

BIO-REMEDIATION OF MINE WATER

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South Africa



our future through science

Introduction

Mining History in South Africa

- Gold Mining started in the late 1880's on the farm Langlaagte, west of Johannesburg
- Soon thereafter coal mining started in the Witbank Area as well as in Northern Natal
- Diamond mining in Kimberley



Chinese coal miners in an illustration of the *Tiangong Kaiwu* [Ming Dynasty encyclopedia](#), published in 1637 by [Song Yingxing](#).

Wealth versus environment

- SA is 5th largest coal producer in the world
- SA is third largest coal exporter
- South Africa gains economic prosperity from the act of mining

- The mining industry contributes negatively to the pollution of the water environment by producing Acid Mine Drainage.



Impact of mining

By act of mining: AMD is formed

Arises from oxidation of pyrite, due to exposure to air and water.



Characteristics of AMD:

High SO₄: Salinity

High acidity: low pH

High metal content



Environmental impact: Water pollution



AMD

Water Pollution through mining in South Africa

Statistics

- Mpumalanga coal field produces 40-50ML/d, may increase to 120ML AMD/d
- Gauteng mining area produces 300-400 ML AMD/d



Anno 2006

Mine water remediation

- Biological treatment
- Chemical treatment
- Physical treatment



Possible ground water contamination

Chemical treatment

CSIR Limestone/Lime Neutralisation

The integrated limestone/lime process is used for treating acid, iron and sulphate-rich water with powder calcium carbonate.

- Acidic mine effluents are neutralised
- Sulphate is removed from 20 g/l to less than 2 g/l
- Metal removal (Ca, Mg), using lime



Full scale implementation of CSIR neutralisation technology

South Africa:

- Ticor, Limestone/lime
- Navigation and Kromdraai mines, Anglo coal (both Limestone)
- Namaqua Sands Limestone/lime
- Zincor (limestone)
- Optimum (limestone)

Botswana

- BCL (limestone)

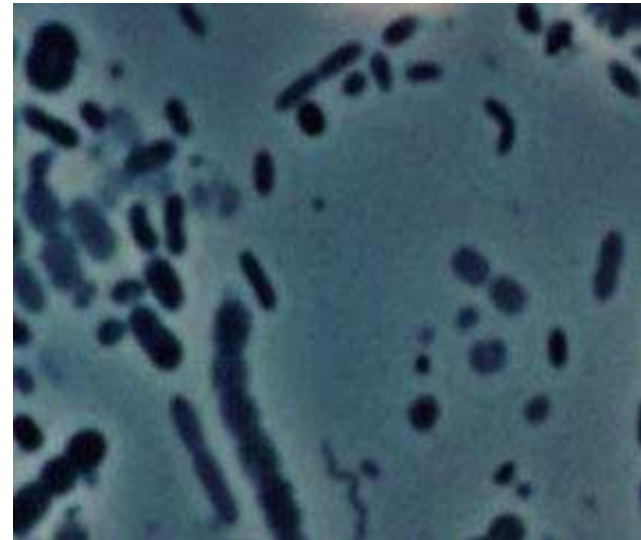
Australia

- Iluka resources (limestone/lime)



Biological Sulphate (SO_4) removal

- Requires Sulphate Reducing Bacteria (SRB)
- Requires a carbon and energy source
- Requires SO_4
- Requires anaerobic conditions



Microorganisms

CSIR developed biological treatment

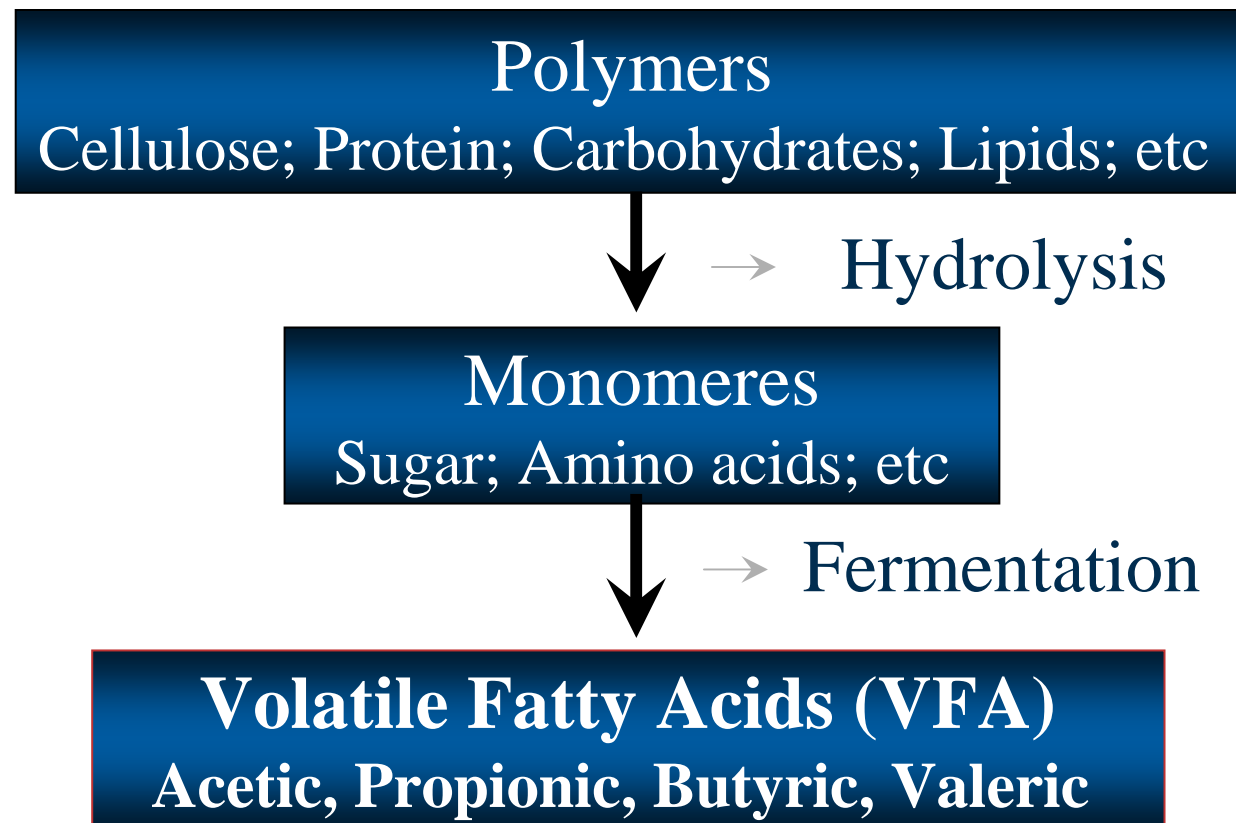
Biological AMD treatment using alternative carbon and energy sources

- Degradation products of cellulose occurring in grass
- Volatile Fatty Acids (VFA) and hydrogen (H_2) production
- VFA and H_2 can function as C+E source for SRB



Grass: potential source of sustainable energy

Biological degradation of cellulose

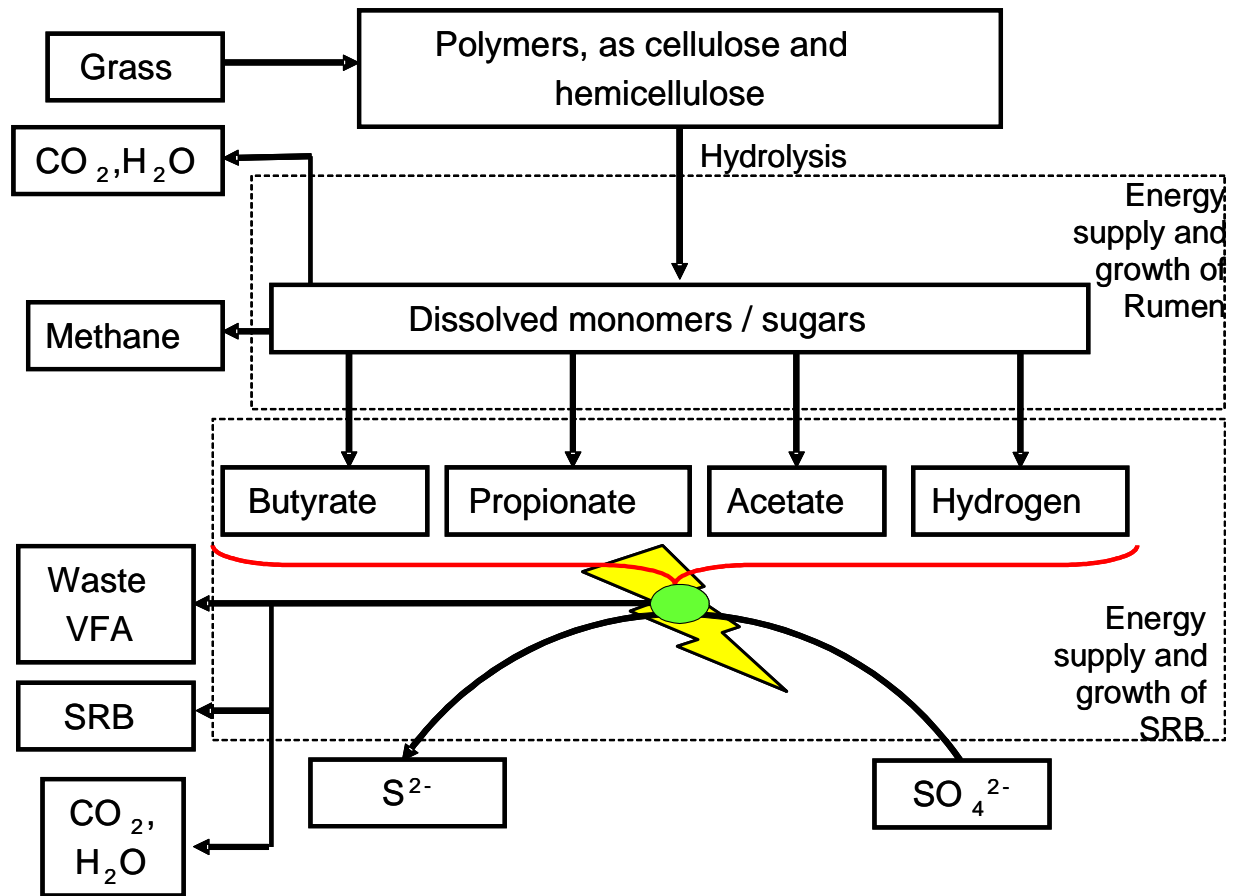


Microorganisms produce VFA and H₂

Use of natural occurring microorganisms

- Rumen fluid from ruminant →
- SRB participate in the degradation of the polymers and monomers to produce VFA





Benefits of biological SO_4 removal

- Sulphate removal to <200 mg/L
- Alkalinity production to increase pH
- Sulphide is the reduction product of sulphate
- Metal removal due to Metal-sulphide precipitation

Biological SO_4 reduction using grass-cellulose as the carbon and energy source

Aim of study

To remove biologically:

- Sulphate from AMD
- Sulphide after sulphate reduction



CSIR developed biological treatment plant

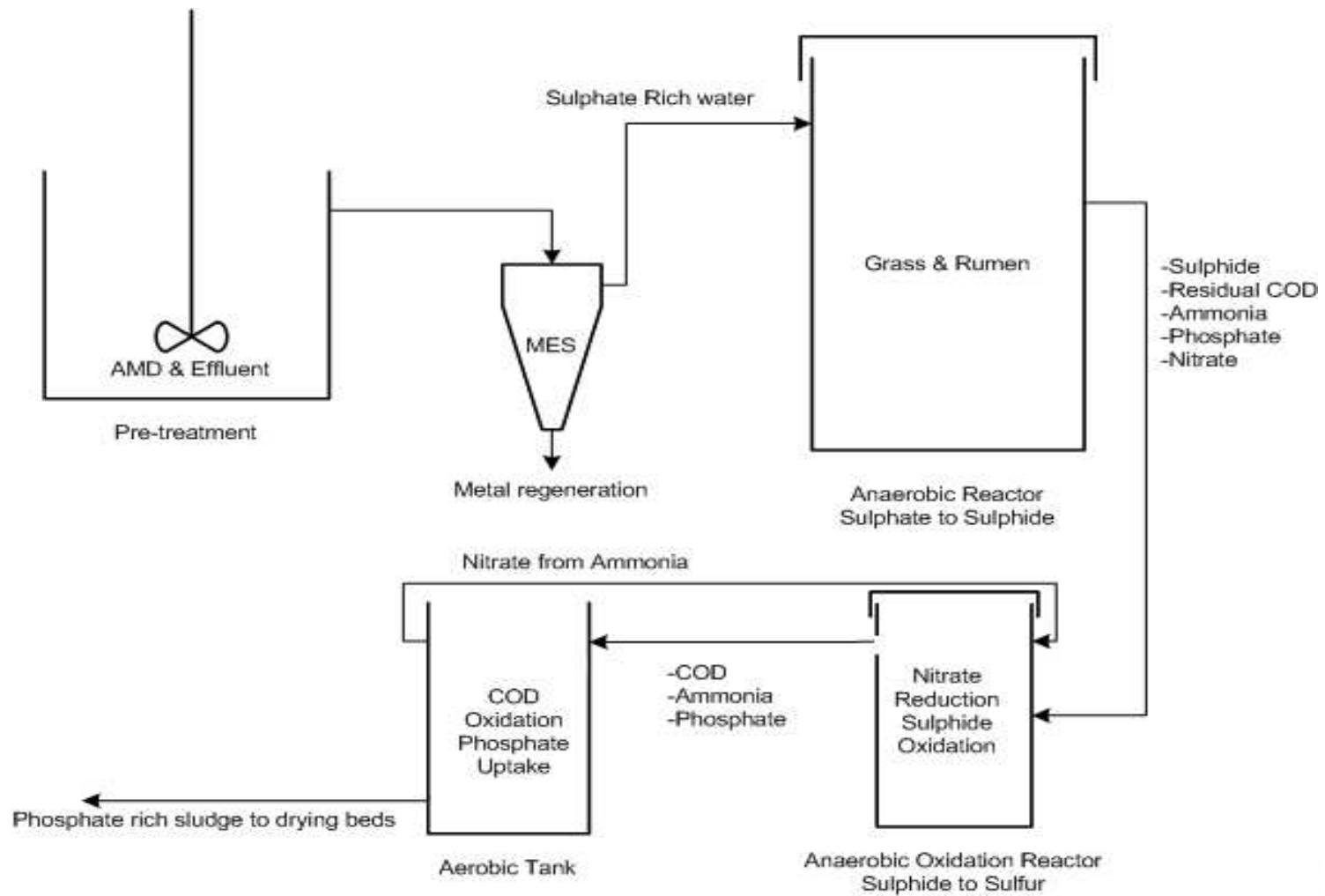
Materials and Methods

Feed water

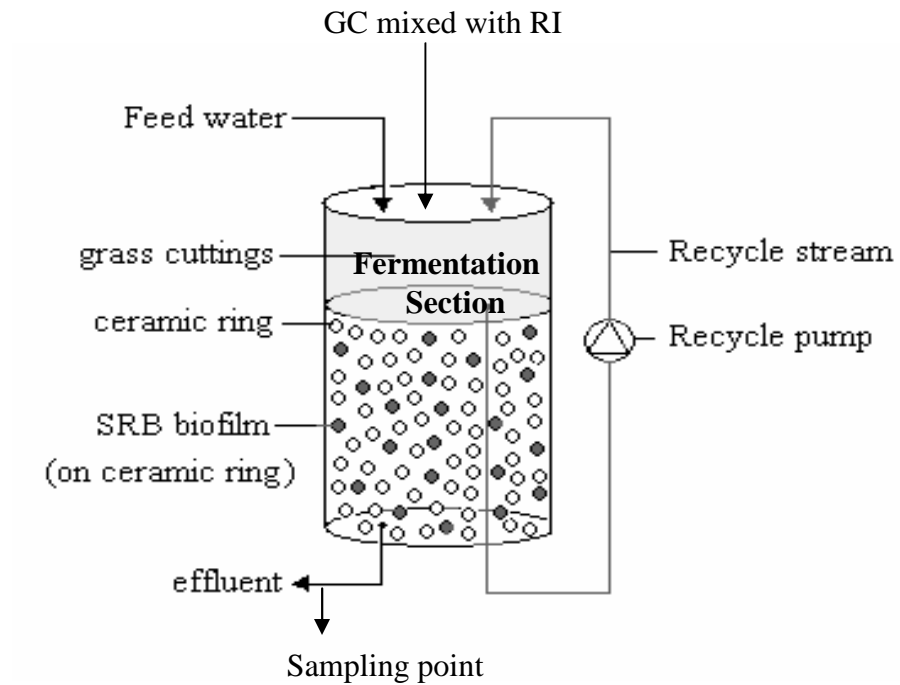
- Pre-treated AMD
- Mix 1 part AMD with 1 part of reactor effluent

Purpose

- Metal removal
- pH increase of AMD



Reactor: Hybrid Reactor System



Schematic overview of HFS reactor system.

Laboratory study: continuous operation of hybrid reactor

Reactor contains

1. Grass cuttings as cellulose source
2. Cellulose degrading microorganisms
3. SRB

Feed water:

Pre-treated AMD for pH increase and metal precipitation

Principle

VFA and H_2 production and utilisation in reactor for biological sulphate removal

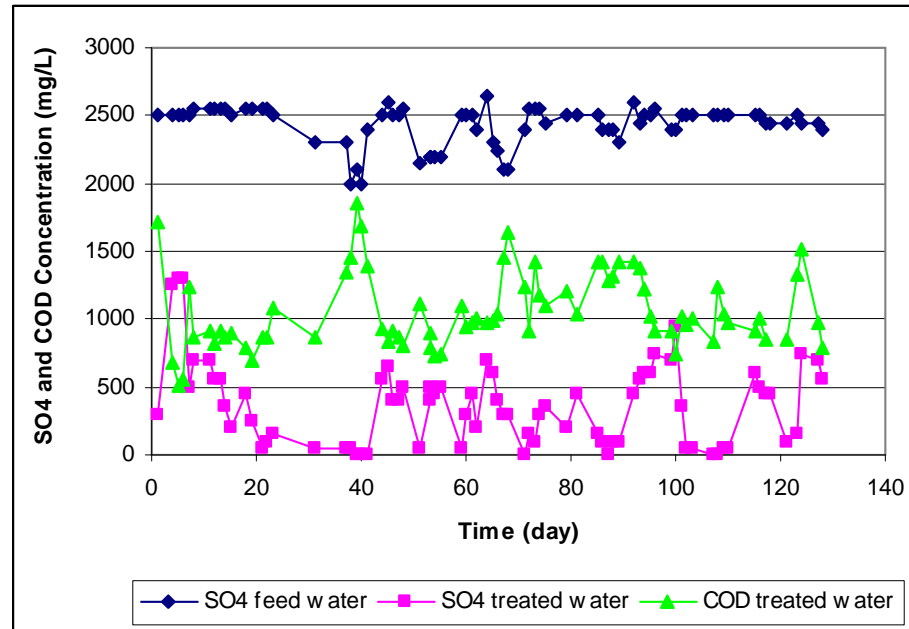


Reactions using VFA and H₂ for SO₄ removal



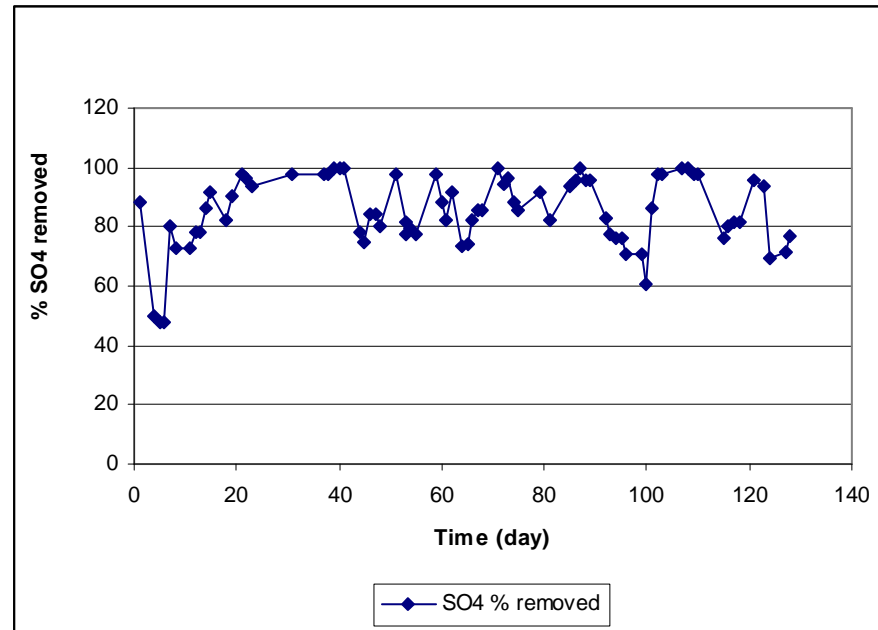
Results

Sulphate removal (mg/L)



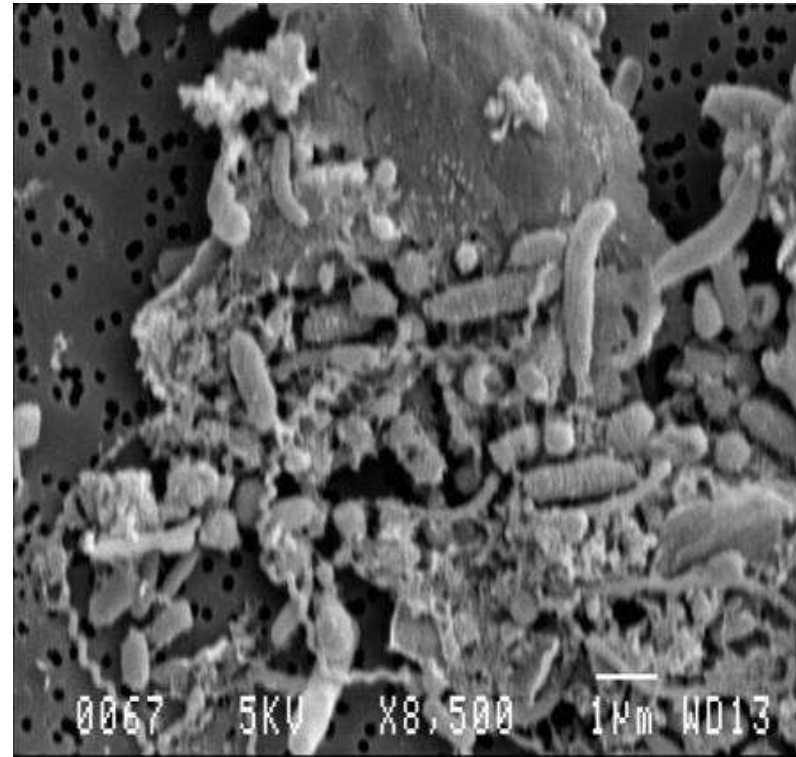
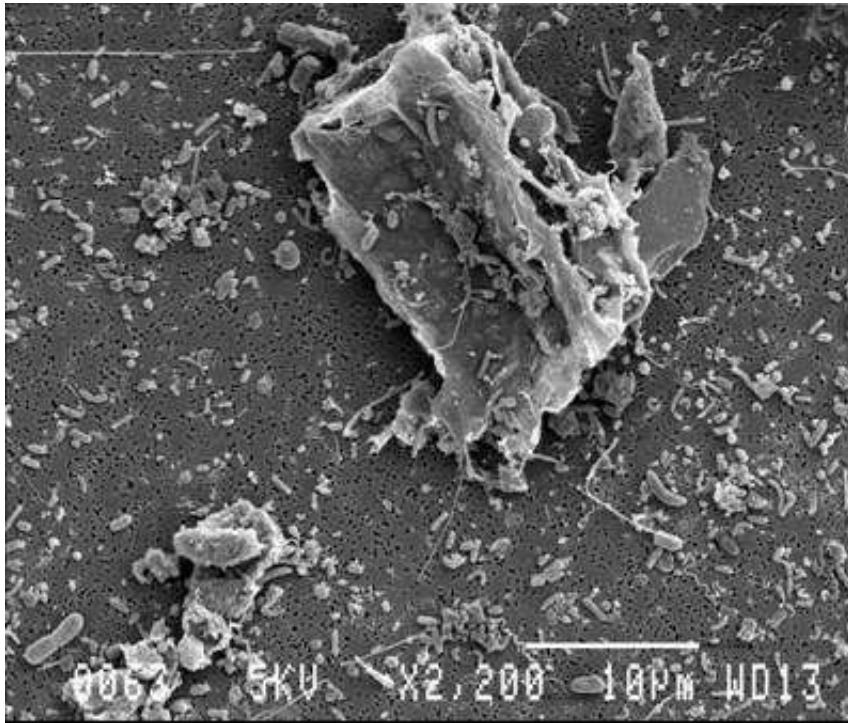
Sulphate removal as function of COD

Percentage Sulphate removal efficiency



Average sulphate removal was 85%

Results scanning electron microscope



Metal removal

Metal	AMD	Pre-treated AMD	Effluent HFS
Aluminium	24	14	<0.09
Chromium	<0.05	<0.05	<0.05
Iron	851	102	0.21
Lead	0.15	<0.03	<0.03
Manganese	48	27	5.9
Nickel	11	4.3	0.04
Zinc	1.4	0.94	<0.06

All units in mg/L

Sulphide oxidation after sulphate reduction

Biological sulphide oxidation using air

- By *Thiobacillus* Species
 - *Thiobacillus thioparus*,
 - *Thiobacillus denitrificans* and
 - *Thiobacillus ferrooxidans* (Chung et al., 1996).
- Producing sulphur or sulphate
 - $\text{HS}^- + \frac{1}{2} \text{O}_2 \rightarrow \text{S} + \text{OH}^-$
 - $\text{HS}^- + 2 \text{O}_2 \rightarrow \text{SO}_4^{2-} + \text{H}^+$

Experimental conditions of a continuous laboratory study

Period (days)	Air supply to reactor (L/min)
29-43	0.2
44-59	0.4
62-81	0.6
83-97	0.8

Results of biological sulphide oxidation after continuous laboratory studies

Air supply (L/min)	S ²⁻ removed (g/d)	Sulphur produced (g/d)	%SO ₄ increase in reactor
0.2	5.13	1.98	8
0.4	4.45	1.96	28
0.6	4.62	2.06	47
0.8	5.77	1.28	57

Conclusions continuous operation SO_4 /grass/rumen reactor and biological sulphide oxidation

- Efficient sulphate removal (Average 85%)
- Metal removal, especially iron
- Biological sulphide oxidation achieved, at low air concentration, otherwise SO_4 is end product rather than Sulphur
- Based on obtained results, pilot scale reactor will be constructed

For more information

www.csir.co.za

Thank You

