

Electronic Supplementary information for

Determination of arsenic in biological samples by slurry sampling hydride generation atomic fluorescence spectrometry using *in-situ* dielectric barrier discharge trap

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Supplementary data

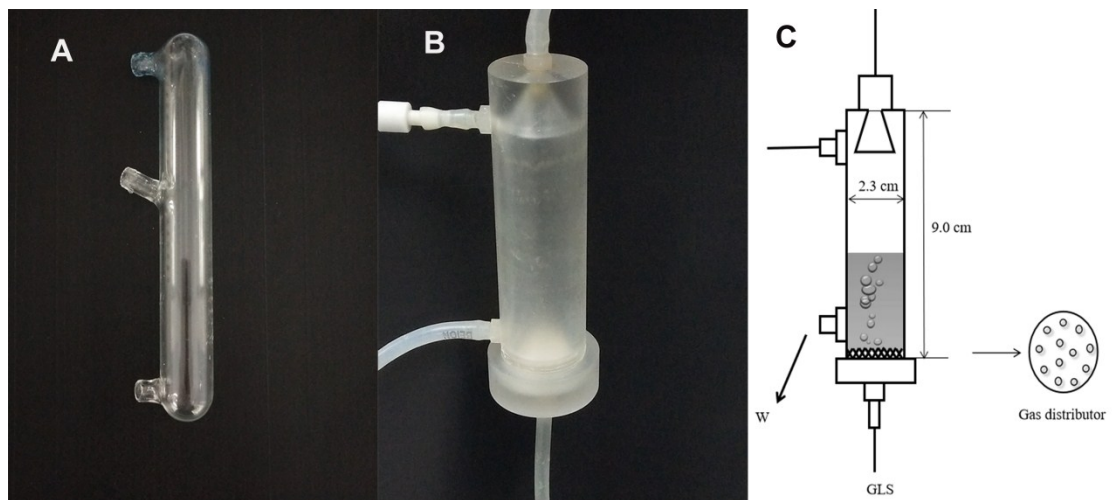


Figure S-1 The pictures of two different GLS apparatuses. Panel A shows the original GLS for the AFS-9130 instrument named GLS-A; Panel B shows the GLS used in this work named GLS-B; Panel C shows the detail structure diagram of GLS-B.

To investigate the effect of different ultrasonic times on arsenic intensity by SLS-HG-*in situ* DBD-AFS, 0.8 g *scenedesmus obliquus* samples were diluted to 20 mL with 5% HCl (v:v) in 50 mL centrifuge tube, and then disrupted for 2 min using a high speed tissue disperser to form a mixture sample, respectively. After an ultrasound water bath at 25 °C for different times, a 2 mL slurry was introduced into the SLS-HG system as manual shaking to keep suspension for the next measurement. The results are shown in Fig.S-2. With the increase of ultrasonic time from 1 to 10 min, the arsenic intensity went up obviously; and the plateau appeared from 10 to 20 min. So, 10 min was chosen as the optimal ultrasonic time for the next experiment.

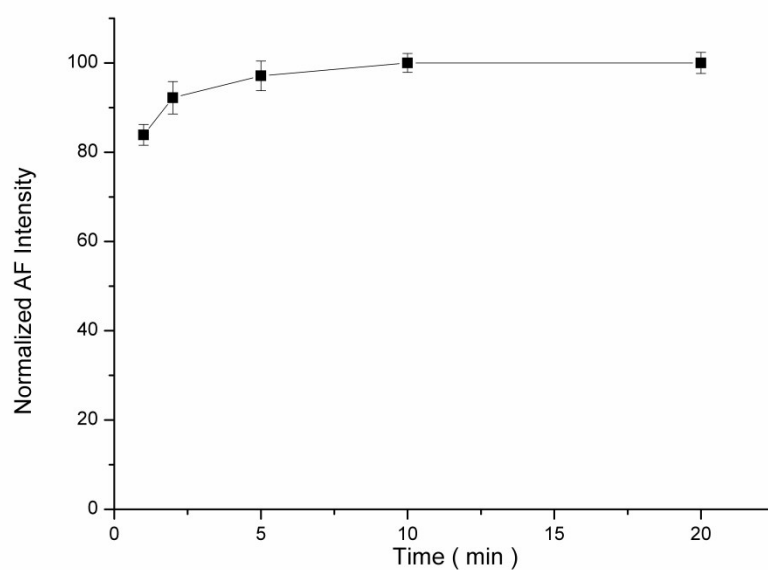


Figure S-2. Effect of different ultrasonic times on detecting arsenic by SLS-HG-*in situ* DBD-AFS. The AF intensity at 10 min is set as 100, and the others are normalized.