

Photocatalytic water-splitting and simultaneous gas segregation from dual-sided photocatalytic membrane.
 Stephen L. Rhoden, Howard D. Mettee and Clovis A. Linkous

Youngstown State University,
 1 University Plaza,
 Youngstown Ohio, 44555.

It is now well understood that efficient solar fuel technologies require increasingly robust semiconductor systems that are capable of generating the necessary photovoltages for transformation of precursors like H_2O and CO_2 into valuable chemicals such as hydrogen and methanol. Heterostructured membranes and co-catalyst systems have shown the most promise in this respect. Also, the separation of H_2 and O_2 becomes a significant safety and efficiency concern in water splitting, if these gases are to be stored and used later for hydrogen fuel cell operation.

CdTe and WO_3 are both capable of utilizing the Sun's abundant insolation. Electrodeposition of thin films of both semiconductors has been performed on either side of a flexible stainless steel substrate (Figure 1); after specific heat treatments, the perforations in the stainless steel plates were filled with Nafion[®], to allow for a sustainable electrolyte pH gradient in each half-cell. Utilizing a 1000 W Xe arc lamp to illuminate both sides, unassisted water-splitting in 0.1 M H_2SO_4 was achieved, producing H_2 on the CdTe photocathode, and the WO_3 photoanode producing O_2 . Typical gas product yields measured using an Inficon GCMS leak detector showed 44 ppm cm^{-2} H_2 and 39 ppm cm^{-2} O_2 . The unexpectedly higher O_2 content was thought to be due to contamination from air. Our current research efforts are to calibrate a self-purging continuous argon flow system. The semiconductors are also being characterized using SEM-EDX, XRD and cyclic voltammetry.

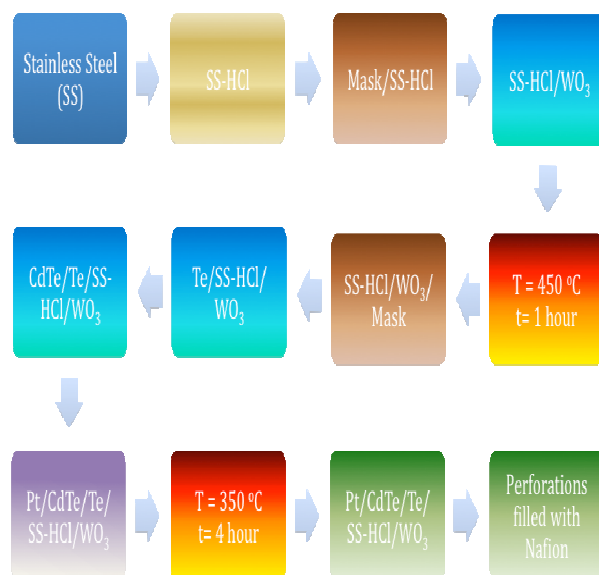


Figure 1. Process fabrication diagram for dual-sided photocatalytic membrane.