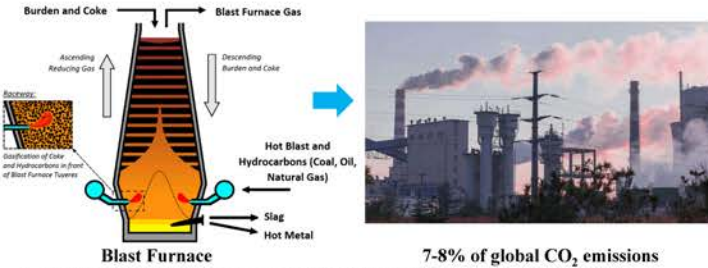


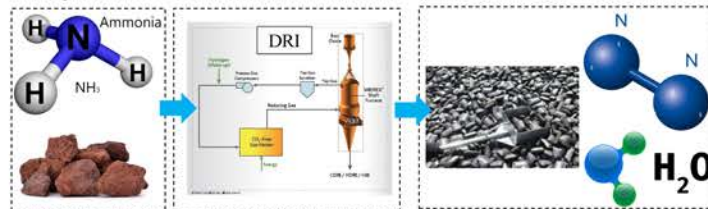
FENEX CRC THEME

Program 2 – Hydrogen Exports and Value Chains

IMPACT AND IMPORTANCE



- Pathways to reduce CO₂ emissions from ironmaking:
 - Ore-carbon agglomerates;
 - Carbon capture and storage (CCS);
 - Direct reduction process (e.g. FINEX, MIDREX) + Electron arc furnace (EAF)
 - Carbon-free or renewable reductant (H₂, NH₃ or biomass gas).
- Green H₂ is an effective carbon free ironmaking process. However, the storage and transportation of H₂ encounter the high cost and safe problems.
- Ammonia (NH₃), a carbon free reductant and hydrogen carrier, offers a path to ironmaking decarbonisation and has mature storage and transportation system.
- NH₃ direct reduction of iron ore is not yet fully-developed.
- Huge iron/steel global market offers an opportunity for Australian iron ore, H₂ and NH₃ export.



PROJECT AIMS

- Investigation of the behaviour and mechanisms of NH₃ direct ironmaking
 - Determine reaction kinetics for the reduction of iron ores with NH₃.
 - Understand reduction behaviour and evolution of phases using characterisation techniques.
 - Inspect the conversion of impurities and physical parameters evolution in iron ore and their effects on DRI process.
- Study the low-temperature pyrophoricity and surface passivation of NH₃ DRI.
- Investigate the reduction mechanisms of typical WA iron ore by single particle modelling.
- Examine the products through benchmarking NH₃/H₂ DRI against other reductants.

LITERATURE REVIEW

- Very limited experimental studies on NH₃ reduction of iron ore.
- NH₃ direct reduction of hematite could began from 430 °C.
- NH₃ would dissociate at high temperatures, as well as the catalysis of iron/iron oxides.
- Produced H₂ contributed further to reduction and could be dominant at high temperatures.
- NO_x formation and diminishment at high temperatures in NH₃ DRI process remains unclear.
- The impurities will affect the reduction process in H₂ DRI, but the effect in NH₃ is unclear
- Iron nitride is unstable at low temperatures but could influence the NH₃ dissociation mode.
- Porosity of produced DRI and reduction temperature are important triggers to pyrophoricity.
- The strong endothermicity of the NH₃ reduction presents a key technical challenge.

RESEARCH METHODOLOGY

- Anti-corrosive thermogravimetric analyser
- Preliminary mechanisms of NH₃/H₂ DRI
- Determination of rate equations for kinetic study
- Fixed-bed reactor
 - Global reduction behaviour of ore gangues
 - Product gas composition analysis
 - Online GC/MS, FTIR – characterisation
 - NO_x analyser – NO, NO₂
 - Solid product characterisation and impurity evolution
 - XRD – Mineral composition and crystal structure
 - Optical microscope and SEM – Morphology
 - TEM – Crystal information
 - CT/BET – Porosity and pore structure
 - ICP-OES – Chemical composition
- Single particle reaction model

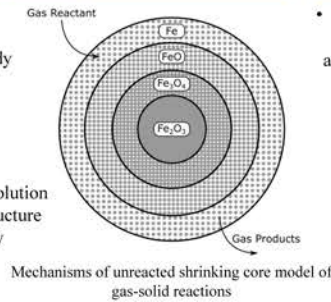


Figure 1 Anti-corrosive thermogravimetric analyser

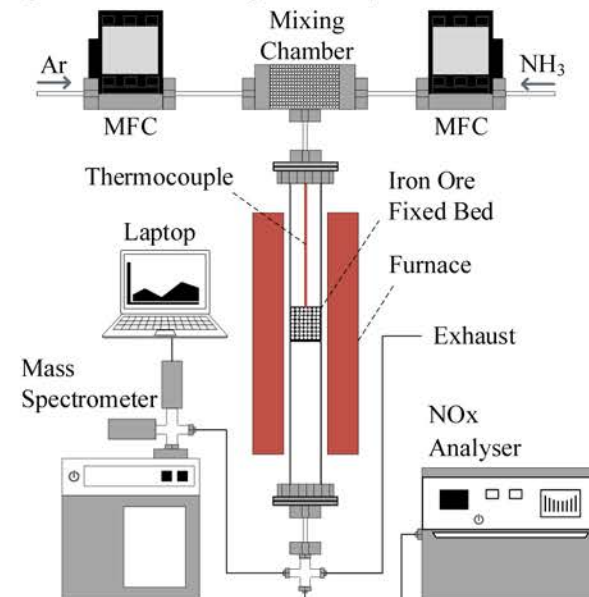


Figure 2 Fixed bed reactor setup

PRELIMINARY WORK

- Reduction degree of WA iron ore using different H₂ concentration and results of numerical simulation

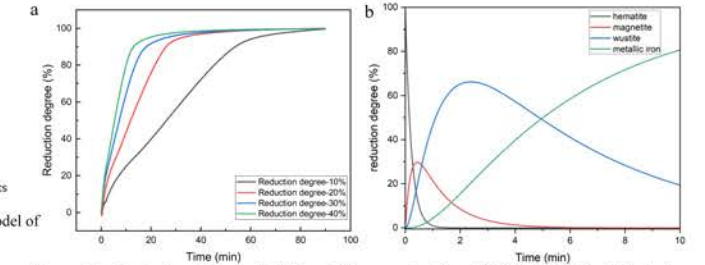


Figure 3 a Reduction degree at different H₂ concentration (700 °C) in TGA; b Reduction degree against time in NH₃ DRI of hematite in COMSOL Multiphysics

- Optical microscope observation

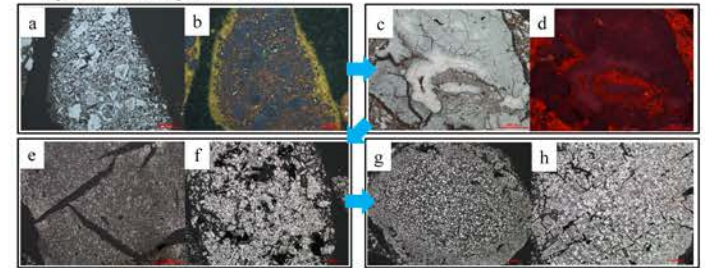


Figure 4 Photomicrographs of samples, a (plain polarized light, PPL) & b (cross polarized light, XPL) raw sample; c (PPL) & d (XPL) control sample; e & f partial reduced sample; g & h fully reduced sample

- Hematite and goethite are the main ore minerals.
- Before the reduction goethite will convert to hematite due to the dehydration forming cracks.
- Metallic iron will form along the shell of the ore and the edge of the cracks.
- The final product is more porous compared with the partial reduced samples.

PROJECT RESEARCHERS

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PROJECT ORGANISATIONS

