Potential Translational Thioflavin T Methodology as a Complement of Cell-Based Assays and after Drug Exposition

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Abstract: Protein aggregation is a common characteristic of several human diseases such as Alzheimer’s disease. Recent evidence has indicated that the aggregation of peptides such as p53 is also marked in cancer cells. The aim of this study was to correlate Thioflavin T (ThT) data with different cellular viability assays (Neutral Red and MTT) in SH-SY5Y neuroblastoma cells and HT-29 colon cancer cells treated with doxorubicin, a classical antineoplastic agent. We also studied the effects of the well-known peptide Aβ42 on the aggregation process in these cells. Our data suggest that both cancer cell lines are responsive to doxorubicin and formed aggregates, highlighting a relationship between ThT and cellular viability methodologies. We observed that lower values of cell viability corresponded with pronounced aggregation. Thus, these results indicated that the ThT methodology used in cells may complement the cell viability assays. In addition, this methodology may be of interest to evaluate the role of protein aggregation in other cancer cells.

Keywords: Thioflavin T; Aβ42; doxorubicin; SH-SYSY cells; HT-29 cells

1. Introduction

In stressful situations, soluble proteins can undergo structural changes and self-assembly, culminating in their aggregation. Protein aggregation has been extensively studied and is pronounced in pathologies such as Parkinson’s disease, Alzheimer’s, and type 2 diabetes [1].

To monitor the in vitro formation of proteins and other aggregation structures, Thioflavin T (ThT) can be used. This molecule is a probe that—after binding to amyloid fibrils, hydrophobic pockets in globular proteins, and other molecules such as DNA oligomers—generates a fluorescent signal at 482 nm after excitation at 450 nm [2–4]. Although the phenomenon of protein aggregation has already been studied in cancer, the evaluation of the formation of aggregates by the ThT technique can generate information regarding cancer diagnosis and carcinogenesis processes [5]. For example, p53 aggregation can be an important factor in both carcinogenesis and tumour progression. The misfolded form of this protein can form amyloid-like fibrils, which ultimately lead to disturbances
in the pro-apoptotic and tumour-suppressing function of p53 [5,6]. Recently, a study also showed that breast cancer cells (MDA-MB-231 cell line) that are damaged at the mitochondrial level accumulate aggregates of p53 that resist degradation by lysosomes [7]. Evidence shows that this probe is also capable of binding nucleic acids [8] and recent studies have found that when using the ThT technique, it is also possible to detect microRNAs, known for being important biomarkers in various types of cancer [9]. Despite the advantages, there are several drawbacks associated with this methodology, particularly the lack of robustness. Additionally, the difficulty in finding optimal concentrations of ThT has generated a lack of consensus regarding the in vitro application of this probe [1]. In this manuscript, our research group aimed to understand whether the phenomenon of aggregation could be connected to cell viability studies (MTT and Neutral Red uptake assays) in the context of cancer. We then performed the ThT methodology on different cancer cell lines, HT-29 (colon cancer) and SH-SY5Y (neuroblastoma), after the application of a drug clinically used in cancer therapy, doxorubicin (DOXO). We then performed a comparison with the results obtained by the MTT and Neutral Red assays. Our results helped us to understand if the phenomenon of aggregation, detected by ThT, was important in cancer, if it was related to the viability of the cells, and if it varied in different types of cancer such as neuroblastoma and colon cancer. We demonstrated, for the first time, that the ThT methodology can be useful for studies involving peptides and cancer cells. In Figure 1, a summary of the methodologies used in this manuscript is represented.

Figure 1. Summary of the methodologies used in this work.

2. Materials and Methods
2.1. Materials
Dulbecco’s Modified Eagle’s Medium (DMEM), foetal bovine serum (FBS), and penicillin-streptomycin mixture were obtained from Millipore Sigma (Merck KGaA, Darmstadt, Germany). Mccoy’s 5A Medium, Thiazolyl Blue Tetrazolium Bromide (MTT; cat. no. M5655), a Neutral Red solution (cat. no. N2889), Thioflavin T (cat. no. T3516), and Hexafluoro-2-propanol (HFIP; cat. no. 105228) were purchased from Sigma-Aldrich (Merck KGaA, Darmstadt, Germany). Doxorubicin hydrochloride (cat. no. 2252) was purchased from Tocris Bioscience (Bristol, UK). The Aβ42 peptide (cat. no. RP20527) was purchased from GenScript (Piscataway, NJ, USA).

2.2. Cell Culture
Human neuroblastoma cell line SH-SY5Y (American Type Culture Collection, Manassas, VA, USA) and human colorectal adenocarcinoma cell line HT-29 (American Type Culture Collection, Manassas, VA, USA) were incubated at 37 °C in a humidified atmosphere (95% air, 5% CO₂). These cells were cultivated in DMEM and Mccoy’s 5A Medium (SH-SY5Y cells and HT-29 cells, respectively) supplemented with 10% FBS and 1% of a
penicillin/streptomycin solution (1000 U/mL; 10 mg/mL). For culture maintenance, both cell lines were cultured in a monolayer and sub-cultured once a week when confluent (75–80%). Before each experiment, the cells were trypsinised (0.25% trypsin-EDTA), centrifuged (1100 rpm, 5 min), and seeded at a density of 1 × 10⁵ cells/mL in 48-well plates, which were attached for a period of 24 h.

2.3. Cell Treatment

DOXO was dissolved in water (filtered) and added to both cell lines at final concentrations of 1 nM, 10 nM, 100 nM, 1 µM, and 10 µM. The cellular viability values were determined by MTT tetrazolium reduction and Neutral Red uptake assays after 24 h and 48 h of treatment with this drug, respectively. Dose-response curves and half-maximal inhibitory concentration (IC₅₀) values for this drug after 24 h and 48 h of treatment, respectively, were obtained through the Neutral Red uptake assay. For the preparation and incubation of the Aβ42 peptide in the cells, the peptide was dissolved in HFIP (1 mg/mL) and incubated overnight at room temperature. The peptide was then frozen (−80 °C) followed by lyophilisation to evaporate the HFIP. After this last process, the peptide was dissolved in water (1 mg/mL) and used in the cells at a concentration of 10 µM as previously described [10,11].

2.4. Cell Viability Assays

After the cells were attached to the 48-well plates, DOXO was applied to the cells for 24 h and 48 h. After the treatment with this agent, the cell viability was evaluated by two distinct assays, MTT and Neutral Red. For the MTT assay, the culture medium was discarded and 200 µL/well of MTT solution (0.5 mg/mL in PBS) was added to the cells. The cells were then incubated for 3 h (light-protected). After that, the MTT solution was removed and 200 µL/well of DMSO was added to the cells. The absorbance values (570 nm) were determined in an automated microplate reader (Tecan Infinite M200, Switzerland). For the Neutral Red uptake assay, the Neutral Red medium (1:100 in the culture medium) was prepared the day before; before being applied to the cells, this medium was centrifugated (10 min, 1800 rpm). After that, the cell culture medium was removed and 250 µL/well of Neutral Red solution was added to the cells followed by a period of incubation of 3 h (light-protected). This solution was then discarded, washed with PBS (300 µL/well), and 250 µL/well of Neutral Red de-stain solution (50% of 96% ethanol, 49% de-ionised water, and 1% glacial acetic acid) was added to the plate. The absorbance at 540 nm was determined in an automated microplate reader (Tecan Infinite M200, Tecan Group Ltd., Männedorf, Switzerland).

2.5. Cell Morphology Visualisation

After the treatments, the SH-SY5Y and HT-29 cell morphology was analysed using a Leica DMI6000B automated microscope (Wetzlar, Germany) coupled to a Leica DFC350 FX camera and LAS X software.

2.6. Thioflavin T Assay

After the cells were attached to the 48-well plates, DOXO and the Aβ42 peptide were applied to the cells for 24 h and 48 h. Both cell lines were then subjected to the Thioflavin T (ThT) assay to assess the cell staining with this dye. To perform this protocol, 300 µL/well of ThT (10 µM in PBS) was added to the cells (without aspiration). The plates were then subjected to a slight agitation for 10 min and the fluorescence (excitation 450 nm, emission 485 nm) was measured using a fluorescence microscope (Olympus IX51, Tokyo, Japan) coupled to an Olympus XM10 digital camera equipped with a FITC filter cube using CellSens software.
2.7. Statistical and Data Analyses

GraphPad Prism 8 software (San Diego, CA, USA) was used to create the graphs, calculate the IC_{50} values, and carry out the statistical analysis. The total cell counts and fluorescence quantification were performed using ImageJ software (version 1.6.1, National Institutes of Health; Bethesda, MD, USA). The results were presented as a mean ± SEM for a minimum of three independent experiments. Statistical comparisons were performed between the control and treatment conditions with a Student’s t-test and a one-way ANOVA test. The differences were statistically significant when the p-value was < 0.05. Dose-response curves were obtained by non-linear regression (curve fit) using the function \( Y = \frac{100}{1 + 10^{(\text{LogIC}_{50} - X) \times \text{HillSlope})}} \).

3. Results and Discussion

3.1. SH-SY5Y Cells and Peptide Aβ42

Our first aim was to observe the natural aggregation in human SH-SY5Y neuroblastoma cells based on morphological changes. The cell morphology of the untreated cells was assessed by phase-contrast microscopy (Figure 2A). Figure 2B represents the untreated cells stained with ThT and displaying background fluorescence.

![Figure 2. Phase-contrast images of untreated human SH-SY5Y cells (A) for 24 h and fluorescence images of untreated SH-SY5Y cells (B). Scale bar: 1020 µm.](image)

Cell-penetrating peptides can be associated with the ability of aggregation and its sequence or cargo, and eventually cell death [12–23]. Human β-amyloid (1–42) peptide is a classical peptide used in studies of aggregation with ThT staining [24] because it is associated with the formation of protein aggregates. As shown in Table 1, the results demonstrated that this phenomenon of aggregation could be associated with the intrinsic properties associated with the protein structure such as the isoelectric point (5.17 for Aβ42). In Figure 3, the ratio of hydrophilic residues/total number of residues was close in the two peptides, representing an advantage.

3.2. Effect of Doxorubicin on the Viability of HT-29 and SH-SY5Y Cells

To explore the phenomenon of aggregation in neuroblastoma and colorectal cancer cells, and to understand if the ThT methodology could be used to complement cell viability assays such as MTT and NR, we started by treating SH-SY5Y and HT-29 cells with increasing concentrations of DOXO, a reference drug used for cancer therapy. This drug intercalates with DNA and inhibits the progression of topoisomerase II, impairing the synthesis of nucleic acids. After an exposition of 24 h and 48 h, the cell viability was evaluated by two different cell-based methods, MTT and Neutral Red (NR). The cell morphology was also evaluated by microscopy.
Table 1. Calculations and estimations on physiochemical properties of peptide Aβ42 using the software Pepcalc.com (peptide property calculator) [25] and the Peptide Calculator from BACHEM [26].

| Peptide  | Aβ42 (Beta-Amyloid Peptide (1–42) (Human)) |
|----------|--------------------------------------------|
| Sequence | DAEFRHDSGYEVHHQKLVFFAEDVGSNKGAIIGLMVGGVVIA (COOH) |
| Number of residues | 42 |
| Molecular weight | 4514.04 g/mol |
| Extinction coefficient | 1280 M$^{-1}$ cm$^{-1}$ |
| Isoelectric point | pH 5.17 |
| Net charge at pH 7 | -2.7 |
| Average hydrophilicity | -0.1 |
| Ratio of hydrophilic residues/total number of residues | 31% |

Figure 3. Representation of net charge and hydrophilicity of peptide Aβ42 (from Peptide Calculator/BACHEM).

For the HT-29 cells, the MTT results for the 24 h treatments demonstrated a lack of significant efficacy of DOXO in reducing the mitochondrial activity for all concentrations between 1 nM and 1 µM (Figure 4A). A significant reduction in the cell viability occurred in HT-29 treated with 10 µM of DOXO with a reduction of approximately 20% of the viable cells. The NR cytotoxicity assay was also used to evaluate the drug cytotoxicity as this technique is based on the ability of viable cells to take up Neutral Red via active transport and incorporate this dye into their lysosomes. The results for the NR assay agreed with the results obtained for the MTT assay and demonstrated the low activity of DOXO in the HT-29 cells except for the higher concentration (10 µM), whose treatment resulted in a decrease of almost 50% of the viable cells (Figure 4B). The dose-response curve for 24 h indicated that DOXO had an IC$_{50}$ value of 1.473 µM (Figure 4C). These results were also observed by microscopy; there was a significant reduction in the cell numbers and changes in the morphology of the HT-29 cells treated with 10 µM DOXO, which became rounder and vacuolised (Figure S1).
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Figure 4. Effects of DOXO on the viability of HT-29 colon cancer cells determined by (A) MTT and (B) NR assays. Based on the results of the NR assay, a dose-response curve was calculated (C). Cells were treated with increasing concentrations of DOXO for 24 h. The results are shown as the percentage of the control and are expressed as the mean ± SEM of three independent experiments (n = 3). * Statistically significant vs. the control at p < 0.05. **** Statistically significant vs. the control at p < 0.0001.

To evaluate if prolonged exposure periods resulted in an enhanced efficacy of DOXO, we treated the HT-29 colon cell line for a period of 48 h with the same concentrations of DOXO as described above. The MTT results demonstrated a significant reduction in the cell viability in the concentrations of DOXO above 100 nM with 10 µM causing a reduction of 50% of the viable cells (Figure 5A). These results also agreed with the results obtained by the NR assay for 48 h (Figure 5B), validating both methodologies for the assessment of drug cytotoxicity. The IC₅₀ for DOXO for 48 h was, as expected, lower than the one obtained for the 24 h treatment with a value of 801.7 nM (Figure 5C), demonstrating that the incubation time was a determinant for DOXO efficacy in the HT-29 cells. In accordance with these results, the cell morphology was also altered compared with the control cells in the treatments with DOXO above 100 nM with the cells becoming more vacuolised and rounder and forming fewer aggregates (Figure S2).

Regarding the SH-SY5Y cells, these cells were responsive to DOXO and, in general, the cell viability was inversely proportional to the concentration values of DOXO at both 24 h (Figures 6 and S3) and 48 h (Figures 7 and S4). This response was more pronounced after 48 h of treatment (Figure 7), and in the Neutral Red assay (Figures 6B and 7B), compared with the MTT assay (Figures 6A and 7A). These results were in agreement with the obtained results for the HT-29 cells and both methodologies for the assessment of drug cytotoxicity were validated. The lowest values of cellular viability were obtained for concentrations greater than 100 nM, which were statistically significant vs. the control. By analysing the obtained dose-response curves, it was possible to observe that with concentrations of 118.5 nM (Figure 6C) and 91.91 nM (Figure 7C), there was a reduction of 50% of the cells. These values of concentration represented the IC₅₀ values of DOXO on the SH-SY5Y cells for 24 h and 48 h, respectively. Analysing Figures S3 and S4, it was possible to verify that...
higher concentrations of DOXO resulted in a smaller number of cells, a greater number of damaged cells, and more rounded morphologies, which are characteristics of non-viable cells. After 48 h of drug exposure and with concentrations greater than 100 nM, this effect was even more pronounced, in agreement with the viability assays.

Figure 5. Effects of DOXO on the viability of HT-29 colon cancer cells determined by (A) MTT and (B) NR assays. Based on the results of the NR assay, a dose-response curve was calculated (C). Cells were treated with increasing concentrations of DOXO for 48 h. The results are shown as the percentage of the control and are expressed as the mean ± SEM of three independent experiments (n = 3). *** Statistically significant vs. the control at p < 0.001.

Taken together, these results supported the antineoplastic activity of DOXO in the HT-29 and SH-SY5Y cell lines.

3.3. Effect of Doxorubicin on the Thioflavin T Staining of HT-29 and SH-SY5Y Cells

We then evaluated the formation of aggregates in HT-29 colorectal cancer cells and SH-SY5Y neuroblastoma cells expressed in relative fluorescence units (RFU; fold change vs. the control cells) using the ThT methodology over 24 h and 48 h with a DOXO treatment. The Aβ42 peptide was also used as a positive control.

Regarding the SH-SY5Y neuroblastoma cells, our results revealed that between the concentrations of 1 nM and 1 µM of DOXO, the fluorescence intensity increased at both 24 h (Figure 8A) and 48 h (Figure 8B). This growing trend was even more pronounced at 48 h of treatment. Of all the concentrations, the treatment with 1 µM of DOXO led to the highest RFU values (Figure 9), especially at 48 h. Due to the interference of the strong reddish solution of 10 µM of DOXO, there was a significant amount of background noise in the fluorescent images and the evaluation of the relative fluorescence could not accurately be ascertained for this concentration.
Figure 6. Effects of DOXO on the viability of SH-SY5Y neuroblastoma cells determined by (A) MTT and (B) NR assays. Based on the results of the NR assay, a dose-response curve was calculated (C). Cells were treated with increasing concentrations of DOXO for 24 h. The results are shown as the percentage of the control and are expressed as the mean ± SEM of three independent experiments \((n = 3)\). * Statistically significant vs. the control at \(p < 0.05\). **** Statistically significant vs. the control at \(p < 0.0001\).

Figure 7. Effects of DOXO on the viability of SH-SY5Y neuroblastoma cells determined by (A) MTT and (B) NR assays. Based on the results of the NR assay, a dose-response curve was calculated (C). Cells were treated with increasing concentrations of DOXO for 48 h. The results are shown as the percentage of the control and are expressed as the mean ± SEM of four independent experiments \((n = 4)\). **** Statistically significant vs. the control at \(p < 0.0001\).
3.3. Effect of Doxorubicin on the Thioflavin T Staining of HT-29 and SH-SY5Y Cells

Figure 8. Effects of treatment with DOXO on the formation of aggregates in SH-SY5Y neuroblastoma cells (A,B) and HT-29 colon cancer cells (C,D) for 24 h (left) and 48 h (right) treatments. Cells were stained with ThT and aggregate formation was quantified in relative fluorescence units. Cells were treated with increasing concentrations of DOXO. Aβ42 peptide was added as a positive control. The results are shown as a fold change vs. the control and are expressed as the mean ± SEM of three independent experiments (n = 3). * Statistically significant vs. the control at p < 0.05. ** Statistically significant vs. the control at p < 0.01. *** Statistically significant vs. the control at p < 0.001. **** Statistically significant vs. the control at p < 0.0001.

Regarding the HT-29 cells, for the 24 h treatment (Figure 8C), there was a concentration-dependent increase in the fluorescence among all concentrations. The highest value of fluorescence was obtained in the treatments with 1 µM of DOXO (Figure 9). For the 48 h treatment (Figure 8D), there was less fluorescence signal per treatment compared with the treatments for 24 h. The relative fluorescence followed the same tendency as the treatment for 24 h with the highest value for the 1 µM DOXO treatment. Regarding the results obtained with the Aβ42 peptide, it led to a higher RFU vs. the control. However, in general, the fluorescence values obtained for this peptide were lower than those obtained for the treatment of the cells with DOXO. Taken together, our results for both cell lines and both periods of treatment demonstrated that DOXO could induce a greater formation of aggregates than the positive control (Aβ42).

Comparing the MTT and NR assays with the ThT methodology, we observed that lower values of cellular viability correlated with greater aggregation both in SH-SY5Y (Figure 10A,B) and HT-29 (Figure 10C,D) for treatments of 24 and 48 h. Our results suggested that the ThT methodology used in the cells could complement cell viability assays such as MTT and NR. Using this methodology, it was possible to hypothesise that an increase in DOXO concentrations (and periods of incubation) led to more significant cellular changes and toxicity, allowing ThT to easily gain access to compromised cells. These results allowed us to establish a relationship between ThT and cellular viability assays.
Figure 9. Images of the formation of aggregates in SH-SY5Y neuroblastoma cells and HT-29 colon cancer cells treated with DOXO (1 µM). Cells were stained with ThT. Aβ42 peptide was added as a positive control. Representative images were obtained in three independent experiments (n = 3). Scale bar: 50 µm.

Figure 10. Relationship between MTT, NR, and ThT results obtained in two different cell lines, SH-SY5Y neuroblastoma (A,B) and HT-29 colon cancer cells (C,D) treated with DOXO for 24 and 48 h, respectively.
Figure 10. Relationship between MTT, NR, and ThT results obtained in two different cell lines, SH-SY5Y neuroblastoma (A,B) and HT-29 colon cancer cells (C,D) treated with DOXO for 24 and 48 h, respectively.

4. Conclusions

Protein aggregation is a frequent characteristic in many neurodegenerative pathologies. Recently, this process was also associated with carcinogenesis. Therefore, the analysis of protein or other structures and aggregation in cancer cells may be a relevant additional parameter for a detailed characterisation of cancer status. Thus, the major aim of the present study was to determine the formation of aggregates in cancer cells (SH-SY5Y cells and HT-29 cells) by a ThT assay and link these data with cellular viability data after treatment with DOXO. Our data indicated that in both cancer cell lines, the increase in the effectiveness of DOXO accompanied a loss of viability of the cells. One of these factors could be associated with aggregation phenomena, which was confirmed by our data obtained with the Thioflavin T method. Our data also revealed that the Thioflavin methodology had a different affinity according to the period of contact of the cells with the drug and the type
of cell. However, in both cancer cell lines in the study, there was a similar tendency of an RFU increase that followed a cellular viability decrease. It is important to mention that the observed fluorescence after the ThT application could be explained by the presence of amyloid aggregates, globular proteins, other compounds such as DNA oligomers, or other non-amyloid structures because, as mentioned in the introductory section, ThT also reflects the presence of other molecules. In summary, with the application of DOXO in crescent concentrations, we observed that lower cell viability values correlated with greater fluorescence values obtained with the ThT methodology. As it has already been extensively described and demonstrated in this manuscript, DOXO is a drug that is cytotoxic but in a different way from the Aβ42 peptide and tends to lead to neurodegeneration/apoptosis in a time-consuming process, which may explain the differences observed between DOXO and the Aβ42 peptide, mainly regarding cytotoxicity [27]. Thus, these data can provide evidence about a connection between the cell viability values by MTT/NR approaches and the fluorescence values by the ThT methodology. Greater cytotoxicity reflects in greater ThT fluorescence. Nevertheless, these results allow the study of aggregation in the context of cancer because there is a strong possibility that DOXO may trigger a few pathways involved in aggregation such as p53 aggregation [28]. Further studies regarding the Thioflavin T technique may be necessary to study the role of protein aggregation in other cancer cells because this technique may be very important to complete other methodologies and provide further details regarding cancer evolution and carcinogenesis processes. Future research should determine the co-localisation of ThT dye and other relevant proteins in the context of cancer such as p53 protein.

Supplementary Materials: The following supporting information can be downloaded at https://www.mdpi.com/article/10.3390/ijtm2020011/s1; Figure S1: Effects of 24 h treatment with DOXO on the morphology of HT-29 cells. Representative images obtained in three independent experiments (n = 3). Scale bar: 50 µM; Figure S2: Effects of 48 h treatment with DOXO on the morphology of HT-29 cells. Representative images obtained in three independent experiments (n = 3). Scale bar: 50 µm; Figure S3: Effects of 24 h treatment with DOXO on the morphology of SH-SY5Y cells. Representative images obtained in three independent experiments (n = 3). Scale bar: 50 µm; Figure S4: Effects of 48 h treatment with DOXO on the morphology of SH-SY5Y cells. Representative images obtained in three independent experiments (n = 3). Scale bar: 50 µm.

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