

開發用具有原子尺度共伴化學效應觸媒並用於氫氣儲存/ 轉化以及氣凝膠細胞損傷修補

Atomic synergetic collaboration catalysts for H₂ storage and evolution toward H₂ aerosol technique in biomedical therapy

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We anticipate to develop a microfluidic system combining nanocatalyst (NCs) and carbon nanosphere (CNS) the direct photocatalytic H₂ production and the slow release of H₂ to the cancel cell treatment. The ROS is an important biophysical signature for the control and the manipulating the function for most types of components in the cancel cells. By interacting with the H₂ molecules, the chemical characteristic of ROS will be changed so that the bio-activity of the cancel cell will be suppressed. With these understandings, based on the physical characteristics of existing materials in our team, two H₂ slow release technologies are proposed for the development of implant medical applications including the hybrid composites of organic-inorganic semiconductor photocatalytic catalysts and the H₂ storage CNS. In the practical scenarios, the former will be sealed in a transparent membrane capsule and the later will be filled with H₂ prior to their deployment in the treating organs. To virtualize the proposed concept, the validation of the conceptual prototype design including the functional test, the process optimization and theoretical materials calculation prediction. Topics include the photocatalytic H₂ production yield on the hybrid composites, the HER performance of the atomic catalysts, the H₂ storage capacity of the CNS as well as the subsequent biomedical responses. The correlation between

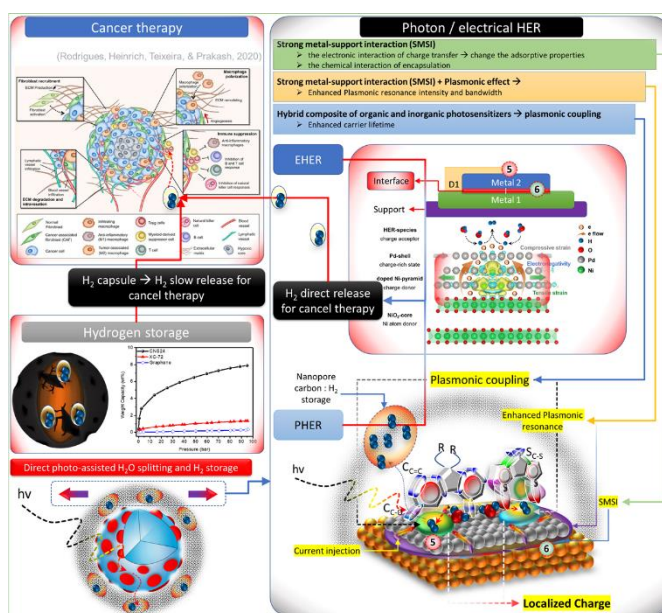


Figure 1 Correlation between CNS, hybrid composites of organic/inorganic semiconductor catalysts and the H₂ treating on cancer therapy

the aforementioned research topics are shown in Figure 1 and the corresponding results are summarized later.

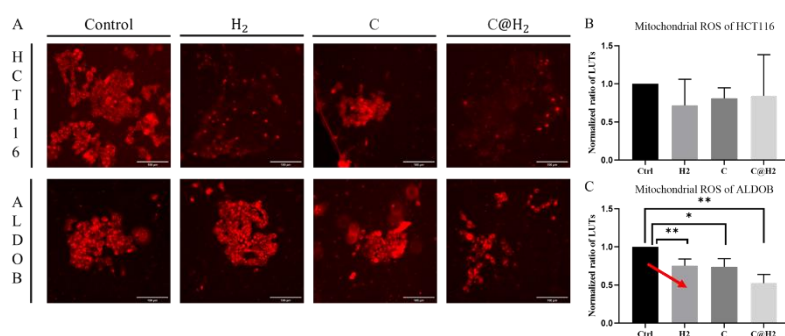


Figure 2 HCT116 and ALDOB cell degradation test results.

In cancer cell treatment, the MitoSOX test reveals that a low level of H₂ ~3ppm reduces the ·O²⁻ radical concentration by 70% and influences the cell degradation mechanism. In addition, the combination of H₂ and 5-FU further degrades the cell by 38% thus showing possibility of such a protocol in the cancer therapy. Slow release of H₂ without outer interference is a key factor in the long-term treatment. Our results indicate that with the insertion of H₂ filled CNS the cell degradation can be further increased by 50% as compared to the H₂ and 5-FU control (Figure 2).

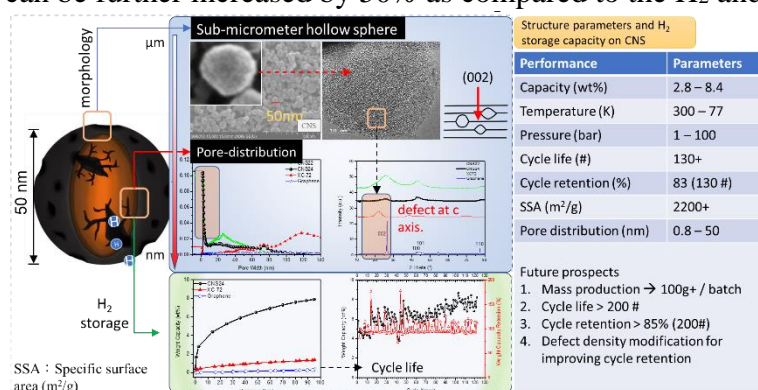


Figure 3 The physical structure parameters and the H₂ storage capacity of CNS.

The H₂ storage capacity, cycle life and the corresponding physical structures of CNS are shown in **Figure 3**. Accordingly, the H₂ capacity is 2.6 and 8.1 wt% respectively at 77K and 300K at 100 bars. These features are dominated by the defect density, the pore ratio, pore distribution and the specific surface area of the carbon material. Our results show that the proposed CNS

possesses the equilibrium physical structure parameters to the state-of-the-art carbon materials with highest H₂ storage capacity. Results of the corresponding topics are summarized in **Table 1**.

Table 1 Summary for the published paper in this project

Topic	SCI paper	Conference paper	in preparation	patent
HER	4			
H ₂ storage			2	1
H ₂ biomedical application	1	2		

Representative References

- [1] J.-R. Zhang, T.-S. Wu, H.-V. Thang, K.-Y. Tseng, **H.-Y. T. Chen*** & Y.-K. Peng*. Chemical state tuning of active sites over single-atom/cluster nanozymes: a balance between their maximum utilization and specific reactivity as peroxidase/oxidase mimetics. *Small*, (2021)..
- [2] Haolin Li, Sheng Dai, Dinesh Bhalothia, Alice Hu*, Jyh-Pin Chou* and **Tsan-Yao Chen***. Interfacial atomic Ni tetragon intercalation in a NiO₂-to-Pd hetero-structure triggers superior HER activity to the Pt catalyst, *Journal of Materials Chemistry A*, 9 (2021) 12019-12028.