Thermal degradation of biological DNA studied by dielectric spectroscopy

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A R T I C L E   I N F O
Keywords:
DNA
Deoxyribonucleic acid
Melting
Dielectric spectroscopy
Thermal stability

ABSTRACT
Dielectric spectroscopy was tested as an alternative tool to study degradation of deoxyribonucleic acid (DNA) in its solid form. The specimens, prepared from biological DNA, were periodically heated and cooled according to a programmed scheme. Simultaneously, their dielectric parameters (permittivity and dielectric loss) were monitored as a function of frequency and temperature. The analysis of Bode plots allowed to determine the upper limit of thermal stability of solid DNA at 120 °C, because heating at higher temperatures resulted in irreversible changes. These changes were identified as denaturation by gel electrophoresis and UV-vis absorption methods.

1. Introduction

The deoxyribonucleic acid (DNA) is a macromolecule of fundamental importance for all living organisms. Many unique chemical and physical properties distinguish DNA from synthetic polymers, due to its ability to maintain the structure of the double-helix.

This fact, since the beginning of the last decade, has motivated the search for new applications in fields outside of the life sciences. In such applications, DNA is used as a bulk material, in its natural form or in a complex with an amphiphilic (surfactant) molecule. These materials in the form of thin layers have been added to the structure of devices classified as organic electronics, optoelectronics and non-linear optics. As a consequence, a significant improvement in performance was observed. An extensive discussion of non-standard DNA applications can be found in recent review articles, for example in [1,2].

DNA is a highly hydrophilic biopolymer [3,4], what directly impacts its morphology [5,6]. In the case of DNA complexes, aliphatic (and so hydrophobic) tails of surfactant molecules extend to outermost directions, making possible to dissolve the complex in some organic solvents [7]. However, even this hydrophobic cocoon does not protect entirely the DNA backbone from water molecules penetrating from the surrounding atmosphere [8–10]. Water molecules confined in either DNA or DNA complex belong to a one of two pools, according to the strength and the character of the binding mechanism. The first shell, referred to as tightly bound water, consists of molecules directly bound via hydrogen bonds to hydrophilic sites on the DNA helix. To the other one, known as loosely bound water, belong water molecules attached to tightly bound waters [11]. These waters can be the source of uncontrolled ionic conduction that may occur extremely detrimental for performance of organic electronics devices. A range of techniques is used to dry DNA, like for example freeze-drying or prolonged equilibration in dry atmosphere. However, even prolonged drying under vacuum in presence of P₂O₅ is insufficient to entirely dry DNA and leaves a few weight percent of water [12]. Thin films of DNA for organic electronics are processed from solution. Such films, even if initially dried, have to be transferred to an evaporating system, where other layers are deposited. During the transfer, the DNA layer risks an extra contamination with atmospheric humidity. For this reason, the film is heated inside the evaporator to remove residual water or solvent. The question arises, what is the actual temperature, that DNA is able to withstand without irreversible changes concerning its principal morphology. The double helix of DNA, where planar base pairs are arranged in equidistant stacks, is believed to determine electrical charge transport [13]. The double helix structure is also vital for effective hosting dye molecules used in DNA-based nonlinear optics [14,15].

The structure of the double helix under the influence of factors such as pH or temperature, undergoes denaturation process [16], also called DNA melting. Its source resides in the disruption of Watson-Crick base-pairs. During the DNA melting occur local openings and reclosing of the double-stranded helix, referred to as breathing fluctuations [16]. In the course of such a process, the DNA molecule is sorely deformed. The double helix is locally stretched, bended, untwisted or compressed [17]. The structural change is crisp, and virtually no stable intermediate state exists between the fully native and fully denatured states. This

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https://doi.org/10.1016/j.polymertesting.2019.106158
Received 6 August 2019; Received in revised form 20 September 2019; Accepted 12 October 2019
Available online 14 October 2019
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In the current study we continue this research with help of dielectric spectroscopy. Each material, on molecular level, contains electric dipoles of different nature. In external electric field, due to interactions with the molecular surroundings, they behave like damped oscillators. The relaxation time is a measure of the time, such an oscillator needs to regain its equilibrium after it has been disturbed. The distribution of relaxation times allows concluding about occurring molecular phenomena. This constitutes the basis of dielectric spectroscopy, that measures permittivity (denoted as $\varepsilon'$) and dielectric loss (denoted as $\varepsilon''$) as function of the field frequency and an environmental factor, most often the temperature [32].

Dielectric spectroscopy has already been applied for DNA testing since pre-Watson-Crick era. However, because of the significance of DNA in bioscience, experiments were mainly limited to aqueous solutions. Nevertheless, the knowledge about relaxation processes occurring in DNA is still incomplete.

An exhaustive review of experimental findings and postulated theoretical models can be found in [33]. The main conclusions presented there can be summarized as follows. DNA double helix does not have a permanent net dipole moment. In the literature, the model qualitatively describing the dielectric DNA response is referred to as Manning-Mandel-Oosawa model. It assumes that once DNA is dissolved in an aqueous solution, some of the negative charge of the molecule is neutralized by cations, especially in saline solutions. In effect a counterion layer forms around the DNA molecules. The counterions are non-localized, what implies that the positive ions can move relatively freely along the DNA surface and within the surrounding hydration shell. Therefore, the dipole moment of DNA occurs as a result of distortion of the counterion layer, forced by external electrical voltage. Typically, in aqueous solution of DNA, three relaxation processes are observed – at several kHz, between 1 MHz and 1 GHz and the last one above 1 GHz. The exact values vary, depending on the material and experimental conditions [33].

However, it is more common to use dielectric spectroscopy limited to low and medium radio frequencies (below a few MHz) as the tool for solid polymers characterization [32].

To interpret results of dielectric spectroscopy it is customary to draw so-called Bode plots that show $\varepsilon'$ and $\varepsilon''$ against frequency in double logarithmic scale. Several typical features of the curves allow a rough classification of the phenomena that are behind them. In vicinity of the frequency corresponding to a relaxation process, $\varepsilon'$ decreases in a step-like manner. A strong increase of $\varepsilon''$ at low frequencies is due to electrode polarization. On the other hand, $\varepsilon''$ exhibits a maximum at the relaxation frequency. Its linear increase with decreasing frequency is explained by ohmic-like conductivity. Maxima appearing in $\varepsilon''$ Bode plot from high to low frequencies by convention labelled by successive letters of the Greek alphabet, starting from the letter α [32].

In the case of linear polymers at moderate temperatures, most often only α-relaxation is observed within this frequency band. This relaxation is associated with micro-Brownian motion of the whole chains [32]. The double strand DNA is a stiff macromolecule of persistence length estimated at c.a. 50 nm [34]. Thus, the expected movements of its segments in the solid would be seriously hampered. Consequently, if a feature similar to α-relaxation occurs in the dielectric spectrum of heated DNA, it can be interpreted as the presence of single chains that have emerged as a result of the decomposition of the double helix.

2. Materials

Low molecular weight DNA (purity 98%) obtained from salmon sperm and other chemicals were provided by Sigma-Aldrich and used without further purification. Stock of Tris-HCl (Tris - (tris(hydroxymethyl)aminomethane) buffer was prepared at 1 M concentration with pH adjusted to 8. DNA was dissolved in the buffer to obtain 5 mg/ml DNA in 10 mM Tris-HCl. For electrophoresis TBE buffer (0.089 M Tris, 0.089 M Boric acid, 0.002 M EDTA) was used.
DNA fibers were freeze-dried at \(-53^\circ\mathrm{C}\), under 0.056 mbar for 18 h and stored in tightly closed container. Next, it was compressed in three tablets (6 mm diameter, thickness c.a. 0.5 mm), each under different loads – 50, 30 and 20 Bar. During this process and transfer to experimental setup, tablets remained in contact with laboratory atmosphere of normal humidity.

3. Methods

3.1. Electrophoresis

Electrophoresis was carried out in 3.5\% (w/v) agarose gel in TBE buffer, following the standard procedure [35] using Mupid-exu apparatus. Reference standards – 50bp and 100bp DNA Step Ladder (Promega) were simultaneously introduced to the gel. The DNA were stained in the gel with a fluorescent dye – ethidium bromide and visualized by illumination with UV light. By this method, molecular mass of the reference and the thermally treated DNA specimens were estimated.

3.2. UV

The absorbance of DNA samples in solution was measured on Spectord 250 plus spectrophotometer (Analytik Jena) in a 10 mm quartz cuvette.

3.3. Dielectric spectroscopy

Dielectric measurements in the frequency range from 1 Hz to 2.5 MHz and testing voltage of 0.3 V were performed using a turn-key Broadband Dielectric Spectrometer (CONCEPT 81, Novocontrol Technologies GmbH) equipped with an automatically controlled cryostat (Novocool Cryosystem V.2.2). Samples were measured in plate capacitor geometry, mounted in proprietary holder BDS 1200. The temperature was varied in 10\^\circ\mathrm{C} steps. Rise or decrease of the temperature was carried out at 10 \degree C/min speed, then the temperature was kept constant for next 8 min. This time was empirically established as sufficiently long for the sample to become thermally equilibrated. The heating program was divided in several heating/cooling cycles as it is illustrated in Fig. 2. In the further text the cycles will be named by descriptions used in the diagram.

4. Results and discussion

4.1. Distribution of DNA molecular mass

DNA extracted from natural cells is a statistic polymer. Bulk quantity consists of shorter and longer macromolecules. Thus, such a DNA is characterized by distributed molecular mass. A range of standard experimental methods is used to determine dispersion of molecular mass in polymers. However, in the case of DNA the most accurate is electrophoresis. A specimen of the tested DNA solution, is introduced to agarose gel. Next, the applied electric field forces the DNA to move. Because shorter macromolecules move faster, the distance travelled in the gel depends on molecular mass.

To calibrate this dependence, a reference sample (so called calibration ladder) of known, discrete molecular mass distribution, travels simultaneously with the studied one. At the end of the protocol, both the ladder and the studied sample are stained with a fluorescent dye and visualized in UV light. A monodispersed molecular mass can be easily identified by reference to the ladder. In the case of a polydispersed molecular mass such a task is more complicated. It can be assumed that staining efficiency is proportional to the number of DNA base pairs that the macromolecule consists of. Hence, the intensity of fluorescence is proportional to the amount of DNA accumulated in the particular region of the gel. Consequently, the polydispersed DNA distributed along the gel can be seen as a streak of gradually varying fluorescence intensity (see the photo in Fig. 3a). The relative intensity at a place chosen in the streak, can be correlated to the located there percent of the total base pairs number of the specimen. Two different ladders in Fig. 3a are in perfect agreement, so positions of their steps can be used to rescale in base pairs units, the distance travelled by the tested DNA. In Fig. 1b are shown fluorescence intensities of 50 bp ladder and the tested DNA as a function of the covered distance, represented in logarithmic scale. Equally spaced peaks from the 50 bp calibration ladder indicate a simple exponential dependence between the distance and the molecular mass. This feature was used to rescale the plot of the tested DNA in terms of relative intensity and base pair number. After simple calculations, there were found for the tested DNA the number average molar mass, \( M_n = 577 \times 10^4 \, \text{g mol}^{-1} \), the mass average molar mass \( M_w = 748 \times 10^4 \, \text{g mol}^{-1} \) and corresponding polydispersity 1.31.

As the first one, the DNA tablet compacted under 50 bar was subjected to the experimental routine. In Figs. 4–11 are shown Bode plots of both \( \varepsilon' \) and \( \varepsilon'' \) recorded during heating and cooling cycles from the 1st to the 4th one. These data will be discussed qualitatively, because of the experiment details. It was not possible to monitor changes of neither its dimensions nor of the quality of the surface in contact with the electrodes. This parameters may have varied, what probably happened in result of the first heating/cooling cycle. Comparison of curves for \( T = 30^\circ\mathrm{C} \) in Fig. 4a and b evidences less intense \( \varepsilon'' \) at lower frequencies, while the value at the opposite end remains similar. Such an effect can be explained by smaller electrode polarization due to a better surfaces accommodation between the sample and the electrode caused by heating.

Similarly, in the course of the measurements, water vapour may have permanently left the sample. Indeed, in traces for \( \varepsilon'' \) shown in Fig. 5a gradually appears a maximum at low frequencies then disappears upon prolonged heating (see Fig. 5b). Such an effect may be associated with the release of excess water. However, a deeper discussion of water dynamics in DNA is beyond the scope of this research. It is worth to underline, that if measured at 30 \degree C, the shape of the curves that \( \varepsilon' \) and \( \varepsilon'' \) adopted after the first heating/cooling cycle remains essentially stable after consecutive heating/cooling to 80 \degree C (Figs. 6 and 7) and next to 120 \degree C (Figs. 8 and 9). As it is shown in Fig. 12 the only difference is a small decrease in values at low frequencies. The relaxation at several kHz known from the literature [33] does not appear.

A closer inspection of Fig. 9a, illustrating variations of \( \varepsilon'' \) on heating up to 120 \degree C, reveals that the general pattern observed up to 80 \degree C imitate that shown in Fig. 7a. At temperature of 110 \degree C and above emerges a shoulder located at c.a. 10 kHz. The cooling process moves this shoulder leftwards (Fig. 7b) and finally, at 30 \degree C, the shape of \( \varepsilon'' \) plot returns to its initial form. The appearance of this shoulder is not
accompanied with any change of $\varepsilon'$ trace that reproduces the scheme observed at heating/cooling up to 80 °C.

The next heating/cooling cycle extended to 150 °C has a completely new character (Fig. 11a). The shoulder appears in $\varepsilon''$ trace again at c.a. 110 °C. Next, at gradually raised temperatures, its maximum moves towards low frequencies, i.e. in the reverse direction that expected for a typical relaxation process. This time, unlike before, the process is accompanied with a distinct step-like change of $\varepsilon'$ (Fig. 10a). As the temperature decreases, the shoulder (Fig. 11b) as well as the step threshold of $\varepsilon''$ (Fig. 10b) move in the direction of lower frequencies, in a way characteristic of a normal relaxation process.

During the cooling this shoulder disappears at c.a 70 °C, then unexpectedly reappears 50 °C (Figs. 11b and 12b). The shoulder persisted at least for a couple of hours after completion of the cooling, when the sample was conditioned a regularly measured at 30 °C. The recorded $\varepsilon'$ traces are also definitely different than that described previously.
The data provided by dielectric spectroscopy may be presented in a more visually convincing manner in terms of the loss tangent (dissipation factor), defined as $\tan\delta = \epsilon''/\epsilon'$. Its value doesn’t depend on sample geometry what is an important asset [32]. Maps, illustrating the dependence of $\tan\delta$ on frequency and temperature are shown in Fig. 13 (for the 3rd heating/cooling cycle) and in Fig. 14 (for the 4th heating/cooling cycle). If one compares Fig. 13a and b, it can be concluded that the two images are very similar by the character, beside of a negligible difference in values of $\tan\delta$. This observation is very true, particularly for lower temperatures.

Fig. 14 contains contrasting images with those shown in Fig. 13. Already in the course of heating, an additional peak, visible as an island on the map, appears in vicinity of $10^5$ Hz (Fig. 14a). During the cooling the peak maintains intensity and shifts towards lower frequencies, as predicted by Arrhenius law. Below 60 °C the peak separates in two parts. The first one continues to move toward lower frequencies, while the other and travels rightwards, ending at $10^2$ Hz, when temperature reaches 30 °C. The picture emerging from the analysis in terms of $\tan\delta$ is consistent with that obtained for permittivity and dielectric loss.
All the reported above observations confirm an important structural change occurring in DNA heated at increased temperatures between 120 °C and 150 °C. This change can be associated with thermal denaturation of DNA double helix. Results obtained for the two other tablets were qualitatively identical, so they will not be discussed.

To confirm the hypothesis, put forward above, tablets, after completion of the heating/cooling routine, were dissolved in Tris buffer and next subjected to electrophoresis together with a reference material. The picture of the stained gel (Fig. 15), evidences the absence of double stranded DNA in thermally treated samples, at least of a length close to the initial (tens of base pairs).

The second assessment was performed with help of UV–vis absorption spectroscopy. The purity of DNA is routinely evaluated by the ratio of absorbance at 260 nm and 280 nm. It is approximately equal to the weighted average of the A260/A280 ratios for the four nucleotides present. Pure DNA preparations have an A260/A280 ratio of greater than or equal to 1.8, depending on actual composition of the nucleic acid. The ratio A260/A230 is used as a secondary measure of nucleic acid purity. Expected values are commonly in the range of 2.0–2.2. If the ratio is appreciably lower in either case, it may indicate the presence of
contaminants that absorb strongly near 280 nm and 230 nm, respectively [16].

Data presented in Table 1 support the conclusion drawn from electrophoretic research. At the end of the heating/cooling treatment, the DNA sample consisted mostly of single strands. Without additional tests, the kinetics of the observed phenomenon cannot be fully and unequivocally understood. However, a purely speculative model can be imagined as follows. Starting at 120 °C, as the temperature rises, DNA helices of increasing length separate. Thus, the average relaxation time increases, what explains the shift of α-relaxation peak towards lower frequencies. During the cooling process, only single strands are present in the sample, so it resembles a linear polymer from morphological point of view. The question about the length of single strands remains open, as well about the origin of the onset peak appearing at the end of the 4th cooling (Fig. 11b).

Fig. 12. Bode plots of (a) $\varepsilon'$, (b) $\varepsilon''$ recorded at $T = 30^\circ C$, shown in semi-logarithmic scale.

Fig. 13. Dependence of tan δ on frequency and temperature. (a) 3rd heating cycle, from 30 °C to 120 °C, (b) 3rd cooling cycle, from 120 °C to 30 °C.

Fig. 14. Dependence of tan δ on frequency and temperature. (a) 4th heating cycle, from 30 °C to 150 °C, (b) 4th cooling cycle, from 150 °C to 30 °C.
5. Conclusions

Dielectric spectroscopy has proven to be an interesting alternative method for studying changes in DNA exposed to elevated temperatures. The reported research, established thermal stability limit at 120 °C for the solid, biological double strand DNA. At higher temperatures the test samples got permanently denatured. No evidence of renaturation was found, at least for a few hours. This conclusion is in line with results obtained for double strand DNA thin films studied by spectroscopic ellipsometry [31]. However, kinetic factor seems to be of vital importance for DNA degradation. In the reported research samples were heated for “sufficiently long” time. In the case of a thermal stress applied for much shorter periods, DNA can resist higher temperatures as it was obtained for double strand DNA thin films studied by spectroscopic ellipsometry [31].

Declaration of competing interest

On behalf of all the authors, I declare that there doesn’t exist any kind of conflict of interest related to the submitted article.

Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.polymertesting.2019.106158.
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