Research Article

Influence of Drug-Carrier Polymers on Alpha-Synucleinopathies: A Neglected Aspect in New Therapies Development

C. Tonda-Turo, M. Herva, V. Chiono, G. Ciardelli, and M. G. Spillantini

1Department of Mechanical and Aerospace Engineering, Politecnico di Torino, Corso Duca degli Abruzzi 24, 10129 Turin, Italy
2Department of Clinical Neurosciences, University of Cambridge, The Clifford Allbutt Building, Cambridge CB2 0AH, UK

Correspondence should be addressed to C. Tonda-Turo; chiara.tondaturo@polito.it

Received 2 October 2017; Revised 15 December 2017; Accepted 25 December 2017; Published 21 February 2018

1. Introduction

Neurodegenerative diseases, such as alpha-synucleinopathies, affect nerve cells function, leading to their death, disruption of neuronal network, and development of disease [1]. Alpha-synucleinopathies such as Parkinson’s disease and Dementia with Lewy bodies are characterized by the presence of intracellular alpha-synuclein filamentous aggregates in degenerating cells in the form of Lewy bodies and Lewy neurites, while in multiple system atrophy they form glial cytoplasmic inclusions [1]. It is now clear that alpha-synuclein aggregates spread from one cell to the other, contributing to the progressive development of pathology.

Although intensively studied for many years, no treatment is currently available to stop progression of the neurodegenerative process. The approved pharmacological treatments available only provide symptomatic relief and a reduction in disease progression. One of the main factors affecting drug efficacy and utilization is the inability to reach the brain at a therapeutic concentration. To overcome this limitation, invasive treatments such as intracranial injection of hydrogels for drug delivery have been developed [2]. The use of a bioresorbable hydrogel loaded with drugs or engineered cells to achieve a direct drug delivery to the brain would ideally provide continuous drug supply at a clinically relevant concentration without the need of surgical intervention for its removal. Synthetic and natural polymers have been therefore investigated for intracerebral release of anti-inflammatory and neurotrophic factors leading to increased neuroprotection [2–4].

As an alternative to intracerebral delivery, the nasal route of administration has recently gained great interest for the treatment of neurodegenerative diseases, since it provides a noninvasive method for rapid drug adsorption bypassing the obstacle created by the blood brain barrier (BBB) [5]. The efficacy of the nasal drug uptake, compared to intraperitoneal route for example, has been demonstrated using different biomolecules, with the results showing that the amount of molecules reaching the brain is more than 5 times higher with the intranasal compared to intraperitoneal delivery [6]. Different formulations of polymers have been tested to maximize the drug uptake from the nasal cavity to the brain. Among them, mucoadhesive hydrogels have been shown to prolong the contact time between the
drug and the nasal mucosa. The drug bioavailability is consequently increased, reducing the number of administrations needed to achieve the targeted therapeutic effect [7].

Synthetic, amphiphilic multiblock copolymers with thermosensitive behavior have been widely applied as injectable drug delivery systems [8–10]. Among them, Poloxamers, a family of triblock poly(ethylene glycol)-poly(propylene oxide)-poly(ethylene glycol) (PEG-PPO-PEG) hydrogels, have been extensively used [8]. Pluronic F-127 is one of the more widely applied Poloxamers in the biomedical field [11]. Chen et al. have recently developed a Pluronic F-127 based gel for the nasal administration of curcumin to the brain, demonstrating superior drug uptake compared to the intravenous administration route [12]. Strappe et al. have developed an injectable Pluronic F-127 gel for the intracranial delivery of a lentiviral vector to the brain [13].

However, despite being widely proposed and sometimes used, hydrogels for the intracerebral and nasal administration of drugs to the brain have not been adequately characterized as the presence of hydrogels and their dissolution products could interfere with the treated pathology. In intracerebral application, hydrogels are in direct contact with the brain tissue and may interact with cells, causing unwanted side effects affecting cell behavior. In intranasal applications, the permeability of the nasal mucosa to hydrogel molecules has not yet been studied; this will be relevant in order to exclude the possibility that lower molecular weight molecules may cross the nasal barrier and reach the brain, with possible side effects.

Up to date, studies analyzing the influence of polymeric drug carriers on neurodegenerative pathologies have not been reported, and experimental methods for such an analysis are consequently missing.

In this work, we investigate the effect of PEG-PPO-PEG hydrogels on alpha-synuclein aggregation using an in vitro cellular model that was previously developed [14], where human neuroblastoma cells are infected with alpha-synuclein filamentous aggregates produced by protein misfolding cyclic amplification (PMCA) [14].

Results show that the presence of PEG-PPO-PEG increases the formation of alpha-synuclein fibrillary aggregates, which could induce unwanted contribution to pathology progression. These data support the need to investigate the effect of hydrogels in the pathological process before their use for drug delivery to the brain.

2. Materials and Methods

2.1. PEG-PPO-PEG Hydrogel Preparation. A PEG-PPO-PEG solution was prepared by dissolving 200 mg of Pluronic F-127 (Sigma-Aldrich) in 1 ml of Roswell Park Memorial Institute (RPMI) medium (Gibco) containing 10% fetal bovine serum (FBS, Invitrogen), 1% of penicillin/streptomycin (Gibco), and 1% L-Glutamine (Gibco) at 4°C and then adding 100 μl of Pluronic F-127 10% solution (Sigma Aldrich). The hydrogel was obtained by pouring 750 μl of the final solution in a silicon mold which was left at 37°C for 30 minutes to achieve complete gelation [12]. Then, a volume of 1.5 ml RPMI medium was added. After 24 h, the supernatant containing PEG-PPO-PEG hydrogel dissolution products was collected and used for in vitro cell testing.

2.2. Cytotoxicity Assay: 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium Bromide (MTT) Assay. MTT assay was performed to evaluate hydrogel cytotoxicity using human neuroblastoma cell line SH-SY5Y (ATCC® CRL 2266™). SH-SY5Y cells were cultured on a 96-multiwell plate at a cell density of 8 × 10⁴ cells/well for 24 h to reach confluence. Then, the culture medium was removed and substituted with the supernatant collected from the PEG-PPO-PEG hydrogel (as prepared as described in par. 2.1) at 1:1 and 1:5 v:v dilution. Positive control (CTRL+) was obtained inducing complete cell death by adding 20% dimethyl sulfoxide (DMSO, Sigma-Aldrich) to medium, while normal medium was used for negative control (CTRL–). After 24 h incubation, the supernatant was carefully removed and the MTT assay was performed following supplier’s instructions. Briefly, a volume of 100 μl MTT solution (Sigma-Aldrich, 5 mg/ml in phosphate-buffered saline (PBS)) was added in each well and the cultures were incubated for 3 h. Then, MTT solution was removed and 100 μl DMSO was added to each well and gently mixed to dissolve the formazan crystals. Absorbance was measured using an UV-Vis plate reader (Tecan Infinite M200PRO) at 570 nm wavelength. Cell viability was calculated as a percentage value compared to CTRL+. Six samples for each condition were used and experiments were performed three times. GraphPad Prism® software was used for one-way or two-way analysis of variance (ANOVA). *p < 0.05 was considered statistically significant.

2.3. Effect of PEG-PPO-PEG Hydrogel on Alpha-Synuclein Fibrils (αFF) Aggregation. SH-SY5Y cells were cultured on glass coverslip in a 24-multiwell plate at a cell density of 2 × 10⁴ cells/well for 24 h prior to infection. Then, fresh media containing 70 nM (1:1000 dilution from the 70 uM stock) of αFF, prepared following PMCA protocol developed by Herva et al. [14], were added and, at the same time, hydrogel supernatant at 1:5 v:v dilution was supplied to the infected cells. After 5 days, cells were fixed with a 4% paraformaldehyde solution (Sigma-Aldrich), permeabilized using 0.003% Triton (Sigma-Aldrich), and then stained using 4′,6-diamidino-2-phenylindole (DAPI, Sigma-Aldrich) 1:10000 diluted in PBS and pentamerform of formyl thiophene acetic acid (pFTAA kindly provided by Klingstedt et al. [15]) 1:10000 diluted in PBS for cell nuclei and αFF detection, respectively.

Cells were photographed under a fluorescence microscope (Leica CTR4000). Six images for sample were taken at 20x magnification and the experiment was repeated three times. Cell nuclei were counted using ImageJ to measure the number of adhered cells, while the amount of cells possessing αFF aggregates (αFF-cells) was quantified, counting the number of cells showing at least one green spot using ImageJ. Data were expressed as the average number of αFF cells with respect to the total number of adhered cells for each image ± standard deviation. Finally, the intensity of green spots within αFF cells was quantified using ImageJ processing tools.
3. Results

3.1. Cytotoxicity Assay: 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium Bromide (MTT) Assay. To determine the toxicity of PEG-PPO-PEG, cells were exposed at two different dilutions (1:1 and 1:5 v:v dilution) of extracts from the hydrogels. Figure 1 shows the viability of SH-SY5Y cells cultured in the presence of the extracts (dissolution products) from PEG-PPO-PEG hydrogels at different dilution ratios. Data were reported as percentage of negative control condition (CTRL−), where cells were cultured using normal medium. A positive control (CTRL+) was also obtained by adding 20% dimethyl sulfoxide (DMSO, Sigma-Aldrich) to medium to induce complete cell death. PEG-PPO-PEG dissolution products showed a good biocompatibility: only a slight reduction in cell viability (reaching 80% compared to CTRL−) was observed for nondiluted solutions (dilution 1:1); while no significant difference in cell viability was measured for 1:5 dilution (>90%) compared to control conditions (100%). The 1:5 dilution was therefore selected to test alpha-synuclein fibrils (αFF) aggregation in the presence of hydrogel dissolution products.

3.2. Effect of PEG-PPO-PEG Hydrogel on Alpha-Synuclein Fibris (αFF) Aggregation. The influence of hydrogel dissolution products on αFF accumulation by the SH-SY5Y cells was measured as a parameter to evaluate the interaction of drug carriers with pathology progression. Results showed that the presence of PEG-PPO-PEG dissolution products caused an increase in the amount of SH-SY5Y cells possessing alpha-synuclein fibrils aggregates (αFF cells) compared to the control (Figure 2). Then, each image was processed to quantify the green intensity ascribed to αFF fibrils within αFF cells. This analysis shows green intensity inside αFF cells from 4 to 10 times higher when αFF cells were cultured with media containing both αFF and PEG-PPO-PEG dissolution products compared to cells cultured in media containing αFF only (Figure 3), confirming the increase in the amount of αFF per cell when medium contains hydrogels dissolution products. Furthermore, when cells were cultured in the presence of both αFF and PEG-PPO-PEG dissolution products, cell clusters formed (Figures 3(e) and 3(f)), foreseeing an accelerated fibrils propagation within closer cells.

4. Conclusion

Drug carriers represent an effective tool to increase the ability of drugs to reach the target tissues. In the last years, the use of both intracerebrally injected and intranasally administered hydrogels for drug delivery to the brain has paved the way for the development of innovative therapies for the treatment of neurodegenerative diseases. When using intracerebrally administered hydrogels, cells are in direct contact with the carrier material and its dissolution products, while in the case of intranasal administration, only an amount of low molecular weight hydrogel dissolution products may pass through the nasal mucosa reaching the brain cells. However, in both cases, the effect of hydrogel dissolution products on the pathology progression requires investigation, particularly in diseases with protein aggregation like alpha-synucleinopathies, where it is known that protein aggregates spread from one cell to another. In this study, an in vitro method was used to analyze the influence of PEG-PPO-PEG hydrogel dissolution products on infection of SH-SY5Y cells with αFF. The experiments showed a relevant increase in the amount of SH-SY5Y cells possessing αFF fibrils (αFF cells) per image and an increase of αFF fibrils inside αFF cells, when cells were infected and cultured in the presence of hydrogel dissolution products. The ability of this hydrogel to enhance αFF aggregates could strongly affect the pathology progression, limiting the efficacy of the treatment inducing an unwanted contribution to aggregates formation. These findings highlight the importance of a more detailed characterization of the polymer carriers used for intracerebral or nasal drug delivery before exploring potential treatments for neurodegenerative diseases. Future work will need to...
Figure 3: Fluorescence microscopy representative images of SH-SY5Y cells after 5 days of culture in normal media ((a) and (b)), in media containing 70 nM of αFF ((c) and (d)), and in media containing both 70 nM of αFF and hydrogel supernatant ((e) and (f)). Cell nuclei are reported in blue (DAPI staining) and αFF fibrils in green (PFA staining). Bars: 50 μm. (g) Green intensity inside αFF cells. Data are reported as percentage compared to green intensity inside αFF cells when SH-SY5Y cells were cultured with media containing αFF only.
be performed to determine in vivo whether the amount of hydrogel dissolution products that reach the brain is also promoting increased alpha-synuclein aggregation and whether there is a difference between intracerebral and intranasal injections. The development of smart polymers able to avoid unwanted effects that could affect the treatment outcomes is a key strategy to open the way for novel therapies based on drug delivery systems for neurodegenerative diseases.

Conflicts of Interest
The authors declare that there are no conflicts of interest regarding the publication of this paper.

Acknowledgments
This work was supported by Smart Injectable Drug-Delivery Systems for Parkinson’s and Alzheimer’s Disease Treatment (PAD-INJ) Project (Compagnia di San Paolo) and the NC3Rs David Sainsbury Fellowship.

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