

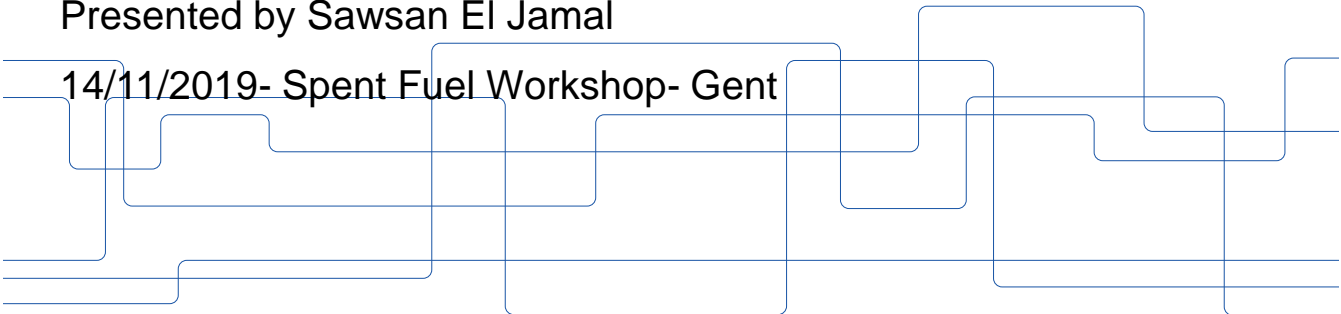


Uranium Carbide, a Potential Nuclear Fuel

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- Purpose:
 - Establish a knowledge base for the behavior some Gen-IV fuel candidates in aqueous systems
 - Understand the release of radionuclides upon accidents and in geological repositories for used nuclear fuel
- The fuel types of main interest :uranium carbides and uranium nitrides:
Compared to standard UO_2 based fuels Uranium Carbide UC has:
 - better thermal conductivity
 - higher fissile material density



Aim

- Elucidate the mechanisms and kinetics of dissolution of the fuel matrixes (UC and UN) in water :
 - dissolution under oxic and anoxic conditions
 - radiation-induced dissolution
 - impact of fission products and solutes in the water
- ➔ To understand the stability of new fuels in aqueous systems



Starting point:

- synthesis of uranium carbides and uranium nitrides
- manufacturing of pellets
- doping pellets with non-radioactive isotopes of fission products as models for irradiated nuclear fuel

Synthesis of UC

Mixing precursors $\text{UO}_2\text{:C}$



Pressing mixed powder into green pellets



Carbo-thermal reduction at 1500C under 5% H_2 in N_2



Characterization of the crushed pellet with X-Ray Powder diffraction (XRD)



Preparing dense pellets by Spark Plasma Sintering



Stability of UC

- Dissolution of UC pellets should be studied experimentally under:
 - H_2O_2 exposure
 - GAMMA irradiation



Model Carbides

- Preliminary studies are based on other carbides such as Titanium Carbide(TiC) to get insights on
 - the manufacturing of these carbides
 - understand the behavior of carbides under oxidative dissolution conditions
- Reactivity of Tungsten carbide is also studied for comparison

Synthesis of TiC

Mixing precursors $\text{TiO}_2\text{:C}$ (1:3,3)

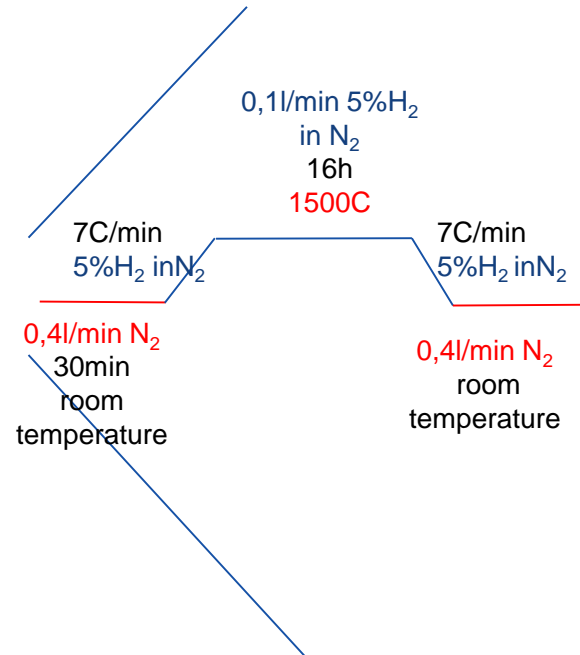
Drying at 80C

Pressing mixed powder into green pellets

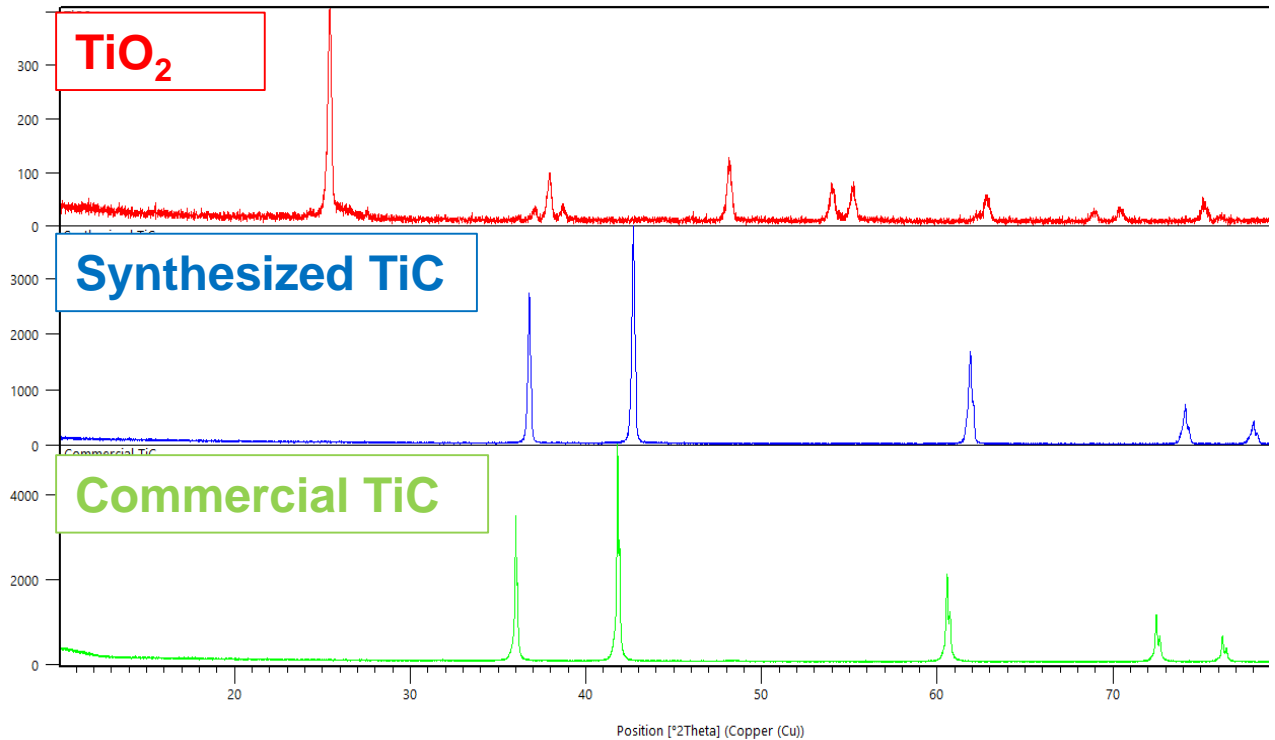
Carbo-thermal reduction for 16h at 1500C
under 5% H_2 in N_2

Characterization of the crushed pellet with X-Ray Powder diffraction (XRD)

Preparing dense pellets by Spark Plasma Sintering



XRD-Diffractogram

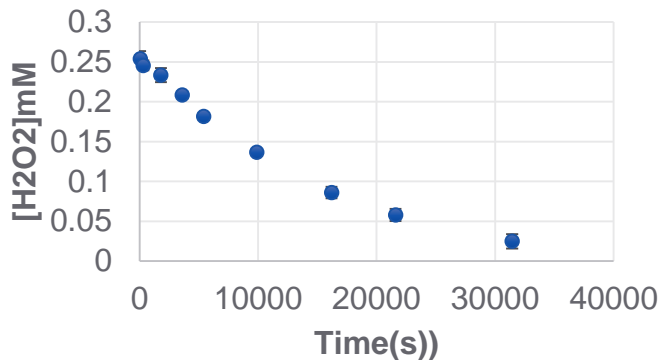


Reactivity of TiC in the presence of H_2O_2

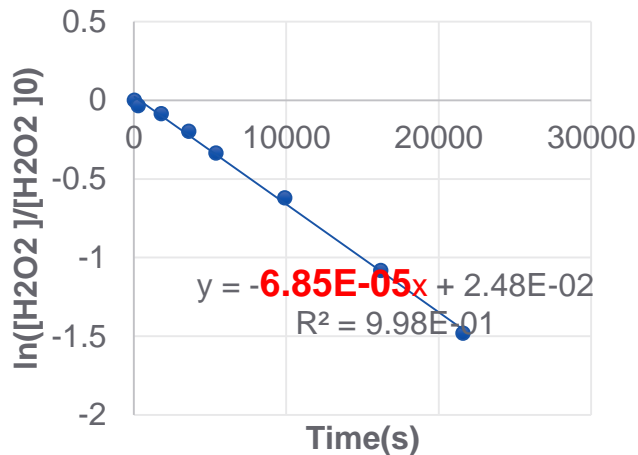
- The Ghormley triiodide method :measure H_2O_2 concentration.
 - The absorbance of I_3^- is subsequently measured spectrophotometrically at 350 nm.
 - A linear correlation between the absorbance and the H_2O_2 concentration was obtained by a calibration curve to calculate $[\text{H}_2\text{O}_2]$
- H_2O_2 decomposition was studied for different amount of of TiC powder (50mg, 0,25g 0,5g and 0,75g) in a 25 ml volume of H_2O .

Reactivity of TiC in the presence of H_2O_2

50mg TiC in 25ml
 H_2O
0,26mM H_2O_2

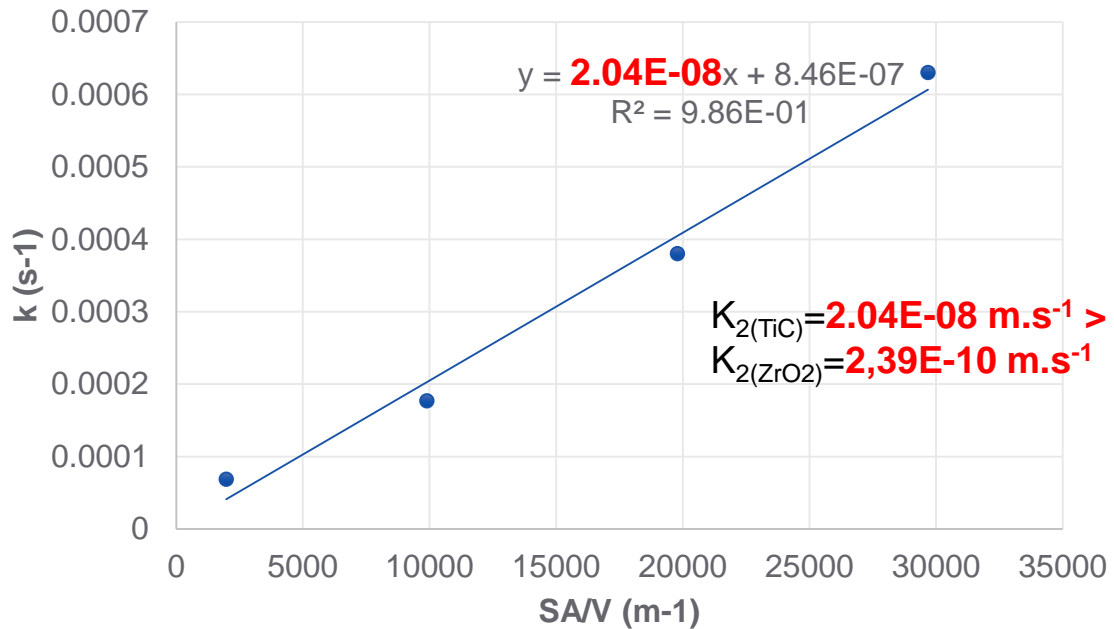


first order rate
constant

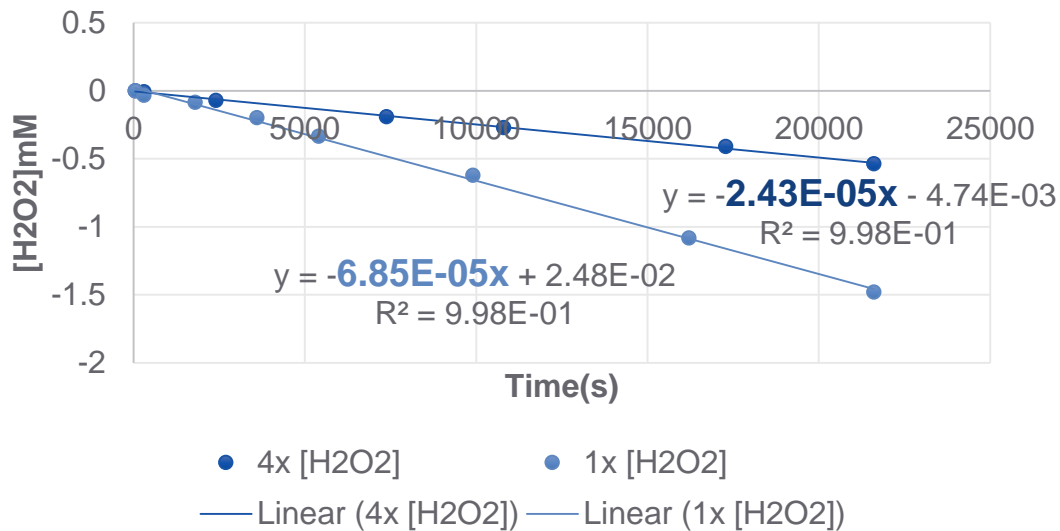


Reactivity of TiC in the presence of H_2O_2

second order rate constant

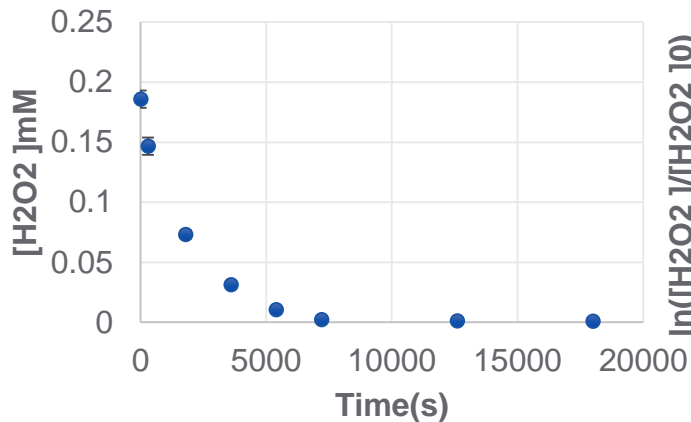


comparing first order rate constant 50mg TiC

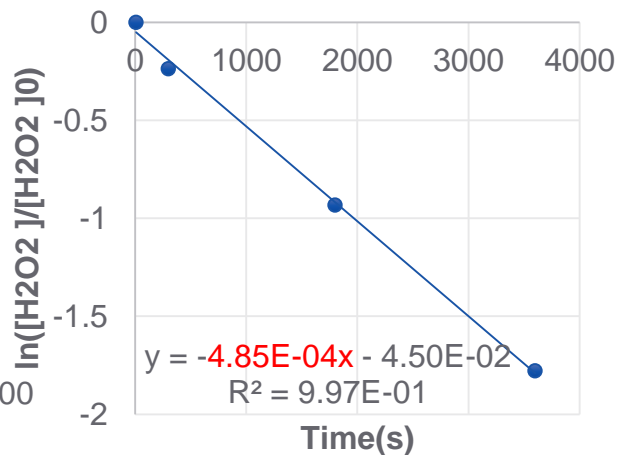


Reactivity of WC in the presence of H_2O_2

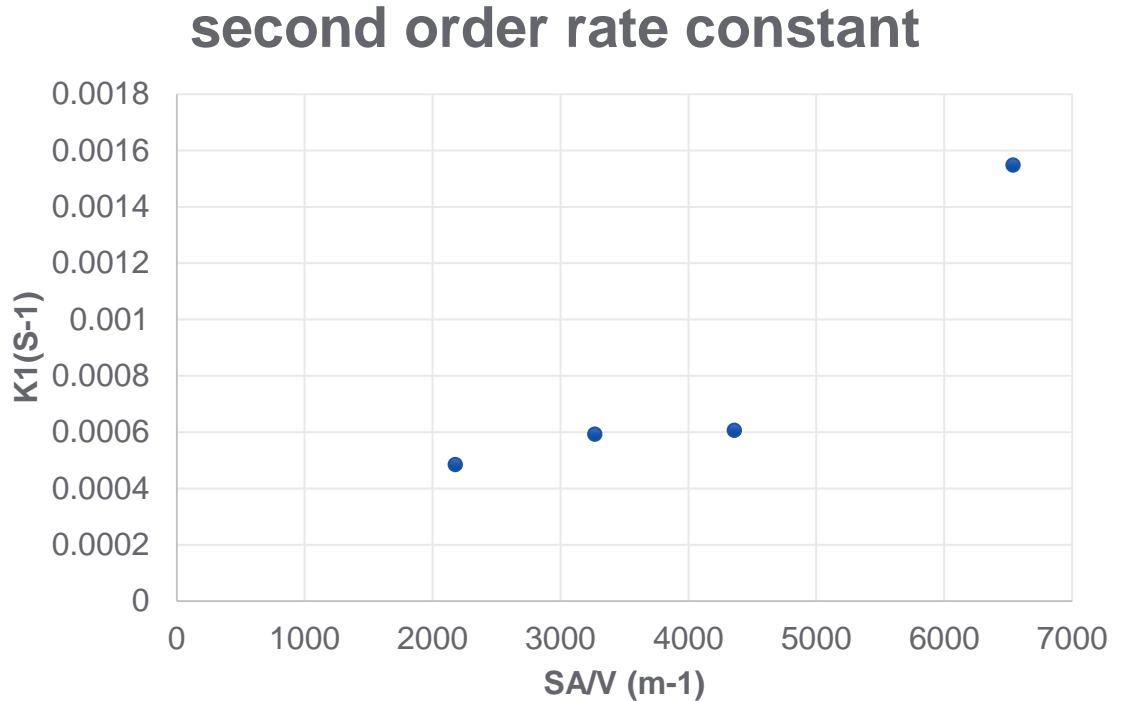
50mg WC in 25 ml H_2O
0,2mM H_2O_2



first order rate
constant



Reactivity of WC in the presence of H_2O_2





Conclusion

- Carbothermal reduction is an effective process in producing carbides
- TiC and WC are reactive once exposed to H_2O_2
- Carbides seems to be more reactive than the oxides



Acknowledgments

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