

RNA Loading on Nano-Structured Hyperbranched β -CyclodextrinSorina Hirbod¹, Shohreh Nafisi^{1,2*}, Howard I Maibach^{2*}¹. Department of Chemistry, Central Tehran Branch, Islamic Azad University, Tehran, Iran². Department of Dermatology, University of California, San Francisco, CA, USA

Abstract

Background: β -Cyclodextrin functionalized hyper-branched polyglycerol (HBCD: β -CD-g-PG), a biocompatible polymer, has recently been proposed for delivery of poorly water soluble compounds.

Methods: The present study examines the interaction of HBCD with RNA, utilizing a constant concentration of RNA and different HBCD/RNA ratios of 1/16 to 1/1, at physiological condition in an aqueous solution. Circular Dichroism (CD), UV-visible, FTIR spectroscopic methods, zeta potential and Dynamic Light Scattering (DLS) were used to analyze the particle formation, particle charge, particle size, aggregation, RNA conformation, binding constant and mode, and the effect of polymer complexation on RNA stability.

Results: The results indicate that the interaction of RNA with HBCD leads to the formation of a linear dendritic supramolecule biopolymer with an overall binding constant of $K_{\text{HBCD/RNA}} = 1.25 \times 10^3$.

Conclusion: The small sized synthesized polymer can be considered as an appropriate system for preventing RNA aggregation and protecting the gene by host-guest interaction.

Avicenna J Med Biotech 2018; 10(1): 15-21

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Received: 18 Nov 2016

Accepted: 13 Feb 2017

Keywords: Aggregation, Gene delivery, Particle size, Polymer

Introduction

Recently, a variety of macromolecule structures such as Cyclodextrins (CDs) and dendrimers have been examined for the formation of stable complexes with genes¹⁻⁶. CDs, cyclic oligosaccharides are well-known structures in supramolecular polymers in which the monomers are held together by reversible and highly directional non-covalent bonds^{7,8}. These biocompatible and nontoxic compounds⁹ possess a cage-like supramolecular structure, with hydrophilic outer surfaces and lipophilic inner cavities. Various hydrophobic drugs can be embedded in their hydrophobic cavities *via* non-covalent interactions¹⁰⁻¹⁴. However, CDs can only cause a minor enhancement in the solubility of hydrophobic drugs^{15,16}.

Recent advances have been focused on functionalizing CDs to obtain small sized macromolecules with higher free cavities and water solubility¹⁷⁻¹⁹. Polyglycerol dendron functionalized cyclodextrins (β -CD-g-PG) are novel carriers which can be synthesized by controlled anionic polymerization of glycidol. The Hyperbranched Cyclodextrins (HBCD) are small sized, well-defined polymers with higher water solubility, oxidative stability and biocompatibility. They exhibit high potential of encapsulating hydrophobic guest mol-

ecules due to the molecules binding to interior or exterior part of their structure²⁰⁻²².

Tao *et al* studied the host-guest interaction between β -CD-g-PG with long alkyl chain adamantines²³. The incorporation of insulin into HBCD polymers significantly caused enhanced drug absorbance across the nasal barrier and decreased blood glucose concentration²⁴. Paclitaxel was effectively encapsulated by HBCD for targeted drug delivery²⁵. Recently, β -cyclodextrine-polyethyleneimine polymers (PEI-CDs) were examined for *in vitro* transport of miRNA (microRNA) and siRNA (small interfering RNA)^{6,26}. CDs have also been studied as topical drug delivery systems. Application of CDs derivatives in transdermal drug delivery has improved drug release/permeation, decreased drug-induced local irritation and has optimized systemic and local dermal drug delivery. Recently, Loftsson *et al* reviewed the role of cyclodextrins on drug delivery through biological membranes²⁷.

Although several experiments report the interaction of CDs with guest molecules^{28,29}, there is no precise information on the molecular aspects of interaction between HBCD and RNA in aqueous solution. Thus, HBCD interaction with RNA was studied in aqueous

solutions with HBCD/RNA molar ratios of 1/16 to 1/1 at pH=6.5-7.5 utilizing CD, UV, and FTIR measurements. The complexes were analyzed to study the particles size and charge parameters. The structural analyses of RNA secondary structures, binding site of HBCD to RNA, and HBCD-RNA binding constant were provided by spectroscopic results.

Materials and Methods

Materials

Sodium methoxide, methanol and acetone were purchased from Merck. β -cyclodextrin, glycidol and Baker's yeast RNA from Sigma Chemical (St. Louis, MO) were employed without further purification. The absorbance band at 280 nm was utilized to check the protein content of RNA solutions. The A260/A280 ratio for RNA was 2.10, indicating that the RNA samples were nearly free from protein³⁰. Other chemicals were of reagent grade and used without further purification.

Synthesis of HBCD polymer

HBCD (β -CD-g-PG) was synthesized by anionic ring opening multi branching polymerization method²⁰. Briefly, β -CD (0.5 g) was deprotonated by dissolving sodium methoxide (14.4 mmol) in dried methanol. Mixture was stirred at room temperature using a polymerization ampule equipped with a magnetic stirrer for 1 hr. Methanol was vaporized *via* vacuum oven at 60°C for 2 hr. Glycidol (92 mmol) was added to deprotonated β -CD and temperature was gradually increased up to 120°C. The mixture was stirred at 120°C for 12 hr and allowed to cool. It was dissolved in pure methanol, and precipitated upon addition of acetone. It was neutralized by filtration over cation-exchange resin. The precipitate was dried *via* vacuum oven at 80°C. Finally, a yellow viscous product was obtained with 87% yield.

Preparation of stock solutions

Homogeneous sodium-RNA solutions (1% w/w; 10 mg/ml) were prepared by dissolving RNA in 10 ml of phosphate buffer (pH=7.4) with occasional stirring at 5°C for 24 hr. Using the molar extinction coefficient of 6600 cm⁻¹ M⁻¹ which was represented as molarity of phosphate groups, the final concentration of RNA stock solution was assessed spectrophotometrically at 260 nm^{31,32}. UV absorbance of the diluted RNA solution (40 μ M) at 260 nm was 0.11 (with the path length of 1 cm), and the final concentration of RNA stock solution was 25 mM in phosphate. For infrared spectroscopic measurements, various amounts of HBCD in phosphate buffer (pH=7.4) were added dropwise to RNA solutions to achieve favorable HBCD contents of 0.625, 1.25, 2.5, 5 and 10 mM with an ultimate RNA concentration of 10 mM. Infrared spectra were recorded 1 hr after the mixing of HBCD solutions with RNA solutions. For UV measurements, various concentrations of HBCD (500-1200 μ M) were used with constant RNA concentration (40 μ M).

Table 1. Zeta potential of RNA, HBCD and RNA-HBCD

Sample [(Concentration (μ M))]	ζ (mV)
RNA (40 μ M)	-8.47
HBCD (1000 μ M)	-1.8
HBCD-RNA	-9.22

FTIR spectroscopy measurements

To record the infrared spectra, a Nicolet FTIR spectrometer (Magna 550) equipped with a liquid-nitrogen-cooled HgCdTe (MCT) detector was used. The spectra of the HBCD/RNA solutions were collected utilizing a cell assembled with ZnSe windows and treated by OMNIC software. The spectra of solutions were recorded after 1 hr. The bands were measured in three individual samples, at the same HBCD and RNA concentrations. 100 scans were collected for each spectrum with resolution of 4 cm⁻¹. To obtain difference spectra [(polynucleotide solution+HBCD solution)-(polynucleotide solution)], a sharp RNA band at 968 cm⁻¹ was used as internal reference^{33,34}. This band is assigned to sugar C-C and C-O stretching vibration and does not change (intensity or shifting variation) upon HBCD complexation to RNA, so it will be deleted upon spectral subtraction.

By considering the reference band at 968 cm⁻¹ (RNA) as a function of HBCD concentrations, the intensity ratios of RNA in-plane vibrations related to phosphate stretching vibrations and A-U and G-C base pairs were measured with an error of $\pm 0.9\%$. In order to determine the ligand binding to RNA bases and backbone phosphate groups, similar intensity variations have been used^{35,36}. After peak normalization, the plot of relative intensity (R) of various RNA in-plane vibrations including 1698 (guanine), 1650 (uracil), 1606 (adenine), 1489 (cytosine) and 1241 cm⁻¹ (phosphate groups) versus polymer concentrations were gained. The peak normalization was performed using $R_i = I_i/I_{968}$ where I_i is the intensity of the absorption peak for pure RNA in the complex with i as ligand concentration, and I_{968} as the intensity of the peak at 968 cm⁻¹ (RNA internal reference) (Figure 1).

Absorption spectroscopy

A LKB model 4054 UV-Visible spectrometer was used to record the absorption spectra; quartz cuvettes of 1 cm were used with various HBCD concentrations of 5×10^{-4} - 12×10^{-4} M and constant RNA concentration of 40×10^{-6} M. Binding constants of HBCD-RNA complexes were calculated as reported³⁷. The interaction between HBCD as ligand [L] and RNA as substrate [S] is assumed to be 1:1 which forms a single complex SL (1:1). Equation (1) describes the relationship between the system variables and parameters with the observed absorbance change per centimeter:

$$(1) \frac{\Delta A}{b} = \frac{S_t K_{11} \Delta \epsilon_{11} [L]}{1 + K_{11} [L]}$$

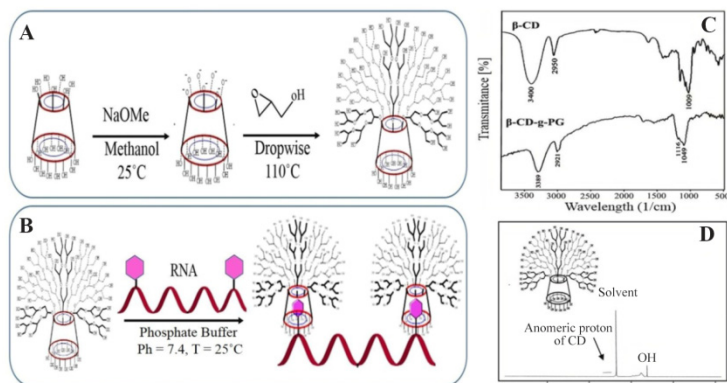


Figure 1. A) HBCD synthesis by anionic ring opening polymerizations. B) HBCD-RNA synthesis by host-guest interactions within the cavity of HBCD polymers and RNA bases. C) FTIR spectra of β -CD and HBCD. D) ^1H -NMR spectra of HBCD in the D_2O as solvent.

$\Delta A = A - A_0$ from the mass balance expression $S_t = [S] + [SL]$, $[S] = S_t / (1 + K_{11}[L])$ is obtained

Equation (1) is the binding isotherm, which indicates the hyperbolic dependence on free ligand concentration. According to the following equation, the linearization of equation (1) yields the double-reciprocal form of plotting the rectangular $\frac{1}{y} = \frac{f}{a} \cdot \frac{1}{x} + \frac{e}{a}$ hyperbola:

$$(2) \frac{b}{\Delta A} = \frac{1}{S_t K_{11} \Delta \epsilon_{11} [L]} + \frac{1}{S_t \Delta \epsilon_{11}}$$

As the double reciprocal plot of $1/\Delta A$ versus $1/[L]$ is linear, the binding constant can be estimated from the following equation:

$$(3) K_{11} = \frac{\text{intercept}}{\text{slope}}$$

CD spectroscopy

Various HBCD concentrations (125, 250, 500, 1000 and 2000 μM) were mixed with constant concentration of RNA (40 μM). The spectra of free baker's yeast RNA and HBCD-RNA adducts were recorded using a Jasco J-720 spectropolarimeter at pH=7.3. A quartz cell with a path length of 1 cm was used for far-UV region (200-260 nm) measurements. Six scans were accumulated at a scan speed of 50 nm/min, with data being collected at every nanometer from 200 to 260 nm. The temperature of sample was kept at 25°C using a Neslab RTE-111 circulating water bath connected to the water-jacketed quartz cuvette. Using Jasco Standard Analysis software, spectra were corrected for buffer signal, and a conversion to Mol CD ($\Delta \epsilon$) was performed.

Determination of size and potential

To determine polydispersity and the average diameter of HBCD-RNA polymer, free HBCD (1000 μM) and free RNA (40 μM), a Dynamic Light Scattering (DLS) particle size analyzer (Brookhaven-model: 90plus USA) was used. A zeta potential meter (Brookhaven-model: ZetaPALS) was used to measure zeta potential of HBCD-RNA complexes, free HBCD (1000 μM) and free RNA (40 μM) at 37°C.

Results

IR and ^1H -NMR of HBCD

HBCD polymer was synthesized by anionic ring opening polymerization method (Figure 2A) which was previously reported^{20,28}. The IR spectral changes of β -CD and HBCD are presented in figure 2B.

Absorption bands at 3380, 1116 and 2921 cm^{-1} are related to O-H, C-O-C and C-H groups, respectively^{20,28}. Broadening of the C-O band in HBCD spectrum comparing with β -CD can be assigned to polyglycerol binding to cyclodextrin functional groups. Similar broadening in the C-O absorbance of HBCD compared with β -CD was assigned to polyglycerol binding to the OH groups of cyclodextrin^{20,28}.

In the HBCD ^1H NMR spectrum (Figure 2C), the signals of cyclodextrin protons have overlapped with polyglycerol signals at 4.1-3.6 ppm. The weak signal at 5.2 ppm is related to anomeric protons of cyclodextrin. The same ^1H -NMR results for HBCD was reported by Adeli *et al*^{28,38}.

Infrared spectra of HBCD-RNA complexes

FTIR spectral studies have identified weak interaction of HBCD with phosphate and bases of RNA at low HBCD concentrations (*via* host-guest) and major participation of bases mainly with guanine, cytosine and backbone phosphate groove at higher concentrations. Evidence for this comes from the spectral changes of free RNA upon HBCD complexation.

At low HBCD concentration ($r=1/16$ to $1/8$), no significant interaction was observed between HBCD and RNA bases as represented by small shifting and intensity changes of RNA in-plane vibrations at 1698 (G, U mainly G), 1650 (U, G, A, C mainly U), 1606 (A),

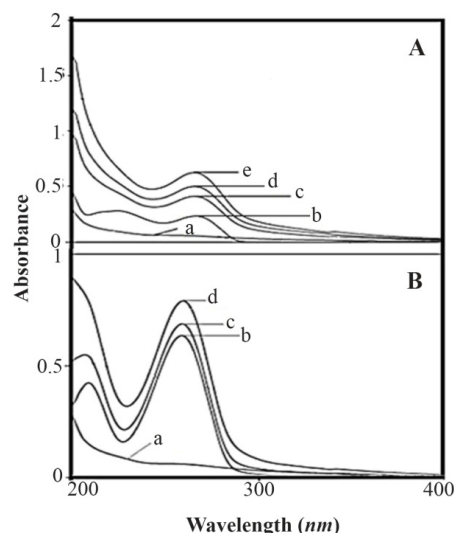


Figure 2. A) UV-visible spectra of (a) HBCD (100 μM); (b) RNA (40 μM), (c-j) HBCD-RNA complexes: (c) 250, (d) 300, (e) 350, (f) 400, (g) 450, (h) 500, (i) 550, (j) 600 μM . B) Plot of $1/(A-A_0)$ vs. $(1/\text{polymer concentration})$ for HBCD and RNA complexes, where A_0 is the initial absorbance of RNA (260 nm) and A is the recorded absorbance (260 nm) at different polymer concentrations (250-600 μM) with constant RNA concentration of 40 μM at pH=7.4.

1489 (C) and 1241 cm^{-1} (asymmetric PO_2 stretch) (Figures 1 and 3). The difference spectra showed negative features at 1724 and 1782 cm^{-1} which is due to the loss of RNA intensity vibrations (Figure 1). This is indicative of a minor, indirect interaction of RNA with HBCD via H-bonding system at low HBCD concentrations. Similarly, the loss of DNA vibrations intensity and negative features at cisplatin-DNA spectrum was assigned to minor and indirect H-bonding interaction of DNA with CD-37³⁹.

At higher HBCD concentrations ($r=1/4$), major shifting of the guanine and cytosine bands (G band at 1698 to 1680 cm^{-1} ($1/1$); C band at 1489 cm^{-1} to 1474 ($r=1/4$) was observed and the intensities of the adenine, guanine, cytosine and uracil bands strongly augmented. The intensity increase was characterized by the presence of several positive features at 1715 (guanine), 1650 (uracil), 1510 , 1490 cm^{-1} (cytosine) and 1245 (phosphate) in the difference spectrum of HBCD-RNA (Figures 1 and 3; the difference spectrum is not shown in figure 1). The spectral changes (intensity and shifting) for the A, G, C and U bases ($r=1/4$) can be assigned to the interaction of HBCD (through OH) with mainly guanine and cytosine and to a lesser extent with the adenine and uracil bases.

At higher HBCD concentrations ($r=1/2$, $1/1$), drastic shifting of the guanine and cytosine bands (guanine band at 1698 to 1706 cm^{-1} ($1/1$); cytosine band at 1489 to 1460 ($r=1/2$), 1477 cm^{-1} ($r=1/1$)) were observed. The spectral changes for the bases ($r=1/2$ and $1/1$) can be attributed to the interaction of OH branches of HBCD with mainly G and C and to a lesser extent with A and U bases (Figures 1 and 3). Similar spectral changes were observed in narigin-RNA complexes⁴⁰.

HBCD- PO_2 interaction was evident from intensity increase and shifting of the PO_2 antisymmetric band at 1241 and symmetric band at 1085 cm^{-1} in the spectra of HBCD-RNA complexes (Figures 1 and 3).

For the phosphate group, shifting of the band at 1241 cm^{-1} to 1243 ($r=1/8$), 1245 ($r=1/4$), 1253 ($r=1/2$) and 1249 cm^{-1} ($r=1/1$) for the complexes can be ascribed to minor phosphate interaction with HBCD at lower concentrations and strong interaction at higher concentrations ($r=1/2$, $1/1$). Similar spectral changes were observed in PAMAM-tRNA complexes³⁶.

Circular dichroism spectra and RNA conformation

CD spectra of RNA and its complexes with different HBCD concentrations are shown in figure 6. CD spectrum of RNA, composed of four major peaks at 209 (negative), 221 (negative), 240 (negative), and 269 nm (positive) (Figure 3) is consistent with previously reported CD spectra of double-helical RNA in a conformation^{38,41}.

At low HBCD concentration (125 , $250\text{ }\mu\text{M}$), no significant shifting was observed; however, the molar ellipticity of the band at 209 nm was strongly enhanced by increasing the polymer concentration (500 to $2000\text{ }\mu\text{M}$). In the spectra of HBCD-RNA complexes, no

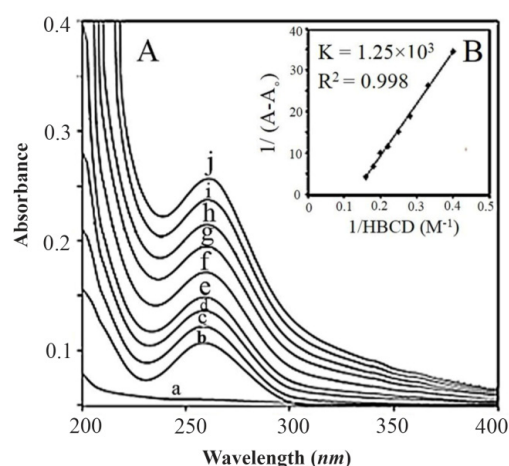


Figure 3. FTIR spectra in the region of $1800\text{--}600\text{ cm}^{-1}$ for (A) Baker's yeast RNA (10 mM) (B) HBCD (C) HBCD-RNA: RNA (10 mM), HBCD (5 mM) (D) HBCD-RNA: RNA (10 mM), HBCD (10 mM) in aqueous solution at $\text{pH}=7$. RNA, HBCD and two complexes at various HBCD/RNA (phosphate) molar ratios ($1/2$, $1/1$) (four top spectra); two difference spectra (bottom two spectra).

shifting was observed for the band at 269 nm (Figure 3) which can be attributed to the presence of a conformation in free RNA and in HBCD-RNA complexes. This is consistent with the infrared results which showed a conformation in free RNA with IR marker bands at 1698 (G), 1241 (PO_2), 861 and 810 cm^{-1} (ribose-phosphate) and in HBCD-RNA complexes at $1689\text{--}1688$ (G), $1241\text{--}1234$ (PO_2), $864\text{--}868$ and $812\text{--}810\text{ cm}^{-1}$ (ribose-phosphate) in the complexes (Figure 4). Besides, minor intensity change of the band at 269 nm and its collapse in the spectrum of HBCD-RNA can be related to the lack of RNA aggregation in the presence of HBCD polymer (Figure 3). This is consistent with DLS results of HBCD-RNA complexes which revealed that RNA aggregation did not occur.

Stability of HBCD-RNA complex

HBCD-RNA binding constant was determined as demonstrated previously in UV-Visible spectroscopy (Figure 5A). The overall binding constants were obtained using UV spectroscopy in a previously reported method³⁷. Concentrations of the complexed HBCD (ligand) were determined by subtracting the absorbance of the free RNA at 260 nm from those of the complexed. The concentration of free ligand was determined by the subtraction of the complexed ligand from total ligand. The data of $1/[\text{ligand complexed}]$ almost proportionally augmented as a function of $1/[\text{free ligand}]$ (Figure 5B). As the double reciprocal plot of $1/(A-A_0)$ vs. $1/(\text{HBCD concentration})$ was linear, binding constant (K) was calculated from the ratio of the intercept to the slope (Figures 4A and 4B); A_0 is the initial absorbance of the free RNA at 260 nm and A is the recorded absorbance of the complexes at various HBCD concentrations. The overall binding constant of HBCD-RNA ($K=1.25\times 10^3\text{ M}^{-1}$) was indicative of a weak stability through attraction forces between HBCD and RNA. Similar bindings

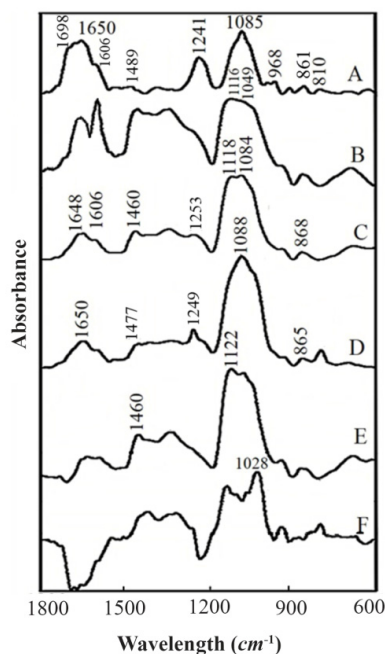


Figure 4. Intensity ratio variations for various RNA in-plane vibrations as a function of HBCD concentration. Intensity ratios for RNA bands at 1698 (G, U), 1650 (U,G,A,C), 1606 (A), 1489 (C,G) and 1241 (PO₂ asymmetric) referenced to RNA band at 969 cm^{-1} .

were reported for G_{3.0} PAMAM dendrimers conjugated to 1-pyrene carboxaldehyde G_{3.0}-PY (as guest) and HBCD (as host) complexes⁴² in which the enhanced λ_{Max} of the guest molecule was attributed to the host-guest interactions. Besides, UV-Vis spectrum of RNA at 260 nm (Figures 5A and 5B) exhibited increased intensity upon the addition of HBCD (spectra c to j) which was assigned to the host-guest interactions between the two segments and formation of HBCD-RNA complex.

RNA conformation

RNA remains in a conformation in HBCD-RNA complexes. The lack of major shifting of the A-RNA marker bands at 815-809 cm^{-1} (phosphodiester), 867-861 (ribosephosphate), 1247-1241 (phosphate), and 1700-1688 (guanine) indicates that RNA remains in a conformation upon HBCD complexation (Figure 1)⁴³⁻⁴⁵.

Size and zeta potential

The size distribution profile of small particles in suspension or (macro) molecules in solution can be determined by Dynamic Light Scattering (DLS)⁴⁶. DLS measurements were applied to estimate the average hydrodynamic diameter of RNA, HBCD and their supramolecular assembly in solution (HBCD-RNA complexes) (Figures 6A-C). The obtained hydrodynamic diameter for free HBCD (2.8 nm at pH=7.4 and at 37°C) is in good agreement with previous literature reports (Figure 6)²⁸. After addition of HBCD to RNA solution, the d_h value of the HBCD-RNA complex increased to 5.7 nm which is larger than the sum of the hydrodynamic di-

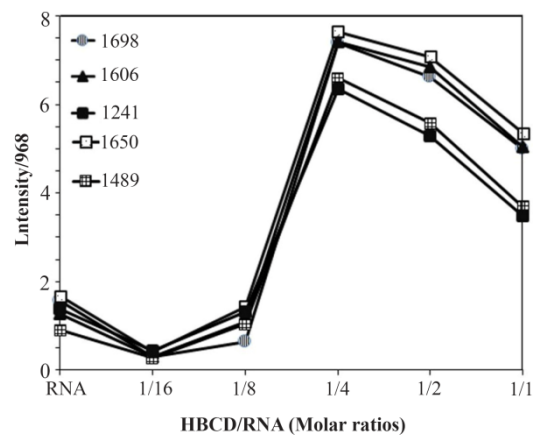


Figure 5. CD spectra at 25°C for (a) free RNA (40 μM in phosphate buffer, pH=7.4), (b-f) HBCD-RNA complexes with different concentrations of HBCD containing: (b) HBCD (125 μM), (c) HBCD (250 μM), (d) HBCD (500 μM), (e) HBCD (1000 μM) and (f) HBCD (2000 μM).

ameters of the pure components. As discussed earlier, by formation of HBCD-RNA complex, RNA remained in a conformation. Therefore, increase in HBCD-RNA dimension can be attributed to an extended conformation of the polyglycerol chains of HBCD into the solvated environment. Based on the interactions between albumin and HBCD, this result could also be explained by the structural properties of HBCD and cyclodextrin hydrophobic cavity, the polar outer shell as well as complex tendency to interact with RNA⁴⁷. RNA zeta potential due to the presence of ionized residues on RNA surface is greatly affected by the concentration and charge of the cationic particles. Indeed, the phosphate backbone of nucleic acids carries one negative charge per residue⁴⁸. By the addition of HBCD ($\zeta = -1.23 \text{ mV}$), the value of ζ for the supramolecular complex (HBCD-RNA) decreases to -9.22 mV . Decrease in ζ -value can be resulted from the intermolecular forces which led to HBCD-RNA supramolecular assembly.

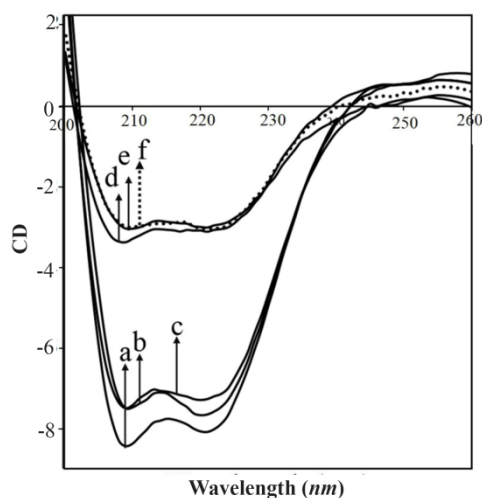


Figure 6. DLS diagrams (phosphate buffer, pH=7.4 at 37°C); A) RNA (40 μM), B) HBCD (1000 μM) C) HBCD-RNA complexes.

Conclusion

An unprecedented type of gene delivery system was successfully developed based on linear dendritic supramolecular biopolymer. RNA-HBCD complexes can be used to gain insight into the new strategies for gene delivery. Our study provided substantial quantitative data on the effect of low and high HBCD concentrations on polynucleotide structure and the binding affinity of RNA to HBCD. According to the spectroscopic results, the following points are considerable: RNA binds HBCD through non-covalent binding (host-guest interactions) between the gene bases as guest and β -cyclodextrin cavities in HBCD as host. The binding constant ($K=1.25 \times 10^3$) indicated weak HBCD interaction with RNA which easily dissociates in aqueous solution. Strong bonds between RNA and HBCD were formed at higher HBCD concentrations. The small sized synthesized complex (5 nm) successfully prevented RNA aggregation and protected RNA by host-guest interactions between the cavity of HBCD polymers and the bases of gene. No RNA conformational change occurred upon HBCD interaction and RNA remained in A-family structure.

Acknowledgement

The authors appreciate Central Tehran Branch, IAU, Tehran, Iran for support of this work.

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