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Periodic cooking of eggs



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Egg cooks are challenged by the two-phase structure: albumen and yolk require two cooking temperatures. Separation or a compromise temperature to the detriment of food safety or taste preference are the options. In the present article, we find that it is possible to cook albumen and yolk at two temperatures without separation by using periodic boundary conditions in the energy transport problem. Through mathematical modeling and subsequent simulation, we are able to design the novel cooking method, namely periodic cooking. Comparison with established egg cooking procedures through a plethora of characterization techniques, including Sensory Analysis, Texture Profile Analysis and FT-IR spectroscopy, confirms the different cooking extents and the different variations in protein denaturation with the novel approach. The method not only optimizes egg texture and nutrients, but also holds promise for innovative culinary applications and materials treatment.

Eggs are one of the most valuable foods¹ on the tables of consumers and in the kitchens of chefs due to their abundant functional properties that make them the funniest and most versatile ingredients to work with. In fact, not only do they contain almost all essential nutrients², which is fundamental for human nutrition^{3,4}, but they are also distinguished by their excellent foaming and emulsification capacity and their ability to coagulate and form gels upon heating^{1,5}.

As a consequence, they are commonly added in a variety of recipes, but are also used alone, and extremely different products can be obtained by changing the conditions under which they are cooked. In fact, tens of cooking methods are available today, including the most basic shell-on egg cooking techniques, of which *à la coque*, hard-boiling and soft-boiling are only the tip of the iceberg. In particular, the interest of chefs is currently attracted by the so called *sous vide* egg or 6X °C egg. This novel cooking method requires immersion of the shell-on egg in water at low and constant temperatures, usually between 60 and 70 °C, for at least 1 h⁶, and gives a very peculiar result, where both albumen and yolk have the same creamy texture.


The problem is that with this cooking technique, the albumen does not fully set (only one albumen protein is capable of doing so at such low temperatures, ovotransferrin⁷). Due to the fact that albumen and yolk have two very different compositions^{4,8,9}, they require two different temperatures for optimal cooking, namely around 85 °C for albumen and around 65 °C for yolk^{5,6,10}. Many chefs have tried to overcome this problem by cooking yolk and albumen separately at the appropriate corresponding temperatures, but this results in a series of too long and complicated steps.

As a response to this, we explore the idea of imposing two different cooking temperatures in two different regions of the egg without necessarily

cracking the shell open. The inspiration comes from previous works of our group, where time-varying boundary conditions (BCs) in the mass transport of the blowing agent are exploited to produce foams with different layers in terms of morphology and/or density¹¹. This same principle can also be exploited to obtain a desired thermal profile inside any material, eggs included. Therefore, we propose here to cook the egg by imposing a periodic time-varying BC in the energy transport problem to build our *ad hoc* thermal profile and repeat it until optimal cooking of both yolk and albumen. On a practical level, the idea is to place the raw shell-on egg alternatively in hot water (T_h) and cold water (T_c) for relatively short periods of time (t_h and t_c , respectively) and repeat these cycles N times until the cooking of both the yolk and the albumen is reached.

In the present article, design of the novel cooking method, namely periodic cooking, is conducted through mathematical modeling of the process, involving concomitantly heat transfer inside the egg and gelation of both egg yolk and albumen. A fine solution of the problem is then found through simulation with a Computational Fluid Dynamics (CFD) software, where imposition of different BCs allows to compare the evolution of the temperatures and the cooking degrees inside the egg obtained with different cooking methods (we will specifically refer to hard-boiling, soft-boiling, *sous vide* cooking and periodic cooking). Finally, validation of the simulation is allowed by cooking intact fresh eggs with all four methods and analyzing the final cooked products through both Sensory Analysis and Texture Profile Analysis (TPA) to gather information about color, consistency, texture, and taste. Additional information are then collected also through FT-IR spectroscopy, hydrogen-1 Nuclear Magnetic Resonance (¹H-NMR) and High Resolution Mass Spectrometry (HRMS) in order to assess the extent of

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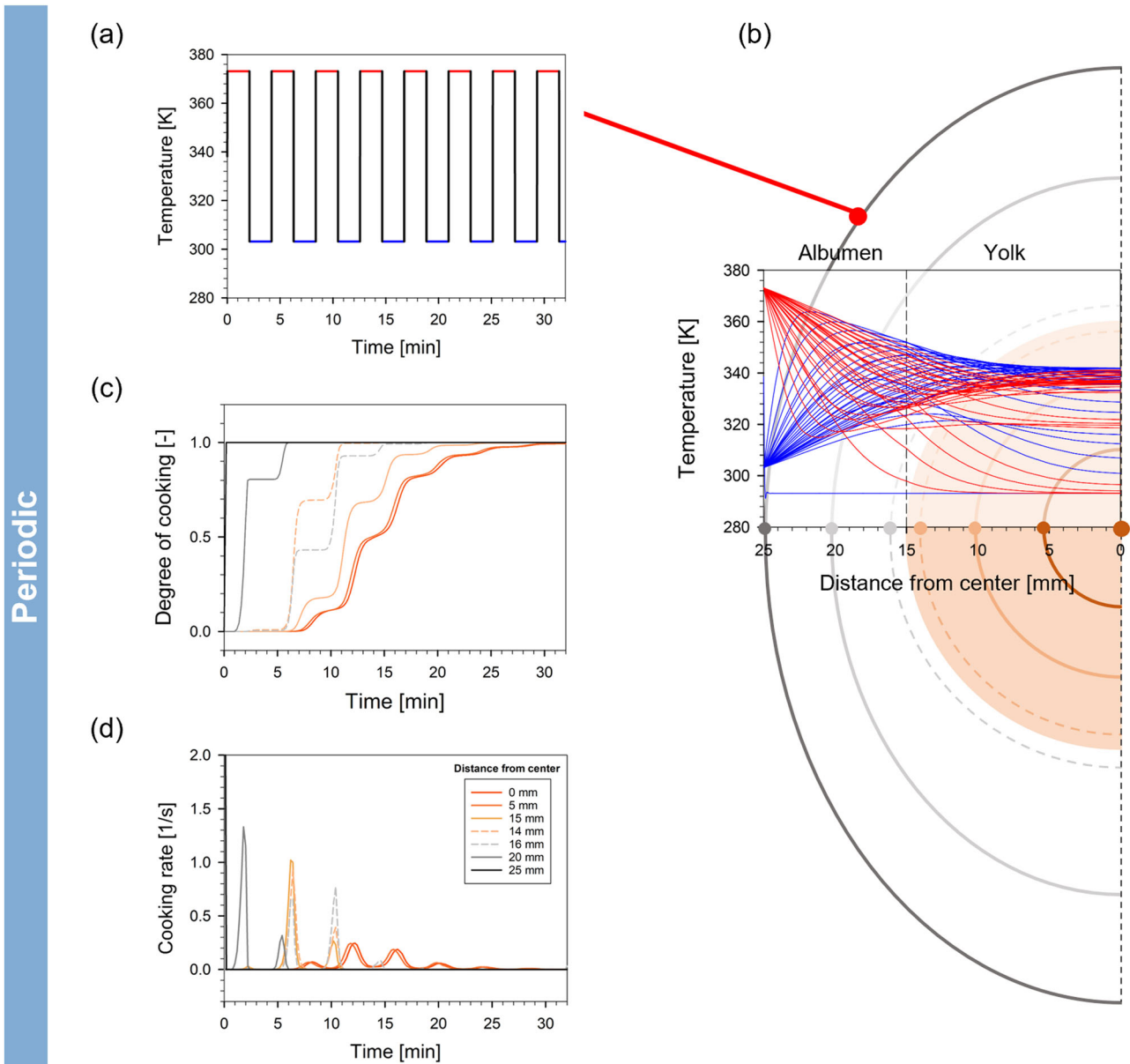


Fig. 1 | Simulation of periodic cooking. Results of the simulation of the cooking of an egg with the periodic cooking method: **a** periodic time-varying BC imposed, **b** evolution of the thermal profile over time, **c** evolution of the degree of cooking over time at different distances from the center of the egg and **d** evolution of the

cooking rate over time at different distances from the center of the egg. The distances from the center selected to construct the graphs of figures **c** and **d** are identified by the lines in figure **b**. The precise legend for figures **c** and **d** is given only in figure **d**.

protein thermal denaturation and to analyze the nutritional profile of the eggs, respectively.

Overall, we found that our cooking method leads to improved texture and nutritional content with respect to traditional shell-on egg cooking techniques, thus elevating the already established concept by which temperature and time have a critical role in the resulting properties of egg parts. The potential of this approach beyond cooking, with possible uses in curing, crystallization, and material structuring, is also foreseen.

Results and discussion

Cooking simulations

Results of the simulation of the cooking of an egg are here shown. Four different cooking techniques were simulated: periodic cooking (Fig. 1), hard-boiling (Fig. 2a, b), soft-boiling (Fig. 2c, d) and sous vide cooking (Fig. 2e, f). For all cooking techniques we show the evolution of the thermal profile over time and distance from the center of the egg, and the

evolution of the degree of cooking over time and distance from the center of the egg.

In the case of hard-boiled and soft-boiled eggs, the temperature grows monotonically over time (Fig. 2a, c). The different cooking time (12 min vs. 6 min, respectively), though, causes differences in the final thermal profile: hard-boiled eggs are at 100 °C in all their parts at the end of the 12 min cooking, while soft-boiled eggs show a lower and non-uniform temperature in almost all their parts at the end of the 6 min cooking, except for the parts adjacent to the shell. This different cooking time also causes a different cooking degree: in hard-boiled eggs the cooking degree reaches the unit in both albumen and yolk (Fig. 2b), while in soft-boiled eggs this happens only in the albumen. On the other hand, in the yolk the cooking degree remains lower, down to a minimum of 0.5 in the center of the egg (Fig. 2d). This is compatible with the cooking outcome of hard-boiling and soft-boiling, since soft-boiled eggs typically show a much runnier yolk compared to hard-boiled eggs.

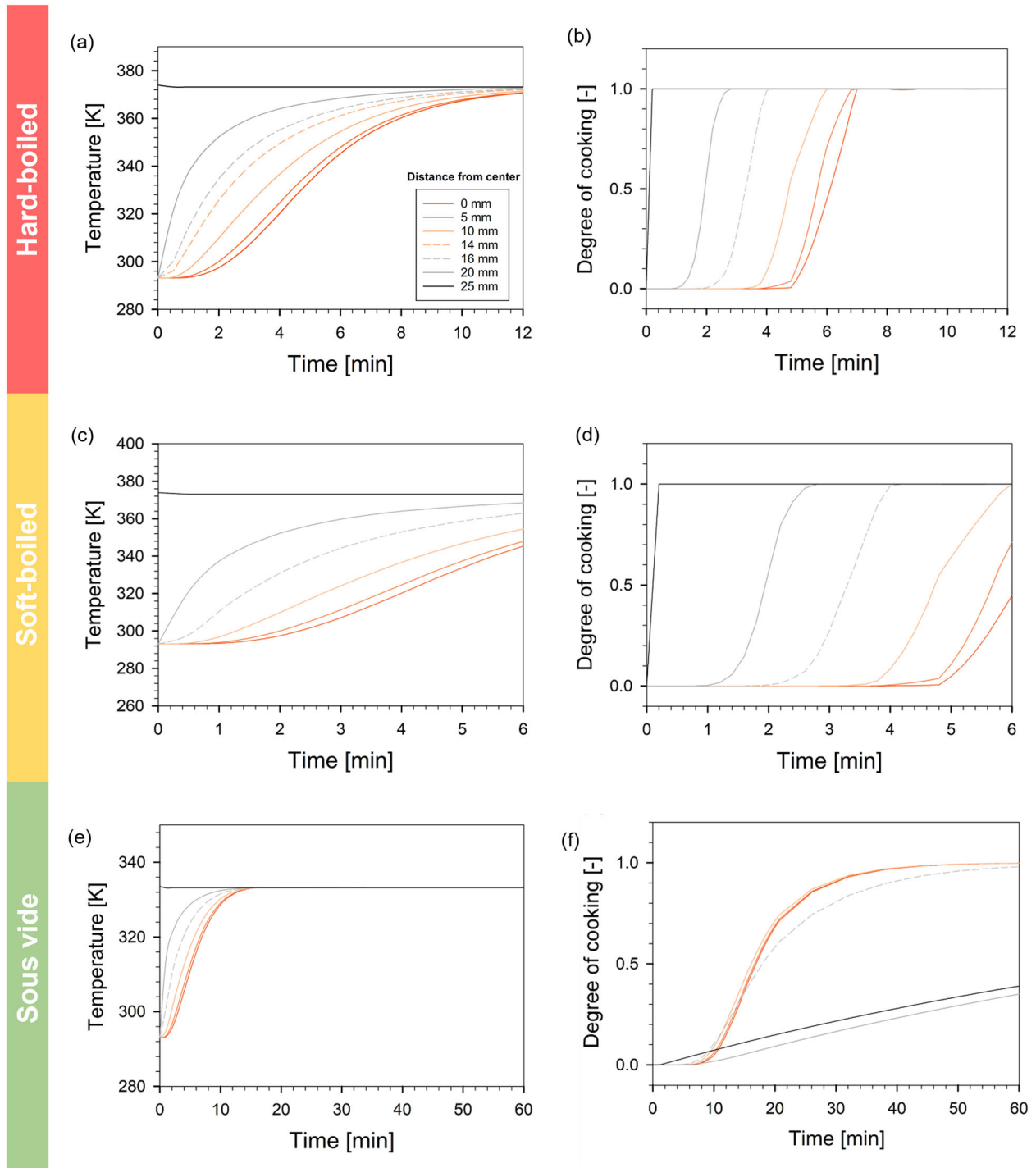


Fig. 2 | Simulation of hard-boiling, soft-boiling and sous vide cooking. Results of the simulation of the hard-boiling (red), soft-boiling (yellow) and sous vide cooking (green) of an egg: **a, c, e** temperature values over time at different distances from the

center of the egg; **b, d, f** degree of cooking values over time at different distances from the center of the egg. The distances selected to construct these graphs are identified by the lines in Fig. 1b. The precise legend for all the figures a–f is given in figure a.

Similar results were found for the sous vide egg in terms of the evolution of temperature over time (Fig. 2e): it grows monotonically until a plateau is reached at 65 °C in all parts of the egg. Nevertheless, the use of a lower cooking temperature causes differences in the final degree of cooking (Fig. 2f): unit is reached inside the yolk, but not inside the albumen, because the egg albumen proteins do not denature and aggregate at such low temperatures. This effect is visible in the simulation thanks to the kinetics equation that we used to model the gelation process.

Very different but satisfying results are obtained from the simulation of the periodic cooking technique (Fig. 1). As expected, temperature

inside the egg does not grow monotonically, in accordance with the imposed BC, but a stationary state at the center of the yolk is reached as well (Fig. 1b): while the albumen alternatively sees temperatures in the range 100–87 °C and 30–55 °C during the hot and cold cycles respectively, the yolk sees a constant temperature of 67 °C, which is around the mean value between T_h and T_c , as predicted. This peculiar thermal profile allows for optimal cooking of the egg in all its parts, and this is confirmed by Fig. 1c in which it is clear that a degree of cooking equal to 1 is reached both in egg yolk and albumen, differently from what happens with the sous vide cooking.

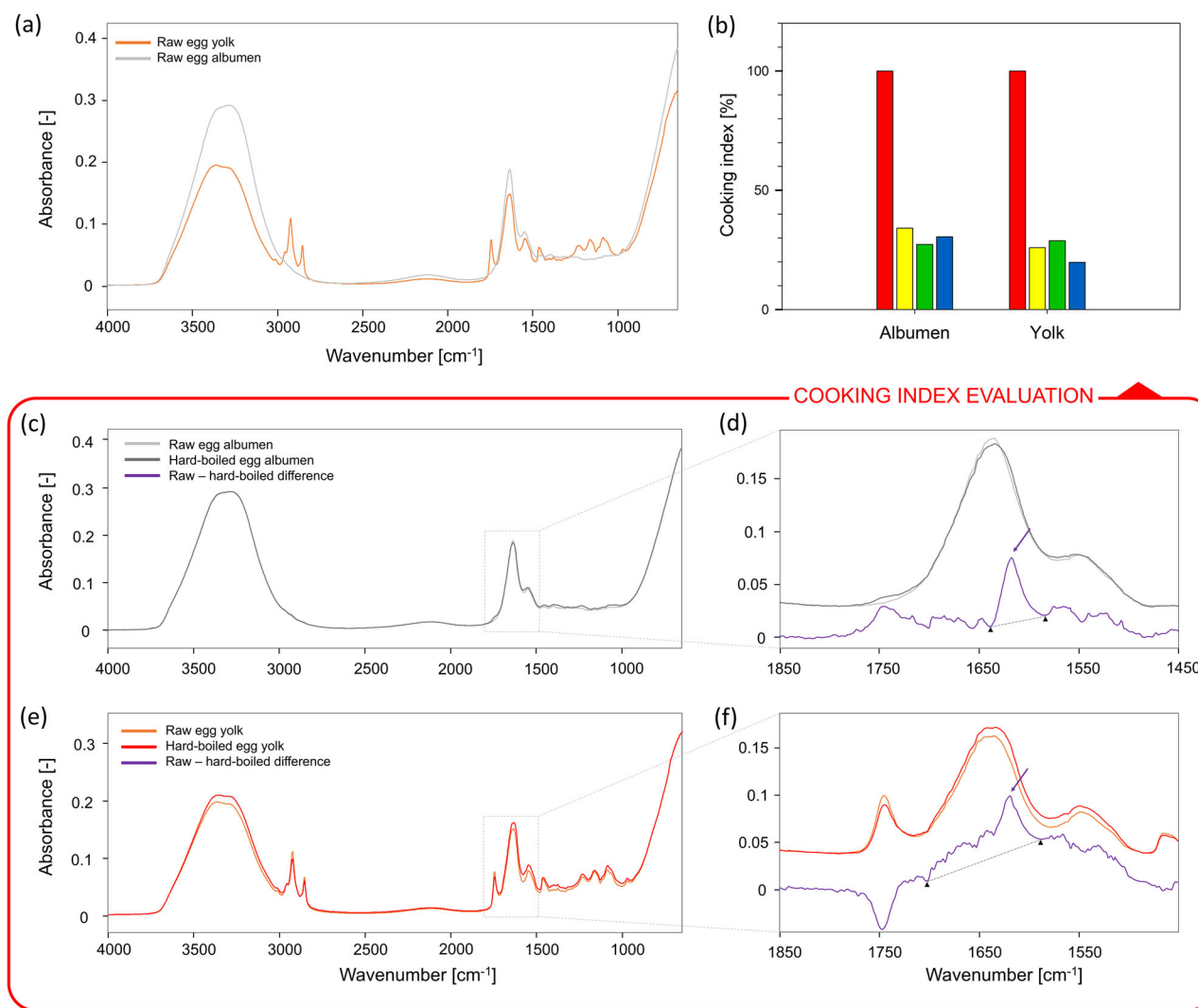


Fig. 3 | Overview of the spectroscopic analysis conducted on raw and cooked eggs. **a** IR spectra of raw egg albumen (light gray) and yolk (orange) over the whole IR range (4000–650 cm^{-1}). **b** Cooking index derived through the FT-IR spectroscopic analysis for all cooking techniques in the case of egg albumen and yolk. The different colors of the bars evidence the different cooking methods: hard-boiling (red), soft-boiling (yellow), sous vide cooking (green) and periodic cooking (blue). **c** IR spectra of raw and hard-boiled albumen (light and dark gray, respectively) and

(e) of raw and hard-boiled yolk (orange and red, respectively) over the whole IR range. A close up on band no. 4 and 5 (amide I region) is displayed for both albumen (d) and yolk (f) with addition of the difference spectrum magnified $\times 5$ (purple) and the baseline used for area quantification in the Amide I region (black dashed line). The purple arrow indicates in both cases the appearance of the aggregation band.

Finally, Fig. 1d shows the evolution of the cooking rates in albumen and yolk. These curves present various peaks over time at all distances from the center, which are clearly linked to the BC imposed: during the cold cycles ($T = T_c$), interruption of the cooking causes the cooking rate to be equal to zero. A final plateau of null cooking rate is eventually reached everywhere when the cooking degree reaches the unit, and this happens at different times depending on the distance from the center. In the case of hard-boiling, no such peaks are detected, but an equal plateau of null cooking rate is reached when the cooking degree reaches the unit, coherently with the BC imposed. A similar evolution of the cooking rate is expected for all other cooking techniques using a constant BC (soft-boiling and sous vide). Details are shown in the Supplementary Information (Supplementary Section A.2 - Cooking simulations, Supplementary Fig. A2).

FT-IR spectroscopic analysis

FT-IR spectroscopy was performed on raw and cooked eggs to verify protein thermal denaturation and aggregation, which knowingly leads to thickening and gelling¹². IR spectra collected for raw egg albumen and yolk are

presented in Fig. 3a. Here, functional groups of lipids, proteins and water components can be detected. The complete list is given in Table 1.

Bands no. 2, 3, 4, 7, 8, 9 and 10 are associated with functional groups of the lipid component. These can be easily detected in the yolk spectrum, while they are not present in the albumen spectrum because egg albumen does not contain fats. Bands no. 5 and 6 represent the amide I and amide II region, respectively. They are attributed to the secondary structure of proteins and they are visible in the spectra of both egg phases, since proteins are present in both albumen and yolk. Finally, band no. 1 is the water band and has a wide range in the spectra of both albumen and yolk.

Amongst these, the amide I region (band no. 5), more resolved in the IR range with respect to the amide II region, is the most important region in our study, because it has a huge potential to give information on the secondary structure of egg proteins. The band at 1636 cm^{-1} , in particular, is the most relevant because it is characteristic of the amide groups involved in the β -sheet structure, which in turn are linked to the denaturation of the proteins: an increase in the β -sheet structure at the sacrifice of the helical structure is interpreted as a result of heat denaturation⁵. It is noted that in some works the characteristic frequency of intermolecular β -sheets is found at

Table 1 | Band assignment for FT-IR spectra of raw eggs

Band number	Wavenumber [cm ⁻¹]	Functional group	Mode of vibration
1	3000–3700	-O-H (H ₂ O)	Stretching
2	2920	-C-H (CH ₂)	Stretching (antisymmetric)
3	2849	-C-H (CH ₂)	Stretching (symmetric)
4	1743	-C=O (ester)	Stretching
5	1590–1720	-C-O, -C-N	Stretching
6	1541	-N-H, C-N (H ₂ O)	Bending, stretching
7	1462	-C-H (CH ₂ , CH ₃)	Bending (scissoring)
8	1230	-C-O, -CH ₂ -	Stretching, bending
9	1159	-C-O, -CH ₂ -	Stretching, bending
10	1085	-C-O	Stretching

Wavenumbers of FT-IR bands of the egg samples with the assigned functional groups and modes of vibration¹.

1620 cm⁻¹^{4,10}. In general, the evaluation of the band at 1636/1620 cm⁻¹, often called “aggregation band”, is a good diagnostic tool for monitoring thermal unfolding of the proteins. It is in our interest, then, to see how the aggregation band changes with different cooking methods, so that we can understand how different cooking methods can affect the level of protein unfolding, i.e., protein denaturation and aggregation and consequent egg texture.

An example of the outcome of such measurement in the specific case of hard-boiled egg yolk and albumen is shown in Fig. 3c, e, completed with a close up on the regions of interest and the corresponding difference spectrum (purple) (Fig. 3d, f). The appearance of the aggregation band, indicating β -sheet formation, is highlighted in both cases with a purple arrow. Details on all the IR spectra collected (egg albumen and yolk cooked with all the different cooking techniques) can be found in the Supplementary Information (Supplementary Section A.2 - FT-IR spectra, Supplementary Figs. A3–A10).

From these spectra, evaluation of a Cooking index, according to the procedure described in the Supplementary Information (Supplementary Section A.1 - Analysis of FT-IR spectra), was possible and this helped understand the different extent of protein denaturation with the different cooking techniques. Results are shown in Fig. 3b for both yolk and albumen.

In the case of yolk, the Cooking index is maximum for hard-boiling (as imposed), followed by, in order, sous vide, soft-boiling and periodic cooking. This is linked to the type of thermal treatment and matches the different textures and consistencies of the cooked products: hard-boiled egg yolk is a solid paste, while sous vide, soft-boiled and periodic egg yolks are gel-like substances with an increasing tendency to flow.

In the case of albumen, the Cooking index is maximum for hard-boiling, followed by, in order, soft-boiling, periodic cooking and the sous vide. Again, this is linked to the type of thermal treatment and matches the different textures: hard-boiled egg yolk is fully set, while this isn't true for soft-boiled, periodic and sous vide. The sous vide albumen, in particular, is completely runny and cooking appears to be not sufficient due to lack of reaching the appropriate albumen cooking temperature (85 °C). This problem appears to be overcome with the periodic cooking technique because the use of the cycles helps to reach a higher temperature in the albumen, allowing for further cooking and setting, while still maintaining the creamy consistency of the yolk. This detail is what makes it possible to have a runny, creamy yolk with a nicely set albumen when using the periodic cooking technique, differently from what happens with other cooking methods. In other words, spectroscopy confirms the simulation results.

Quantitative descriptive analysis

The quantitative descriptive analysis is an objective sensory analysis that allows to define with greater accuracy and detail, with respect to what the

consumer is able to do, the sensory profile of a product. Results of this analysis are shown in Fig. 4, where specific significant differences are highlighted.

As it is visible, the main differences between the 4 different products concern not only texture-related properties, but also several characteristics related to taste area.

Compared to the periodic, both the albumen and yolk of the hard boiled sample differ mainly for its less wet, more adhesive and more powdery/sandy consistency when pressed between tongue and palate. When analyzing the taste, the egg albumen is sweeter and more stringent, the yolk is less sweet and both egg albumen and yolk have less umami taste (although it is perceived at a weak intensity).

The soft boiled sample, differently from the periodic, has a shinier/brighter surface of the albumen and it is drier in the mouth and less sweet. The yolk is less dense/bodied and wetter, less sweet as well as less salty.

Finally, compared to the periodic sample, the sous vide albumen is shinier and clearer/transparent. It is also definitely softer, wetter and more soluble during tasting. On the other hand, the yolks are very similar to each other and no significant differences emerge.

In conclusion, the periodic egg is more similar to the soft boiled when analyzing the texture of its albumen, while it is very similar to the sous vide sample when considering its yolk. This confirms the effectiveness of periodic cooking in delivering two very different textures and tastes in the egg albumen and yolk, respectively: the yolk is most similar to that cooked at a constant temperature of 65 °C, while the albumen is most similar to that cooked at 100 °C, in full agreement to simulation results and expectations.

Texture profile analysis

TPA parameters of both egg albumen and yolk are presented in Table 2. In general terms, the biggest differences are found in hardness (also displayed in the box plot of Fig. 5) and chewiness (or gumminess).

In the albumen, hardness is higher in the hard-boiled sample and progressively decreases in the soft-boiled, sous vide and periodic samples, while chewiness is higher in the hard-boiled and soft-boiled samples, and progressively decreases in the periodic and, most significantly, in the sous vide sample. This is consistent with the results of the sensory evaluation, which shows the same trend in the values of softness, wetness and meltability.

In the yolk, hardness is higher in the hard-boiled sample and progressively decreases in the periodic, soft-boiled and sous vide samples, while chewiness (here partially gumminess because most of the samples are semi-solids) is higher in the hard-boiled sample and progressively decreases in the soft-boiled, periodic and sous vide samples, the latter bearing more similarities with each other. Again, this is consistent with the results of the sensory evaluation, which shows that the highest difference in terms of texture (density/body, wetness, solubility, pastiness, adhesiveness, powderiness) is found in the hard-boiled sample, while the remaining are more similar to each other.

Overall, TPA confirms that different cooking methods are able to deliver different textures, but, unlike the sensory analysis, it is less capable of highlighting specific differences between the soft-boiled, sous vide and periodic samples probably due to sampling complexity.

¹H-NMR and UHPLC Q-Orbitrap HRMS based analyses

To evaluate the effect of the various cooking methodologies proposed in this study on the nutritional profile of the eggs, an untargeted study has been performed on the albumens and yolks by using ¹H-NMR spectroscopy. The samples were prepared and the spectroscopic data processed as described in the Supplementary Information (Supplementary Section A.1). All the NMR spectra were analyzed by Principal Component Analysis (PCA) to specifically explore the differences in albumens and yolks across the investigated class. The primary purpose of PCA is to reduce the complexity of a dataset containing many interrelated variables, allowing for a visual representation of the main sources of variance in the data. PCA achieves this by transforming the original correlated variables into a smaller set of uncorrelated

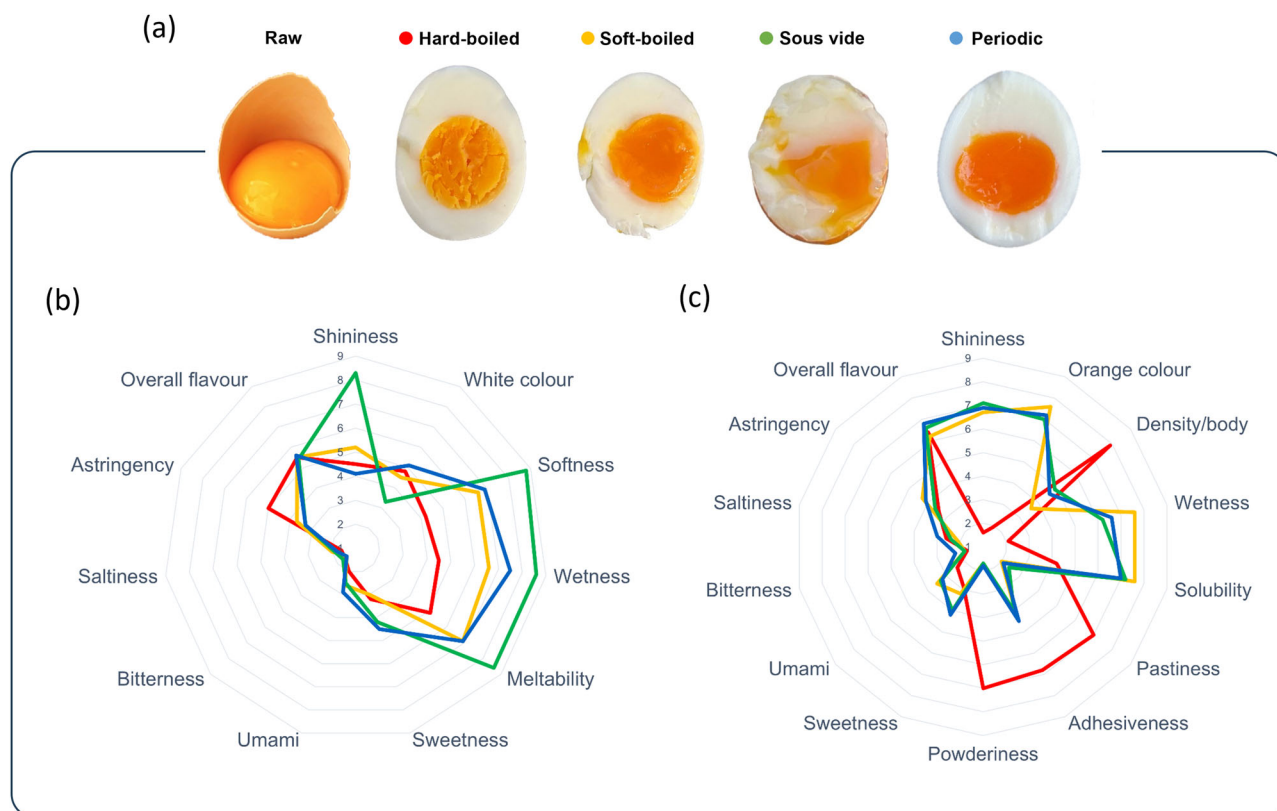


Fig. 4 | Sensory analysis evaluation. **a** Photographs of the raw, hard-boiled (red), soft-boiled (yellow), sous vide (green) and periodic (blue) eggs. Results of the Sensory Analysis performed on **b** albumen and **c** yolk. Significant differences ($p = 95\%$) are identified between: hard-boiled and periodic yolk (in Astringency, Softness, Wetness, Meltability, Sweetness and Umami); soft-boiled and periodic yolk (in Shininess, Wetness and Sweetness); sous vide and periodic yolk (in Shininess, White

color, Softness, Wetness and Meltability); hard-boiled and periodic albumen (in Shininess, Orange color, Density/body, Wetness, Solubility, Pastiness, Adhesiveness, Powderiness, Sweetness and Umami); soft-boiled and periodic albumen (in Density/Body, Wetness, Sweetness and Saltiness). No significant differences are found between sous vide and periodic yolk.

variables called principal components (PCs). These PCs form the axes of a new coordinate system, with the first principal component (PC1) representing the highest variance, the second (PC2) representing the next highest, and so on. PCA results are typically displayed in two types of plots: a “scores plot”, which shows groups samples based on their similarity, and a “loadings plot”, which highlights the variables that contribute to the differences between samples along each principal component. In this study, however, PCA results are presented using a biplot (Fig. 6), which combines the scores plot and the loadings plot into one. This approach facilitates the visualization of both samples and variables within a single plot, improving interpretability. Notably, PCA biplot of the $^1\text{H-NMR}$ -based metabolomic data (Fig. 6a) revealed that, consistent with previous works, PC1, accounting for 84.31% of the total variation, clearly distinguished yolks from albumens¹³. In addition, the distribution of the variables on the PCA biplot suggested that most of the detected amino acids were more abundant in the yolk compared

to the albumen (left side of the PCA biplot reported in Fig. 6a). Conversely, albumens were richer in formate and sugars (right side of the PCA biplot reported in Fig. 6a).

Among yolk amino acids there are leucine (Leu), isoleucine (Ile) and valine (Val), known as branched-chain amino acids (BCAAs). BCAAs, which are primarily metabolized in brain, kidney and muscle, can provide nitrogen groups for the nitrogen-containing substances biosynthesis (e.g. creatinine, creatine, glutathione, pyridine and carnitine) and are recognized as gluconeogenic amino acids, providing a source of glucose¹⁴. Moreover, isoleucine can promote myogenesis and intramyocellular fat deposition, thereby increasing muscle mass¹⁵. BCAAs also play a crucial role in psychological disorders, with studies showing an inverse relationship between dietary BCAAs and the likelihood of depression and anxiety¹⁶.

Concerning albumen composition, it’s worth to notice that lysine is an essential amino acid involved in the synthesis of crucial proteins such as nucleoproteins, and hemoglobin. Lysine (Lys) also promotes bone growth and development in children enhancing calcium absorption and increasing appetite¹⁷. In addition, albumens are richer in tryptophan, an essential amino acid that cannot be synthesized by living organisms and needs to be obtained from daily diet. Tryptophan, an aromatic amino acid, is key element for brain functioning, due to its role as a precursor for the neurotransmitter serotonin. Lower levels of tryptophan are often associate with depressive mood¹⁸. Therefore, the consumption of both egg yolk and albumen can provide more energy and may help prevent brain diseases¹⁹.

Interestingly, PC2 (12.02% of total variance) shows a separation of the egg samples according to the cooking method (Fig. 6a). Specifically, both the yolks and albumens cooked using the periodic method

Table 2 | TPA parameters of cooked egg albumen and yolk

	Albumen				Yolk			
	HB	SB	SV	P	HB	SB	SV	P
Hardness	14.37	9.46	1.33	4.73	22.13	3.04	1.59	4.34
Cohesiveness	0.30	0.47	0.56	0.57	0.45	0.67	0.78	0.55
Springiness	0.30	0.40	0.57	0.38	0.19	0.47	0.55	0.32
Gumminess	–	–	0.90	–	–	1.85	1.08	2.20
Chewiness	1.34	1.89	–	1.03	1.84	–	–	–

Instrumental texture measurements in egg albumen and egg yolk cooked with the 4 different methods: Hard-boiled (HB), Soft-boiled (SB), Sous vide (SV) and Periodic (P). Data are evaluated as the mean of 4 replicates.

(blue samples) are positioned in the upper part of the plot, well separated from the samples cooked using other methods. This suggests that the periodic cooking approach affects the egg metabolome in a peculiar way, deserving further investigation through Mass Spectrometry (MS)-based analysis. Particularly, we decided to quantify polyphenols and amino acids only in the yolk, which is known to contain the vital nutrients that characterize the egg.

The PCA model obtained with the UHPLC Q-Orbitrap HRMS data is reported in Fig. 6b.

Excitingly, the PCA biplot showed that the separation between the periodic and all the remaining groups is even more pronounced compared to that observed in the NMR-based analysis. Interestingly, the periodic yolk group is characterized by higher content of all the investigated polyphenols (the most abundant and broadly distributed class of bioactive molecules) compared to the other samples. The association between polyphenol consumption and human health has been explored²⁰, and it has been observed that a diet rich in polyphenols appears to be protective and prevent the onset

of several diseases. Particularly, flavonoids are the main class of polyphenols in the analyzed extracts. It's worth to notice that among the detected flavonoids there is an isoflavone named daidzein, whose concentration in eggs depends on the diet of laying hens. It is also used as a dietary supplement for laying hens. This compound has gained popularity as a dietary supplement, especially for animals in the post-estrogenic period, offering a natural and safe alternative to estrogen-like compounds²¹. Liu et al.²² assessed that daidzein improved egg quality by controlling lipid metabolism in layers and enhancing the antioxidant capacity of egg yolks. Moreover, isoflavones like daidzein are biologically active compounds with estrogenic properties, that may have a modest beneficial effect for human health, particularly on menopausal symptoms and may reduce the incidence of prostate and breast cancer²⁰. In addition, the investigation identified also the presence of some phenolic acids among the extracts' polyphenols, such as the ferulic and chlorogenic acid.

Overall, these results strongly suggest that the periodic cooking method have a better advantage over conventional cooking methods in terms of nutritional content.

Conclusions

It appears that the design of the novel cooking method, namely periodic cooking, was carried out successfully. Designing of the process and subsequent simulation proved fundamental in the understanding of the physics behind the cooking of an egg and allowed us to finely tune the process

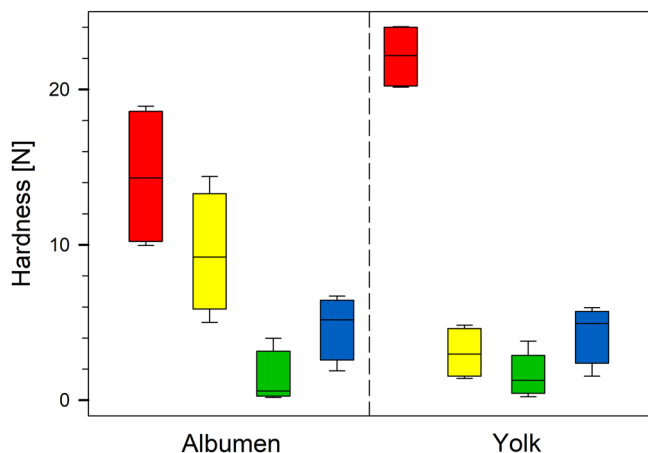


Fig. 5 | Hardness of egg albumen and yolk as measured with TPA. Hardness evaluated through TPA for albumen and yolk. The color legend is the same of Fig. 4 and indicates the different cooking techniques: hard-boiled (red), soft-boiled (yellow), sous vide (green) and periodic (blue). Each box of the box plot displays (bottom to top) minimum, first quartile, median, third quartile, and maximum.

Table 3 | Acronyms of metabolomic analysis

Asp	Aspartic acid	Met	Methionine	Arg	Arginine
Phe	Phenylalanine	TPC	Total Phenolic Content	QA	Quinic acid
Lac	Lactate	Asn	Asparagine	Pro	Proline
Gln	Glutathione	Trp	Thryptophan	Ala	Alanine
Val	Valine	For	Formate	Seco	Secoisolariciresinol
Glu	Glutamic acid	Tyr	Tyrosine	Hesp	Hesperidin
Ile	Isoleucine	Ser	Serine	Lys	Lysine
Leu	Leucine	Thr	Threonine	DPPH, ABTS	Antioxidant activity assays

Full list of acronyms used in Fig. 6.

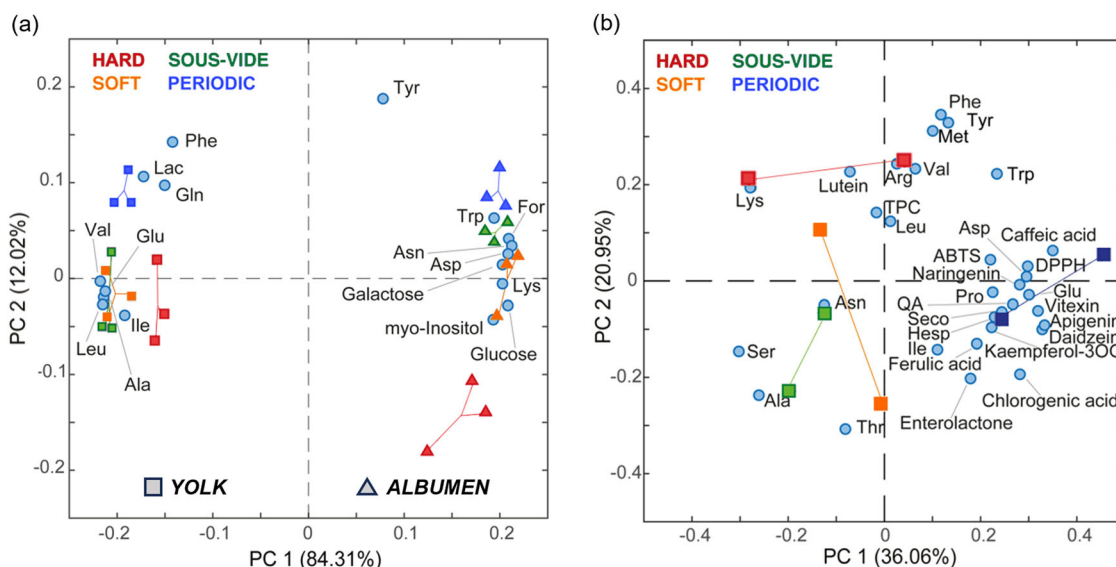


Fig. 6 | Nutritional aspects of cooked eggs. PC1 vs. PC2 biplot of the PCA model calculated using (a) both albumen and yolk extracts analyzed by ¹H-NMR and (b) yolk extracts analyzed by MS analysis and spectrophotometric assays. Acronyms used in the graphs are reported in Table 3.

parameters with respect to our objective, i.e., having two different, optimal, temperatures in the two phases of the egg. Simulated results proved to be consistent with the results of cooking experiments, and the FT-IR spectra measurements corroborated them, proving the different extent of protein denaturation (and consequent thickening and gelation) with different cooking methods (namely, hard-boiling, soft-boiling, sous vide and periodic cooking). Analysis of the color, texture, and consistency of all the egg products was only the final proof of a successful cooking experiment that might inspire new fancy recipes, proving how knowledge of the science behind simple problems can improve even the slightest bits of our daily life, like the simple act of eating an egg. Studying the nutritional profile of such eggs only helped validating this final statement, as periodic cooking clearly stood out as the most advantageous cooking method in terms of egg nutritional content. An even higher impact on human diet is here implied: not only we are able to reach a perfectly diverse texture in a two-phase food product, but we are also able to preserve, in both these phases, a higher nutrient amount, providing a useful tool to boost poor dietary habits. The possibility of exploiting this process outside the kitchen realm is also foreseen: curing, crystallization, and structuring of materials are only a few of the applications for which a thoroughly designed periodic thermal treatment is at hand.

Materials and methods

Materials

Fresh hen eggs were purchased at a local supermarket (Naples, Italy) and stored at room temperature in a cool and dry place. These were used for all experiments except for Sensory Analysis. Fresh hen eggs for the Sensory Analysis were provided by Ovomont S.r.l., Castelvetere sul Calore (AV), Italy.

Methods

Egg cooking methods. Cooking experiments of shell-on eggs were performed with a heater and a kitchen pan. The pan was filled with tap water and placed on the heater to reach the boiling or the desired temperature. Constant monitoring of the water temperature was possible thanks to a food thermometer, which was immersed in water as the cooking proceeded. Four different cooking techniques were tested, namely hard-boiling, soft-boiling, sous vide or 6X °C and periodic cooking. Hard-boiled eggs were placed in boiling water for 12 min; soft-boiled eggs were placed in boiling water for 6 minutes; sous vide eggs were placed in water at 65 °C for 1 h and periodic eggs were placed alternatively in boiling water ($T_h = 100$ °C) for $t_h = 2$ min and water at $T_c = 30$ °C for $t_c = 2$ min, for a total cooking time of 32 minutes, which corresponds to the repetition of the hot and cold cycles for a total of $N = 8$ times. In the case of periodic eggs, a bowl filled with water kept at 30 °C was used for the cold cooking cycle.

After each cooking experiment, the eggs were cooled down under running water and then the shell was cracked open. Pictures of a cross section of each egg were taken (See Fig. 4a).

Mathematical modeling and simulation. To model the heat transfer phenomenon and concomitant thickening and gelling of egg yolk and albumen during cooking of the egg in hot water, the following assumptions were made:

1. Egg albumen and yolk are homogeneous and isotropic during cooking;

2. The initial temperature is constant and the same in all eggs ($T = 20$ °C);
3. Thermal conductivity and density of both egg phases are a function of temperature;
4. Presence of an air cell, natural convection and moisture transfer inside the egg are neglected. The derived model is here presented:

$$\rho_i c_{p_i} \frac{\partial T_i}{\partial t} + \nabla \cdot (-k_i \nabla T_i) = 0 \tag{1}$$

$$\frac{\partial X_i}{\partial t} = R_i \tag{2}$$

Equation (1) is the conduction only energy transfer equation, while Eq. (2), coupled with it, describes the evolution of the degree of cooking $X(t)$ over time. R_i represents the gelation rate and is thus expressed:

$$R_i = A_i e^{\frac{-E_{a_i}}{RT_i}} (1 - X_i) \tag{3}$$

Where ρ_i , c_{p_i} and k_i are the thermal properties of egg yolk and albumen and $\frac{1}{\tau_i} = A_i e^{\frac{-E_{a_i}}{RT_i}}$ is the rate constant of the gelation process, whose dependence on the absolute temperature is given by the Arrhenius equation, A_i and E_{a_i} being the pre-exponential factor and the activation energy, respectively. More specifically, all the parameters used in the system, both thermal properties and kinetics data, are listed and explained in Table 4. It is evident that two different sets of parameters are used for the yolk and the albumen, respectively. This is due to their different composition and the different denaturation processes happening in the two phases.

From a quick analysis of this system we are able to provide an estimate of the parameters of our cooking method. From the values of thermal conductivity, specific heat and density, we can find, for both albumen and yolk, thermal diffusivities $a_i = \mathcal{O}(10^{-5} \text{ m}^2 \text{ s}^{-1})$, and, considering a length scale $L = \mathcal{O}(10^{-2} \text{ m})$, the characteristic time for heat transport $\mathcal{O}(10^1 \text{ s})$, which gives us an idea of t_h and t_c . T_h is the optimal cooking temperature for the albumen, while T_c is chosen such that the mean between these two values is the optimal cooking temperature for the yolk (in case $t_h = t_c$). Finally, the number of cycles N is related to the kinetic constants, being the largest between $\frac{\tau_{\text{yolk}}}{t_h}$ and $\frac{2 \cdot \tau_{\text{albumen}}}{t_h}$. A first design of the novel cooking method was made exploiting these considerations.

Simulation with a CFD program was then used for a finer resolution of the problem for all the cooking techniques previously listed (hard-boiling, soft-boiling, sous vide and periodic cooking). The geometry of the egg was designed as a 2D axisymmetric oval with 7 and 5 cm axis and with a sphere (radius 1.5 cm) in the geometric center as the yolk. The mesh was Mapped at the edges, where higher changes are expected, and Free triangular elsewhere. Initial conditions were $X_i(t = 0) = 0$ and $T(t = 0) = 20$ °C for all simulations. As for the BCs, a no flux condition was imposed in all simulations, while three different BCs were used on temperature to be able to simulate all the cooking techniques ($T = 100$ °C for hard-boiling and soft-boiling, $T = 65$ °C for sous vide cooking and Fig. 1a for periodic cooking).

The system of differential equation was solved by the “finite element” method in the “Direct UMFPAK” mode at time intervals of 0.5 min. The total process time was 12 min for hard-boiled eggs, 6 min for soft-boiled eggs, 1 h for sous vide eggs and 32 min for periodic eggs.

This simulation can be easily adapted to the variations in the quality and size of the egg. An estimation of the corresponding changes is quickly

Table 4 | Physical properties and kinetics data of eggs

	k_i [$\text{Wm}^{-1}\text{K}^{-1}$]	ρ_i [kgm^{-3}]	c_{p_i} [$\text{Jkg}^{-1}\text{K}^{-1}$]	A_i [s^{-1}]	E_{a_i} [J]
Yolk	$0.00087 + 0.395$	$-0.0023T^2 - 0.1386T + 1037.3$	3120	$2.72 \cdot 10^{50}$	$3.443 \cdot 10^5$
Albumen	$0.00137 + 0.5125$	$-0.00417^2 - 0.01157 + 1043.3$	3800	$4.85 \cdot 10^{50}$	$4.185 \cdot 10^5$

Physical properties (thermal conductivity k_i , density ρ_i , specific heat c_{p_i}) and kinetics data (pre-exponential factor A_i and activation energy E_{a_i}) of both yolk and albumen. Data were taken from ref. 23 and ref. 12.

given by the scaling arguments mentioned above: the characteristic time for heat transport (L^2/a) expresses the relationship with the characteristic dimension of the object (L), while T_h , T_c and the kinetic constants express the relationship with egg quality.

Characterization techniques. To characterize the cooked products obtained with all the cooking methods hereby examined (hard-boiling, soft-boiling, sous vide cooking and periodic cooking), we have used a series of characterization techniques, here mentioned:

- FT-IR spectroscopy, to assess the extent of protein denaturation;
- TPA, to gather information on the texture of egg albumen and yolk;
- Quantitative Descriptive Analysis, to get insights on color, consistency, texture and taste of the cooked products;
- Metabolomic Analysis (specifically $^1\text{H-NMR}$ and HRMS), to investigate the nutritional profile of the eggs.

Moreover, we also carefully verified that in periodic cooking the thermal load was sufficient to avoid microbiological issues. All the details about the characterization techniques and the microbiological considerations can be found in the Supplementary Information (Supplementary Subsection A.1).

Data availability

The data that support the findings of this study are available online on figshare at 10.6084/m9.figshare.26397538.

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Competing interests

The authors declare no competing interests.

Additional information

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