when evaluating HPLC-DAD and HPLC-MS/ MS methodologies can be summarized in the hexagon pictogram [8], as shown in Fig. 2. By using a 0–4 penalization scale, the variables of the methods are organized in six equilateral triangles and quantified with a final qualification mark according to penalty points ranges [8]. The conclusions obtained when comparing the penalization scores from Fig. 2 are the following: the Sunset Yellow analysis by means of HPLC-MS/MS based method is the worst environmentally friendly analysis (2 versus 0) due to the intrinsic characteristics of the technique. However, HPLC-DAD method offers advantages in terms of environmental impact and better cost-effectiveness relation. Therefore, it can be concluded that HPLC-DAD method provides satisfactory analytical performance for the determination of Sunset Yellow in animal feed and chicken samples, as already stated in Ref. [11].

In addition to HPLC-DAD and HPLC-MS/MS analytical techniques, the evaluation of the SY analysis has also been carried out when considering capillary electrophoresis with UV detection (CE) [12], polyclonal antibody-based indirect enzyme-linked immunoabsorbent assay (ELISA) [13], photoacoustic spectroscopy [14], and first derivative spectrophotometry (FDS) [14]. The final penalization score for each method is indicated in the corresponding hexagon pictogram, in Fig. 3. The figures presenting more data concerning evaluation of the selected methods are shown in the Electronic Supplementary Material (ESM), in Fig. S1.

When comparing the results between the methods, it should be mentioned that photoacoustic spectroscopy (PAS) presents better adaptation of the figures of merit than the other methods (compare penalization score FM-1 equal to 2 and FM-2 equal to 1 with 3/2 or 3/1 for CE and ELISA methods, respectively). PAS showed high sensitivity and satisfactory precision, together with its non-destructive character. Also, it allowed the simultaneous determination of food dyes, among them SY, with a very good agreement between the values determined by using first derivative spectrophotometry (FDS). Therefore, these results indicated the potential of photoacoustic spectroscopy as an analytical method in the analysis of food dyes, where no preliminary separation step is required.

As regards toxicity and safety variables, CE and ELISA analytical methods show the worst penalization score (4/3 in comparison to 1/2 from PAS/FDS). This can be understood by the fact that both methodologies imply a sample pretreatment that requires the use of several chemical reagents and materials. For instance, diamino moieties functionalized silica nanoparticles (dASNsPs) are employed as both adsorbents in preconcentration of SY colorant by the dispersive solid-phase microextraction (DSPME) process, and pseudostationary phases (PSPs) in capillary electrophoresis (CE) separation. On the other hand, ELISA method showed high sensitivity, simplicity and rapidity for the detection of SY, although it is the most expensive method because of the wide variety of chemicals and equipment needed in the analysis.

With the aim of ranking the analytical procedures from the most sustainable (0 score) to the least (4 score), the arithmetic mean (Sₘ) of each method is indicated in Table 2. The results obtained are comparable to those already explained by using the hexagon pictogram for each method.

5. RGB algorithm

5.1. Working principle

The RGB model develops and extends the concept of "greenness" of an analytical method by the other primary colors assigned to other basic attributes of a method, as a result of which the resultant color of a method is determined by the contribution of the Red, Green and Blue components [9]. Red (R) color is assigned to analytical performance expressed by validation criteria, which are a measure of the quality of analytical result, green (G) to safety and environmental friendliness, and blue (B) to practical efficiency and productivity.

The intensity of a given primary color is expressed by the CS parameter (Color Score) on the scale of 0–100, distinguishing three ranges: <33.3% - the range of a general lack of acceptance for the attribute under consideration, >33.3% and <66.6% - the range of acceptance but not satisfaction, and >66.6% - satisfaction for a given attribute. The above ranges allow to significantly simplify the use of the RGB model for the assessment of analytical methods and distinguish the limited number of resultant/final colors of a method, presented in Fig. 4.

A method’s color is the qualitative parameter that is easy to estimate and interpret. Another parameter called the “method brilliance” (MB) is dedicated to a more thorough quantitative assessment. MB is calculated as the weighted geometric mean of three CS values corresponding to the respective primary colors, with “W” weights, selected by the user. As a result, MB has no direct correlation with color, as it allows for assigning different weights to the given primary attributes, e.g. greater for red (analytical performance) than greenness, etc. In addition, recognizing MB as the geometric mean makes it more sensitive to extremely low values of CS, which may constitute bottlenecks of the whole method and affect its general utility.

To determine the CS value for a given primary color, at least three criteria adequate for the given attribute should be selected, for example: precision, accuracy and sensitivity for R, reagent toxicity, amount of waste and other hazards for G, and cost of analysis, sample throughput and sample consumption for B. Then, appropriate weights should be assigned to the selected criteria (w), independent of the weights assigned to the primary colors (W).

The next step is to assess the given criterion by an appropriate score on a scale of 0–100, e.g. precision expressed by the RSD value, using the two proposed reference points, Lowest Acceptable Value (LAV) and Lowest Satisfactory Value (LSV). LAV is a value from which the result can be considered “only” acceptable (e.g. RSD = 5%), while LSV is a value from which the result is satisfactory (e.g. RSD = 2%). The value obtained for a given criterion equal to LAV should be awarded the score 33.3, while to LSV values – 66.6. Extreme values, i.e. 0 and 100, are given when the criterion is completely unacceptable (e.g. RSD > 25%) or the best of the available
methods in a given area (e.g. RSD <0.25%). The relationship be-
 tween the value received for a given criterion (here RSD) and the
 score to be placed does not always have to be linear over its entire
 range, and should be adapted to the specifics of a given criterion.

To facilitate the assessment process, the awarded scores can be
rounded off and graded every 5 points, taking into account the
additionally mentioned values 33.3 and 66.6 when the result of the
method equals LAV and LSV, respectively. Finally, the CS values for a
given primary color are calculated as the weighted geometric mean
of the scores awarded, with the weights assumed (w), similar to the
MB value calculated as the geometric mean of the CS values taken
with weights (W).

A special algorithm was designed to evaluate methods using the
RGB model, based on a standard Excel spreadsheet. The spread-
sheet is available on-line as the original publication’s supplement
[9]. It should be noted that the proposed model is flexible and al-
 lows the user to adjust the assessment specification to his subject-
ive preferences: weights assigned to given primary colors – W,
selection of appropriate criteria for a given primary attribute,
weights of given criteria - w, and LAV and LSV values which play the
role of reference points. This flexibility is good because it allows a
method to be assessed in terms of the planned application and the
resulting expectations, it allows the reverse option, i.e. predicting
the best method application according to the assessments received.
with different sets of guidelines, and allows for assessment according to defined internal standards adopted e.g. in a given laboratory due to its specificity. In addition, the flexibility of the model allows one to re-evaluate the method according to other guidelines. In another scenario, the assessment can be made in an objective manner, according to strictly regulated guidelines adopted by a wider group of analysts. It can be assumed that proposing such standards increasing objectivity of assessments is a matter of the near future.

5.2. Exemplary analysis

5.2.1. Algorithm specification

The assessment using the RGB algorithm was focused on the critical analysis of an overall analytical potential of individual methods. The word “method” was defined as an entire procedure including the preparation of sample material and instrumental analysis, assuming the initial availability of all necessary reagents ready for use in the laboratory. The choice of given assessment parameters and their significance (weights) was subjective, but was supported by an informal discussion among the widest possible group of employees of the Department of Analytical Chemistry at the Faculty of Chemistry, Jagiellonian University in Krakow, and people from friendly analytical laboratories.

The red attribute (analytical performance) was treated with the highest relative weight $W = 5$, which reflects the analysts’ general expectations that an effective method should primarily ensure a good quality of the analytical result. The blue attribute (productivity and practical efficiency) was treated as the second most important, with the relative weight $W = 4$, because as we assume, the next general expectation of analysts relates to the practical aspects of the analytical procedure, which can often be another limiting factor. The green attribute (compliance with the principles of green analytical chemistry, i.e. environmental friendliness and safety) was treated with the weight $W = 3$. This choice still reflects the strong emphasis on “greenness”, but does not give it priority or equal significance to the red or blue attributes, indicated in informal discussions as generally more important. A detailed

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**Fig. 3.** Hexagon pictograms for the analytical methods: a) CE, b) ELISA, c) PAS and FDS.

**Fig. 4.** Nine resultant colors of a method predicted by the RGB model.
discussion of the individual criteria selected for evaluation in the red, green and blue areas is given in the ESM.

5.2.2. Evaluation results

The completed Excel worksheets presenting the assessment results of the six considered methods are shown in Figs. 5–10. To shorten the length of the main text, the detailed discussion of the scores assigned to the individual criteria and resulting CS values is shown in ESM. Below is presented only the discussion of the methods’ resulting colors and MB values (main evaluation outcomes).

5.2.2.1. Resultant color. The RGB model offers two ways to express an overall assessment of the method, qualitative — using the resultant color, and quantitative — using the method’s brilliance value (MB) in the range 0–100%. The best qualitative assessment is white color, which requires having all three CS values over the satisfaction level (Fig. 4). This color is however lacking among the evaluated methods, thus none of them is fully complete. The best method in this sense is HPLC-DAD [11], with the magenta color, and FDS [14] — characterized as cyan. The former one has two primary colors, red and blue, and it lacks green. Thus, in overall, it both ensures good quality of analytical results and is practically/economically-effective. Therefore, it seems to be worth recommending when other more ecological methods are not available.

Interestingly, the FDS method has gained another equipotent secondary color of the RGB model (cyan), also indicating the possession of two primary colors, but in this case, they are green and blue. This shows that the strengths and weaknesses of these methods are different. The FDS method lacks analytical performance compared to HPLC-DAD, but is more environmentally-friendly and safe for users in general. It can therefore be a good alternative if frequent routine tests are performed, without stringent requirements regarding the quality of the quantitative results.

The methods that have gained only one primary color are HPLC-MS/MS [11], ELISA [13] and PAS [14]. The HPLC-MS/MS method has been classified as red, i.e. its missing attributes are green and blue. It is worth remembering, however, that despite the high Csred value, above the 66.6% threshold, the two criteria were rated quite poorly — precision and, above all, accuracy (RE above 30%), see Fig. 5 and ESM for more details. Although this method undoubtedly makes up for these deficiencies by other features, such as low LOD and “other red aspects”, the low reliability of the assays can be a bottleneck and exclude successful application of the method in some situations. The ELISA method, in turn, gained red color due to its great precision, accuracy, linearity and by far the lowest LOD value. Its big drawback may be the lack of multi-component analysis, which is simple in the case of separation methods. Therefore, in this respect, HPLC-MS/MS and ELISA complement each other well.

The lack of other primary colors, however, informs about their other limitations, which may be crucial in some situations, e.g. costs or complexity of the entire procedure. Another example is the PAS method, considered green, which lacks red and blue colors to be complete. Its use may be justified in the situation when few analyzes are carried out with relatively low requirements regarding the quality of results, e.g. at universities, for educational and didactic purposes related to the problem of adding food colors.

The only method which has not obtained any primary color is CE [12], marked as gray (colorless). This demonstrates its lack of clearly strong points, hence in the qualitative sense, it can be considered the weakest of the whole group. However, as described in the next paragraph, this does not preclude its competitiveness in relation to other methods, because color is not a good measure of overall potential, but rather a simple indication of predispositions and potential areas for improvement, easy to encode graphically, remember and interpret.

5.2.2.2. Brilliance value. An appropriate measure of the global method potential is MB, which is calculated as a weighted geometric mean from the corresponding CS values. In addition to the quantitative nature, enabling a more accurate assessment than using color, it depends on the weights (W) assigned to the given primary attributes (here W = 5 for red, 3 for green and 4 for blue). Therefore, it allows adjusting the specification of assessment (algorithm) to the subjective preferences of evaluator, or the more objective rules accepted by a wider group of users.

Considering the individual MB values, HPLC-DAD proved to be the best method from the whole group, with MB = 68.1%. This result, in combination with the obtained magenta color, indicate the undoubted high potential of this method in various respects, with one small drawback in the form of fairly average greenness (see Fig. 7 and ESM). The second method, which may come as a surprise when analyzing the obtained colors, is CE, with the value of MB = 60.6%. Although this method was previously considered as gray, i.e. without clearly strong points, the high MB value shows its good inner balance, i.e. maintaining all three attributes at a fairly high level, but slightly below the threshold that guarantees obtaining color (66.6%). This indicates its fairly wide applicability, confirmed by its high position in the ranking. The spectroscopic methods, FDS (MB = 59.4%) and PAS (MB = 58.6%), as well as HPLC-MS/MS (MB = 58.3%), were rated slightly lower than CE. The differences between these three methods are small. Nevertheless, as shown by the respective colors, the advantages and weaknesses of these methods are different, especially when it comes to the red and green aspects. In this regard, the HPLC-MS/MS (red) is complementary to spectroscopic methods (green).

The ELISA method was rated the lowest, with MB = 52.7%. This result is actually not very low, it still remained above 50%, i.e. above the general “average”, but indicates clear deviations from the other methods. Despite the highest Csred value among all methods and the highest weight assigned to red (W = 5), the low-rated green and blue attributes, associated mainly with the complexity of an entire experimental procedure, influenced the lowest position in the general ranking.

6. Comparison of algorithms

The purpose of this section is to compare the evaluation outcomes obtained using the three tools described above. This test was carried out in two variants. In the first, the selection of the evaluation criteria and the weights assigned to them was done by the authors of the particular approach. It was not previously agreed between the co-authors, so it differed significantly between the algorithms (M.T. was responsible for TOPSIS, for HEXAGON A–B, and P–C–F, while for RGB P–N and P–K). In the second variant, the specificity of the algorithms has been unified, taking the same criteria for assessment as possible, while maintaining the mathematical and visual distinctiveness of each tool.

In addition, a new measure of the overall method efficiency, Averaged Method Efficiency Index (AMEI) expressed in percentages was proposed, which results from assessments obtained using each of the three algorithms:

\[ \text{AMEI} = \sqrt{100C^r + 100 \left(1 - \frac{4-S_m}{4}\right) MB} \]  

(8)
**Fig. 5.** Outcomes of the HPLC-DAD method evaluation using the RGB algorithm.

| REDNESS (analytical performance) | W=5 | Precision (RSD) | Accuracy (RE) | LOD | Linearity (R²) | Other aspects |
|----------------------------------|-----|----------------|--------------|-----|----------------|---------------|
| CS: 74.8%                        |     | 15%            | 25%          | 500 ng/mL | 0.980          | acceptable    |
| LAV=33.3                        |     | 5%             | 10%          | 50 ng/mL  | 0.995          | acceptable    |
| Result                          | 4.45% | 12.5%        |              | 74.8 ng/mL | 0.999          | very good     |
| Score (0-100)                   | 70   | 70             | 80           | 65       | 85             | 95            | 95            | 90            | 90            |

| GREENNESS (safety and eco-friendliness) | W=3 | Waste amount | Toxicity of chemicals | Other occupational hazards | Other aspects |
|-----------------------------------------|-----|--------------|-----------------------|----------------------------|---------------|
| CS: 58.2%                               |     | moderate     | satisfactory          | 1 hazard                   | satisfactory  |
| LAV=33.3                                |     | acceptable   | satisfactory          |                           |               |
| LSV=66.6                                |     | satisfactory | satisfactory          |                           |               |
| Result                                  |     |              |                       |                           |               |
| Score (0-100)                           | 50  | 50           | 50                    | 50                        | 50            | 80            | 80            | 88.6          | 88.6          |

| BLUENESS (productivity / practical effectiveness) | W=4 | Cost of analysis | Time of analysis | Sample consumption | Other aspects |
|--------------------------------------------------|-----|-----------------|-----------------|---------------------|---------------|
| CS: 68.2%                                        |     |                 |                 |                     |               |
| LAV=33.3                                        |     |                 |                 |                     |               |
| LSV=66.6                                        |     |                 |                 |                     |               |
| Result                                           |     |                 |                 |                     |               |
| Score (0-100)                                    | 66.6| 66.6           | 66.6            | 66.6                | 66.6          | 75            | 75            |

**FINAL COLOR:** **MAGENTA**

- REDNESS: ±33.3%
- GREENNESS: ±266.6%
- BLUENESS: ±33.3%

Short annotation: 68.1magenta
Long annotation: 68.1magenta(74.8/red-68.2/green-68.2/blue)

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**Fig. 6.** Outcomes of the HPLC-MS/MS method evaluation using the RGB algorithm.

| REDNESS (analytical performance) | W=6 | Precision (RSD) | Accuracy (RE) | LOD | Linearity (R²) | Other aspects |
|----------------------------------|-----|----------------|--------------|-----|----------------|---------------|
| CS: 68.8%                        |     | 15%            | 25%          | 500 ng/mL | 0.980          | acceptable    |
| LAV=33.3                        |     | 5%             | 10%          | 50 ng/mL  | 0.995          | acceptable    |
| Result                          | 7.73% | 31%         |              | 2.1 ng/mL  | 0.999          | excellent     |
| Score (0-100)                   | 60   | 60             | 80           | 65       | 85             | 95            | 95            | 90            | 90            |

| GREENNESS (safety and eco-friendliness) | W=3 | Waste amount | Toxicity of chemicals | Other occupational hazards | Other aspects |
|-----------------------------------------|-----|--------------|-----------------------|----------------------------|---------------|
| CS: 53.0%                               |     | moderate     | satisfactory          | 5 hazards                   | acceptable    |
| LAV=33.3                                |     | acceptable   | satisfactory          |                           |               |
| LSV=66.6                                |     | satisfactory | satisfactory          |                           |               |
| Result                                  |     |              |                       |                           |               |
| Score (0-100)                           | 50  | 50           | 50                    | 50                        | 50            | 80            | 80            | 86.6          | 86.6          |

| BLUENESS (productivity / practical effectiveness) | W=4 | Cost of analysis | Time of analysis | Sample consumption | Other aspects |
|--------------------------------------------------|-----|-----------------|-----------------|---------------------|---------------|
| CS: 51.1%                                        |     |                 |                 |                     |               |
| LAV=33.3                                        |     |                 |                 |                     |               |
| LSV=66.6                                        |     |                 |                 |                     |               |
| Result                                           |     |                 |                 |                     |               |
| Score (0-100)                                    | 33.3| 33.3           | 66.6            | 66.6                | 66.6          | 66.6          | 50            | 50            |

**FINAL COLOR:** **RED**

- REDNESS: ±33.3%
- GREENNESS: ±266.6%
- BLUENESS: ±33.3%

Short annotation: 58.3red
Long annotation: 58.3red(68.8/red-53.0/green-51.1/blue)
Fig. 7. Outcomes of the CE method evaluation using the RGB algorithm.

Fig. 8. Outcomes of the ELISA method evaluation using the RGB algorithm.
**Fig. 9.** Outcomes of the PAS method evaluation using the RGB algorithm.

| CS. 51.0% | **REDNESS (analytical performance)** | W=5 | Precision (RSD) | Accuracy (RE) | LOD | Linearity (R²) | Other aspects |
|-----------|--------------------------------------|-----|-----------------|--------------|------|----------------|---------------|
| LAV=33.3  | 19%                                  | 25% | 500 ng/mL       | 0.980        | acceptable |
| LSV=50.6% | 5%                                   | 10% | 50 ng/mL        | 0.965        | satisfactory |
| Result    | 1.89%                                | ??  | 28 ng/mL        | 0.990        | below acceptable |
| Score (0-100) | 90 | 90 | 33.3 | 33.3 | 70 | 70 | 66 | 66 | 30 | 30 |

| CS. 68.2% | **GREENNESS (safety and eco-friendliness)** | W=3 | Waste amount | Toxicity of chemicals | Other occupational hazards | Other aspects |
|-----------|-----------------------------------------------|-----|--------------|------------------------|---------------------------|---------------|
| LAV=33.3  | acceptable                                    | acceptable | 5 hazards | acceptable |
| LSV=50.6% | satisfactory                                  | satisfactory | 2 hazards | satisfactory |
| Result    | satisfactory                                  | satisfactory | 0 hazards | acceptable |
| Score (0-100) | 68.8 | 68.6 | 68.6 | 68.6 | 68.6 | 100 | 100 | 50 | 50 |

| CS. 62.3% | **BLUENESS (productivity / practical effectiveness)** | W=4 | Cost of analysis | Time of analysis | Sample consumption | Other aspects |
|-----------|----------------------------------------------------------|-----|------------------|------------------|-------------------|---------------|
| LAV=33.3  | acceptable                                               | acceptable | acceptable | acceptable |
| LSV=50.6% | satisfactory                                              | satisfactory | satisfactory | satisfactory |
| Result    | over satisfactory                                         | over satisfactory | acceptable | satisfactory |
| Score (0-100) | 75 | 75 | 75 | 75 | 75 | 33.3 | 33.3 | 66.6 | 66.6 |

| **FINAL COLOR:** | REDNESS | GREENNESS | BLUENESS |
|------------------|---------|----------|----------|
| **GREEN**        | ≥33.3%  | ≥33.3%   | ≥33.3%   |
| **BRILLIANCE**   | (MB):   | (MB):    | (MB):    |
|                  | 58.6%   | 58.6%    | 58.6%    |

Short annotation: 58.6green
Long annotation: 58.6green(51.0/5red-68.2/3green-62.3/4blue)

**Fig. 10.** Outcomes of the FDS method evaluation using the RGB algorithm.

| CS. 49.5% | **REDNESS (analytical performance)** | W=5 | Precision (RSD) | Accuracy (RE) | LOD | Linearity (R²) | Other aspects |
|-----------|--------------------------------------|-----|-----------------|--------------|------|----------------|---------------|
| LAV=33.3  | 15%                                  | 25% | 500 ng/mL       | 0.980        | acceptable |
| LSV=50.6% | 5%                                   | 10% | 50 ng/mL        | 0.965        | satisfactory |
| Result    | 1.89%                                | ??  | 28 ng/mL        | 0.990        | below acceptable |
| Score (0-100) | 90 | 90 | 33.3 | 33.3 | 70 | 70 | 66 | 66 | 30 | 30 |

| CS. 68.2% | **GREENNESS (safety and eco-friendliness)** | W=3 | Waste amount | Toxicity of chemicals | Other occupational hazards | Other aspects |
|-----------|-----------------------------------------------|-----|--------------|------------------------|---------------------------|---------------|
| LAV=33.3  | acceptable                                    | acceptable | 5 hazards | acceptable |
| LSV=50.6% | satisfactory                                  | satisfactory | 2 hazards | satisfactory |
| Result    | satisfactory                                  | satisfactory | 0 hazards | acceptable |
| Score (0-100) | 66.6 | 66.6 | 66.6 | 66.6 | 66.6 | 100 | 100 | 50 | 50 |

| CS. 67.3% | **BLUENESS (productivity / practical effectiveness)** | W=4 | Cost of analysis | Time of analysis | Sample consumption | Other aspects |
|-----------|----------------------------------------------------------|-----|------------------|------------------|-------------------|---------------|
| LAV=33.3  | acceptable                                               | acceptable | acceptable | acceptable |
| LSV=50.6% | satisfactory                                              | satisfactory | satisfactory | satisfactory |
| Result    | very good                                               | over satisfactory | acceptable | over satisfactory |
| Score (0-100) | 90 | 90 | 90 | 75 | 75 | 75 | 33.3 | 33.3 | 75 | 75 |

| **FINAL COLOR:** | REDNESS | GREENNESS | BLUENESS |
|------------------|---------|----------|----------|
| **CYAN**         | ≥33.3%  | ≥33.3%   | ≥33.3%   |
| **BRILLIANCE**   | (MB):   | (MB):    | (MB):    |
|                  | 59.4%   | 59.4%    | 59.4%    |

Short annotation: 59.4cyan
Long annotation: 59.4cyan(49.9/5red-68.2/3green-67.3/4blue)
where $C_i^*$ is the closeness to the ideal solution obtained by TOPSIS, $S_{av}$ is the average score (from 0 to 4) obtained by HEXAGON and MB is the brilliance value obtained by RGB model.

AMEI was defined as a weighted geometric mean, to make it more sensitive to possible extreme outcomes that may indicate serious limitations overlooked by the other algorithms. It allows one to rank the compared methods globally from the best to the worst, treating each algorithm used with the same importance. This parameter is therefore useful in expressing the potential of the method in the most general way, eliminating to some extent the subjectivity and specifics of each approach.

Comparison of the compatibility between the individual assessment outcomes, including: TOPSIS, HEXAGON, RGB and AMEI approaches, is shown in Figs. 11 and 12. They show the values of respective quantitative parameters indicating the overall potential of given methods as a percentage agreement with the ideal situation, i.e. the best possible value of a given parameter ($C_i^*$, $S_{av}$ and MB). Fig. 11 refers to the variant in which the selection of assessment criteria was different, while Fig. 12 refers to the second variant in which the selection of criteria was agreed as much as possible. To facilitate the analysis of differences between these two variants, the individual criteria used in each algorithm and their relative weights are given in Tables 3 and 4.

As can be seen in Fig. 11, the adoption of different criteria for applying the given algorithms results in the quite different outcomes. The largest differences are observed in the position of HPLC-DAD and CE methods, while significantly smaller and indicating better compatibility, for HPLC-MS/MS and ELISA methods that were poorly assessed by each model, as well as for FDS and PAS that were well rated in any case. This results directly from the guidelines followed by the evaluators of the methods, which are manifested in the selection of the parameters to be evaluated and their relative significance. For example, the low position of HPLC-DAD and the high CE according to the TOPSIS model results from a great emphasis on aspects of environmental friendliness and solvent consumption, while the different results obtained by the RGB model for these methods result from the lower relative weight of these criteria and the consideration of other aspects omitted in TOPSIS, e.g. accuracy, linearity or cost of analysis.

On that account, it can be assumed that this situation, combined with a different mathematical structure and other details, is not surprising. On the other hand, the obtained AMEI values indicate an averaged potential of the given methods taking into account all

Fig. 11. Comparison of the evaluation results obtained using all discussed approaches, according to the different specifics of each algorithm (TOPSIS, HEXAGON and RGB). AMEI presents the averaged outcomes.

Fig. 12. Comparison of the evaluation results obtained using all discussed approaches, according to the unified specifics of each algorithm (TOPSIS, HEXAGON and RGB). AMEI presents the averaged outcomes.

### Table 3

| TOPSIS | HEXAGON | RGB |
|--------|---------|-----|
| Criterion | $W_{tot}$ | Criterion | $W_{tot}$ | Criterion | $W_{tot}$ |
| LOD | 1/8 | Figures of merit 1 (sample treatment, method characteristics and calibration) | 1/6 | RSD (precision) | 10/120 |
| RSD (precision) | 1/8 | Figures of merit 2 (quality control and accuracy) | 1/6 | Accuracy | 10/120 |
| Amount of organic solvent | 1/8 | Toxicity/Safety | 1/6 | LOD | 10/120 |
| Amount of organic solvent$^a$ | 1/8 | Carbon footprint | 1/6 | Linearity ($R^2$) | 10/120 |
| toxicity (hazard) | | | | | |
| Amount of sample | 1/8 | Residues | 1/6 | Other “red” aspects | 10/120 |
| Solid waste | 1/8 | Cost of analysis | 1/6 | Waste amount | 9/120 |
| Other analytes | 1/8 | | | Toxicity of chemicals | 9/120 |
| (multicomponent analysis) | | | | | |
| Time of analysis | 1/8 | | | | |

$W_{tot}$ is the relative weight of a given criterion in respect to all other criteria listed.

$^a$ In the case of RGB algorithm $W_{tot}$ includes both “W” and “w” (see Section 4.1).
algorithms with the same significance. They indicate CE, FDS and PAS as globally better than HPLC-DAD, HPLC-MS/MS and ELISA.

The comparison of Figs. 11 and 12 gives a clear confirmation of the role played by the selection of criteria for the final evaluation results. In the second variant, much better agreement between individual algorithms was achieved, despite the differences in the assessment method — similarity to the best alternative in TOPSIS, awarding penalty points in HEXAGON and awarding score values based on LAV and LSV reference points in RGB. In addition, other mathematical rules and visualization of outcomes remained different as well. For instance, as far as HEXAGON and RGB present outcomes in a quite pictorial way, TOPSIS is limited to only one table presenting similarity to ideal solution. At the same time, the obtained compliance confirms the usability of each algorithm, and the minor differences observed for some methods should be attributed to maintaining the aforementioned separateness in certain aspects. It is also worth noting that the scores obtained for the methods have different range for the given algorithms. The biggest differences between the methods are observed for TOPSIS, while the smallest for RGB, which results directly from the mathematical structure.

Our intention was not to indicate the most appropriate approach to method evaluation, but to make it easier for the reader to choose the optimal tool in a given case. For example, if the graphic design is an important element facilitating the analysis, the HEXAGON and RGB methods are recommended. Conversely, if a user values simplicity, MCDa would be optimal. Moreover, if the assessment of the criteria according to the awarded penalty points seems convenient, as in the case of Eco-scale [3], HEXAGON will be the best choice. If a user values an overall flexibility of the assessment process, he should choose RGB, if minimum effort, TOPSIS method. To provide readers with additional support, the main advantages and drawbacks of each algorithm are gathered in Table 5. Moreover, the Excel spreadsheets used for evaluating the presented methods, which can be used by readers as a template for other assessments, are provided for each tool in ESM.

7. Conclusions

It can be concluded from this work that the global assessment of the analytical method is not simple, although desirable for many reasons. In this work we illustrated and compared the three different approaches that can be used for this purpose, each with a different specification, each worth considering as a valuable auxiliary tool. Despite some differences in the mode of operation, mathematical structure and form of presentation of results, it can be assumed that the results of the assessment of selected methods using these tools will be similar, provided that agreement on the choice of parameters to be assessed and their significance is ensured. What is unavoidable, however, some differences should always be expected. In this regard, it is also worth considering the approach of applying all three algorithms, followed by the calculation of AMEI, a new measure of method’s general efficiency. It quantifies and expresses the averaged overall potential of a given method, and offers yet another perspective of comparing alternative solutions. It may also be interesting to apply another MCDa method, different than TOPSIS, which might occur more effective in a given situation. The aim of this work was not to indicate absolutely better or worse approaches to the assessment of methods, because, as the authors agree, each of them offers its own
advantages, but also some limitations. We hope, however, that choosing the optimal option in a given situation will be easier, and the interest in carrying out a comprehensive assessment of analytical methods, going beyond the usual validation criteria, will be greater than before and will translate into new opportunities. In the future, we plan to use the described algorithms to evaluate other analytical methods, to further explore their capabilities, develop and improve. For example, a new version of the RGB algorithm will be presented soon, it will be simplified in terms of the selection of variables, allowing for faster evaluation and easier interpretation of results. In case of ambiguities and questions regarding the presented tools, we encourage readers to contact their authors directly.

CRediT authorship contribution statement

Paweł Mateusz Nowak: Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Software, Supervision, Visualization, Writing - original draft, Writing - review & editing.
Pawel Koscielniak: Conceptualization, Supervision.
Marek Tobiszewski: Conceptualization, Data curation, Formal analysis, Investigation, Writing - original draft, Writing - review & editing.
Ana Ballester-Caudet: Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Software, Visualization, Writing - original draft, Writing - review & editing.
Pilar Campíns-Falcó: Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Software, Supervision, Visualization, Writing - original draft, Writing - review & editing.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.trac.2020.116065.

Electronic Supplementary Material (ESM)

Additional description of algorithms and assessment details, additional outcomes of assessment, Excel worksheets used for evaluations with the integrated algorithms (as a template for modification and further use by readers).

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