β-CD Dimer-immobilized Ag Assembly Embedded Silica Nanoparticles for Sensitive Detection of Polycyclic Aromatic Hydrocarbons

Eunil Hahm1, Daham Jeong1, Myeong Geun Cha2, Jae Min Choi2, Xuan-Hung Pham2, Hyung-Mo Kim2, Hwanhee Kim2, Yoon-Sik Lee3, Dae Hong Jeong2, Seunho Jung1 & Bong-Hyun Jun1

We designed a β-CD dimer on silver nanoparticles embedded with silica nanoparticles (Ag@SiO2 NPs) structure to detect polycyclic aromatic hydrocarbons (PAHs). Silica NPs were utilized as a template for embedding silver NPs to create hot spot structures and enhance the surface-enhanced Raman scattering (SERS) signal, and a thioether-bridged dimeric β-CD was immobilized on Ag NPs to capture PAHs. The assembled Ag NPs on silica NPs were confirmed by TEM and the presence of β-CD dimer on Ag@SiO2 was confirmed by UV-vis and attenuated total reflection-Fourier transform infrared spectroscopy. The β-CD dimer@Ag@SiO2 NPs were used as SERS substrate for detecting perylene, a PAH, directly and in a wide linearity range of 10^{-7} M to 10^{-2} M with a low detection limit of 10^{-8} M. Also, the β-CD dimer@Ag@SiO2 NPs exhibited 1000-fold greater sensitivity than Ag@SiO2 NPs in terms of their perylene detection limit. Furthermore, we demonstrated the possibility of detecting various PAH compounds using the β-CD dimer@Ag@SiO2 NPs as a multiplex detection tool. Various PAH compounds with the NPs exhibited their distinct SERS bands by the ratio of each PAHs. This approach of utilizing the assembled structure and the ligands to recognize target has potential for use in sensitive analytical sensors.

Polycyclic aromatic hydrocarbons (PAHs) consist of several benzene rings and are carcinogens, teratogens and mutagens1. The sources of these compounds are diverse as they are byproducts of incomplete combustion of fossil fuels or organic substances. Various PAH detection methods have been developed. Generally, PAHs in water, soil and solid waste are analyzed after extraction and concentration by chromatography such as, liquid chromatography with fluorescence detection (LC/FLDs)3, high-performance liquid chromatography (HPLC) with UV and fluorescence4–5, gas chromatography coupled with a flame ionization detector (GC/FID)6 and mass spectrometry (GC/MS)7,8. The PAH detection limits of these techniques are in the ppb range. However, the chromatography method is hampered by the prolonged sample preparation required9. Also, application is problematic because of their incompatibility with column chromatography. Surface-enhanced Raman spectroscopy (SERS) has been proposed to overcome these problems10,11.

SERS is an ultra-sensitive analytical technique12,13, which can detect very low concentrations of the target compound14,15 and determine structural information16. Over the past decade, a range of multidirectional strategies for detecting chemicals on SERS substrates have been investigated, utilizing single-metal NPs with a modified surface and various shapes17–20 and sol-gels21 as sensors. Further, metal NPs held together in solution can generate hot spots22,23, which have been used for SERS detection. However, aggregation of metal NPs in solution is difficult to control due to their instability and uncontrollability. Assembled Ag SERS substrates comprising a silica core and
Ag NPs could overcome the weaknesses mentioned above, affording high sensitivity. However, Ag NPs on the surface of silica core particles have poor affinity to PAHs.

Cyclodextrins (CDs), which have hydrophobic internal holes, are excellent receptors of PAHs and can be placed on the metal surface via the hydrophobic effect. In particular, bridged dimeric β-CDs were reported to exhibit significantly enhanced binding and recognition of target materials due to cooperative binding of two neighboring CDs, compared to native CDs. However, these bridged dimeric β-CDs are not frequently utilized for sensing applications and their use for detection of PAHs has not been investigated extensively to date.

In this study, we used β-CD dimers-immobilized Ag assembled silica NPs (β-CD dimer@Ag@SiO₂ NPs) as a SERS substrate to detect PAHs. The assembled Ag structures enhanced the SERS signals and the β-CD dimers functioned as affinity ligands for PAHs. The structure of the β-CD dimer@Ag@SiO₂ NPs was analyzed by transmission electron microscopy (TEM), UV-vis spectroscopy and attenuated total reflection-Fourier transform infrared spectroscopy (ATR-FTIR). Sensitivity of β-CD dimer@Ag@SiO₂ NPs for detecting PAHs was compared with that of Ag@SiO₂ to determine the role of the β-CD dimer. Additionally, the possibility of multiplex detection was tested using mixtures of several PAHs, such as phenanthrene, pyrene and benzo(a)pyrene.

Results and Discussion

Characterization of β-CD dimer@Ag@SiO₂ NPs structure. The structure of β-CD dimer@Ag@SiO₂ NPs, used as a SERS substrate for detection of PAHs, is shown in Fig. 1. Silica NPs of ~180 nm diameter were prepared according to the Stöber method and utilized as a template to generate assembled Ag NPs on silica NPs. Silica NPs were first thiolated with 3-mercaptopropyl trimethoxysilane (MPTS) to introduce thiol groups to their surface.

Ag NPs were embedded on silica NPs by reduction of silver nitrate by octylamine with polyvinylpyrrolidone (PVP) in ethylene glycol. Thioether-bridged β-CD dimer, which was prepared as reported previously, was immobilized on Ag NPs by means of the affinity of thiol groups for silver. Here, β-CD dimers were used to capture PAHs so that PAHs can be located on the surface of the Ag NP through the host-guest complexation. The assembled Ag NPs on silica NPs were designed to create hot spots to enhance the SERS signals of target molecules.

Figure 2A shows TEM images of the Ag@SiO₂ NPs. These NPs were uniform in size (ca. 206 ± 10 nm) and well dispersed without aggregation. The size of Ag NPs embedded on the surface of silica NPs was 8.4 ± 3.7 nm. β-CD dimers were immobilized on the surface of the assembled Ag NPs on the silica NPs to capture PAHs via host-guest interactions. To investigate the optical properties of these particles, UV–vis spectra were recorded (Fig. 2B). The UV-vis spectrum of the assembled Ag@SiO₂ NPs (line f) showed a broad band from 322 to 800 nm with a maximum peak at 430 nm, indicating the presence of aggregates of Ag NPs on the surface of silica NPs. Following assembly of β-CD dimer on the surface of the Ag@SiO₂ NPs, UV-vis spectrum of the β-CD dimer@Ag@SiO₂ NPs show a slightly shifted localized plasmon resonance band from 430 nm to 464 nm (line e). The spectra of β-CD dimer@Ag@SiO₂ NPs with various concentration of perylene from 1 × 10⁻² to 1 × 10⁻⁴ M in line a–d were showed a peak of 452 nm, which is assigned as perylene. The increasing of absorbance peaks by perylene were relatively weak, compared to the absorbance peak of β-CD dimer@Ag@SiO₂, however the intensity of absorbance peaks was increased by raising the concentration of perylene. We believe that when perylene was caught by β-CD dimer@Ag@SiO₂, the presence of perylene on the surface of β-CD dimer@Ag@SiO₂ may have caused the combination effect of the extinction coefficient of our materials to visible light that increased the intensity of the absorbance peak.

To confirm assembly of β-CD dimers on the surface of the Ag@SiO₂ NPs, attenuated total reflection-Fourier transform infrared spectroscopy (ATR-FTIR) was performed. The IR spectra of Ag@SiO₂ NPs and β-CD dimer@Ag@SiO₂ NPs are shown in Figure S1. The IR spectra of the Ag@SiO₂ NPs and the β-CD dimer@Ag@SiO₂ NPs differed. The intensity of the band at 2923 cm⁻¹, assigned to C-H stretching vibration, increased due to the β-CD dimers. Also, a new band at 2349 cm⁻¹ was assigned to the β-CD dimers. Band shifts from 1048 cm⁻¹ to 1044 cm⁻¹ and from 1656 cm⁻¹ to 1685 cm⁻¹ were observed due to the assembly
of β-CD dimer on Ag@SiO₂ NPs. In particular, a characteristic band at 708 cm⁻¹ was assigned to C-S stretching vibration bands of the β-CD dimers. Our result is consistent with a previous report. Therefore, we demonstrated that β-CD dimers were successfully assembled on the Ag@SiO₂ NPs.

**Detection of perylene using β-CD dimer@Ag@SiO₂ NPs.** β-CD can interact with PAHs in its hydrophobic cavity and form various host–guest complexes. Unlike native CDs, dimeric β-CDs exhibit significantly enhanced binding and recognition of target materials due to cooperative binding of two neighboring CDs. Therefore, we assumed that the β-CD dimer-modified Ag@SiO₂ NPs would facilitate enrichment of PAHs on the metal surface, and PAH–CD complexes could be detected by surface-enhanced Raman spectroscopy.

SERS spectra of β-CD dimer@Ag@SiO₂ NPs treated with 1 × 10⁻² M to 1 × 10⁻¹ M perylene for 30 min are shown in Fig. 3. The β-CD dimer@Ag@SiO₂ NPs were mixed with perylene and the SERS signals were measured in a capillary tube. SERS intensity increased with perylene concentration, particularly the peaks at 1293 cm⁻¹ and 1576 cm⁻¹. The peak at 1293 cm⁻¹ was assigned to the C-C stretching on bridges and the peak at 1576 cm⁻¹ was assigned to the C-C stretching on benzene rings of perylene. Perylene at up to 10⁻² M was detected by β-CD dimer@Ag@SiO₂ NPs treated with the capillary method and 10⁻⁴ M using the drop-cast method (Figure S2). To confirm the β-CD dimer effect, the Ag@SiO₂ NPs lacking β-CD dimers were mixed with perylene. The SERS spectra of the Ag@SiO₂ NPs in the presence of perylene are shown in Figure S3. The normalized SERS intensity at 1293 cm⁻¹ of the β-CD dimer@Ag@SiO₂ NPs and Ag@SiO₂ NPs with perylene in various concentrations are shown in Fig. 3B. The SERS signals of the β-CD dimer@Ag@SiO₂ with the same concentration of perylene were markedly higher than those of Ag@SiO₂ NPs lacking β-CD dimers. A similar phenomenon appeared at 1567 cm⁻¹ (Figure S4). Therefore, we concluded that β-CD dimers played an important role in capturing and enriching PAHs on the surface of the Ag@SiO₂ NPs. Although the detection limit of perylene using the SERS signals was equivalent to those of traditional methods, such as isotopic dilution capillary column gas chromatography/mass spectrometry (GC/MS) and high-performance liquid chromatography (HPLC), the SERS technique is more cost-effective and less time-consuming.

**Multiplex detection of PAHs using β-CD dimer@Ag@SiO₂ NPs.** To utilize the multiplex detection capability of the β-CD dimer@Ag@SiO₂ NPs for PAH compounds, phenanthrene, benzo(a)pyrene, pyrene and perylene were mixed with β-CD dimer@Ag@SiO₂ NPs.

Figure 4A shows the SERS spectra of phenanthrene, benzo(a)pyrene, pyrene and perylene (10⁻⁴ M each) mixed with β-CD dimer@Ag@SiO₂ NPs. The SERS bands of the β-CD dimer@Ag@SiO₂ NPs in the presence of various PAHs differed from those of β-CD dimer@Ag@SiO₂ NPs only. In particular, pyrene showed a characteristic peak at 1240 cm⁻¹ and perylene a characteristic peak at 1293 cm⁻¹. Benzo(a)pyrene had a characteristic peak at 1234 cm⁻¹. Phenanthrene exhibited somewhat weaker intensity than the other PAHs, and its characteristic peak was at 836 cm⁻¹. These results demonstrate that β-CD dimer@Ag@SiO₂ NPs can interact with various PAHs and show distinct SERS bands. More experiments to provide the sensor selectivity were performed. Each organic compound (perylene, naphthalene, toluene, isopropyl acid and ethylene glycol) at the concentration of 10⁻⁴ M was added in β-CD dimer@Ag@SiO₂ NPs solutions and those solutions were compared with β-CD dimer@Ag@SiO₂ NPs solution. Perylene showed its characteristic peak, while other solutions that had added organic compounds did not show their peaks in Raman spectra.

Figure S5 (A) showed the Raman spectra of those solutions. The presence of interferences showed insignificant changes of spectra compared to that of no targeted sample. To investigate the interference of organic compounds in detail, toluene was chosen as a model for investigating the effect of interference at high concentration. The perylene/toluene ratios were adjusted to 1:1 (10⁻⁴ M : 10⁻⁴ M) and 1:10 (10⁻⁴ M : 10⁻³ M). β-CD dimer@Ag@SiO₂ NPs was added in a mixed solution of perylene and toluene. At a perylene/toluene ratio of 1:1, the mixture...
showed the characteristic peak of perylene at 1567 cm$^{-1}$. However, at a perylene/toluene ratio of 1:10, the mixed solution appeared in the peaks of toluene (10$^{-3}$ M) at 622, 785 and 1030 cm$^{-1}$ and perylene (10$^{-4}$ M) at 1567 cm$^{-1}$ in Raman spectra before washing. After washing, these peaks of toluene disappeared. The intensity at 1567 cm$^{-1}$ in Figure S5 (B) showed insignificant change in the presence of toluene. This result demonstrated that β-CD dimer@Ag@SiO$_2$ NPs can be a selective sensor.

To investigate the multiplex detection capability, we used pyrene and perylene as model PAHs for SERS detection. β-CD dimer@Ag@SiO$_2$ NPs were mixed with a mixture of pyrene (10$^{-3}$ M) and perylene (1 × 10$^{-3}$ M)
to $1 \times 10^{-4}$ M). Figure 4B shows an average Raman spectrum of the $\beta$-CD dimer@Ag@SiO$_2$ NPs in the presence of pyrene and perylene. As the perylene concentration increased, the intensities of the characteristic peaks of perylene at 1293 cm$^{-1}$ and at 1567 cm$^{-1}$ also increased, while the typical peak of pyrene at 1240 cm$^{-1}$ did not change markedly. Therefore, $\beta$-CD dimer@Ag@SiO$_2$ NPs could be used for multiplex detection of pyrene and perylene.

To demonstrate the stability of our sensor, we measured the intensity of perylene at 1567 cm$^{-1}$ after several washing steps. Acetonitrile, which was mentioned in the previous report for extracting PAHs in HPLC technique$^{41,42}$, was used as a solvent for washing.

Figure S6 showed the Raman spectra of $\beta$-CD dimer@Ag@SiO$_2$ in the $10^{-4}$ M perylene solution after each washing (Figure S6 (A)). The characteristic peak at 1567 cm$^{-1}$ was plotted in Figure S6 (B). After it was washed 5 times, the intensity of the perylene peak at 1567 cm$^{-1}$ was insignificantly changed. Perylene with the $\beta$-CD dimer@Ag@SiO$_2$ was stable after washing 5 times.

We prepared $\beta$-CD dimer-immobilized Ag assembly embedded silica NPs ($\beta$-CD dimer@Ag@SiO$_2$ NPs). The Ag@SiO$_2$ NPs were uniform in diameter (ca. 206 ± 10 nm) with 8.4 ± 3.7 nm Ag NPs on their surface. The $\beta$-CD dimer@Ag@SiO$_2$ NPs were able to detect perylene selectively at concentrations of $1 \times 10^{-2}$ to $10^{-2}$ M. The limit of detection was $10^{-4}$ M. The $\beta$-CD dimer@Ag@SiO$_2$ NPs can be used for multiplex detection of PAHs compounds, such as pyrene, perylene and so on. This method of combining an assembled structure and the ligands to recognize target materials has considerable potential for detection of chemicals or biomolecules.

Method

Preparation of assembly Ag-embedded silica nanoparticles (Ag@SiO$_2$ NPs). Approximately 180 nm Ag assembly-embedded silica NPs were synthesized using a method reported previously$^{24,43}$. First, silica NPs were prepared using the Stöber method and their surface was modified with thiol groups. Forty milliliters of 99.9% EtOH, 3 mL NH$_4$OH, and 1.8 mL TEOS were mixed and the solution stirred vigorously for 20 h at 25°C. Then, the mixture was centrifuged, washed with 95% EtOH, dispersed in absolute EtOH, and the volume adjusted to 50 mg mL$^{-1}$ silica NPs solution. Two hundred microliters of MPTS and 50 μL NH$_4$OH were added to 4 mL silica NPs solution (50 mg mL$^{-1}$) and the mixture was stirred vigorously for 12 h at 25°C. The solution was centrifuged, followed by washing with EtOH to remove excess reagent. Second, Ag NPs were embedded on the surface of thiolated silica NPs by reducing silver nitrate with octylamine and PVP in ethylene glycol. Five milligrams of PVP in 25 mL EG and 26 mg AgNO$_3$ in 25 mL EG were added to a thiolated silica NP (30 mg) suspension (final AgNO$_3$ concentration, is 15 mM). After thorough mixing, 41.4 μL octylamine were added and the resulting suspension was stirred for 1 h at 25°C. The resulting suspension was centrifuged, washed several times with EtOH, and dispersed in 3 mL absolute EtOH to obtain a 10 mg mL$^{-1}$ Ag@SiO$_2$ solution.

Synthesis of $\beta$-cyclodextrin dimer. $\beta$-Cyclodextrin dimer was synthesized according to a reported method$^{32}$. Mono-6-O-p-toluenesulfonyl-$\beta$-CD (1.00 g, 0.78 mmol), synthesized from $\beta$-CD by modifying a 6-hydroxy group on the primary side, was dissolved in anhydrous DMF (40 mL) with sodium iodide (1.20 g, 8 mmol) under nitrogen stirring for 5 h at 90°C. Acetone (25 mL) was added to obtain a precipitate, and the solvent was removed through reduced pressure (rotary evaporator) after filtration and washing. The resulting solid was dissolved in demineralized water (50 mL) and excess acetone was added to induce precipitation. After filtering and drying, the precipitate, which is 6-iodo-6-deoxy-$\beta$-CD (0.88 g, 0.71 mmol), was dissolved in 8 mL of dry DMF. The resulting solution was reacted with sodium sulfide (27.64 mg, 0.35 mmol) for 15 h at 80°C. The reaction mixture was precipitated with 200 mL of acetone after concentration under reduced pressure. The precipitates collected on a glass filter were dried under reduced pressure. Final $\beta$-CD was purified using a Bio-Gel P6 column after dissolving the precipitates in water. The dried final product was obtained with a yield of 13.5% from $\beta$-CD.

Preparation of $\beta$-CD dimer@Ag@SiO$_2$ NPs. To $\beta$-cyclodextrin dimer solution (5 mM in distilled water) was added 5 mg of assembled Ag embedded silica NPs. The suspension was stirred vigorously for 12 h at 25°C. The suspensions were then centrifuged and washed with EtOH, and dispersed in 1 mL absolute EtOH.

Interactions of $\beta$-CD dimer@Ag@SiO$_2$ NPs with PAHs. A perylene stock solution (100 mM) was prepared using dichloromethane as the solvent. Subsequently, a series of PAH solutions of various concentrations were obtained by dilution with dichloromethane. The prepared $\beta$-CD dimer@Ag@SiO$_2$ NPs (0.5 mg in 500 μL of EtOH) were mixed with 500 μL of dichloromethane solution containing various concentrations of PAHs and shaken for 30 min under ambient conditions.

Raman measurements. To evaluate the sensitivity of the synthesized SERS materials, the SERS signals were measured in a capillary tube using a DXR™ Raman Microscope system (Thermo Fisher Scientific, USA). The SERS signals were collected in a back-scattering geometry using a ×10 objective lens. A 532 nm diode-pumped solid-state laser was used as the photo-excitation source, with 10 mW laser power at the sample. Selected sites were measured randomly, and all SERS spectra were integrated for 5 s. The spot size of the laser beam was ~2 μm.

References

1. Conney, A. H. Induction of microsomal enzymes by foreign chemicals and carcinogenesis by polycyclic aromatic hydrocarbons: GHA Clowes Memorial Lecture. Cancer research 42, 4875–4917 (1982).
2. Plaza-Bolaños, P., Frenich, A. G. & Vidal, J. L. M. Polycyclic aromatic hydrocarbons in food and beverages. Analytical methods and trends. J Chromatogr A 1217, 6303–6326 (2010).
3. Nassar, H. F. et al. Atmospheric concentrations of polycyclic aromatic hydrocarbons and selected nitrated derivatives in Greater Cairo, Egypt. Atmospheric Environment 45, 7352–7359 (2011).
4. Hollosi, L. & Wenal, T. Development and optimisation of a dopant assisted liquid chromatographic-atmospheric pressure photoionisation-tandem mass spectrometric method for the determination of 15 + 1 EU priority PAHs in edible oils. J Chromatogr A 1218, 21–31 (2011).
5. Purcaro, G., Moret, S., Bučar-Miklavič, M. & Conte, L. S. Ultra-high performance liquid chromatographic method for the determination of polycyclic aromatic hydrocarbons in a passive environmental sampler. Journal of separation science 35, 922–928 (2012).
6. Olatunji, O. S., Fatoki, O. S., Opeolu, B. O. & Ximba, B. J. Determination of polycyclic aromatic hydrocarbons [PAHs] in processed meat products using gas chromatography–Flame ionization detector. Food chemistry 156, 296–300 (2014).
7. Moret, S. & Conte, L. S. Polycyclic aromatic hydrocarbons in edible fats and oils: occurrence and analytical methods. J Chromatogr A 882, 245–253 (2000).
8. Zielinska, B. & Samy, S. Analysis of nitrated polycyclic aromatic hydrocarbons. Anal Bioanal Chem 386, 883–890 (2006).
9. Schmidt, H. et al. Detection of PAHs in seawater using surface-enhanced Raman scattering (SERS). Mar Pollut Bull 49, 229–234 (2004).
10. Graham, D. & Goodacre, R. Chemical and bioanalytical applications of surface enhanced Raman spectroscopy. Chem Soc Rev 37, 883–884 (2008).
11. Halvorson, R. A. & Vikesland, P. J. Surface-enhanced Raman spectroscopy (SERS) for environmental analyses. Environ Sci Technol 44, 7749–7755 (2010).
12. Nie, S. & Emory, S. R. Probing single molecules and single nanoparticles by surface-enhanced Raman scattering. science 275, 1102–1106 (1997).
13. Kneipp, K. et al. Single molecule detection using surface enhanced Raman scattering (SERS). Physical review letters 78, 1667 (1997).
14. Li, J., Chen, L., Lou, T. & Wang, Y. Highly sensitive SERS detection of As3+ ions in aqueous media using glutathione functionalized silver nanoparticles. ACS Appl Mater Inter 3, 3936–3941 (2011).
15. Liang, H., Li, Z., Wang, W., Wu, Y. & Xu, H. Highly surface-roughened “flower-like” silver nanoparticles for extremely sensitive substrates of surface-enhanced Raman scattering. Adv Mater 21, 4614 (2009).
16. Tripp, R. A., Dhuy, R. A. & Zhao, Y. Novel nanostructures for SERS biosensing. Nano Today 3, 31–37 (2008).
17. Jain, P. K., Lee, K. S., El-Sayed, I. H. & El-Sayed, M. A. Calculated absorption and scattering properties of gold nanoparticles of different size, shape, and composition: applications in biological imaging and biomedicine. The Journal of Physical Chemistry B 110, 7238–7248 (2006).
18. Dondapati, S. K. et al. Label-free biosensing based on single gold nano stars as plasmonic transducers. ACS Nano 4, 6318–6322 (2010).
19. Nuse, G. J. et al. Label-free plasmonic detection of biomolecular binding by a single gold nanorod. Anal Chem 80, 984–989 (2008).
20. Larsson, E. M., Alegret, J., Käll, M. & Sutherland, D. S. Sensing characteristics of NIR localized surface plasmon resonances in gold nanorings for application as ultrasensitive biosensors. Nano Lett 7, 1256–1267 (2007).
21. Murphy, T., Schmidt, H. & Kronfeldt, H. D. Use of sol-gel techniques in the development of surface-enhanced Raman scattering (SERS) substrates suitable for in situ detection of chemicals in sea-water. Applied Physics B 69, 147–150 (1999).
22. Fan, M. & Brolo, A. G. Silver nanoparticles self assembly as SERS substrates with near single molecule detection limit. Phys Chem Chem Phys 11, 7381–7389 (2009).
23. Bosnick, K. A., Jiang, J. & Brus, L. E. Fluctuations and local symmetry in single-molecule rhodamine 6G Raman scattering on silver nanocrystal aggregates. The Journal of Physical Chemistry B 106, 8096–8099 (2002).
24. Kim, J.-H. et al. Nanoparticle probes with surface enhanced Raman spectroscopic tags for cellular cancer targeting. Anal Chem 78, 6967–6973 (2006).
25. Szejtli, J. Introduction and general overview of cyclodextrin chemistry. Chem Rev 98, 1743–1754 (1998).
26. Xie, Y. et al. Selective SERS detection of each polycyclic aromatic hydrocarbon (PAH) in a mixture of five kinds of PAHs. J Raman Spectrosc 42, 945–950 (2011).
27. Breslow, R. & Zhang, B. Cholesterol recognition and binding by cyclodextrin dimers. J Am Chem Soc 118, 8495–8496 (1996).
28. Zhao, Y. et al. Molecular selective binding of aliphatic oligopeptides by bridged bis (β-cyclodextrin) s with aromatic diamine linkers. J Mol Struct 930, 73–77 (2009).
29. Li, P., Liu, Y., Wang, X., Tang, B. A new FRET nanoprobe for trypsin using a bridged β-cyclodextrin dimer–dye complex and its biological imaging applications. Analyst 136, 4520–4525 (2011).
30. Zhang, Y.-M., Chen, H.-Z., Chen, Y., Ding, F. & Liu, Y. Molecular binding behaviors of triazole-bridged bis (β-cyclodextrin) s towards cinchona alkaloids. New J Chem 37, 1554–1560 (2013).
31. Yang, L. et al. Bridged β-cyclodextrin-functionalized MWNTs with higher supramolecular recognition capability: The simultaneous electrochemical determination of three phenols. Biosensors and Bioelectronics 68, 617–625 (2015).
32. Cho, E., Jeong, D., Paik, H.-D. & Jung, S. Solubility Enhancement of Flavonols in the Inclusion Complex with Thioether-bridged Dimeric β-Cyclodextrins. Bull Korean Chem Soc 35, 2487 (2014).
33. Szczepanski, J., Chapo, C. & Vála, M. Visible and infrared spectra of matrix-isolated perylene 3G Raman scattering on silver nanocrystal aggregates. The Journal of Physical Chemistry B 106, 8096–8099 (2002).
34. Rao, C., Venkataraman, R. & Kasturi, T. Contribution to the infrared spectra of organosulphur compounds. Canadian journal of chemistry 42, 36–42 (1964).
35. Nakajima, A. Effect of ethanol on the inclusion complex formation between. BETA.-cyclodextrin and pyrene. Bulletin of the Chemical Society of Japan 57, 1143–1144 (1984).
36. Xie, Y. et al. Sensing of polycyclic aromatic hydrocarbons with cyclodextrin inclusion complexes on silver nanoparticles by surface-enhanced Raman scattering. Analyst 135, 1389–1394 (2010).
37. Cho, E. et al. Novel magnetic nanoparticles coated by benzene-and β-cyclodextrin-bearing dextran, and the sorption of polycyclic aromatic hydrocarbon. Carbohydr Polym 133, 221–228 (2015).
38. Choi, J. M. et al. Hydroxypyropyl cyclic (5–(1–2)–5–d)-glucans and epichlorohydrin β-cyclodextrin dimers as effective carbohydrate-solubilizers for polycyclic aromatic hydrocarbons. Carbohydrate research 401, 82–88 (2015).
39. Shinohara, H., Yamaikita, V. & Ohno, K. Raman spectra of polycyclic aromatic hydrocarbons. Comparison of calculated Raman intensity distributions with observed spectra for naphthalene, anthracene, pyrene, and perylene. J Mol Struct 442, 221–234 (1998).
40. Lu, Y., Yao, G., Sun, K. & Huang, Q. β-Cyclodextrin coated SiO2@Au@Ag core–shell nanoparticles for SERS detection of PCBs. Phys Chem Chem Phys 17, 21149–21157 (2015).
41. Manoli, E. & Samara, C. Polycyclic aromatic hydrocarbons in waste waters and sewage sludge: extraction and clean-up for HPLC analysis with fluorescence detection. Chromatography 43, 135–142 (1996).
42. Papadopoulos, A., Paton, G. I., Reid, B. J. & Semple, K. T. Prediction of PAH biodegradation in field contaminated soils using a cyclodextrin extraction technique. Journal of Environmental Monitoring 9, 516–522 (2007).
43. Kim, H.-M. et al. Large scale synthesis of surface-enhanced Raman scattering nanoprobes with high reproducibility and long-term stability. J Ind Eng Chem 33, 22–27 (2016).

Acknowledgements

This work was supported by the Konkuk University in 2014.
Author Contributions
E.H. and B.-H.J. conceived and designed the experiments. E.H., D.J. and M.G.C. et al. performed the experiments. E.H., X.-H.P., J.M.C., H.-M.K. and H.K. et al. analyzed the data. E.H., X.-H.P., Y.-S.L., D.H.J., S.J. and B.-H.J. wrote the manuscript.

Additional Information
Supplementary information accompanies this paper at http://www.nature.com/srep

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Hahm, E. et al. β-CD Dimer-immobilized Ag Assembly Embedded Silica Nanoparticles for Sensitive Detection of Polycyclic Aromatic Hydrocarbons. Sci. Rep. 6, 26082; doi: 10.1038/srep26082 (2016).

This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/