## Supporting Information

## S1-SAXS data of templating microemulsions

The SAXS measurements were performed on a SAXSess high-flux small-angle X-ray scattering instrument (Anton Paar, Austria), attached to a PW3830 X-ray generator (PANalytical) with a sealed-tube anode ( $\mathrm{Cu} \mathrm{K} \alpha$ wavelength of 0.1542 nm ). The generator was operated at 40 kV and 50 mA . The SAXSess camera was equipped with a line collimator block and all measurements were performed at vacuum conditions for an intense and monochromatic primary beam with low background. A semitransparent beam stop was used to enable the measurements of an attenuated primary beam for the exact definition of the zero scattering vector and transmission correction. Vacuum-tight refillable quartz capillaries ( 1 mm diameter, sample volume $\leq 100 \mu \mathrm{~L}$ ) were used in order to determine the size and shape of the w/o- microemulsion containing the metal salts and the reducing agent. All experiments were performed at $T_{\text {wefb. }}$. The sample temperature was controlled with a thermostatted sample holder unit (TCS 120, Anton Paar). The 2-D scattered intensities were recorded on a CCD detector (Princeton Instruments) and were converted via SAXSQuant software (Anton Paar) to one dimensional scattering curves as a function of the magnitude of the scattering vector $q=(4 \pi / \lambda) \sin (\theta / 2)$, where $\theta$ is the total scattering angle. All intensities were transmission-calibrated by normalizing the attenuated primary intensity at $q=0$ to unity and were corrected the background scattering from the capillary and the solvent (octane).

SAXS was used in order to measure the diameter of the w/o- microemulsions containing 13 mM of $\mathrm{H}_{2} \mathrm{PtCl}_{6}+13 \mathrm{mM}$ of $\mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}, 13 \mathrm{mM}$ of $\mathrm{H}_{2} \mathrm{PtCl}_{6}+13 \mathrm{mM}$ of $\mathrm{Bi}\left(\mathrm{NO}_{3}\right)_{3}$ and 320 mM of $\mathrm{NaBH}_{4}$ at $T_{\text {wefb. }}$. The droplet diameter of the w/o- microemulsion containing the metal salts and the reducing agent are presented in Table A1. In Figure A1 the scattering curves of w/o- microemulsions are presented in a double logarithmic plot. In all samples, an initial slope of zero was observed indicating that the w/o- microemulsions have a globular shape. The diameter of the droplets was determined by Guinier extrapolation (extrapolation to zero angle, $q=0.06-0.4 \mathrm{~nm}^{-1}$ ) assuming that the droplets are homogeneous spheres.

Table S1. Diameter of the w/o-microemulsions at $w_{\mathrm{A}}=0.08$ containing the metal salts and the reducing agent determined by SAXS.

| Metal Salts | w/o- microemulsion diameter / nm |
| :---: | :---: |
| $13 \mathrm{mM} \mathrm{H}_{2} \mathrm{PtCl}_{6}: 13 \mathrm{mM} \mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}$ | $26.2 \pm 1.0$ |
| $13 \mathrm{mM} \mathrm{H}_{2} \mathrm{PtCl}_{6}: 13 \mathrm{mM} \mathrm{Bi}\left(\mathrm{NO}_{3}\right)_{3}$ | $26.5 \pm 1.0$ |
| $320 \mathrm{mM} \mathrm{NaBH}_{4}$ | $25.0 \pm 1.0$ |



Figure S1. (a) SAXS curves of the w/o- microemulsion containing ( $\cdots$ ) $13 \mathrm{mM} \mathrm{H}_{2} \mathrm{PtCl}_{6}$ : $13 \mathrm{mM} \mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2},(-) 13 \mathrm{mM} \mathrm{H}_{2} \mathrm{PtCl}_{6}: 13 \mathrm{mM} \mathrm{Bi}\left(\mathrm{NO}_{3}\right)_{3}$ and $(--) 320 \mathrm{mM} \mathrm{NaBH}_{4}$ at $w_{\mathrm{A}}$ $=0.08$.

## S2-EDX analysis

Table S2. EDX analysis of the atomic composition of samples 1a, 1b, 1c, 1d, 1e (s. Table 1).

| $\begin{aligned} & \text { Sample 1a } \\ & / \mathrm{nm}^{2} \end{aligned}$ | $\begin{gathered} \mathrm{Pt}: \mathrm{Pb} \\ \text { atomic ratio } \end{gathered}$ | $\begin{gathered} \text { Sample 1b } \\ / \mathbf{n m}^{2} \end{gathered}$ | Pt:Pb <br> atomic ratio | $\begin{aligned} & \text { Sample 1c } \\ & / \mathbf{n m}^{2} \end{aligned}$ | $\begin{gathered} \mathrm{Pt}: \mathrm{Pb} \\ \text { atomic ratio } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 20 | 12:88 | 5 | 9:91 | 100 | 55:45 |
| 20 | $72: 28$ | 200 | 65:35 | 10 | 72: 28 |
| 200 | 68:32 | 10 | $20: 80$ | 20 | $63: 37$ |
| 20 |  | 10 |  | 20 |  |
| 5 | 74:26 | 10 | $70: 30$ | 20 | 73: 27 |
| 20 | 69:31 | 20 | 59:41 | 10 | 62:38 |
| 20 | 66:34 |  |  | 15 | 66:34 |
|  |  |  |  | 50 | 9:91 |
| Sample 1d/ | $\begin{gathered} \mathrm{Pt}: \mathrm{Pb} \\ \text { atomic ratio } \end{gathered}$ |  | Sample 1e/ $\mathrm{nm}^{\mathbf{2}}$ |  | Pt:Pb <br> atomic ratio |
| 100 |  | 45 | 10 |  | 50: 50 |
| 10 |  |  | 15 |  | 50:50 |
| 20 |  | 35 | 5 |  | 39:61 |
| 20 |  | 37 | 10 |  | 59:41 |
| 10 |  | 35 | 10 |  | 61:38 |
| 15 |  | 34 | 20 |  | 52: 48 |
| 20 |  | 36 | 20 |  | 50 : 50 |
| 50 |  |  | 30 |  | 61:38 |
|  |  |  | 20 |  | 47: 53 |
|  |  |  | 20 |  | 48:52 |
|  |  |  | 20 |  | 40:60 |
|  |  |  | 50 |  | 5:95 |

Table S3. EDX analysis of the atomic composition of samples $2 \mathrm{a}, 2 \mathrm{~b}$ and 2 c (s. Table 1).

| Sample 2a <br> $/ \mathbf{n m}^{2}$ | $\mathbf{P t}: \mathbf{B i}$ <br> atomic ratio | Sample 2b <br> $/ \mathbf{n m}^{2}$ | $\mathbf{P t}: \mathbf{B i}$ <br> atomic ratio | Sample 2c <br> $/ \mathbf{n m}^{2}$ | $\mathbf{P t}: \mathbf{B i}$ <br> atomic ratio |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 10 | $81: 19$ | 10 | $46: 54$ | 20 | $40: 60$ |
| 10 | $75: 25$ | 10 | $50: 50$ | 10 | $21: 79$ |
| 10 | $87: 13$ | 20 | $56: 44$ | 15 | $38: 62$ |
| 10 | $85: 15$ | 5 | $49: 51$ | 8 | $43: 57$ |
| 10 | $87: 13$ | 10 | $52: 48$ | 20 | $64: 36$ |
| 20 | $81: 19$ | 20 | $47: 53$ | 10 | $60: 40$ |
| 20 | $63: 37$ | 20 | $50: 50$ | 15 | $43: 57$ |
| 10 | $86: 14$ | 3 | $50: 50$ | 10 | $64: 36$ |
| 5 | $90: 10$ |  |  | 10 | $59: 41$ |

