

Advanced Wastewater Treatment Technologies



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Introduction

- Industrial wastewater- variety of pollutants with varying concentrations and properties
- No single technology available to treat all industrial wastewaters
- Technology selection- based on type of pollutants, concentration of pollutants and treated water quality requirement
- Treatments: A combination of Physical, physico-chemical and biological processes

Physical Processes

- Sedimentation
- Filtration
 - Surface filtration
 - Sieves, cloth filters, membrane filters
 - Deep filtration
 - Sand filter
 - Other media filters
- Evaporation
 - Natural evaporators
 - Vacuum Evaporators
 - Mechanical evaporators
- Gas Transfer

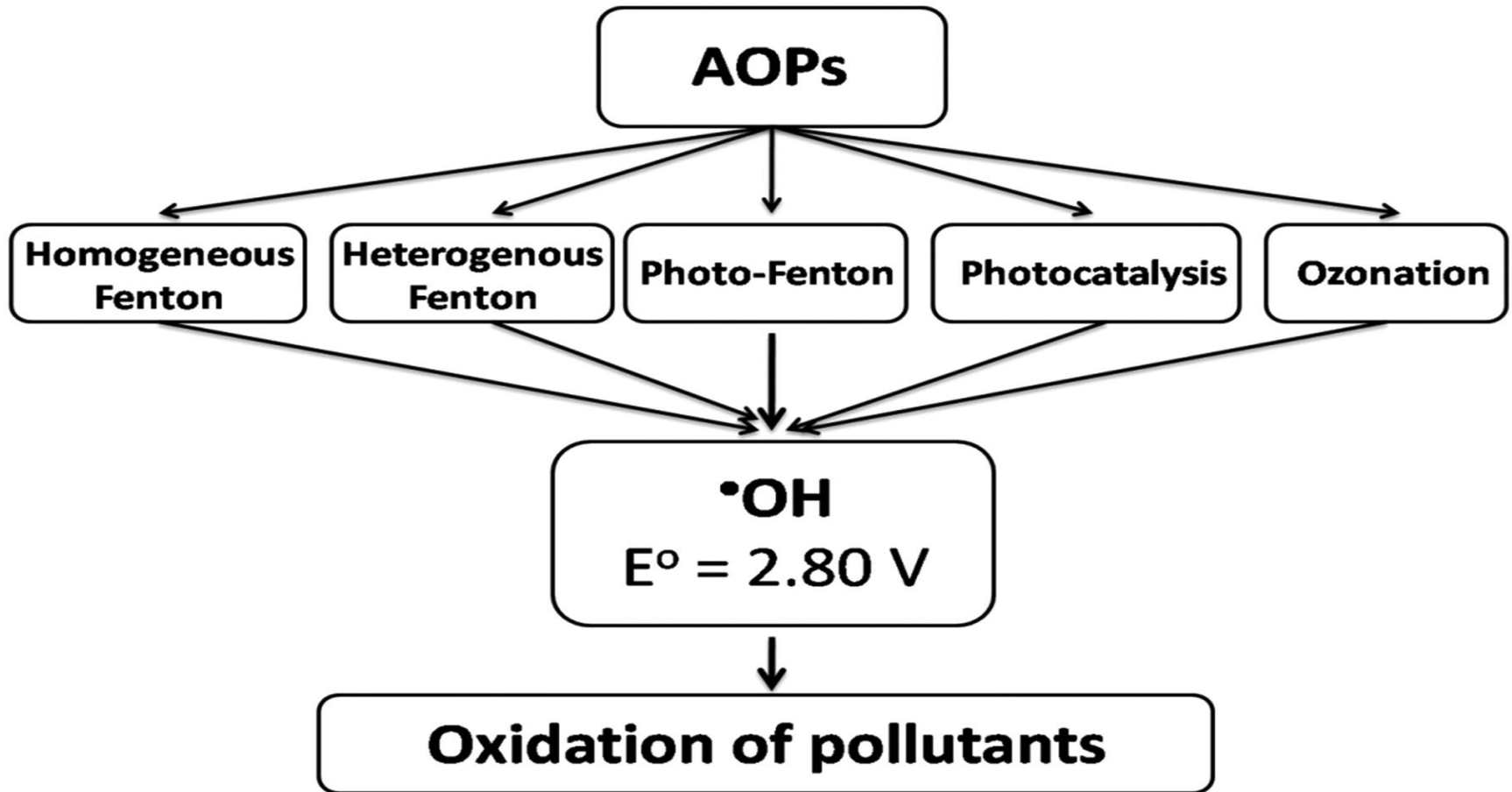
Physico-Chemical Processes

- Coagulation and flocculation, electro coagulation
- Adsorption- New and tailor made adsorbents
- Ion Exchange -
- Precipitation
- Membranes –RO, CDI, electro dialysis
- Oxidation Reduction
 - Advanced Oxidation

Biological Processes

- Aerobic
- Anaerobic
- Biological processes can be modified by using enriched microbes for selective complex organic wastes.

Advance oxidation processes(AOPs)



- Volume and treatment time is considerably reduced.



PLASMA

OH°

H°

O°

O_2

HO_2

H_2O_2

O_3

UV

High
electric
field

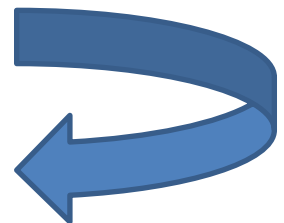
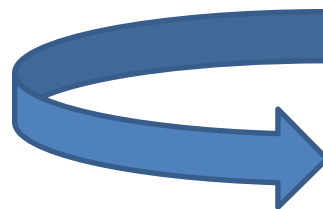
Intense
wave

Reactive
species

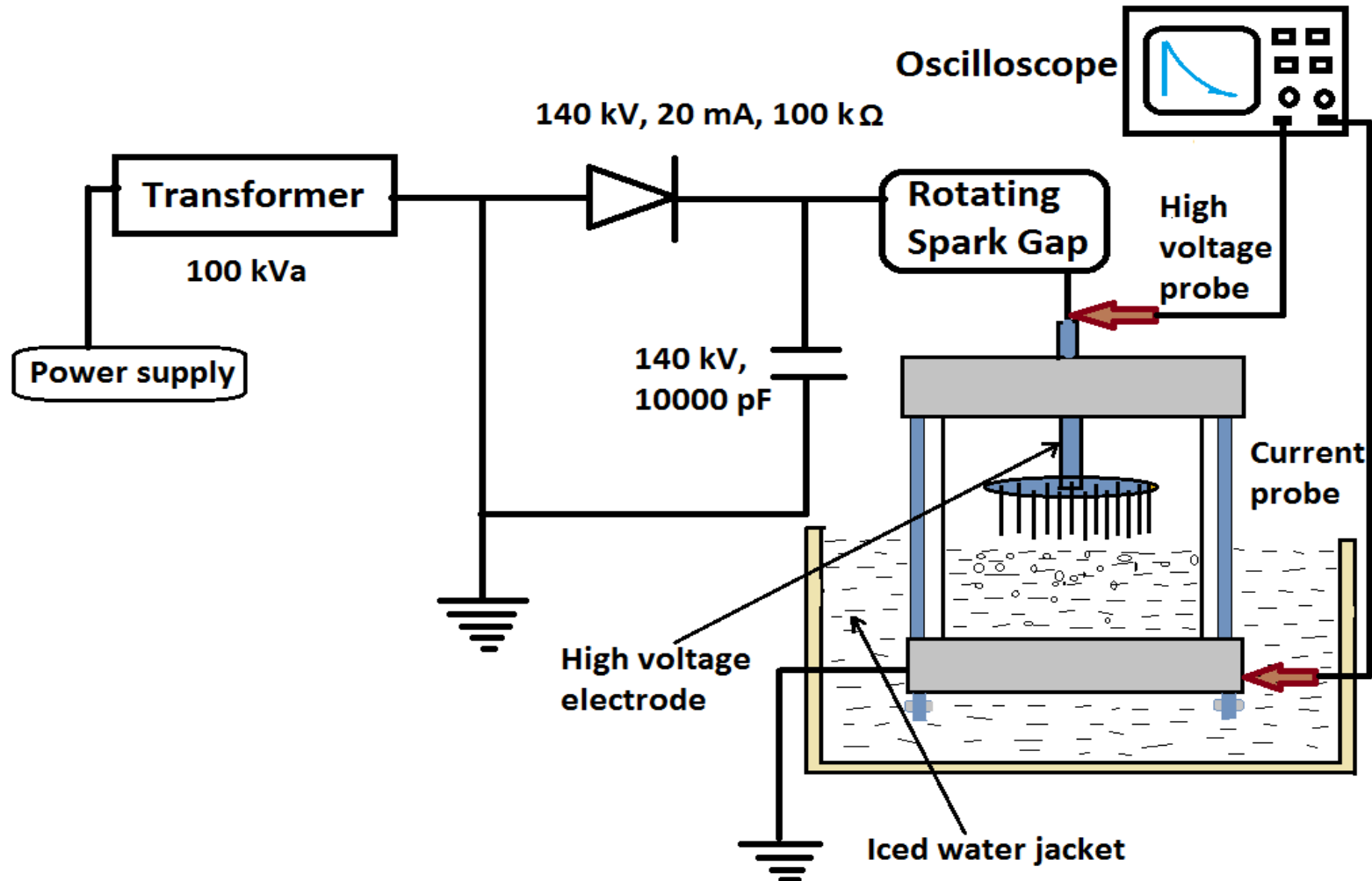
Advance
Oxidation

3.03	F_2
2.80	OH
2.42	O
2.07	O_3
1.78	H_2O_2
[V]	

Oxidation potential

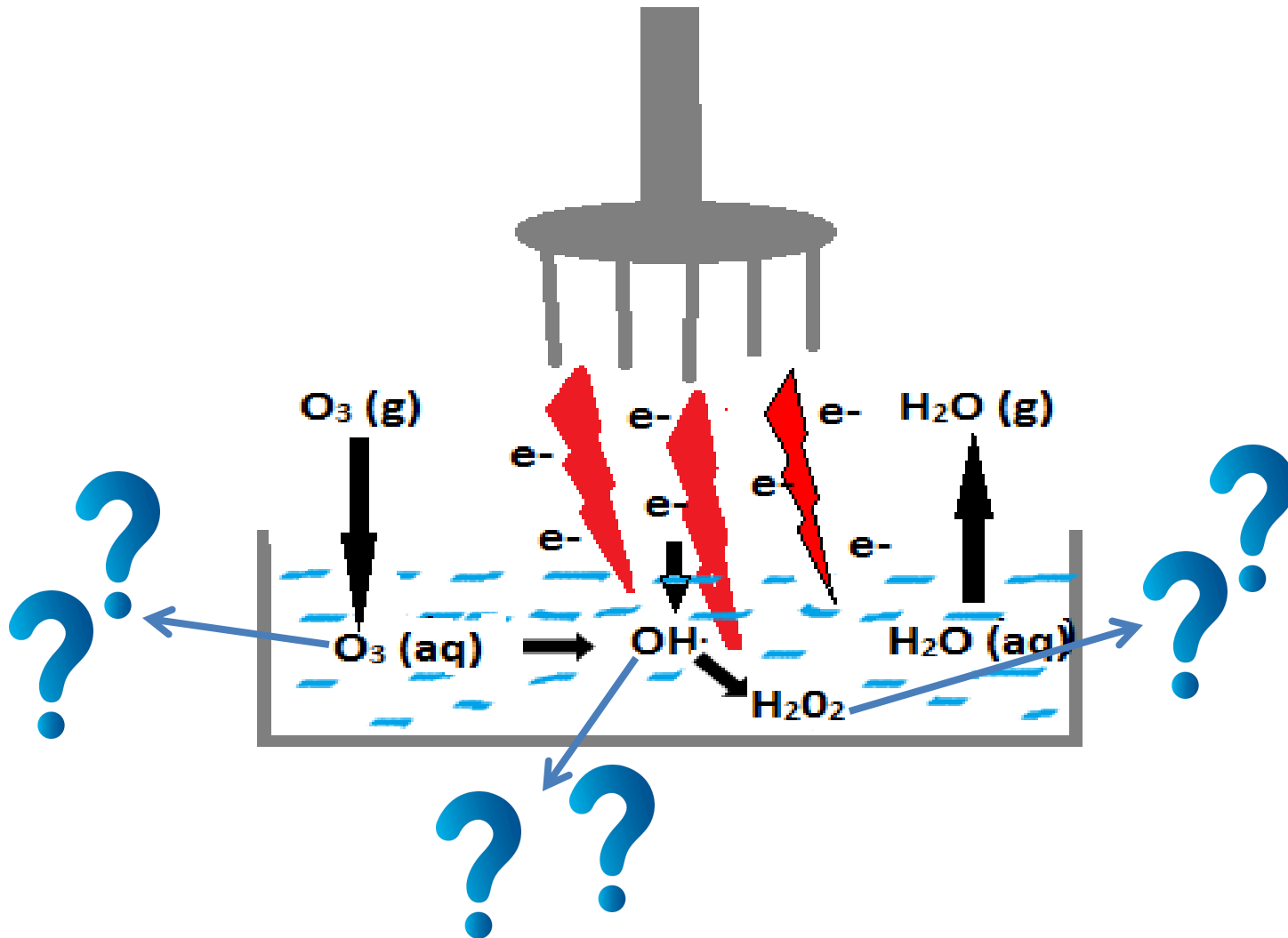


Circuit Diagram for the Reactor set-up

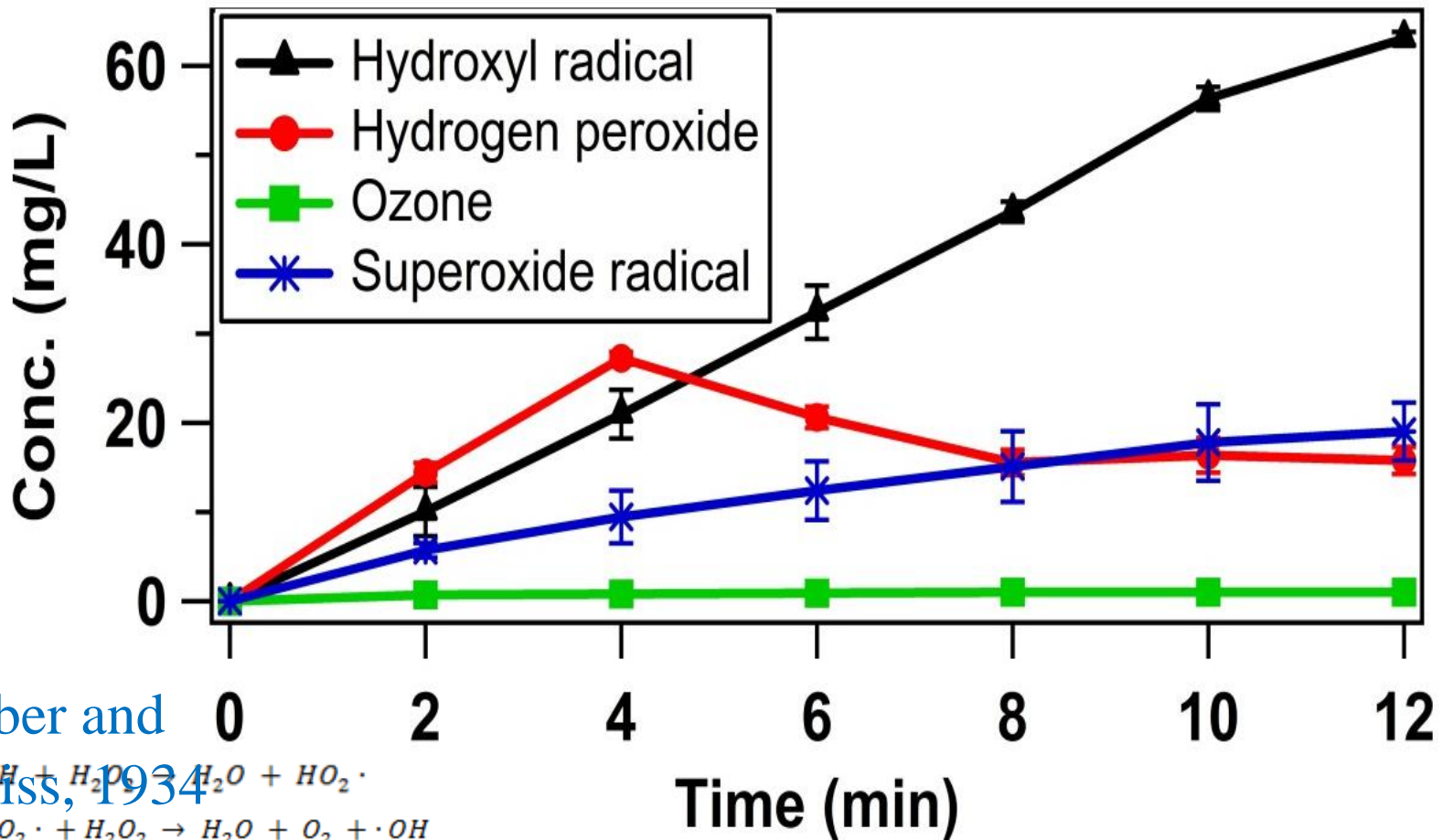


Singh Raj Kamal, Babu V., **Philip Ligy**, Sarathi R., (2016), Disinfection of Water Using Pulse Power Technique: A Mechanistic Perspective, RSC Advances, 6, 11980 – 11990.

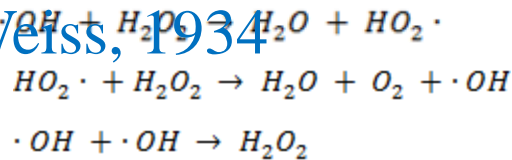
Quantification of ROS



Trend of ROS formation

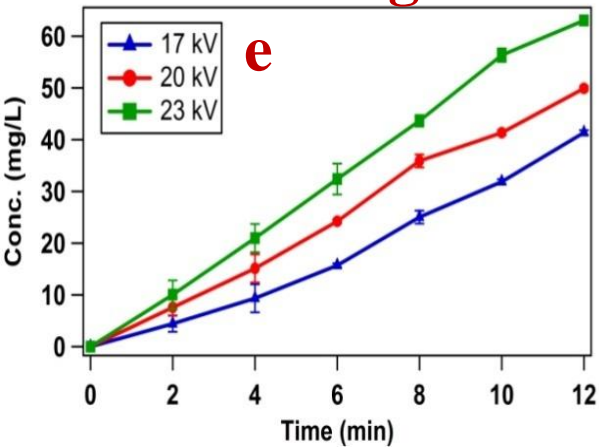


Haber and Weiss, 1934

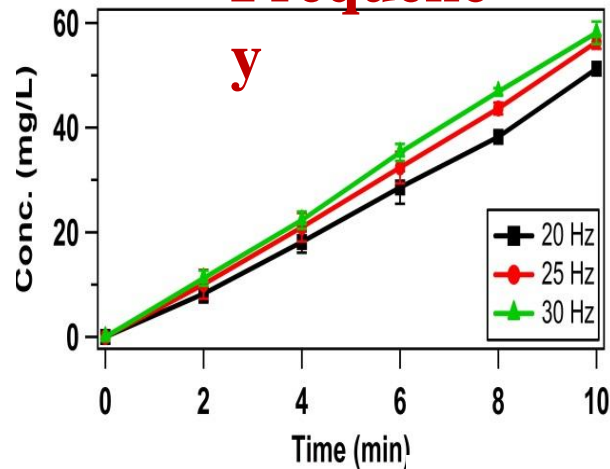


Effects of system parameters on $\cdot\text{OH}$ formation

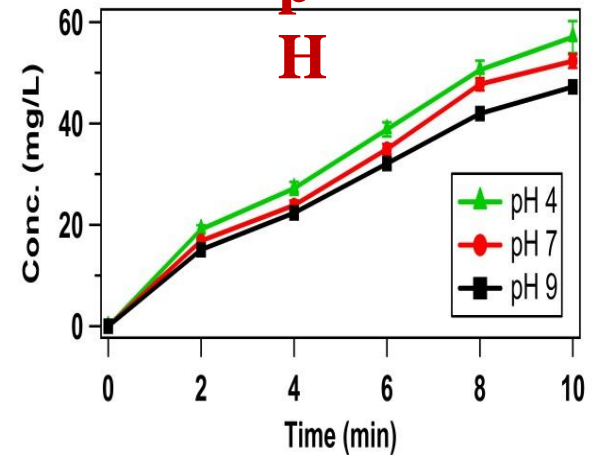
Voltage



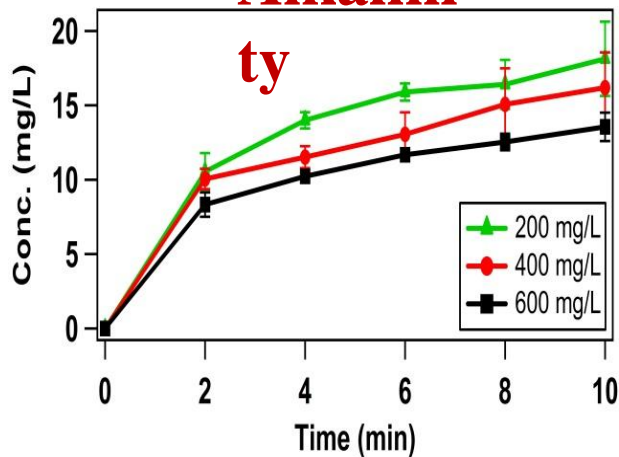
Frequency



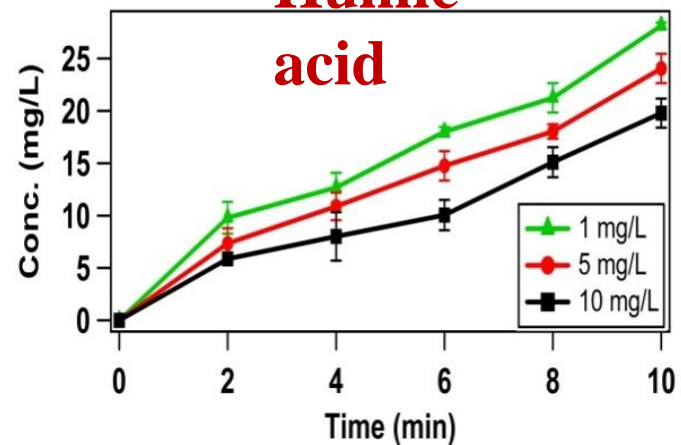
pH



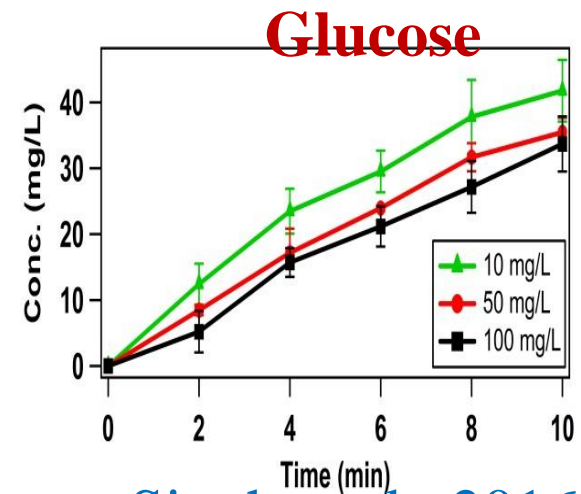
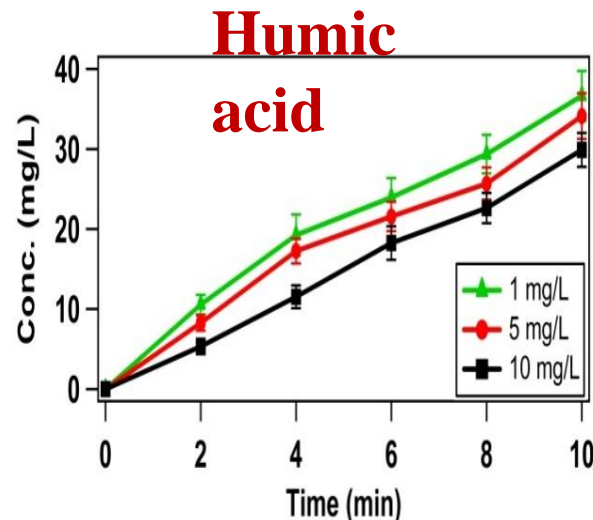
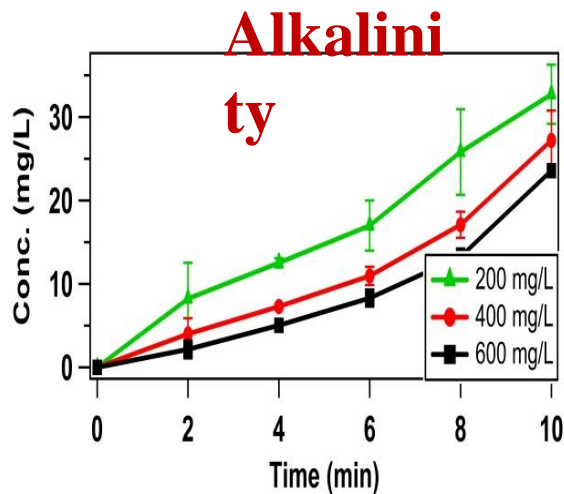
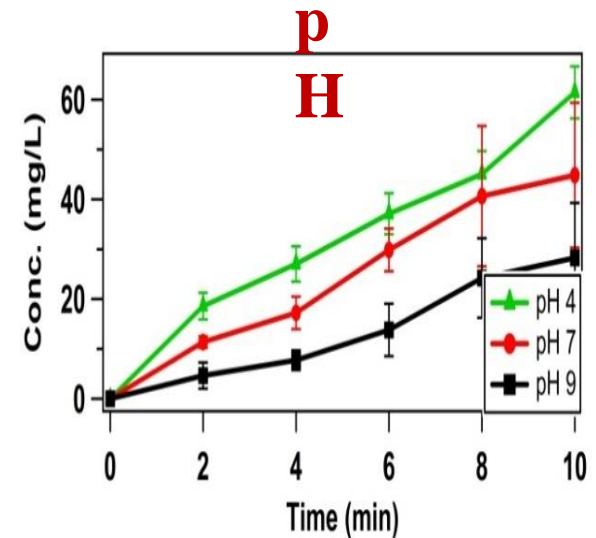
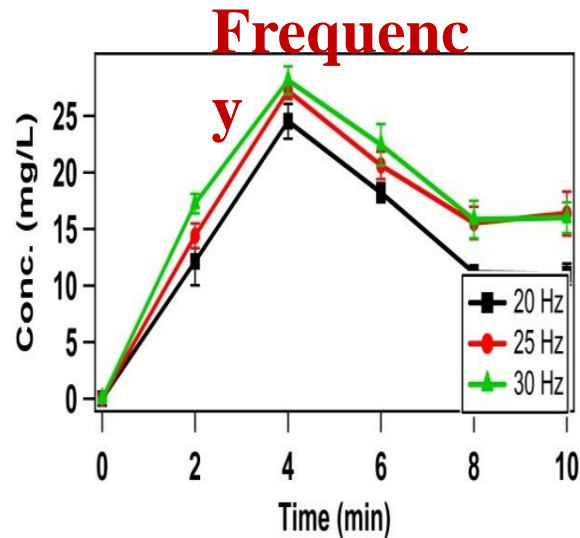
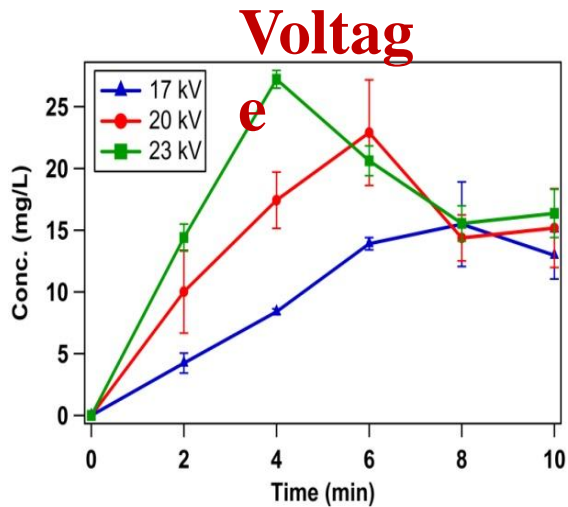
Alkalinity



Humic acid



Effects of system parameters on H₂O₂ formation



Kinetics study of ROS formation

Voltage (kV)	Rate of Reaction for OH radical (mol L ⁻¹ s ⁻¹)	Rate of Reaction for H ₂ O ₂ (mol L ⁻¹ s ⁻¹)	Rate of Reaction for O ₂ ²⁻ (mol L ⁻¹ s ⁻¹)	Rate of Reaction for O ₃ (s ⁻¹)
17	3.1	1.0	0.4	0.195
20	4.0	2.0	0.6	0.225
23	5.3	3.4	1.0	0.28

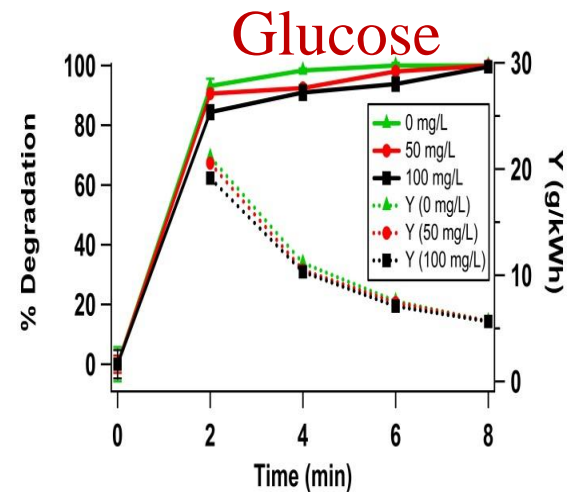
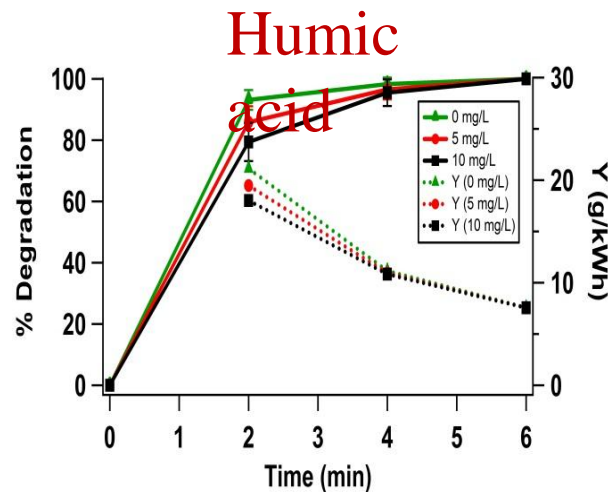
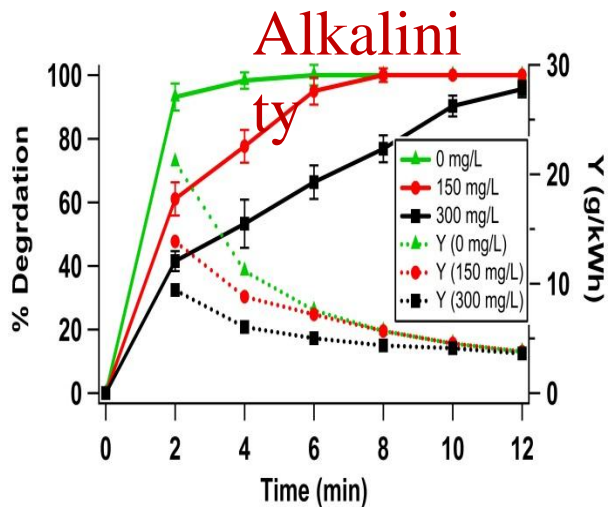
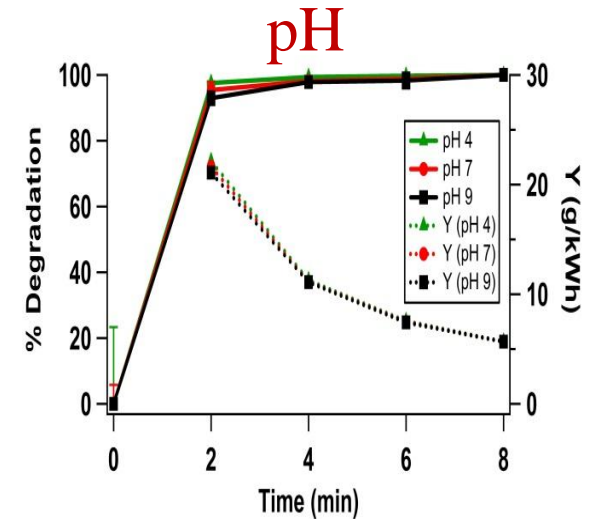
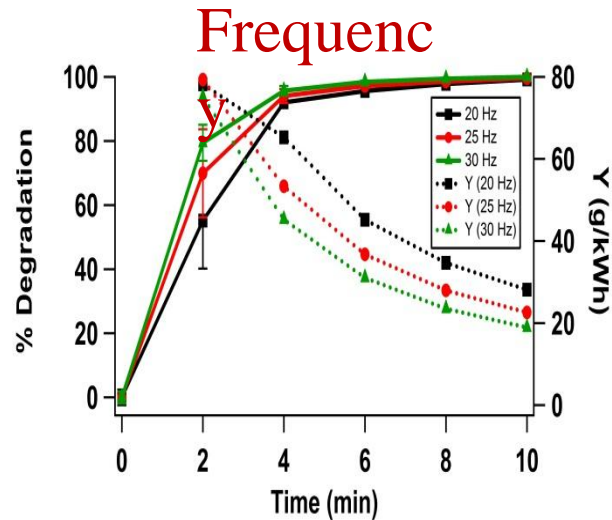
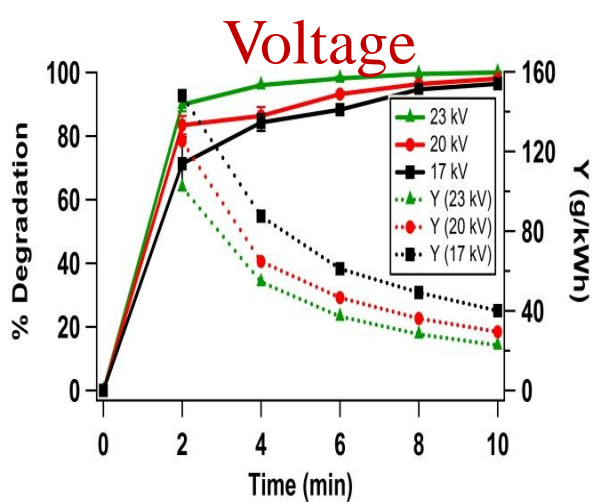
Note: The unit for r_{OH} , $r_{\text{H}_2\text{O}_2}$ and r_{O_3} is $10^{-6} \text{ mol L}^{-1} \text{ s}^{-1}$, $10^{-6} \text{ mol L}^{-1} \text{ s}^{-1}$ and $10^{-6} \text{ mol L}^{-1} \text{ s}^{-1}$.

3. Methylene Blue Degradation Study

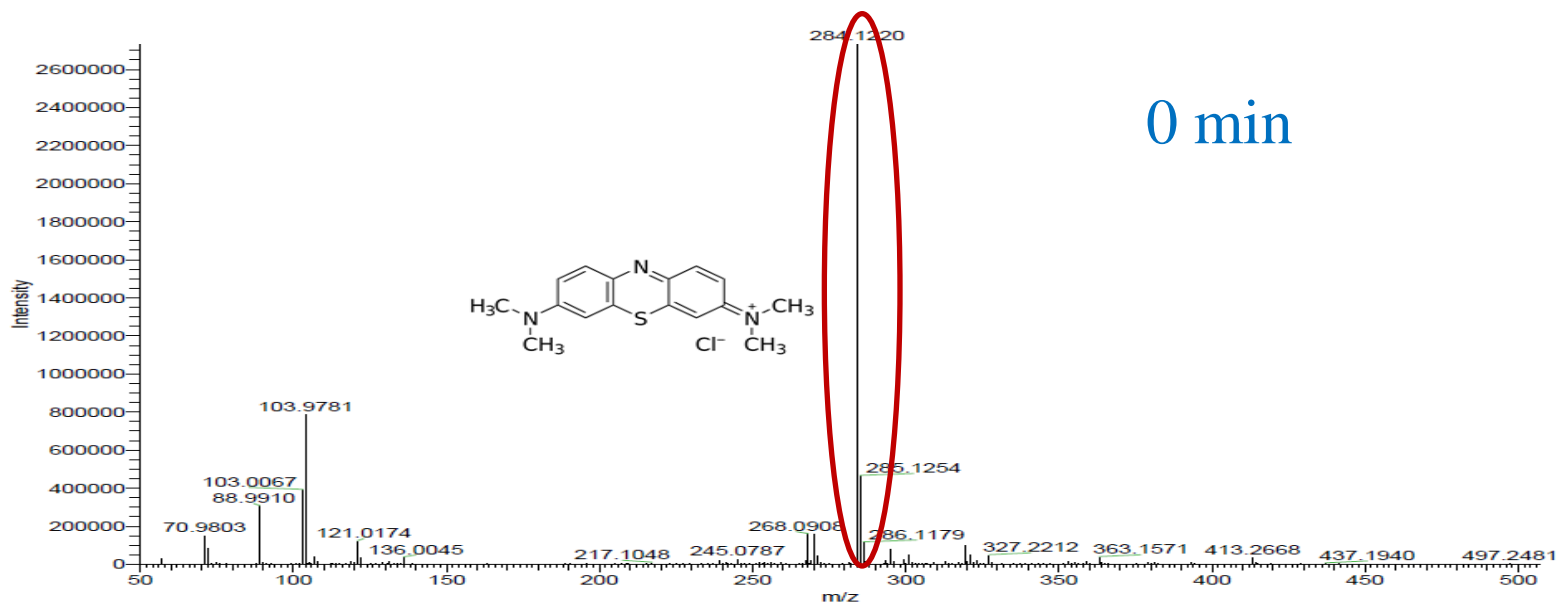
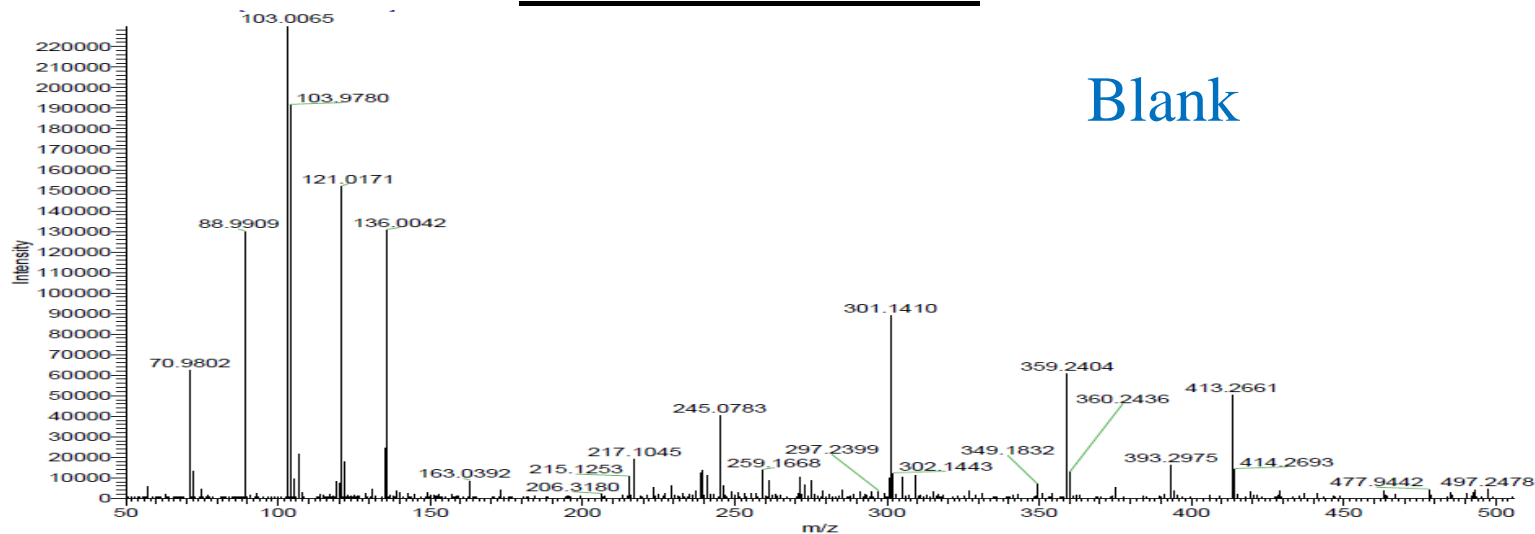


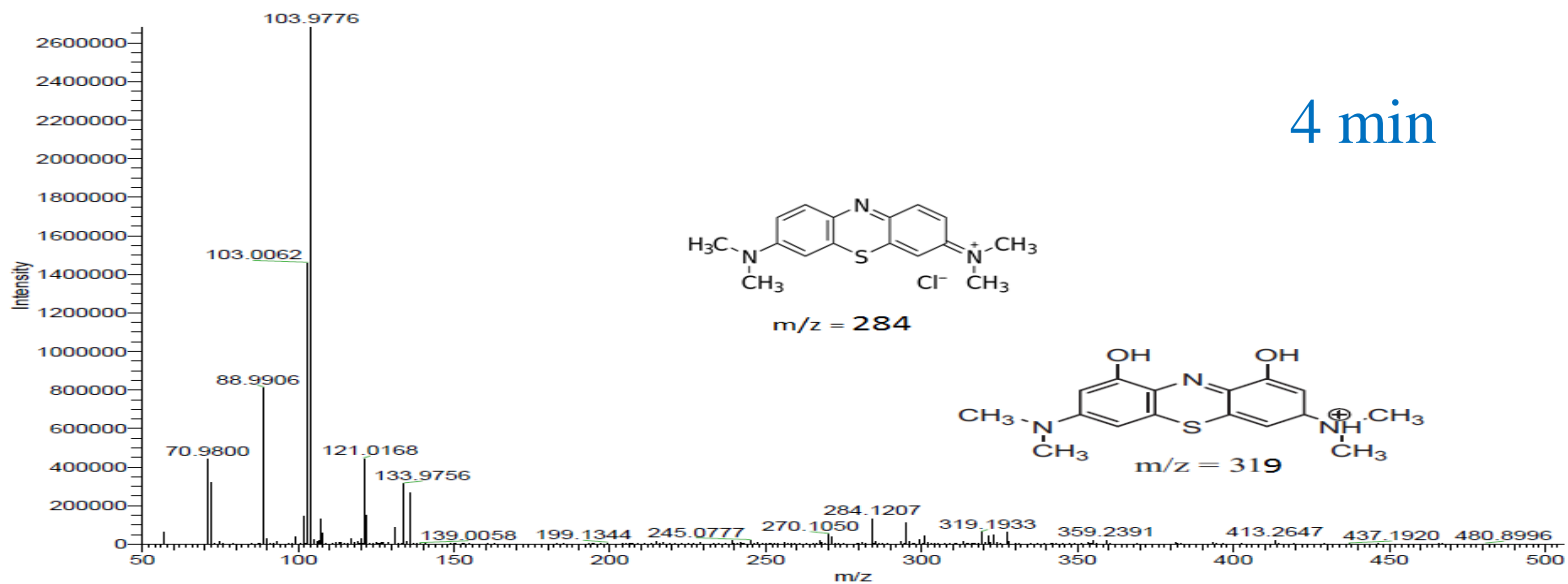
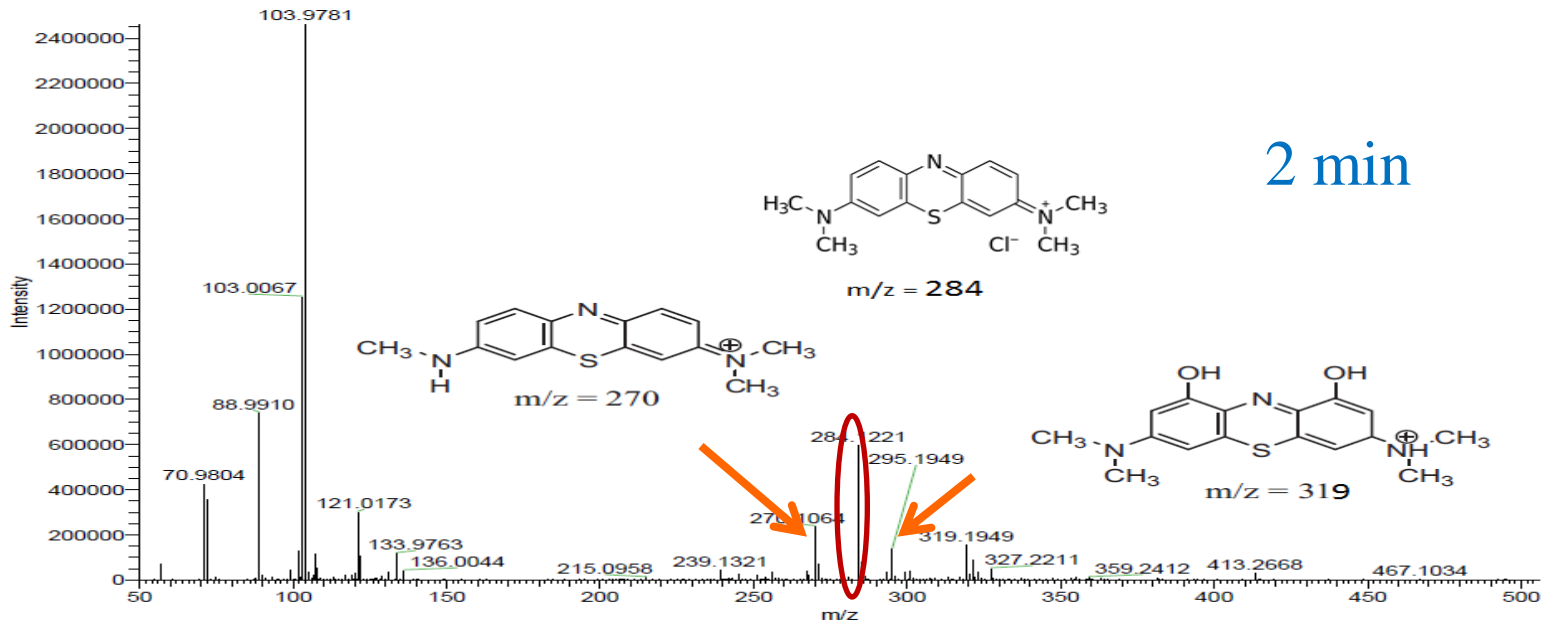
Singh Raj Kamal, Babu. V., Philip Ligy, Sarathi R., (2016), Applicability of Pulsed Power Technique for the Degradation of Methylene Blue, Journal of Water Process Engineering, 11, 118 – 129.

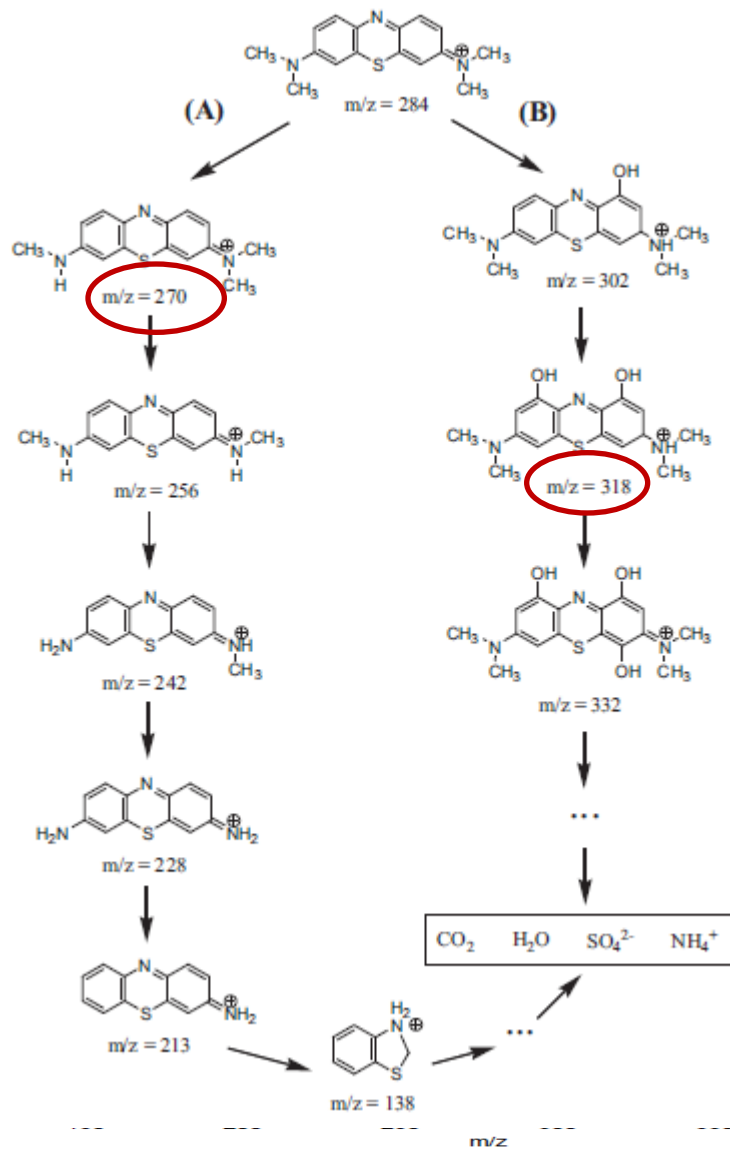
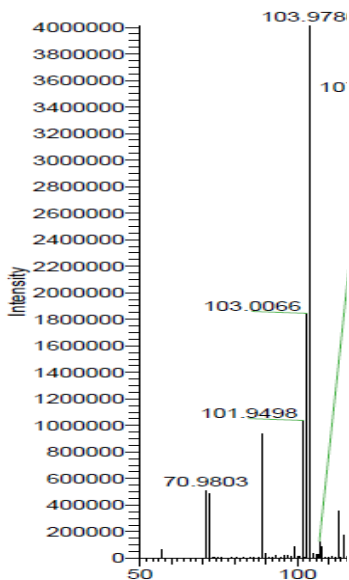
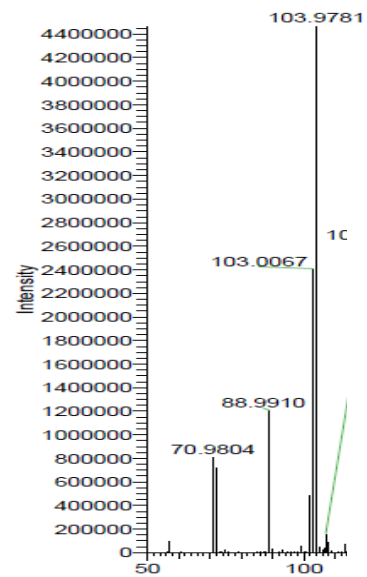
Effects of system parameters



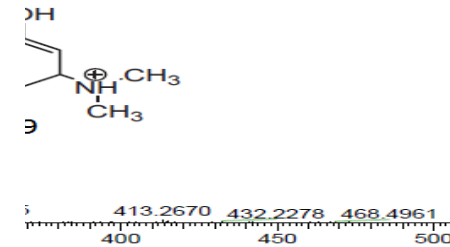
Mass Spectra for Methylene blue and its intermediates



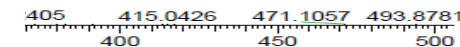




6 min



10 min



Benetoli et
al., 2012

Singh et al., 2016c

Technical Achievements

Rapid degradation and mineralization of Methylene blue (dye)

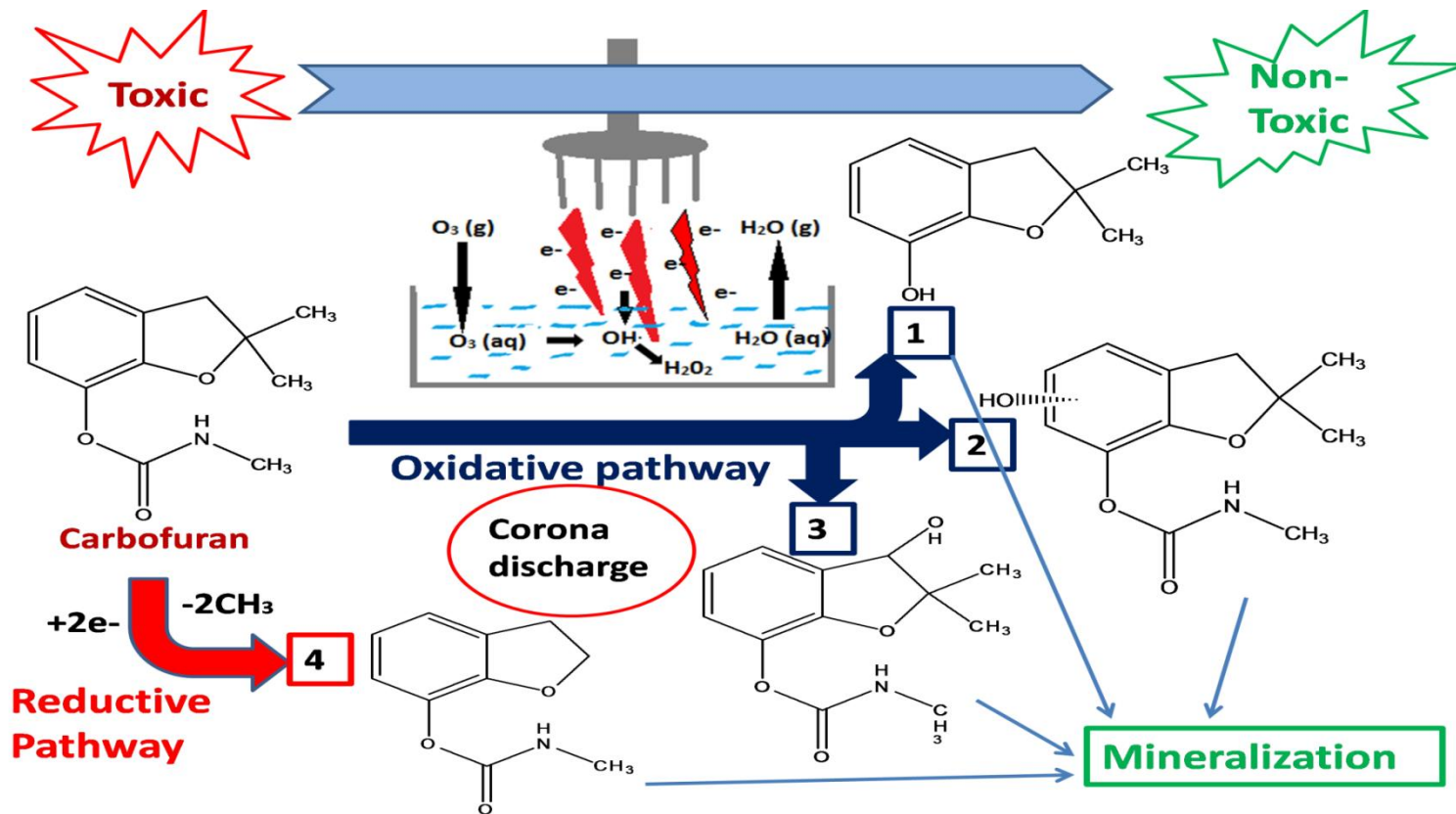
- **Description:**

Investigation of PPT efficiency for the degradation of dye.

- **Novelty:**

- Different **ROS** such as $\cdot\text{OH}$, H_2O_2 , O_3 and $\text{O}_2\cdot^-$ **quantification** in different environmental conditions.
- Effect of different **system parameters** on treatment efficiency.
- Under PPT, methylene blue **degradation pathway** was proposed.

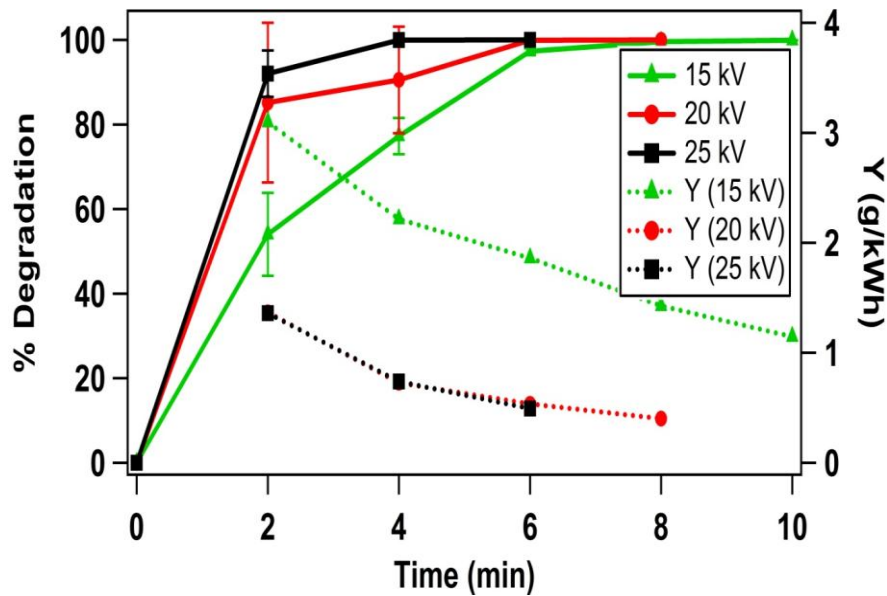
4.ECs degradation study



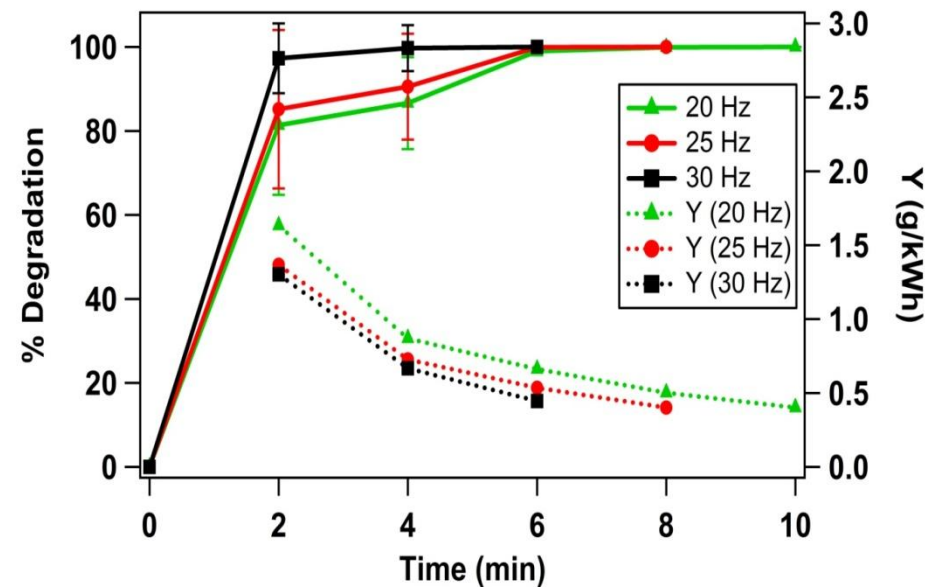
Singh Raj Kamal, **Philip Ligy**, Sarathi R., (2016), Rapid removal of carbofuran from aqueous solution by pulsed corona discharge treatment: Kinetic study, oxidative, reductive degradation pathway and toxicity assay, Ind. Engg. Chem. Res., Accepted manuscript.

Pesticide - Carbofuran

Initial Concentration – 1ppm

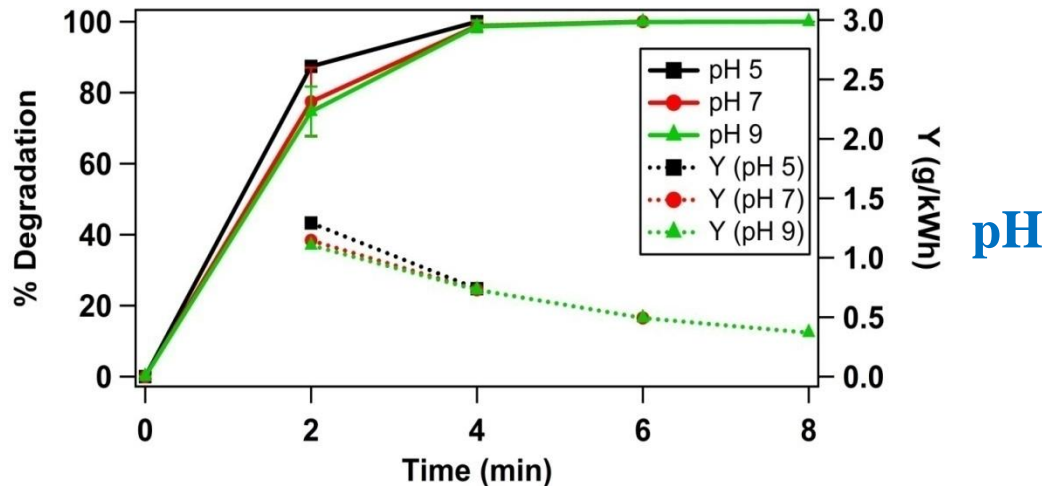
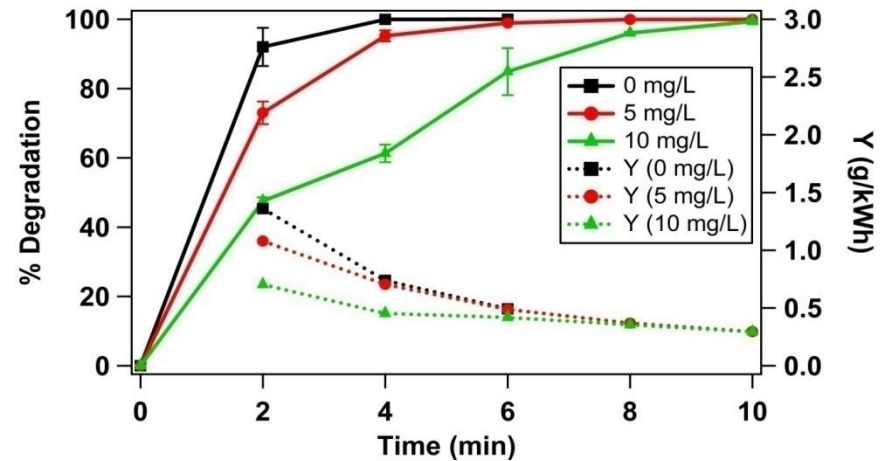
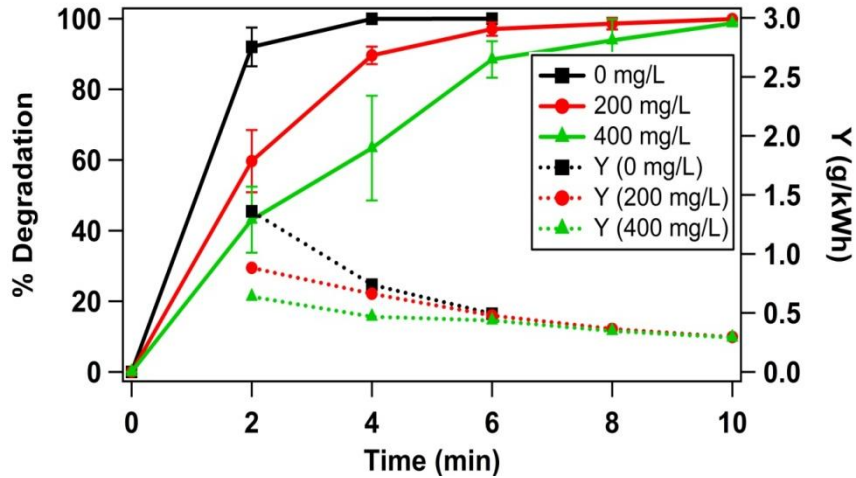


Voltage effect



Frequency effect

Effects of Environmental Parameters



Effect of Initial Carbofuran Concentration

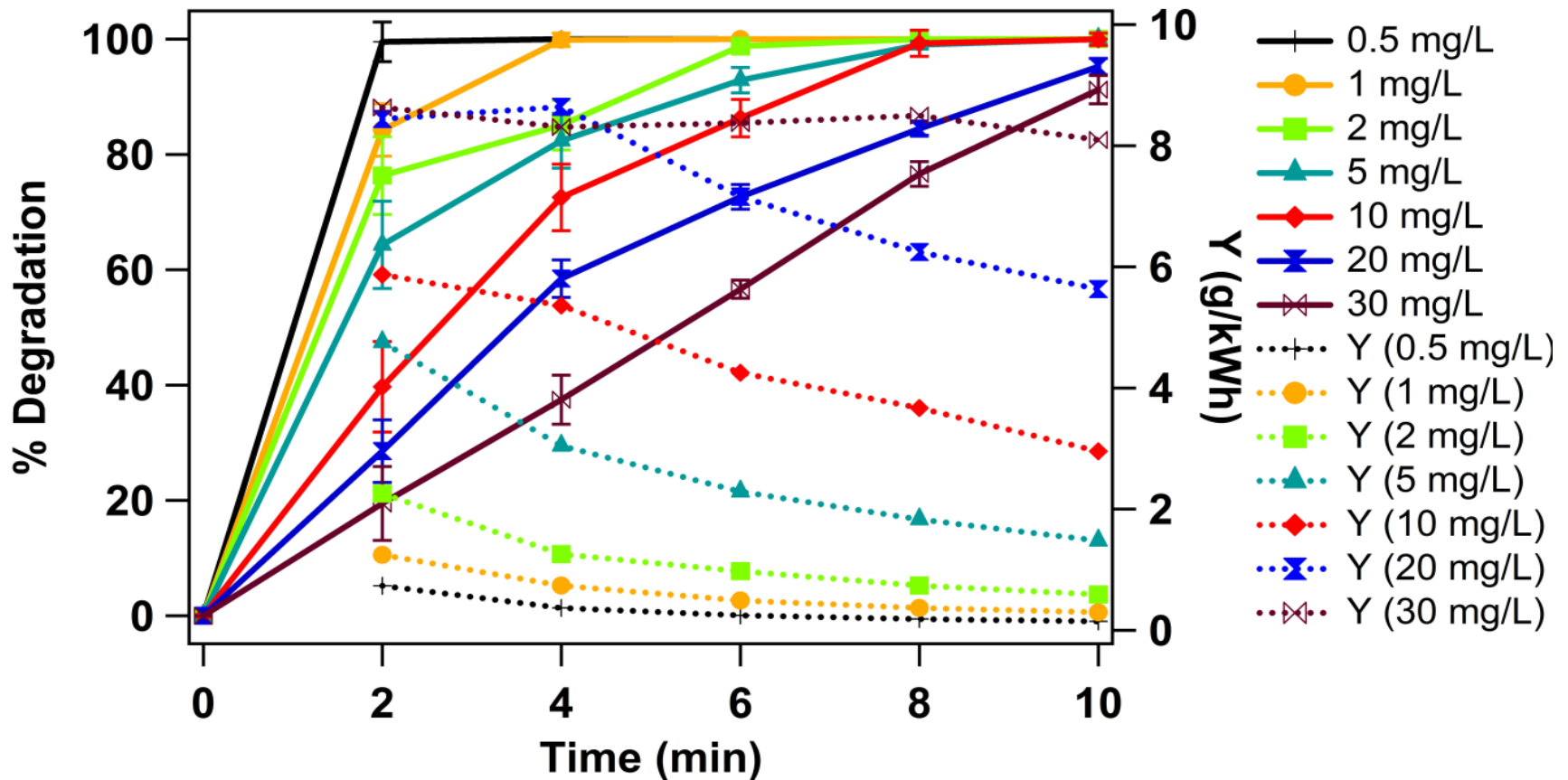
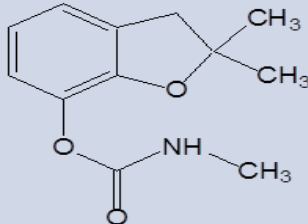
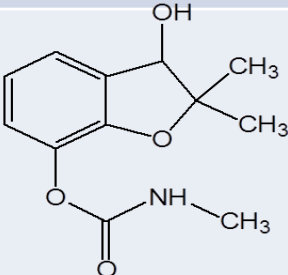
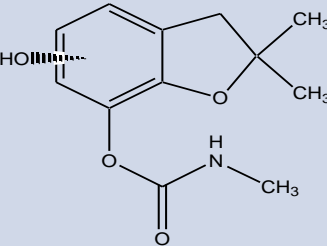
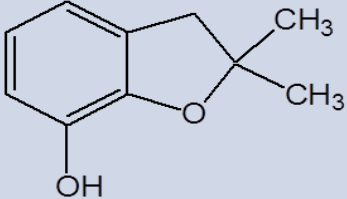
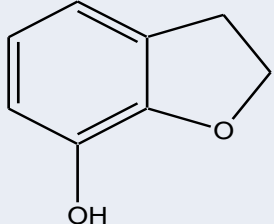
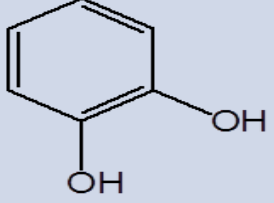
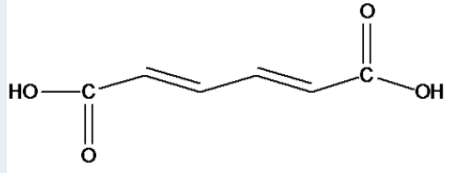
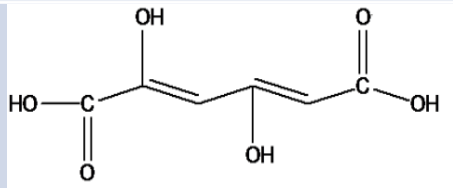


Table – Degradation kinetics of carbofuran degradation

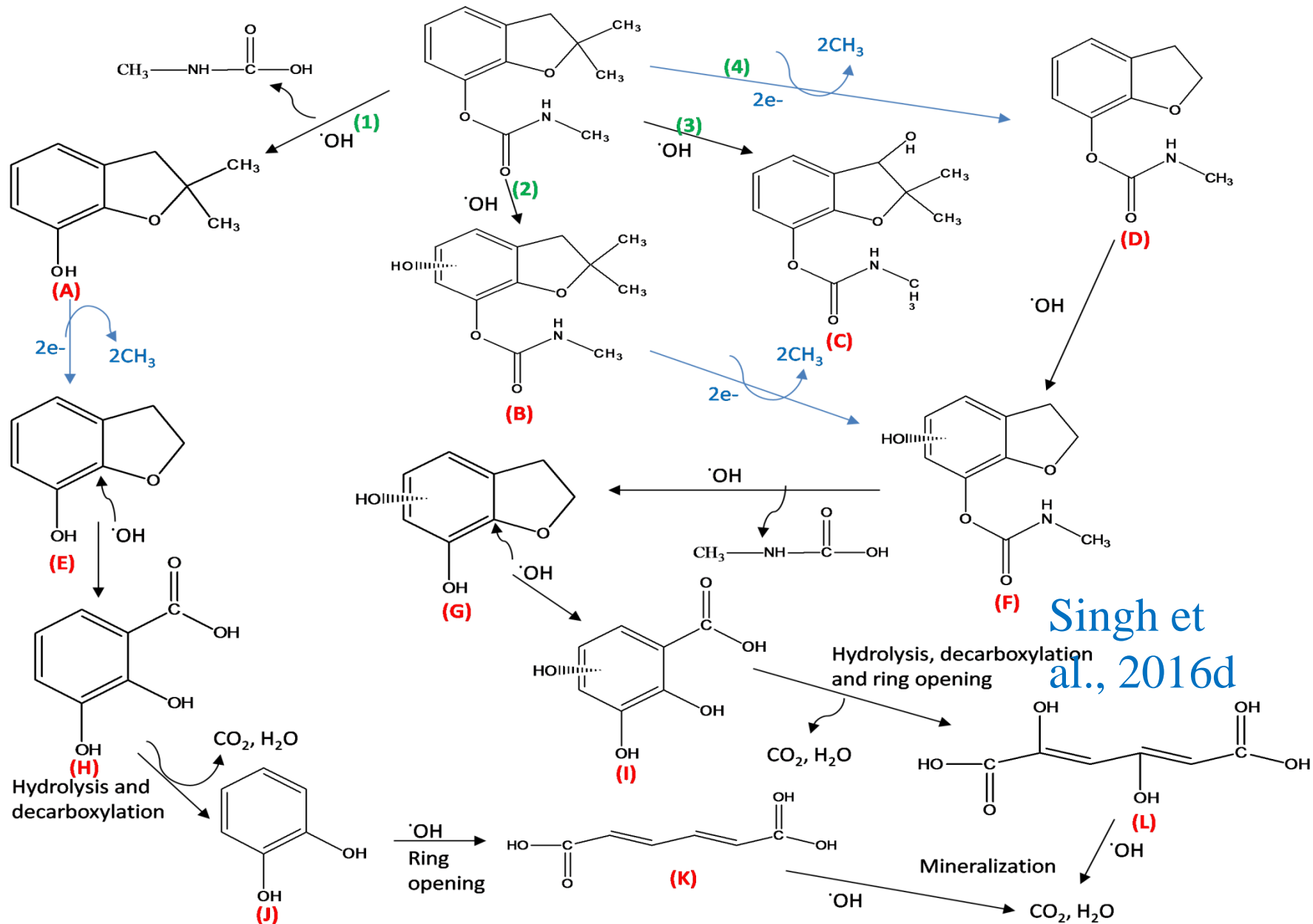
Initial concentration (mg/L)	First order rate constant (min⁻¹)	R²	t_{1/2} (min)
0.5	2.68	1.00	1.0
1	1.71	0.93	1.2
2	0.82	0.97	1.3
5	0.57	0.97	1.6
10	0.61	0.92	2.5
20	0.23	0.91	3.5
30	0.32	0.95	5.6

Main Carbofuran Intermediates – LC/MS analysis

Compound	Molecular mass (m/z) with Na ⁺ adduct	Actual molecular mass (m/z)	Chemical structure
Carbofuran	244	221	
A	260	237	
B	232	209	

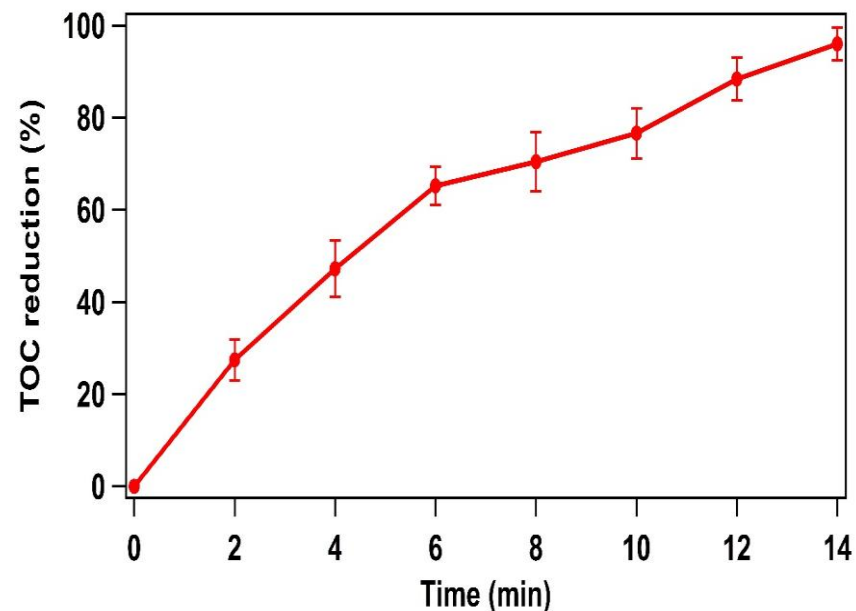
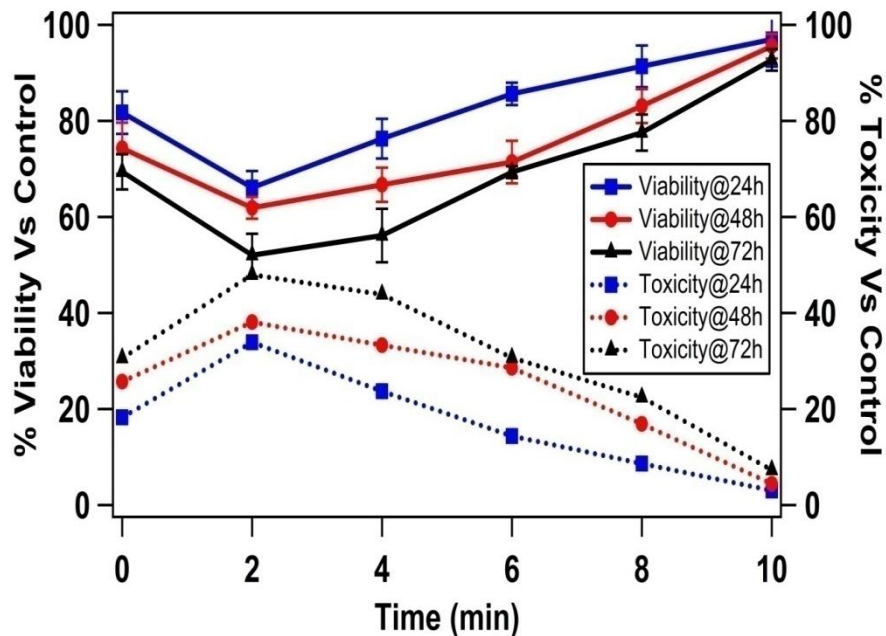
Compound	Molecular mass (m/z) with Na ⁺ adduct	Actual molecular mass (m/z)	Chemical structure
C	187	164	 <p>Chemical structure of 2,2-dimethyl-3,4-dihydro-1H-inden-5-ol, showing a benzene ring fused to a five-membered ring containing an oxygen atom and two methyl groups, with a hydroxyl group on the benzene ring.</p>
D	159	136	 <p>Chemical structure of 3,4-dihydro-1H-inden-5-ol, showing a benzene ring fused to a five-membered ring containing an oxygen atom, with a hydroxyl group on the benzene ring.</p>
E	232	209	 <p>Chemical structure of 1,3-dihydroxybenzene (resorcinol), showing a benzene ring with two hydroxyl groups at the 1 and 3 positions.</p>
F	133	110	 <p>Chemical structure of trans,trans-hexa-2,4-diene-1,6-dioic acid, showing a six-carbon chain with two carboxylic acid groups and two trans double bonds.</p>
G	166	143	 <p>Chemical structure of trans,trans-2,5-dihydroxyhexa-2,4-diene-1,6-dioic acid, showing a six-carbon chain with two carboxylic acid groups, two trans double bonds, and two hydroxyl groups.</p>

Proposed pathway for Carbofuran degradation

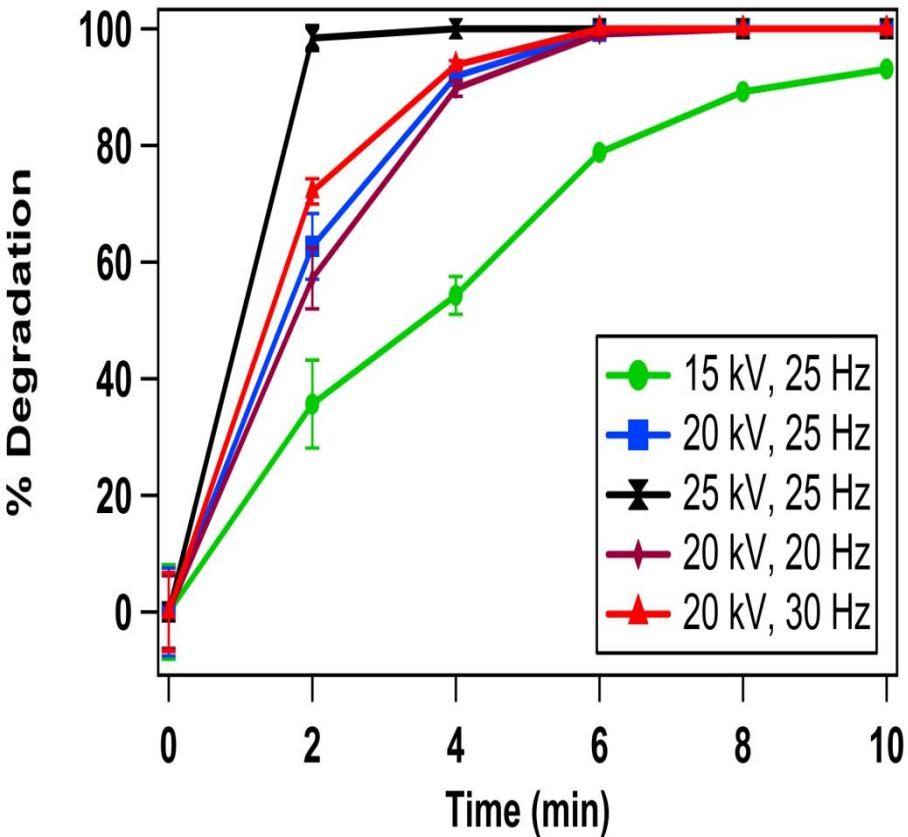


Eco-toxicity assay for Carbofuran degradation

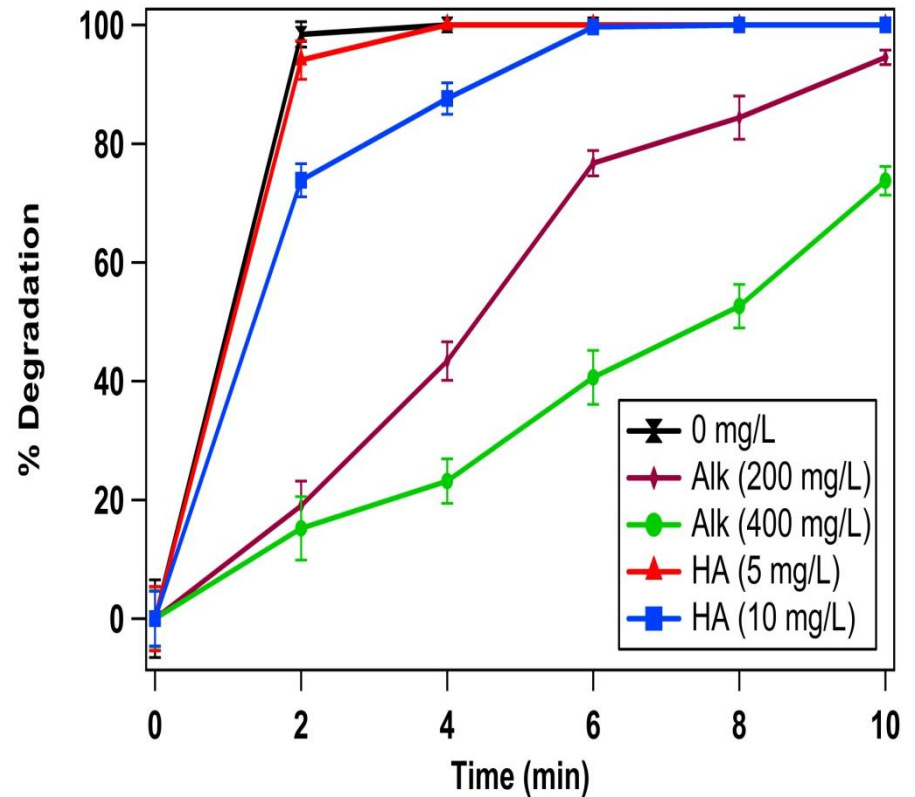
- Model Micro-alga – *Chlorella Vulgaris*



Pesticide: 2-4-D



Voltage and Frequency effect

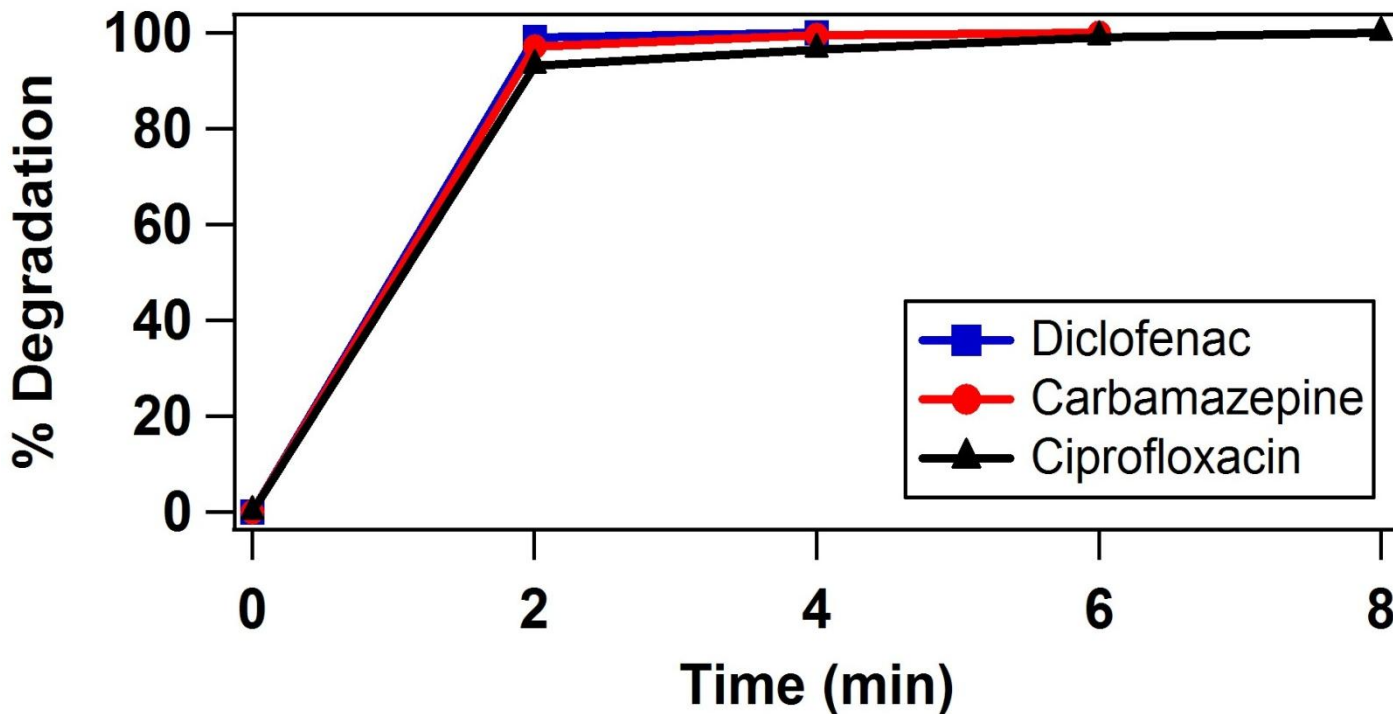


Effect of radicals quencher

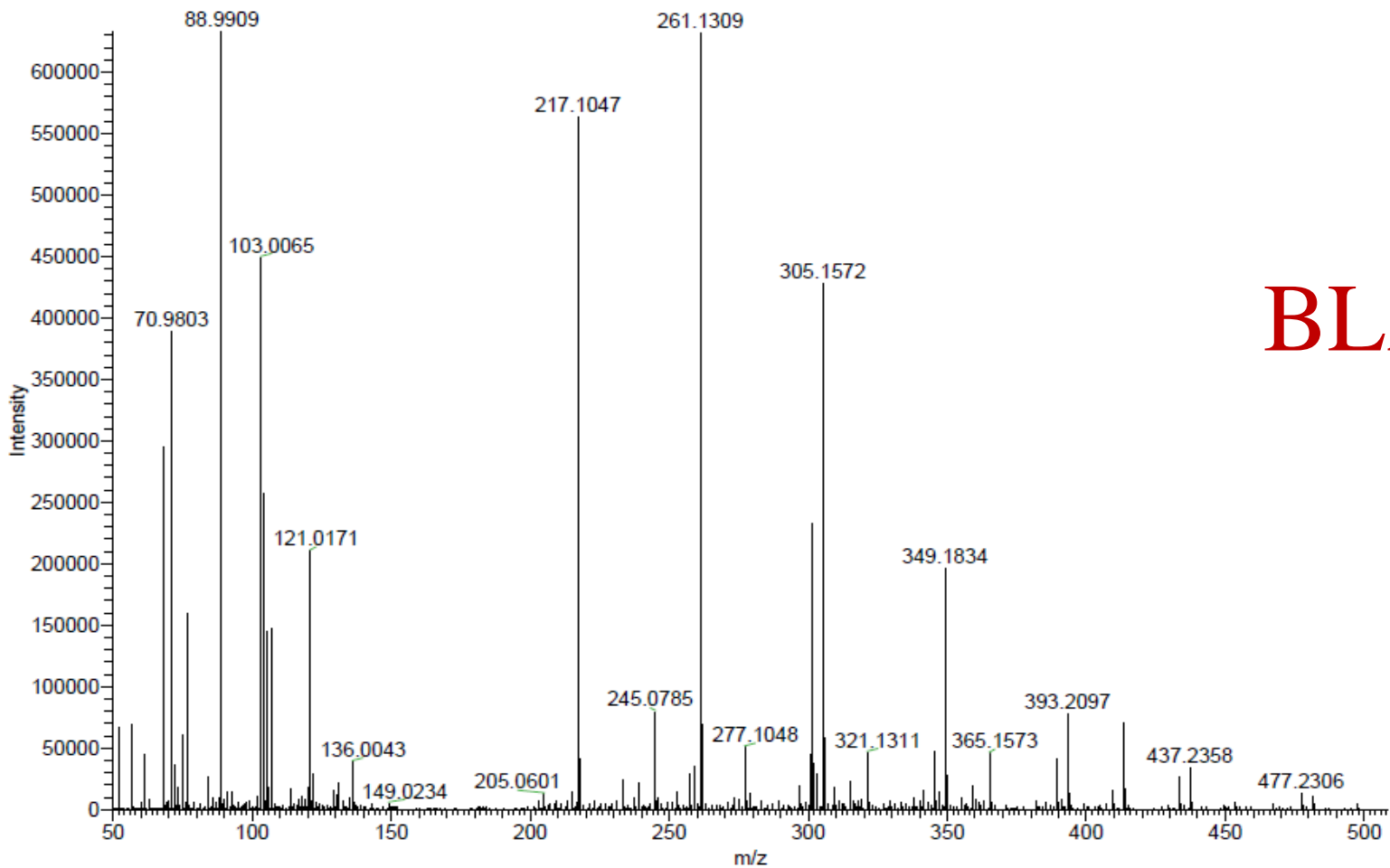
Pharmaceuticals Active Compounds (PACs)

PACs – **Diclofenac, Carbamazepine and Ciprofloxacin – 1ppm**

Voltage – **25 kV** and Frequency – **25 Hz**



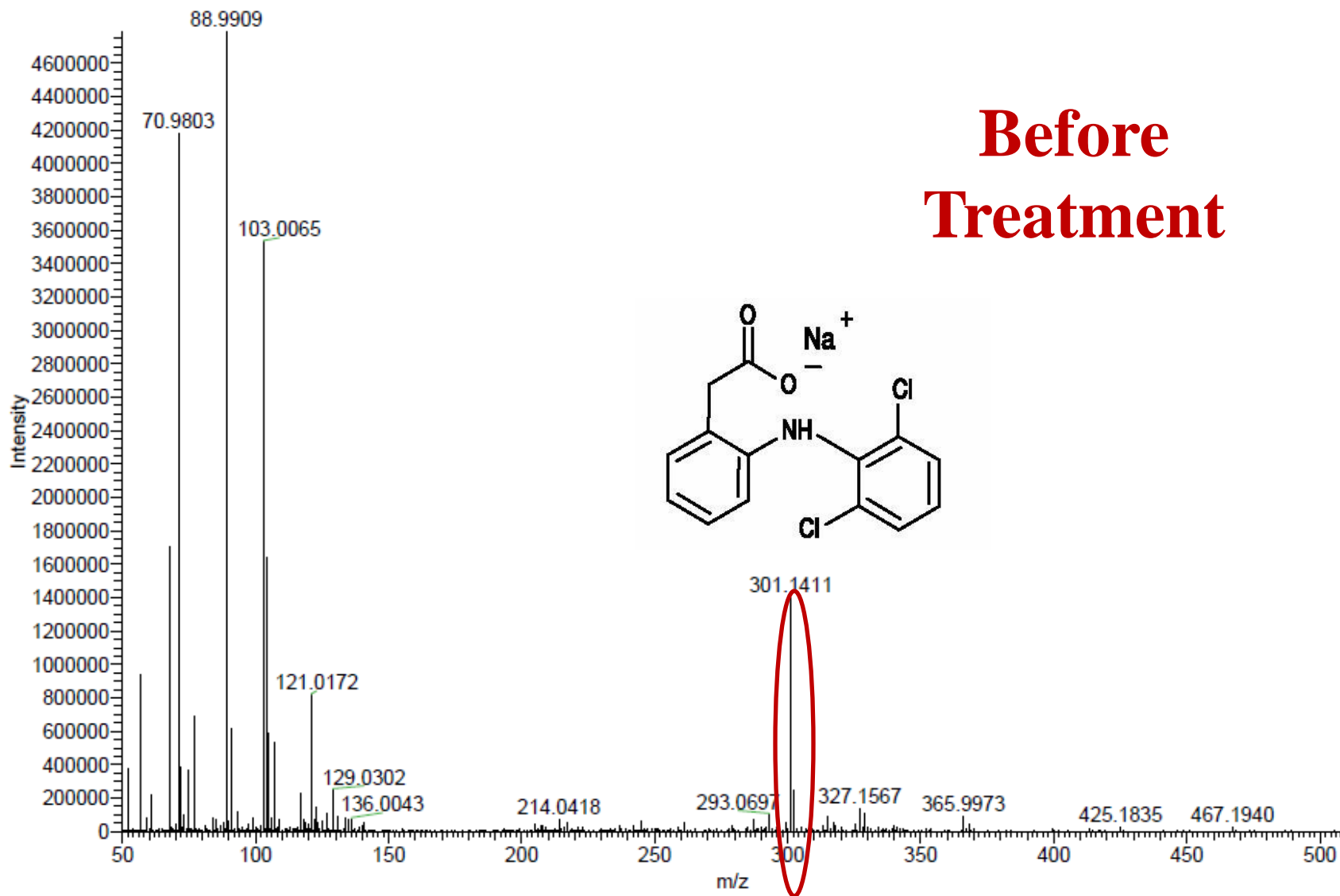
LC-MS Analysis



BLANK

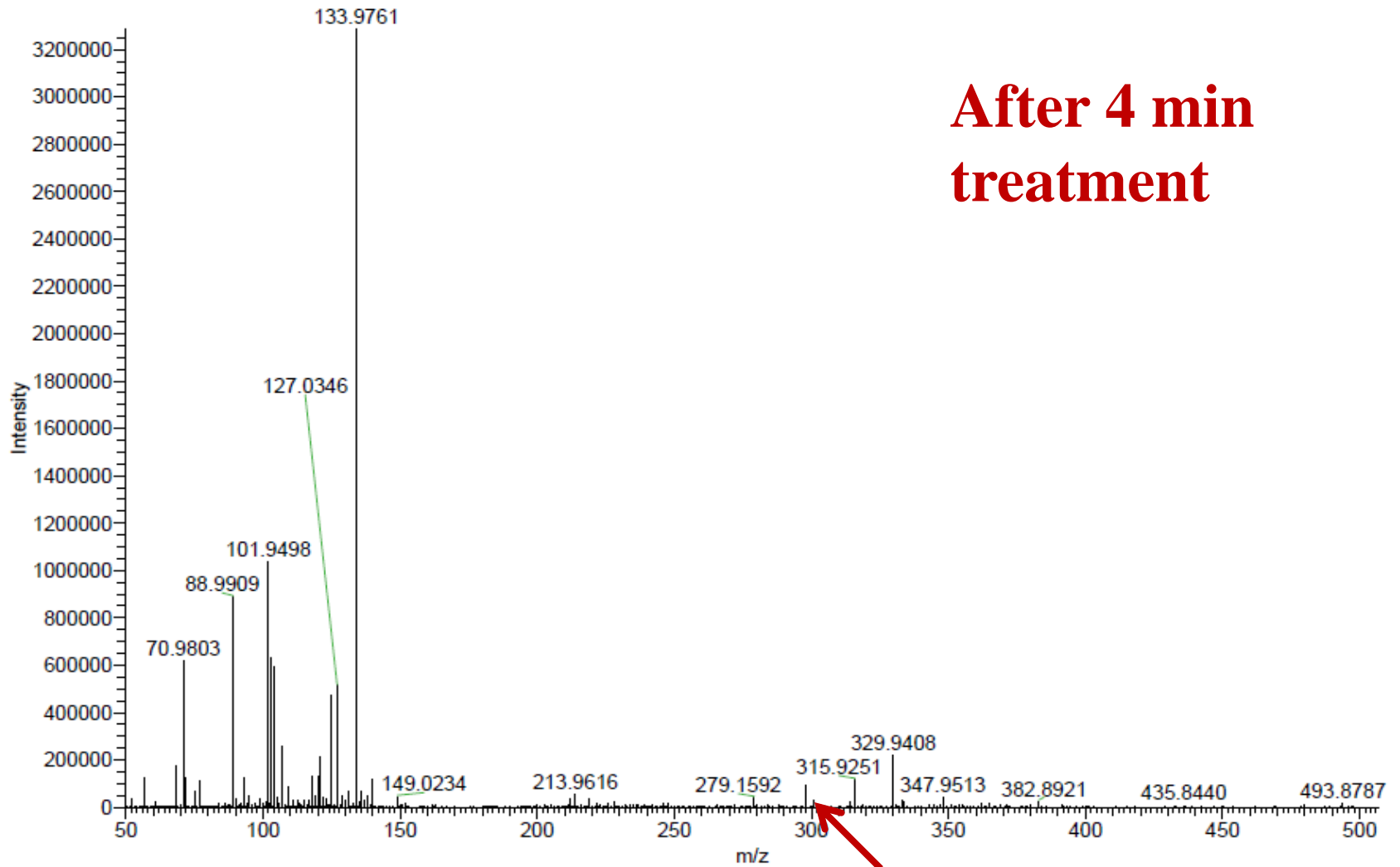
Diclofenac (DCF)

**Before
Treatment**

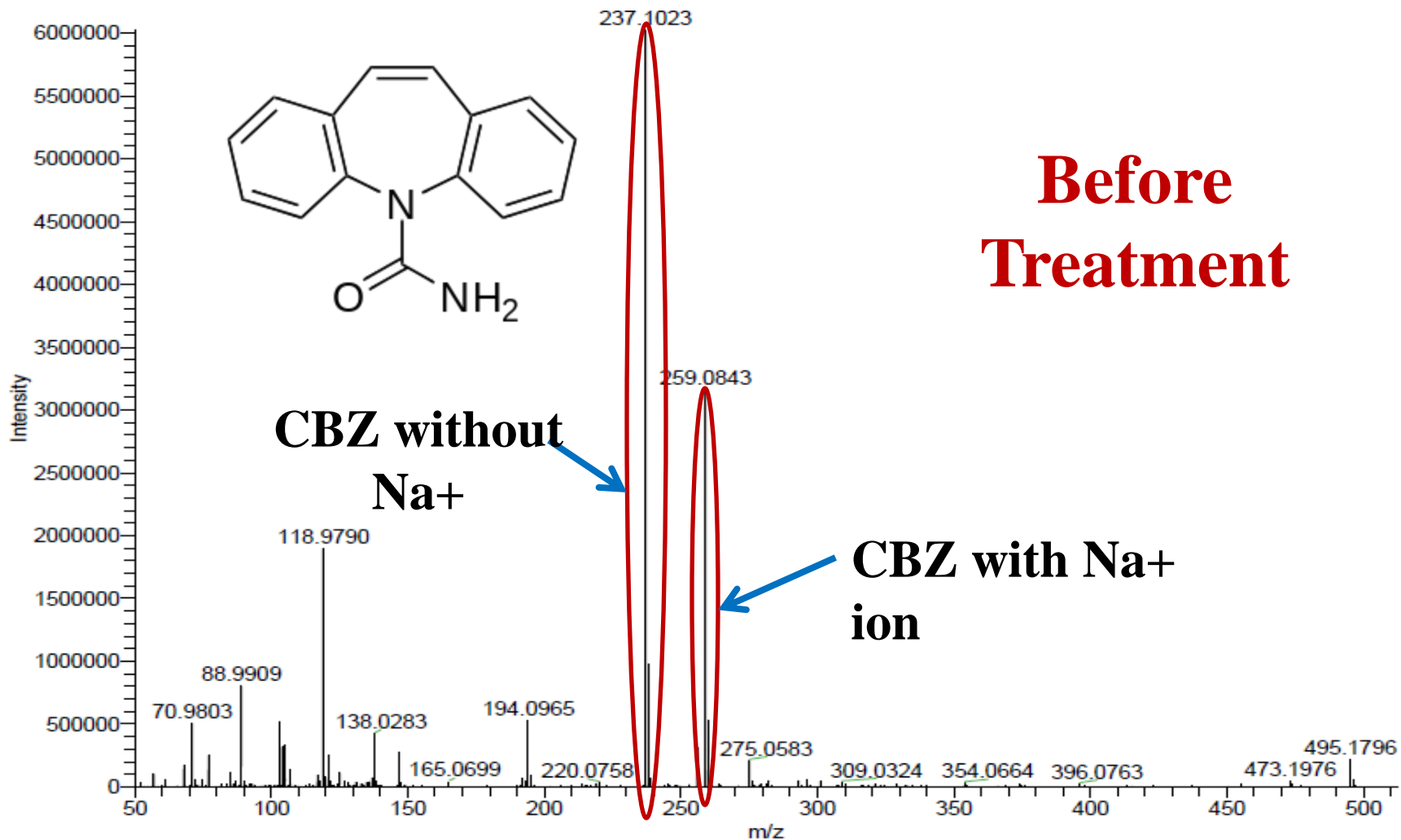


Diclofenec

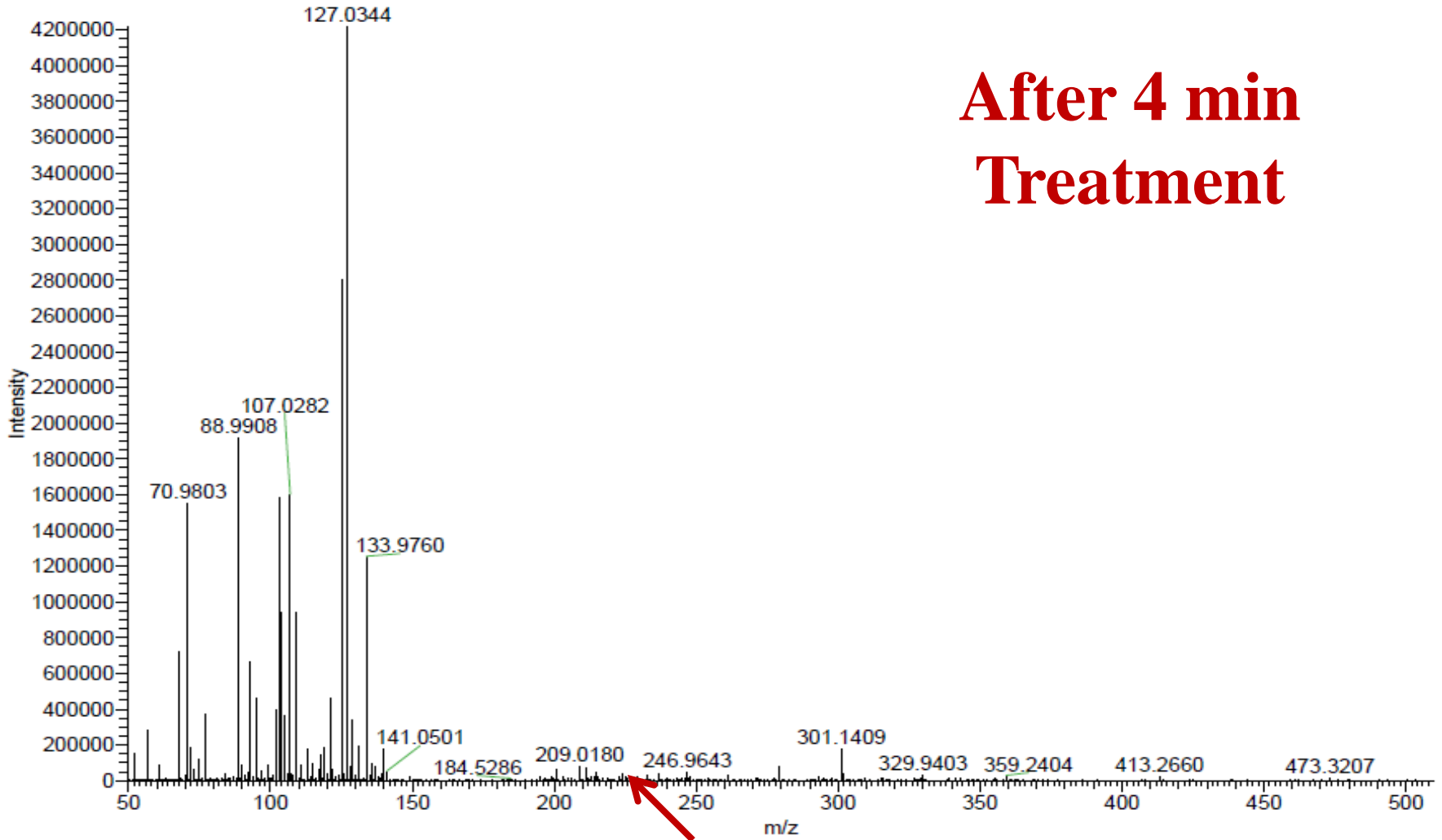
**After 4 min
treatment**



Carbamazepine (CBZ)

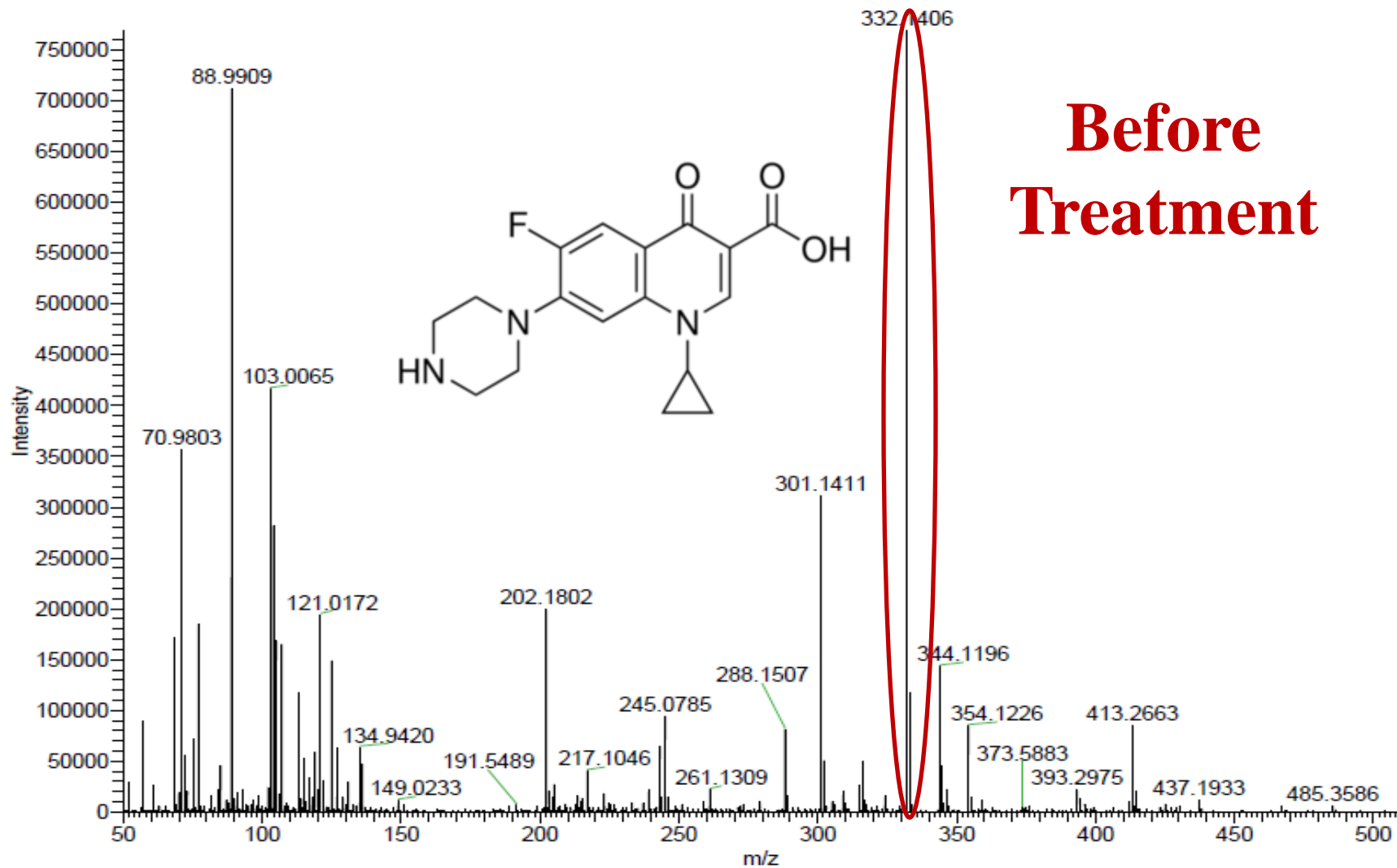


Carbamazepine

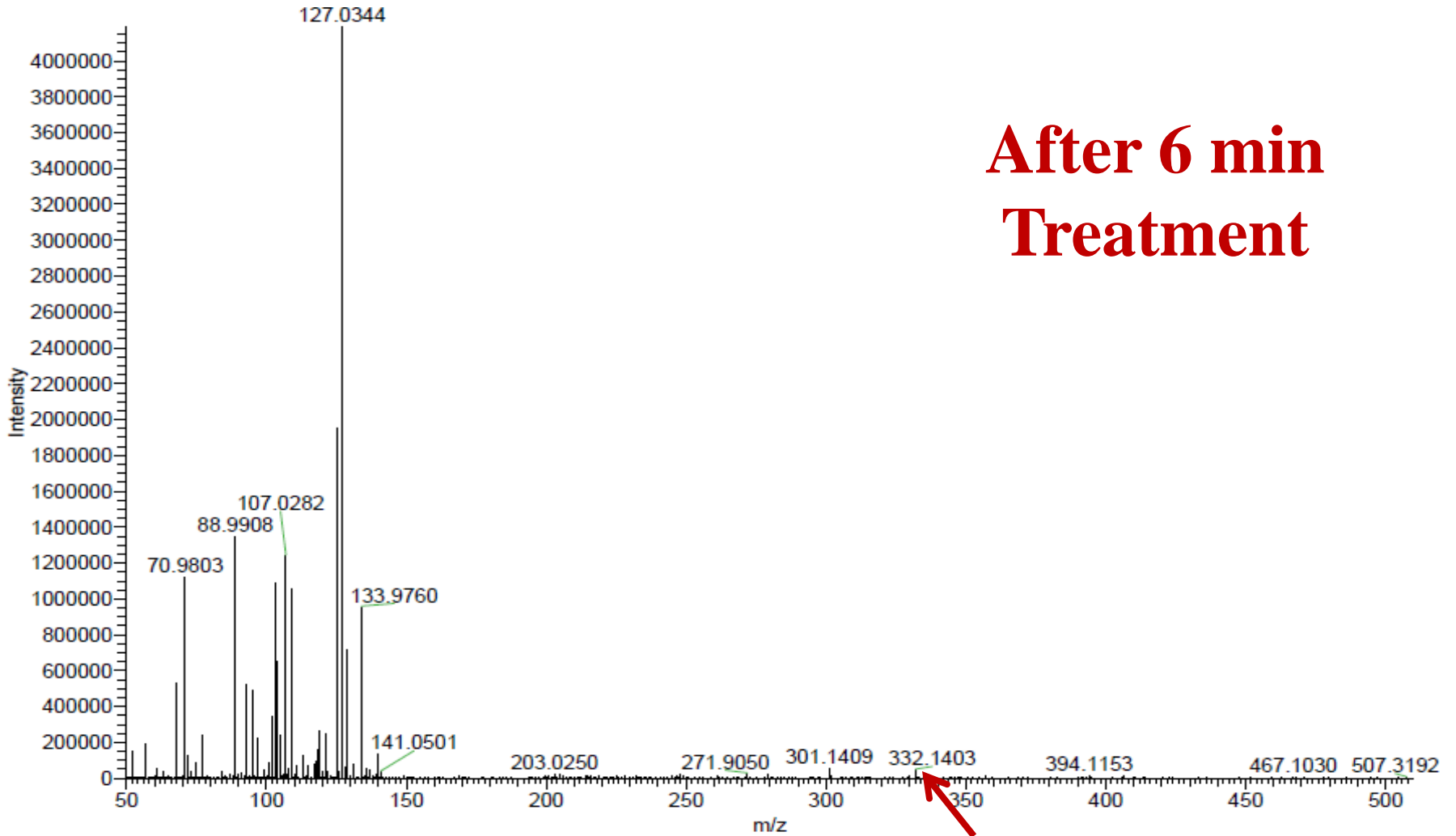


**After 4 min
Treatment**

Ciprofloxacin (CPF)

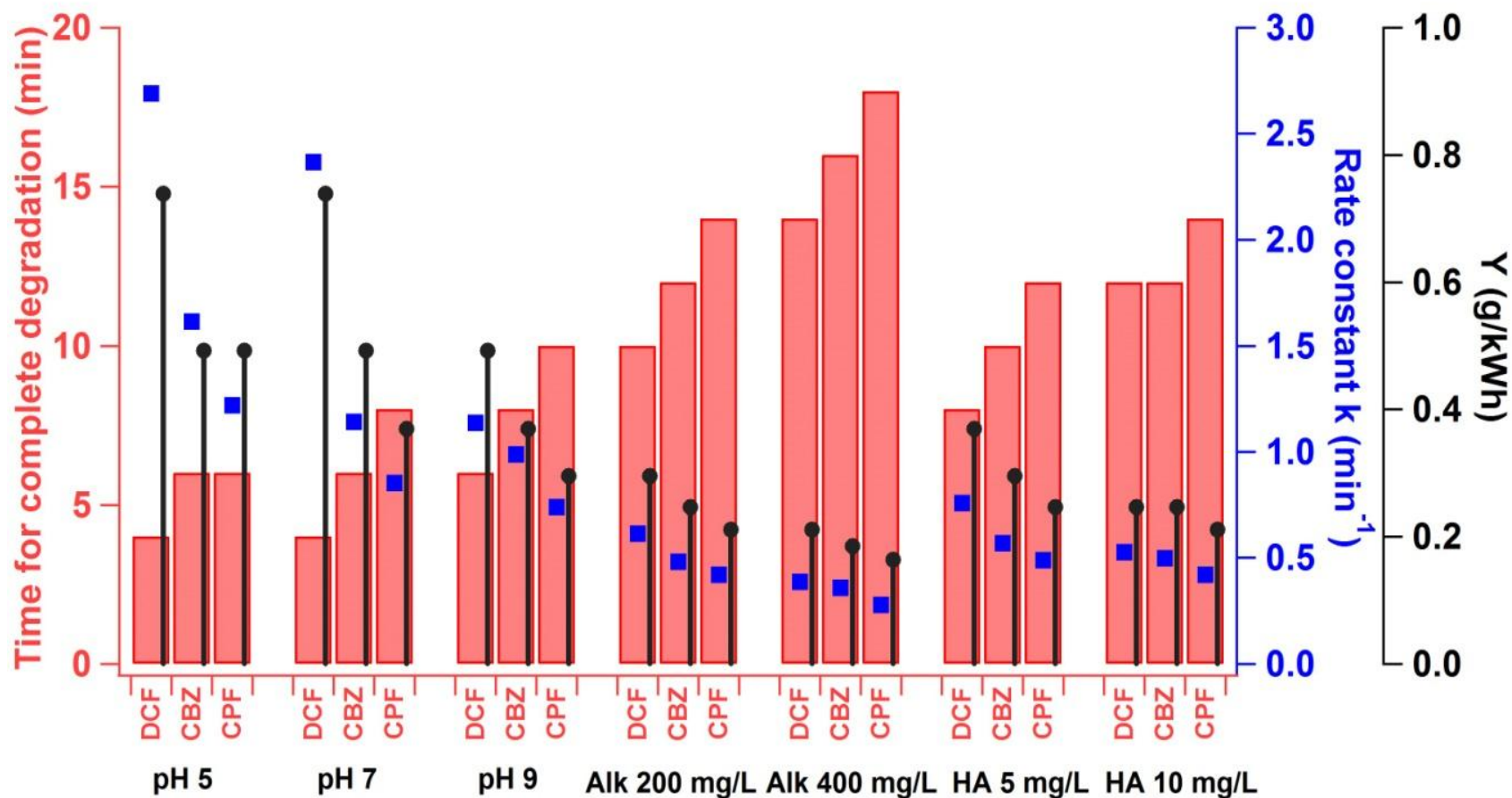


Ciprofloxacin

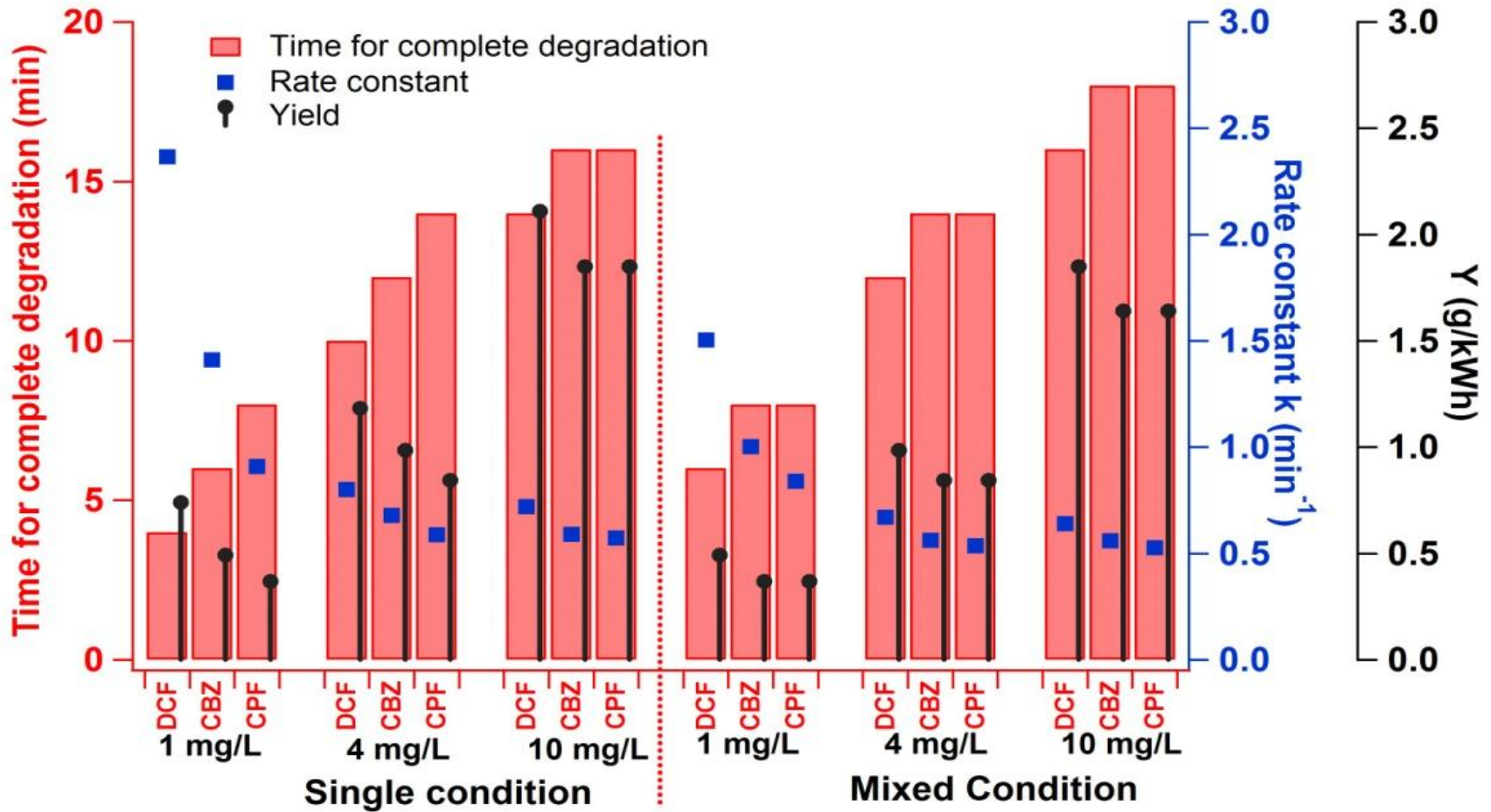


**After 6 min
Treatment**

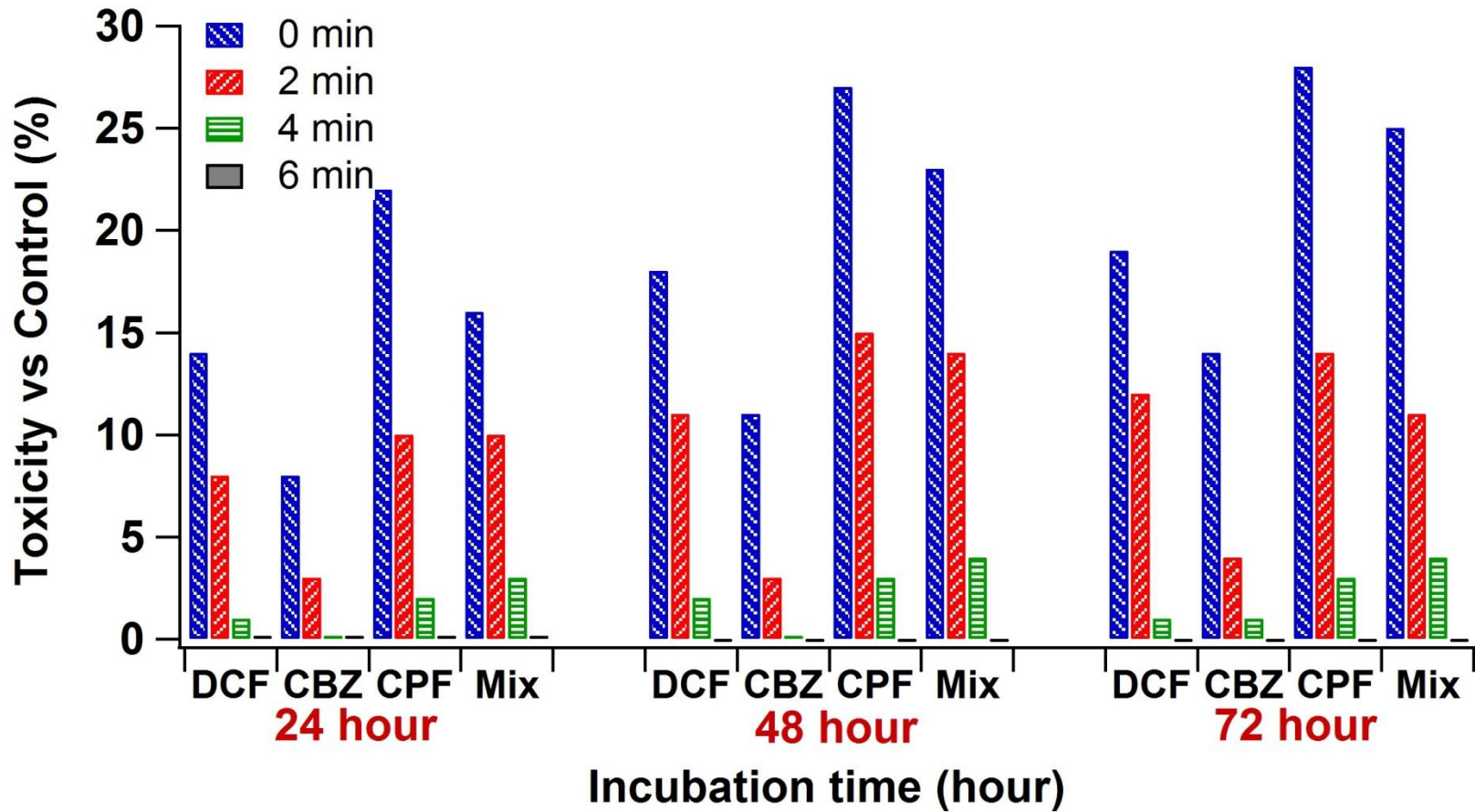
Effect of pH and radical scavengers



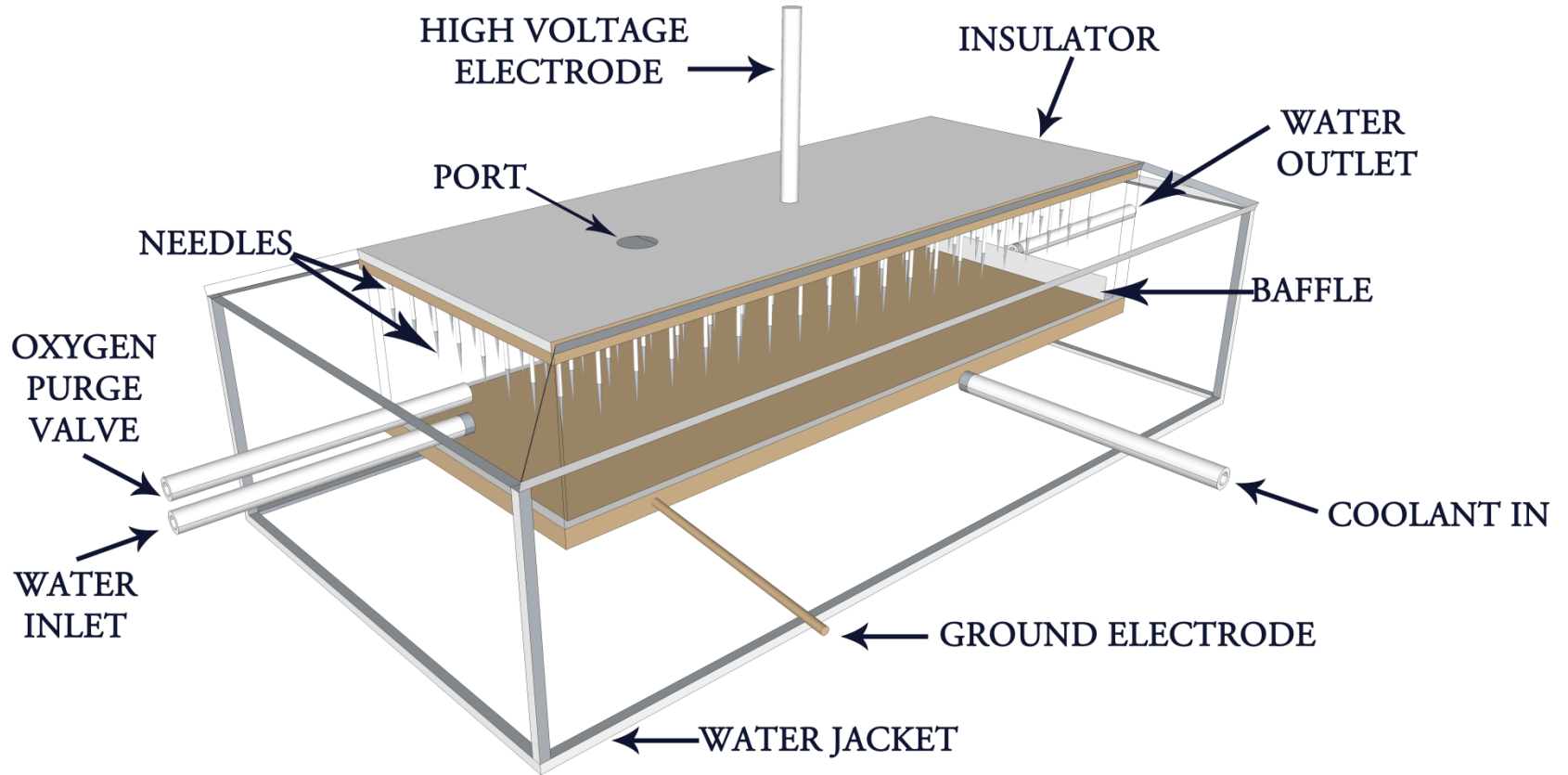
Single and mixed pollutant degradation



Toxicity assay



5. Continuous Reactor study

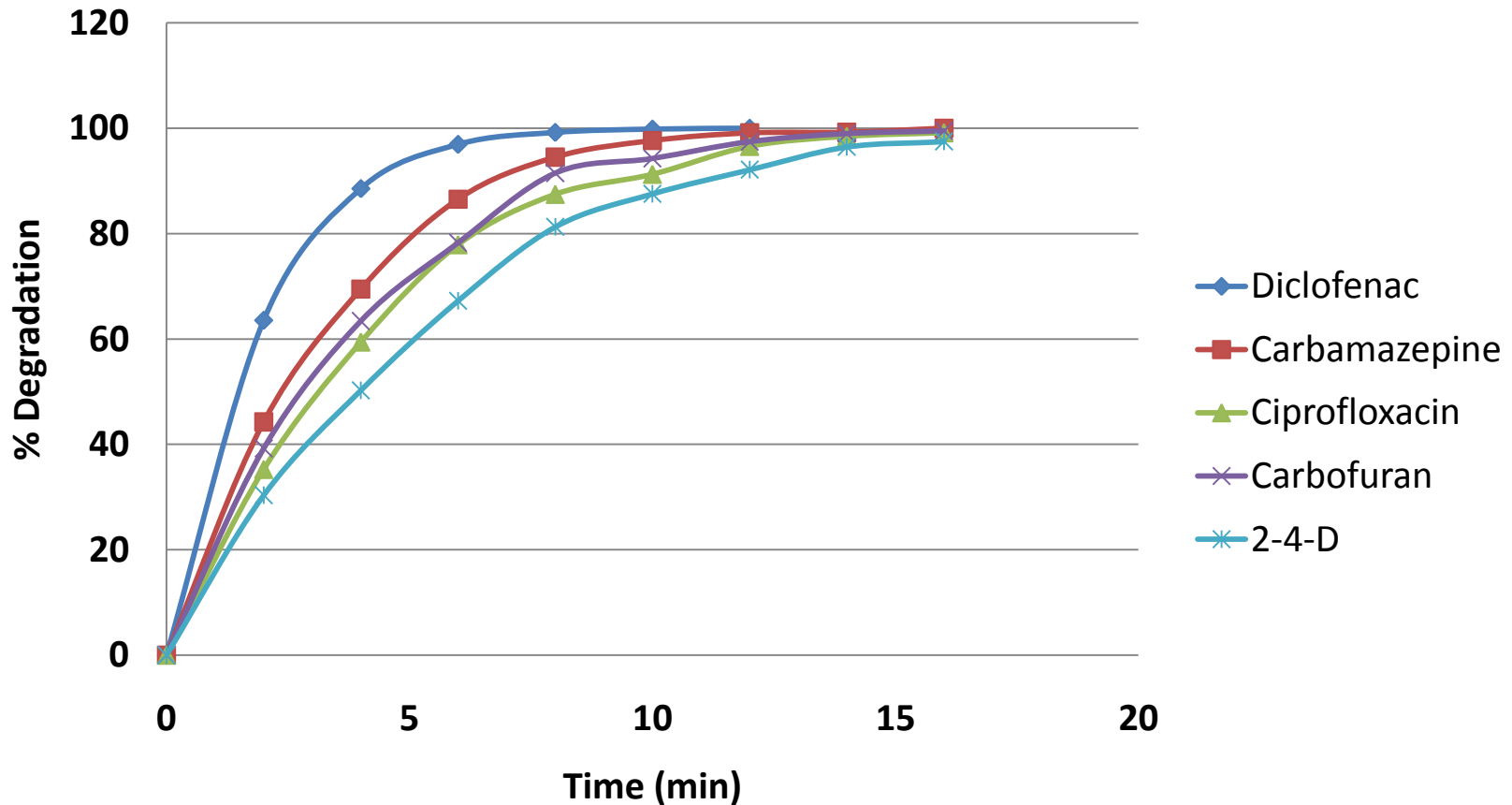


Continuous Reactor

- Reaction volume – 29 cm x 19 cm x 0.5 cm
- Spacing between the needles = 2 cm (optimized in batch reactor)
- Optimized flow rate = 10 mL/min

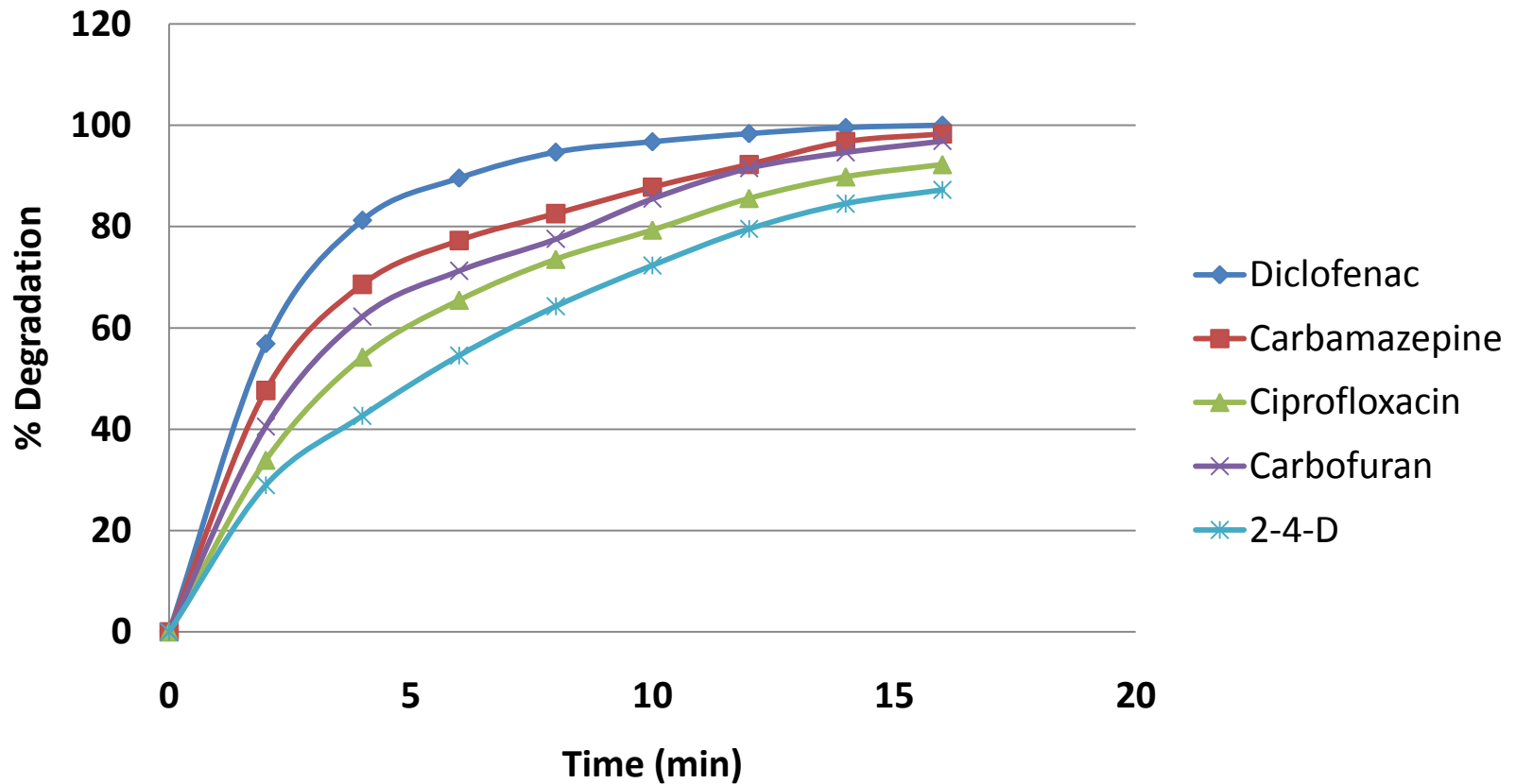
ECs degradation study

- Initial concentration – 1ppm



ECs degradation study

- Initial conc. – 10 ppm



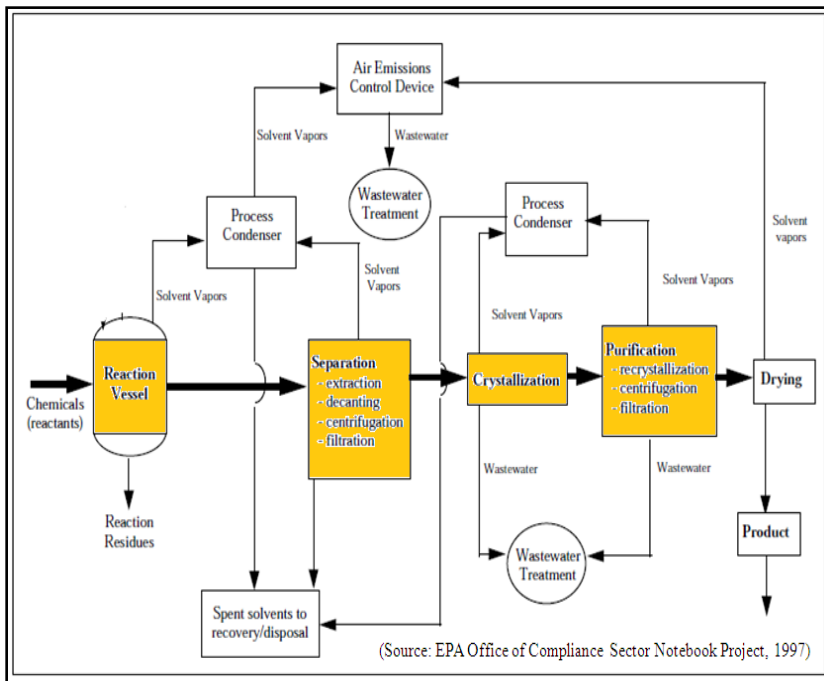
Summary/Conclusion

- Different **ROS** such as $\cdot\text{OH}$, H_2O_2 , O_3 and $\text{O}_2^{\cdot-}$ **quantification** in different environmental conditions.
- Effect of different **time mode** on disinfection efficiency.
- **Combined effect** of system parameters on disinfection efficiency and **empirical model** development.
- Understanding the bacterial **disinfection mechanism** in PPT.
- Study on **dye degradation and its fate** in PPT process.

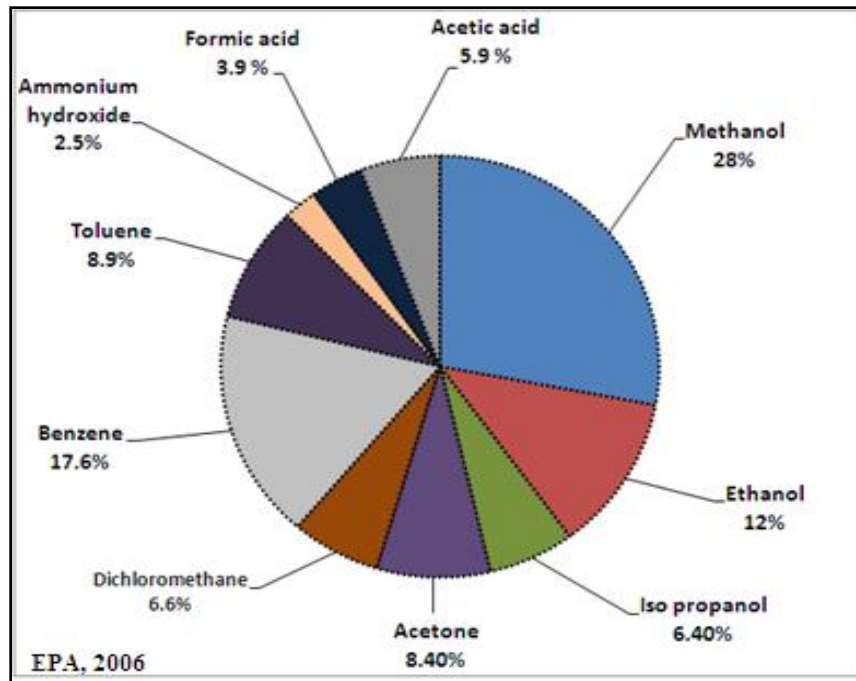
Summary/Conclusion

- Complete degradation of ECs (Carbofuran, 2-4-D, DCF, CBZ and CPF) was achieved within **4 to 6 min** treatment time in batch study.
- **Environmental parameters** significantly affect the degradation efficiency.
- Possibility of **Reductive pathway in plasma technology** – not only oxidative pathway.
- **Complete mineralization and detoxification of ECs** was achieved.
- Continuous reactor – design and efficiency was evaluated.

Development and Performance Evaluation of a Hybrid Treatment System for the Complete Treatment of Pharmaceutical Wastewater



PROCESS IN PHARMACEUTICAL INDUSTRY



DISTRIBUTION OF VOLATILE SOLVENTS IN WASTEWATER

MAJOR PROBLEM : VOC emissions during the treatment of pharmaceutical wastewater.

LIMITATION OF EXISTING TREATMENT SYSTEMS

- Only focus on removal of organic pollutants
- Emission of VOC is not accounted

Saravanane et al., 2001;
 Ince et al., 2002
 Raj and Anjaneyulu, 2005

“Reduction of VOC emission from the treatment units”

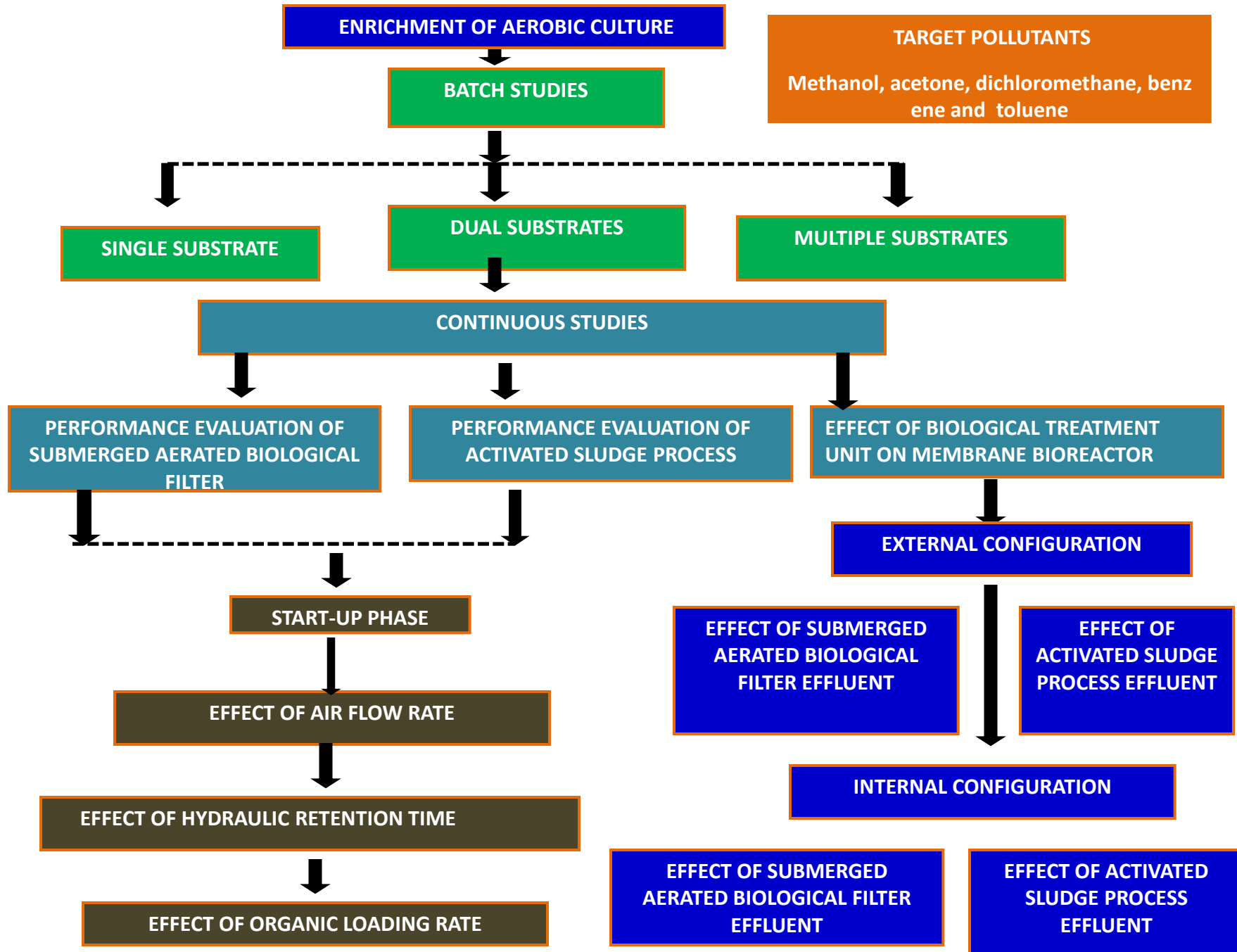
LIMITATIONS IN EXISTING TREATMENT SYSTEM FOR REMOVAL OF VOC

- | | |
|-------------------------------------------------------------------------------------|--------------------------------------|
| <input type="checkbox"/> Individual pollutant study | Cattony et al., 2005 |
| <input type="checkbox"/> Degradation of VOC at low concentration | Quesnel and Nakhla, 2005 |
| <input type="checkbox"/> No focus on the reduction of VOC emission from bioreactors | Ozdemir et al., 2010
Dawery, 2013 |

Presence of high biomass is reported to reduce VOC emissions

- Submerged aerated biological filter (SABF) and Membrane bioreactor (MBR)**
(Cheng , 2009; Min and Ergas, 2006)
- Need to evaluate the performance **SABF** to treat mixture of VOC
- Effect of operational parameters like air flow rate, hydraulic retention time (HRT) and organic loading rate (OLR) on VOC emission is an area to be explored
- Feasibility of **Membrane bioreactor** as a post treatment unit and its potential to reduce VOC emission is an area to be explored

METHODOLOGY



POLLUTANTS IN UNTREATED PHARMACEUTICAL WASTEWATER IN INDIA

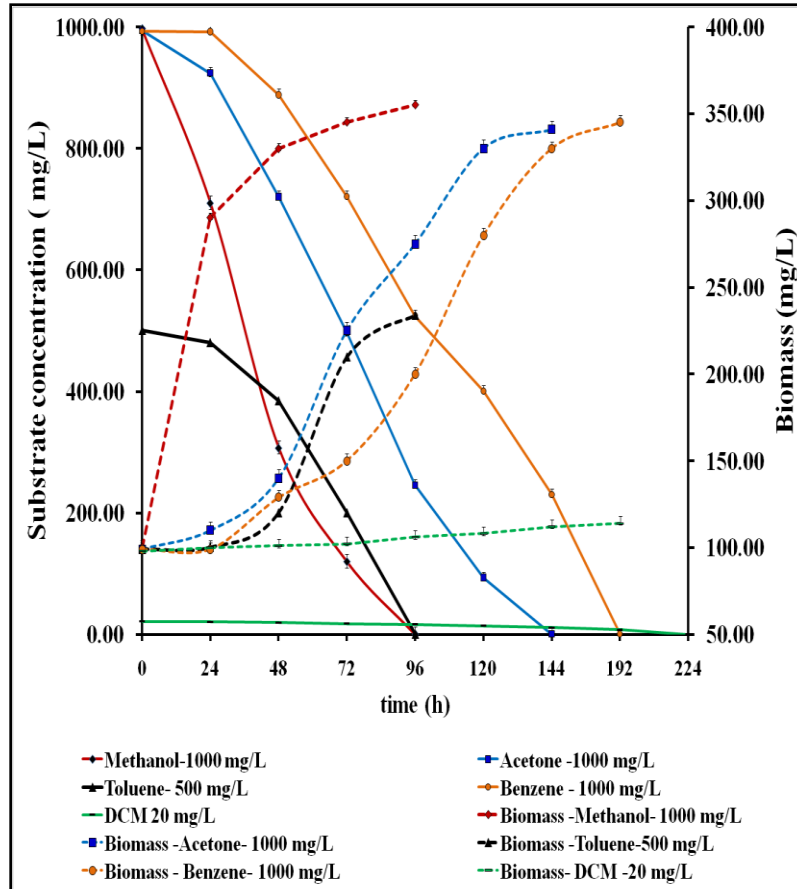
Methanol (2500–3000 mg/L) , Acetone (500 mg/L – 1000 mg/L), Benzene and toluene (400–700 mg/L), Dichloromethane (120 - 380 mg/L)

(Gupta et al., 2005, Virnig et al., 2003)

Studies with single substrate biodegradation		
Methanol, acetone , benzene	Toluene	Dichloromethane (DCM)
100, 300, 500, 700 and 1000 mg/L	100, 300 and 500 mg/L	10 and 20 mg/L
Dual substrate interaction studies with dichloromethane		
Low concentration studies	High concentration studies	
Methanol- DCM / Acetone- DCM Benzene- DCM/ Toluene- DCM Non chlorinated solvent ~ 100 mg/L , DCM ~ 50mg/L	Methanol- DCM / Acetone- DCM Benzene- DCM/ Toluene- DCM Non chlorinated solvent ~ 1000 mg/L, DCM ~ 50mg/L	
Multiple Substrate Interaction studies		
In the absence and presence of Dichloromethane	Mixture of methanol, acetone, benzene dichloromethane and toluene at equal concentration (50, 100, 200 mg/L)	

BATCH BIODEGRADATION RESULTS

SINGLE SUBSTRATE DEGRADATION



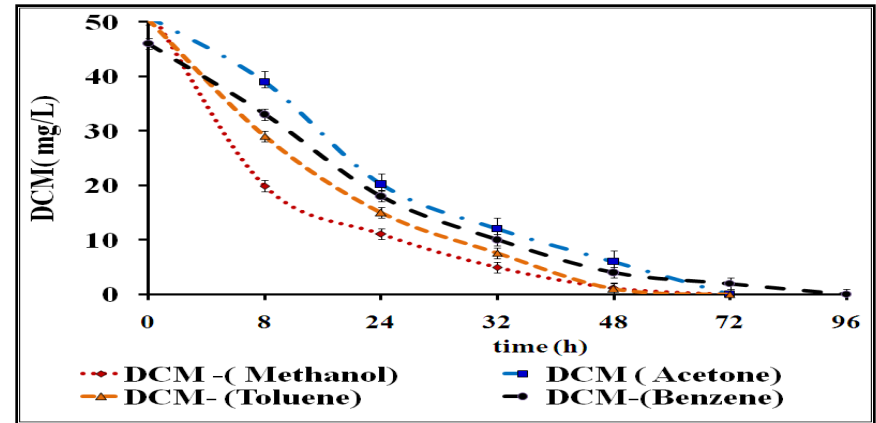
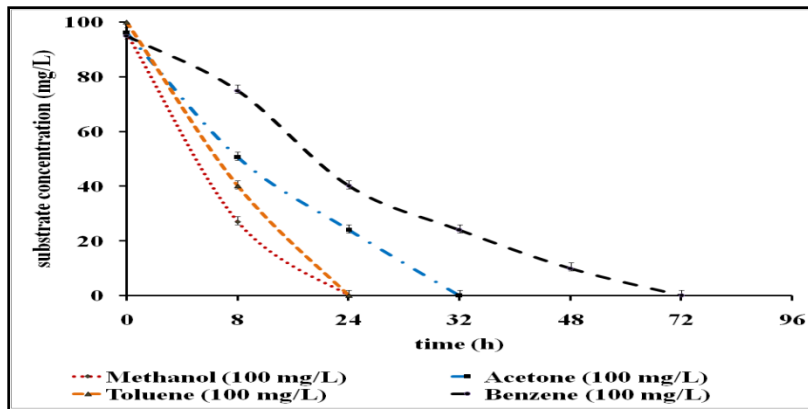
- ❑ Degradation of Non chlorinated pollutants were faster
- ❑ Dichloromethane was observed to be recalcitrant to biodegradation
- ❑ *Burkholderia kururiensis* and *Bacillus cereus* were predominant species.
- ❑ Monod inhibition model predicted single pollutant biodegradation

Priya, V.S., Philip, L. (2013). Biodegradation of Dichloromethane along with other VOCs from Pharmaceutical wastewater. *Applied Biochemistry and Biotechnology*. 169, 1197–1218.

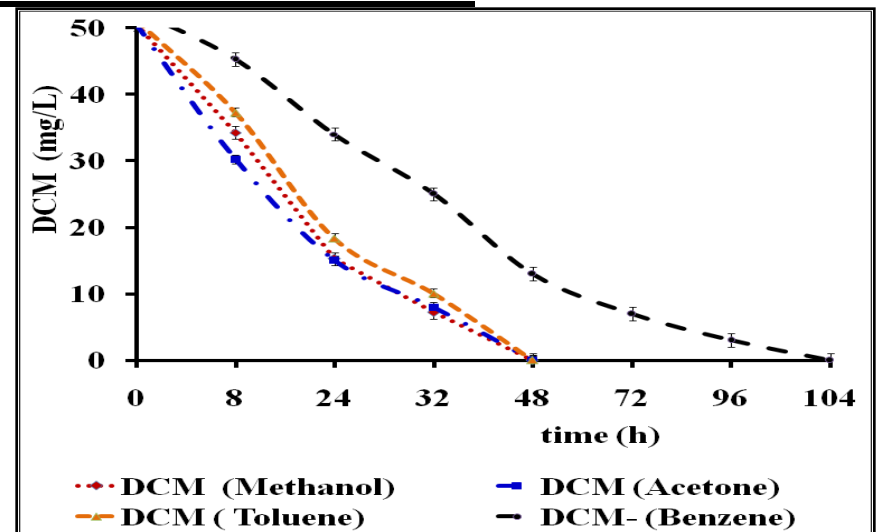
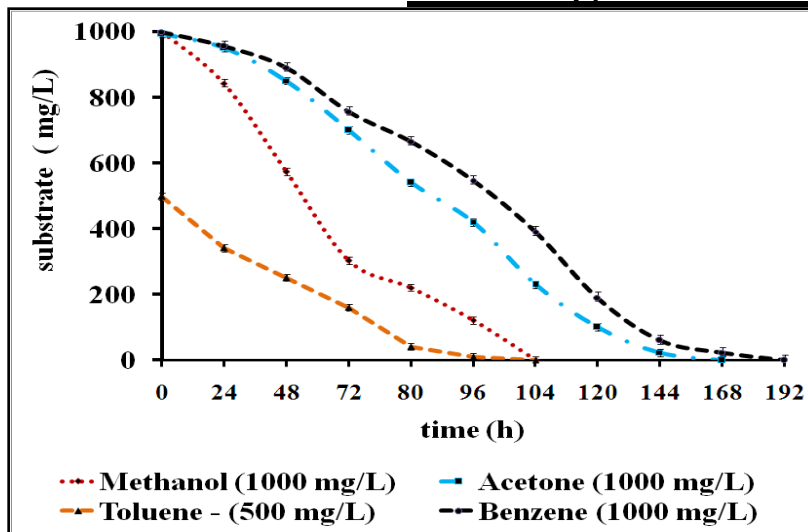
DUAL SUBSTRATE INTERACTION

STUDIES

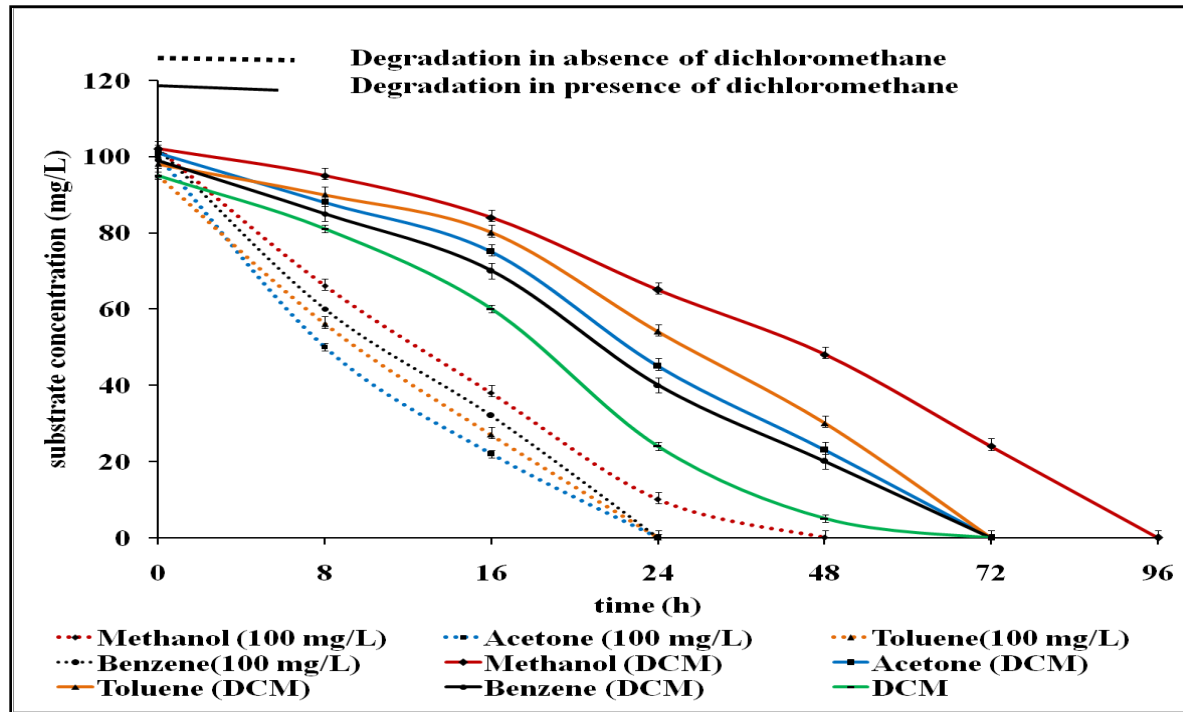
Degradation of 50 mg/L of dichloromethane in the presence of 100 mg/L of non chlorinated solvents



Degradation of 50 mg/L of dichloromethane in the presence of 1000 mg/L of non chlorinated solvents



MULTIPLE SUBSTRATE INTERACTION STUDIES



- ❑ **Absence of DCM** : All the non chlorinated solvents were degraded much faster compared to their degradation in a single pollutant system.
- ❑ **Presence of DCM** : Presence of DCM prolonged the degradation of all the non chlorinated solvents
- ❑ **Enhanced degradation of dichloromethane in the presence of other solvents**

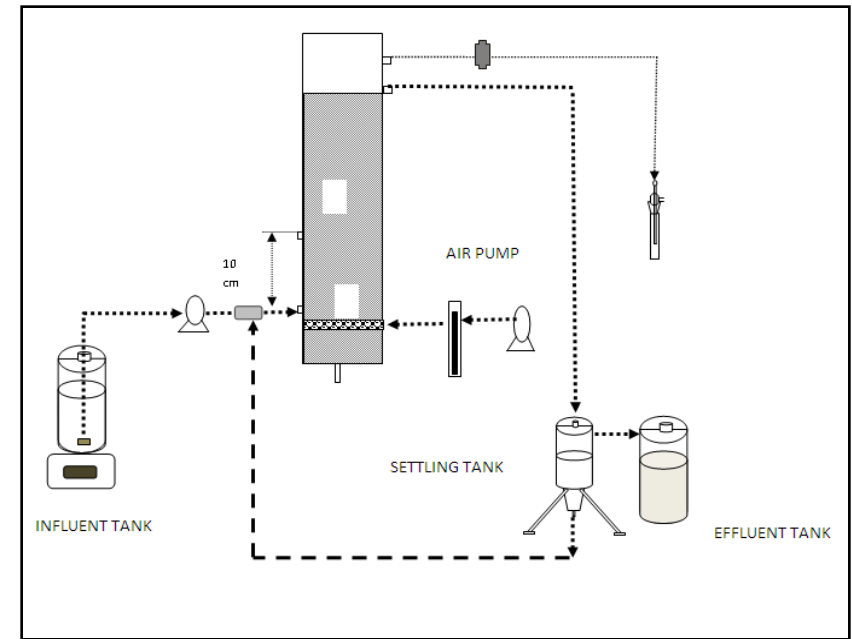
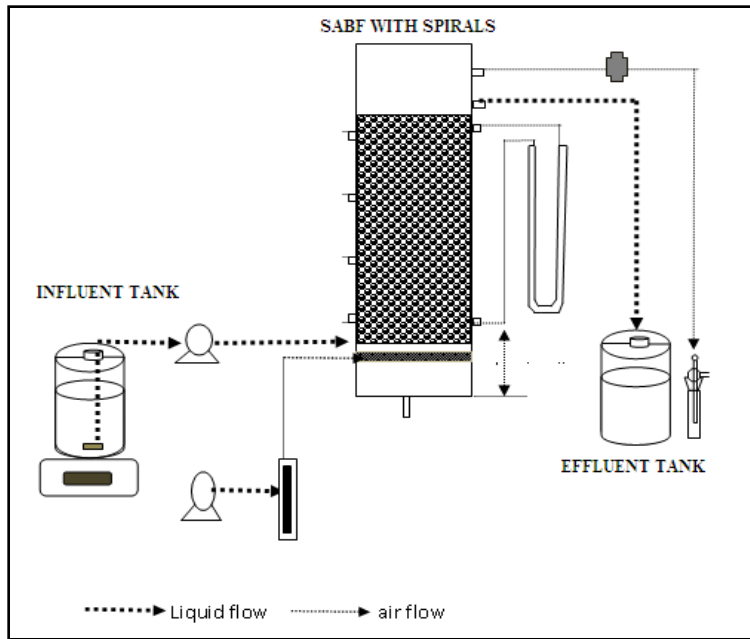
CONCLUSION FROM BATCH STUDIES

- ❑ All the target pollutants were degraded in the aerobic conditions
- ❑ **First report on the enhanced degradation of dichloromethane in the presence of other non chlorinated pollutants**
- ❑ Low concentrations (100 mg/L) of non chlorinated solvents did not interfere with the DCM degradation
- ❑ High concentrations of non chlorinated solvents (1000 mg/L) enhanced the DCM degradation and a severe competition between the chlorinated and the non chlorinated solvents was observed.
- ❑ In multiple substrate system also, presence of DCM prolonged the degradation of the other non chlorinated solvents.

Priya, V.S., Philip, L. (2013). Biodegradation of Dichloromethane along with other VOCs from Pharmaceutical wastewater. *Applied Biochemistry and Biotechnology*. 169, 1197–1218.

DEGRADATION STUDIES IN CONTINUOUS BIOREACTORS

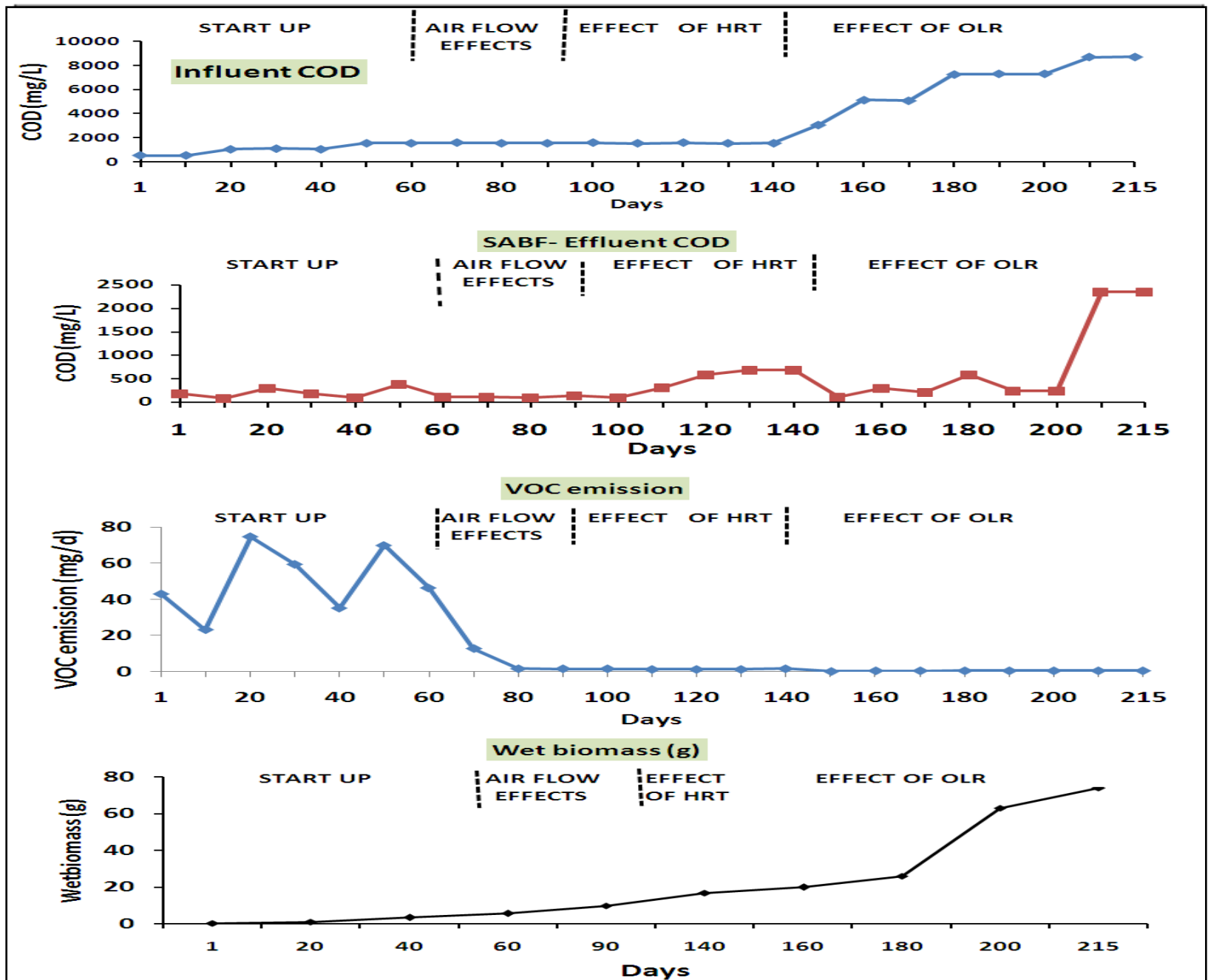
CONTINUOUS BIOREACTORS



SUBMERGED AERATED BIOLOGICAL FILTER

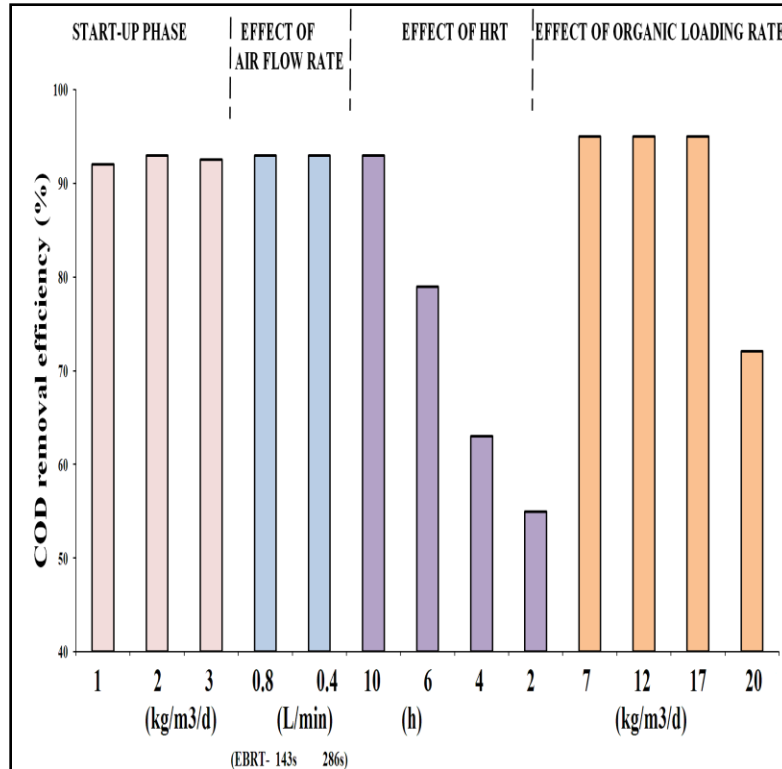
ACTIVATED SLUDGE PROCESS

CONTINUOUS BIOREACTORS

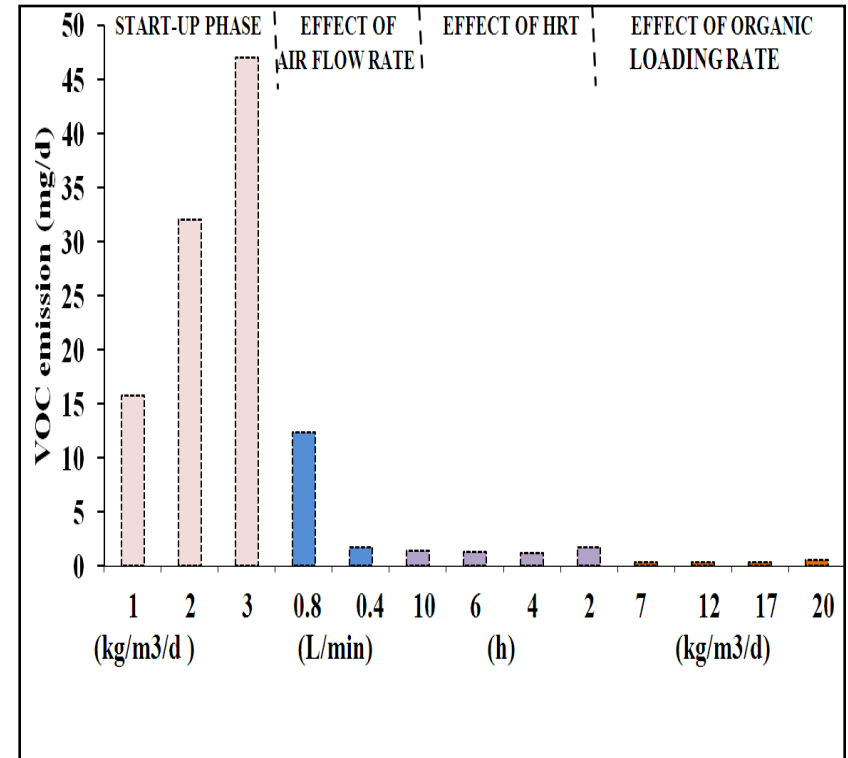


PERFORMANCE OF SUBMERGED AERATED BIOLOGICAL FILTER (SABF) UNDER DIFFERENT OPERATING CONDITIONS

RESULTS



COD REMOVAL FROM SABF

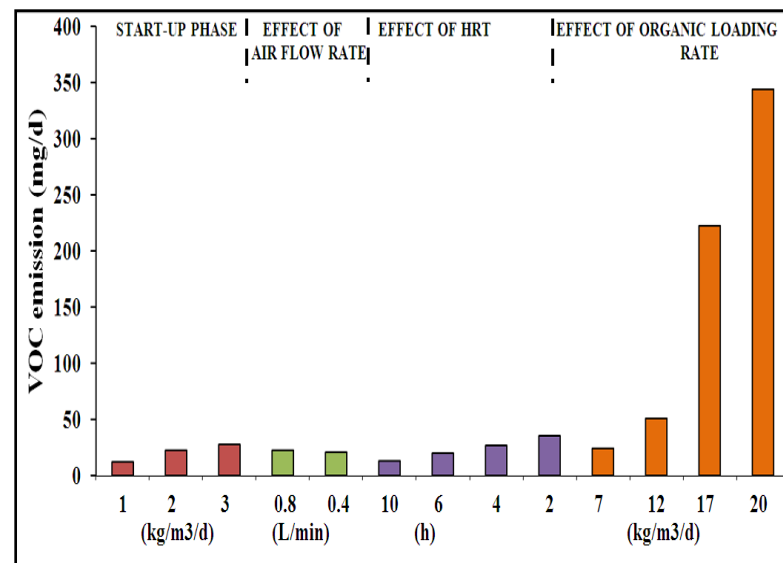
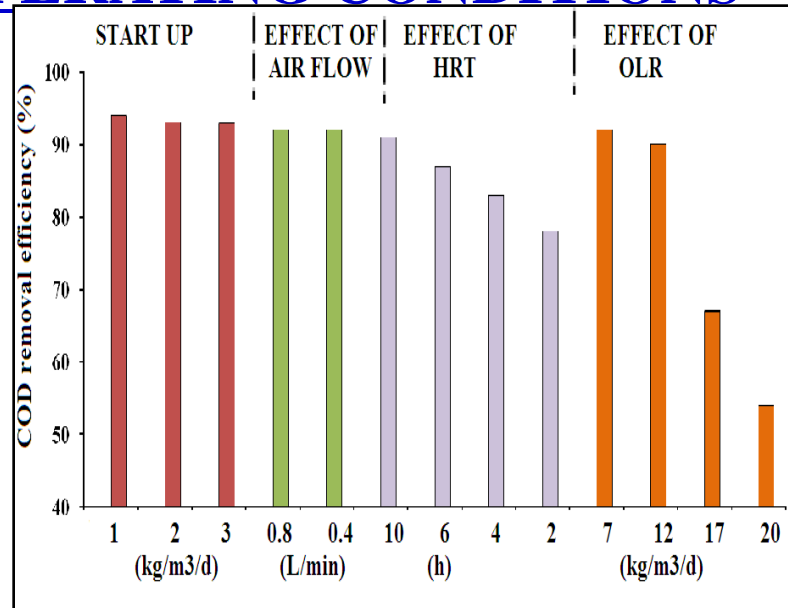
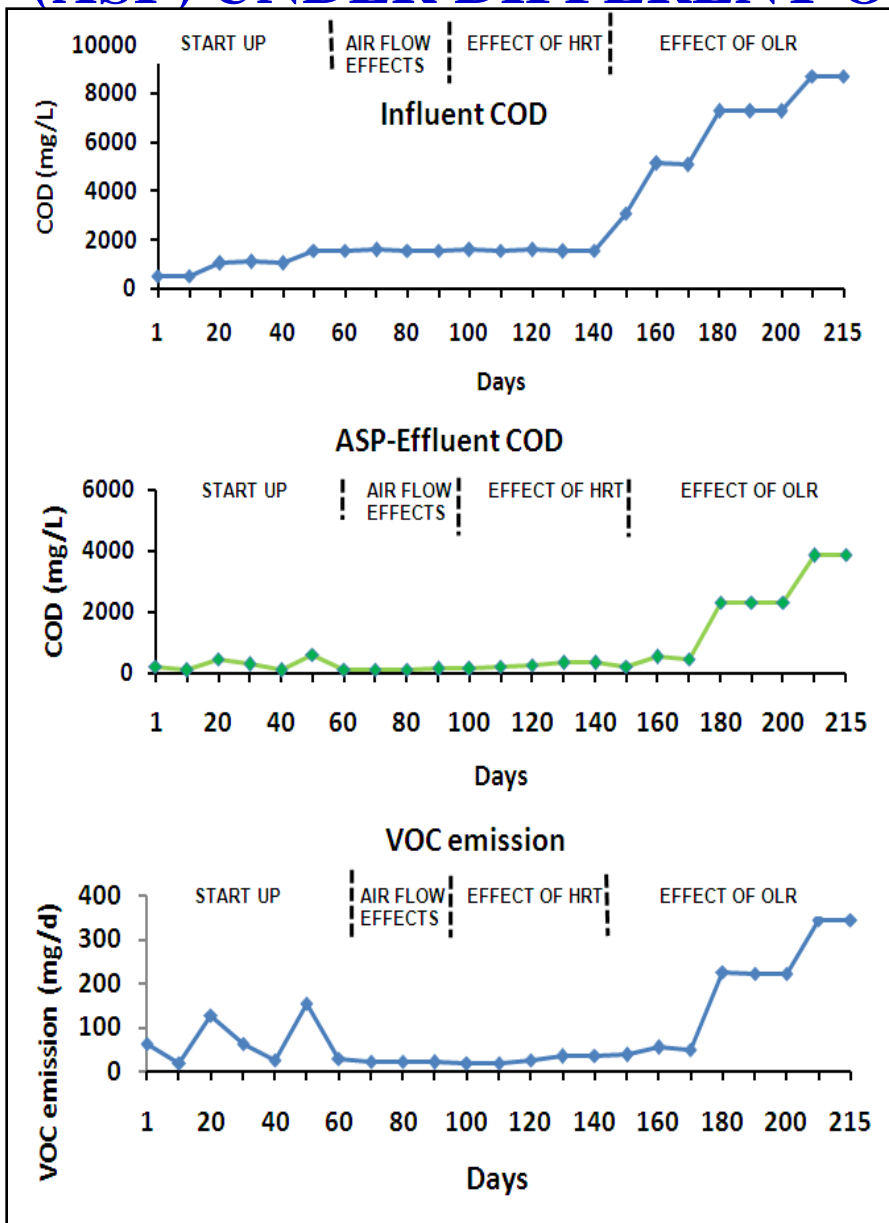


VOC EMISSION FROM SABF

Priya.V.S., Philip,L. Treatment of Volatile Organic Compounds in Pharmaceutical Wastewater using Submerged Aerated Biological Filter (Accepted in Chemical Engineering journal)

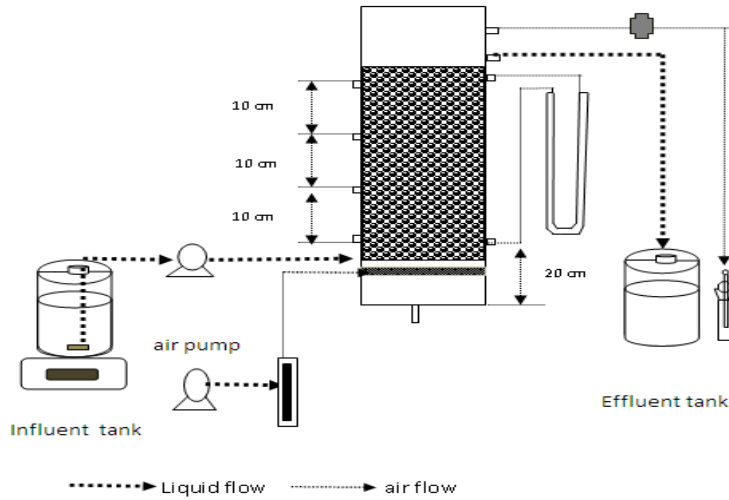
PERFORMANCE OF ACTIVATED SLUDGE PROCESS (ASP) UNDER DIFFERENT OPERATING CONDITIONS

RESULTS



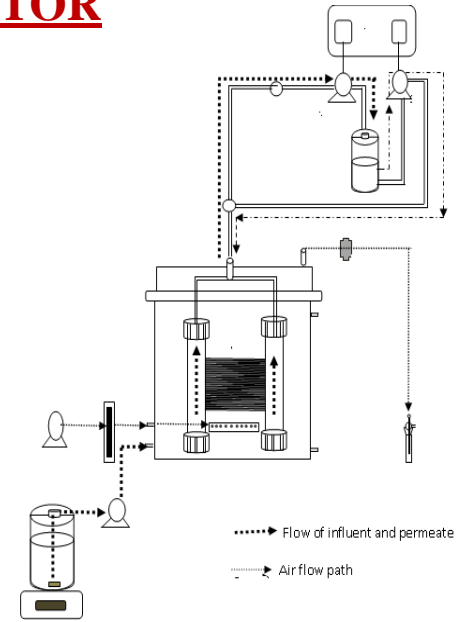
PERFORMANCE EVALUATION OF MEMBRANE BIOREACTOR

SABF COMBINED WITH MEMBRANE BIOREACTOR

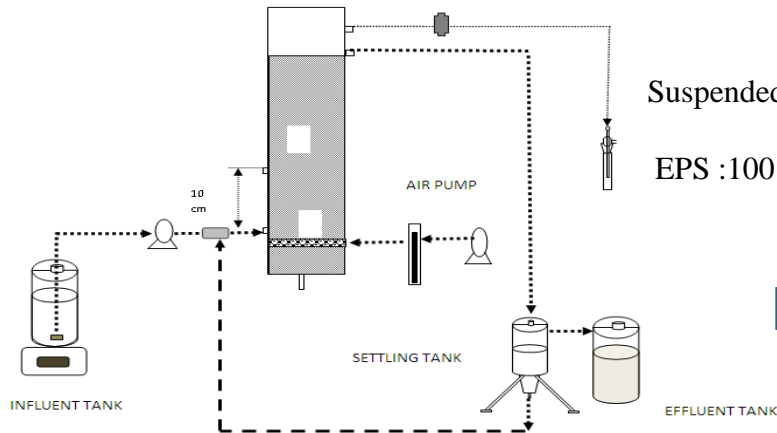


Suspended solids :60mg/L

EPS :12mg/g

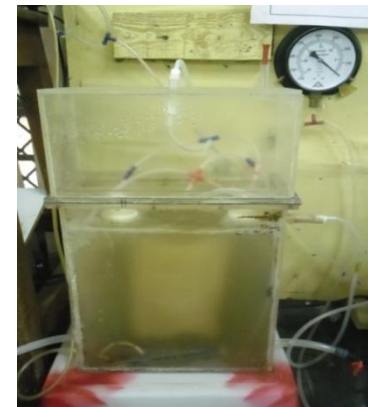


ASP COMBINED WITH MEMBRANE BIOREACTOR



Suspended solids :2000mg/L

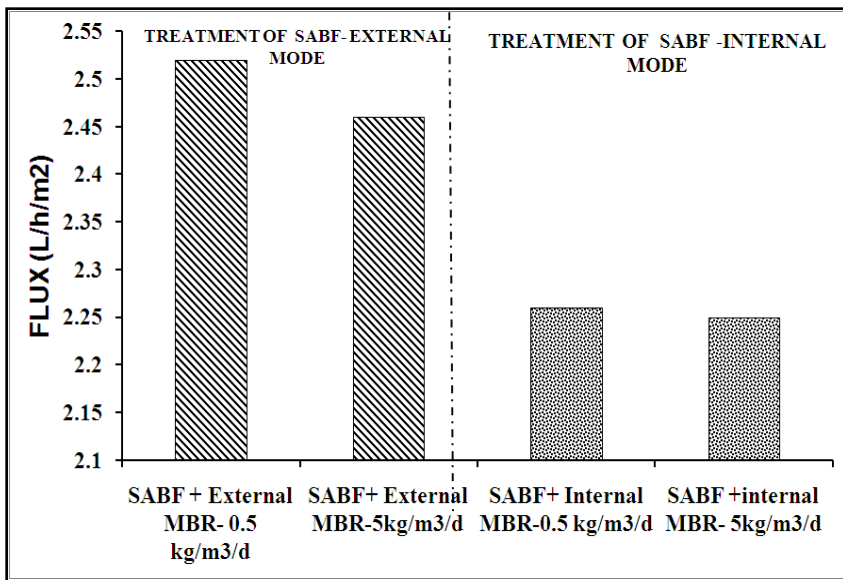
EPS :100 mg/g



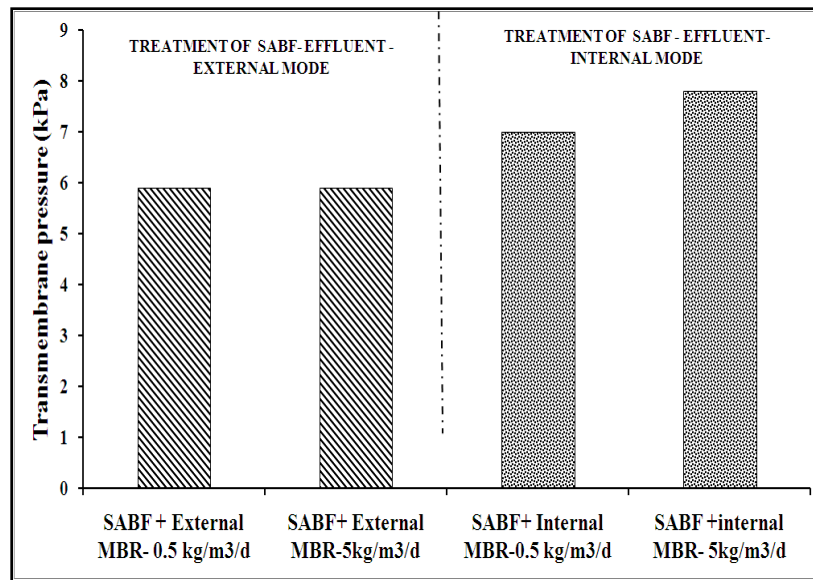
MEMBRANE BIOREACTOR

TREATMENT OF EFFLUENT FROM SUBMERGED AERATED BIOLOGICAL FILTER USING MEMBRANE BIOREACTOR

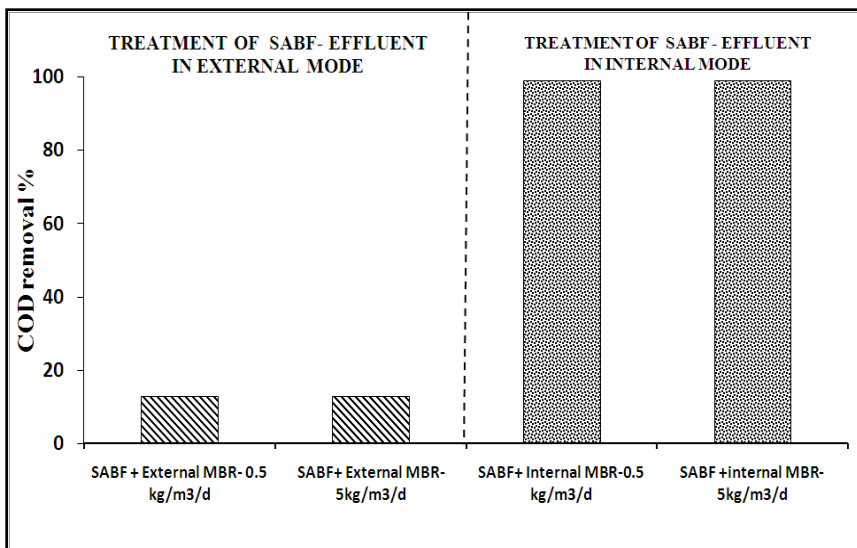
RESULTS



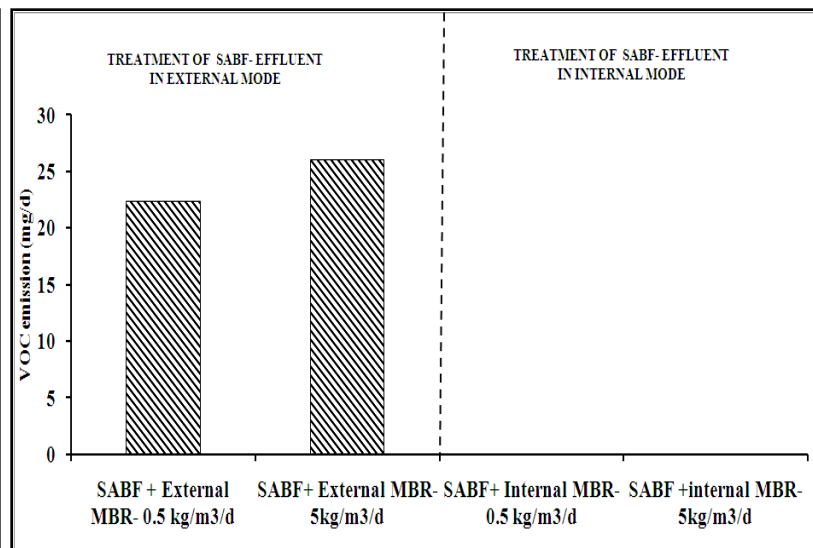
VARIATION IN FLUX



VARIATION IN TRANSMEMBRANE PRESSURE



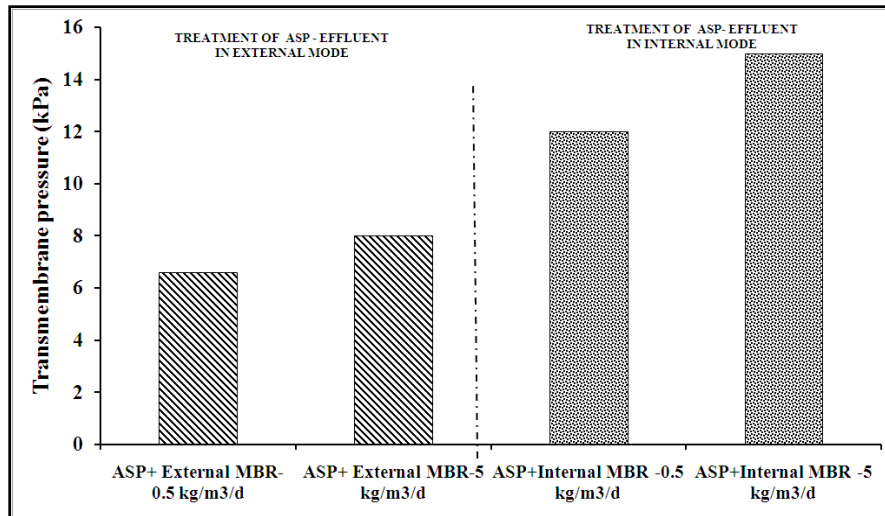
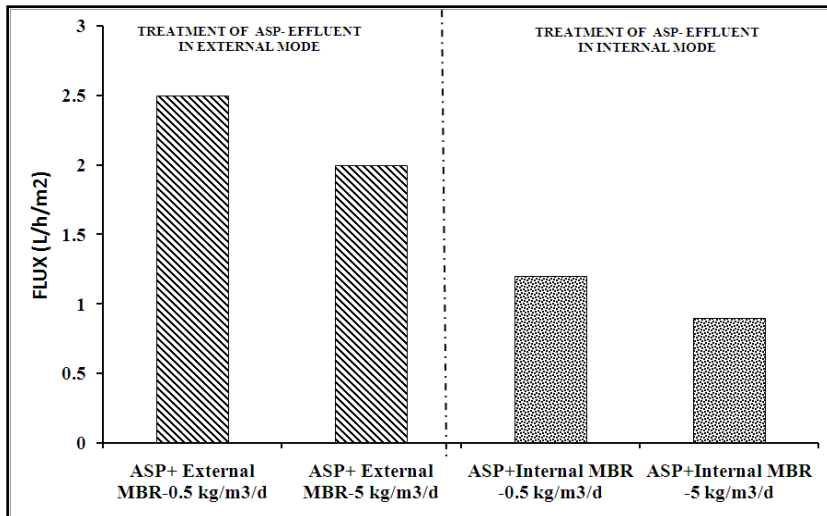
VARIATION IN COD REMOVAL



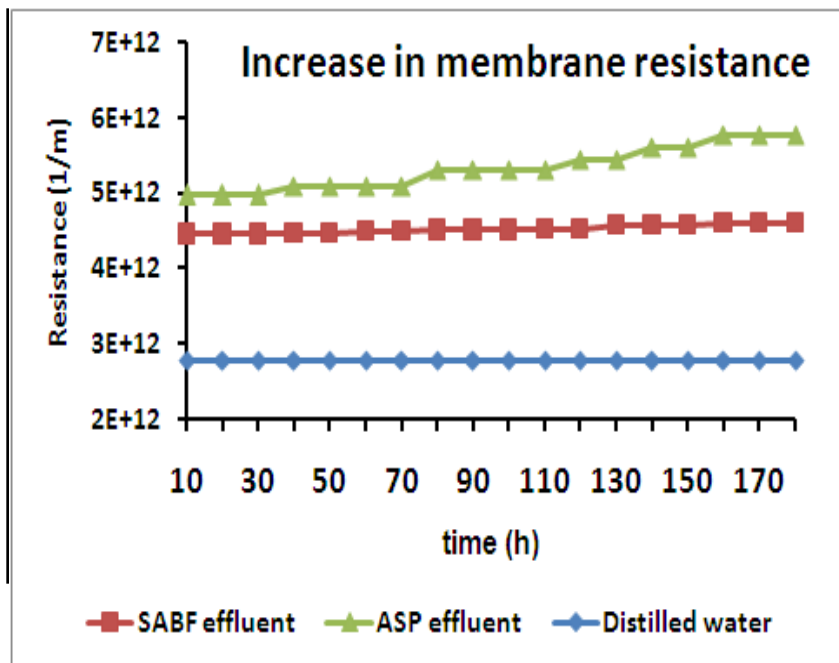
VARIATION IN VOC EMISSION

TREATMENT OF EFFLUENT FROM AERATION TANK OF ACTIVATED SLUDGE PROCESS USING MEMBRANE BIOREACTOR

RESULTS



VARIATION IN FLUX



VARIATION IN TRANSMEMBRANE PRESSURE

Performance of Hybrid treatment system			
Condition	Permeate COD (mg/L)	% COD reduction	VOC emission (mg/d)
SABF+ External MBR	622	13	25
SABF + Internal MBR	0.04	99	nil

CONCLUSIONS

- ❑ Submerged aerated biological filter were more resistant to higher organic loading rate than compared to activated sludge process.
- ❑ Limited mass transfer of VOC to the gas phase at low air flow rate reduced VOC emission from submerged aerated biological filter
- ❑ Optimization of operating conditions such as air flow rate, hydraulic retention time and organic loading rate reduced the VOC emissions from submerged aerated biological filter
- ❑ Effluent from SABF were effectively treated using membrane bioreactor.
- ❑ Complete removal of VOC from SABF effluent was achieved while adopting internal MBR configuration.
- ❑ Flux reduction and TMP rise were more significant during the treatment of ASP effluent
- ❑ SABF can be coupled along with the MBR operated under internal configuration for the complete removal of VOC from the pharmaceutical wastewater.

**BIOREMEDIATION OF Cr(VI)
CONTAMINATED SOIL AND GROUND
WATER SYSTEMS**

MOTIVATION

**TamilNadu Chromate Chemicals Limited,
Ranipet, Vellore District , Tamilnadu.**



Chromium waste

Disposal area: 5 acres (2 hectares)

2×10^5 Tones of waste)





Chromium Leachate in Ground Water

Cr(VI) Concentration in open wells/bore wells in and around TCCL (Prepared by IIT Madras)

● BDL mg/l

TCCL

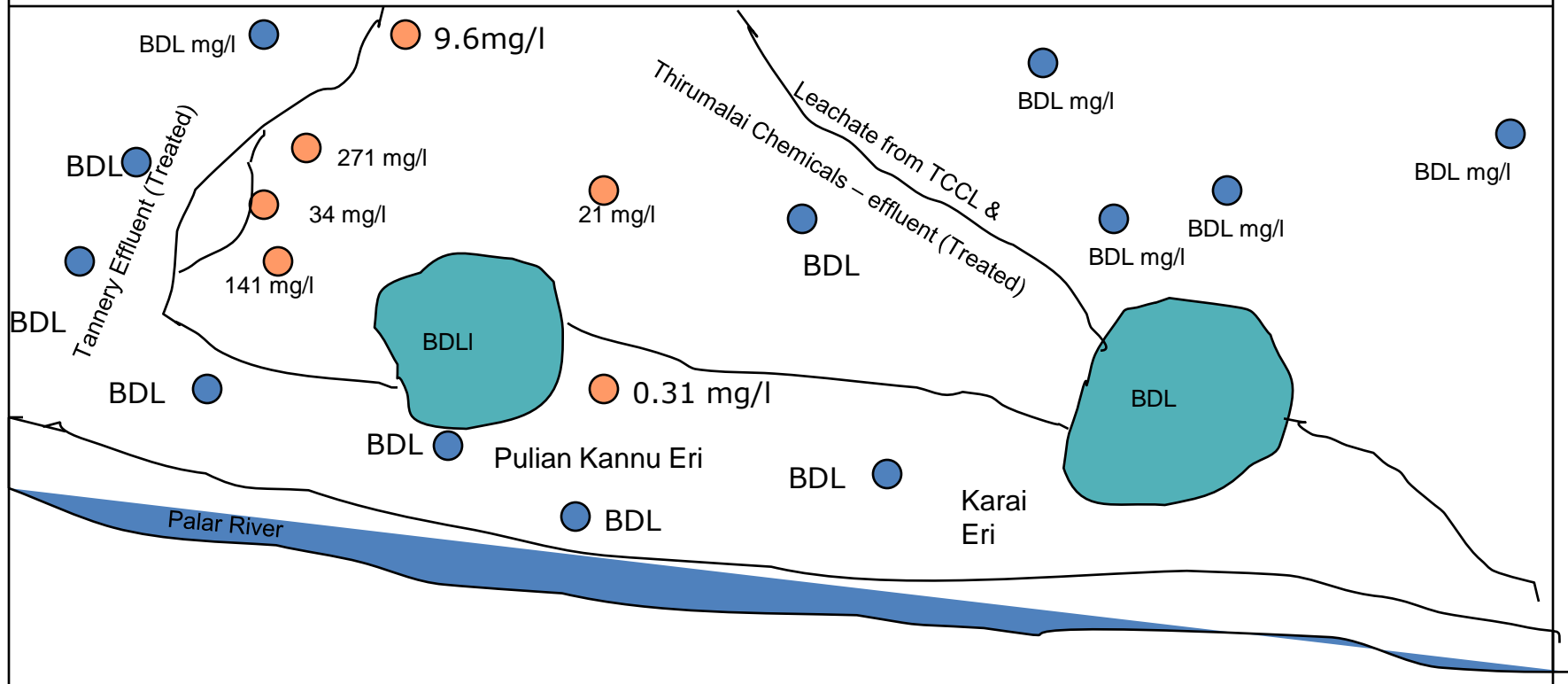
178 mg/l

SIPCOT Service Road

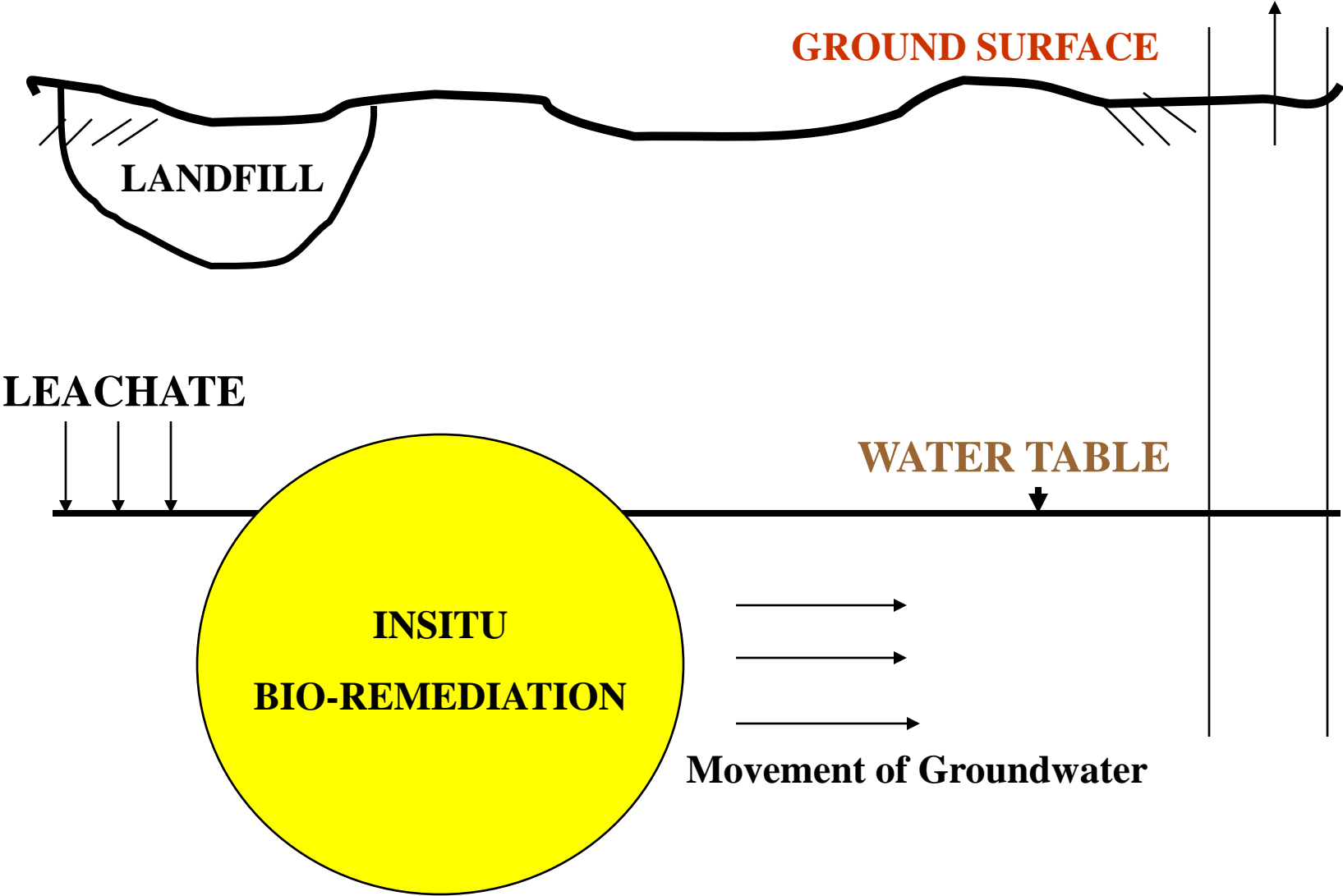
To Banglore

NH - 4

To Ranipet/Chennai

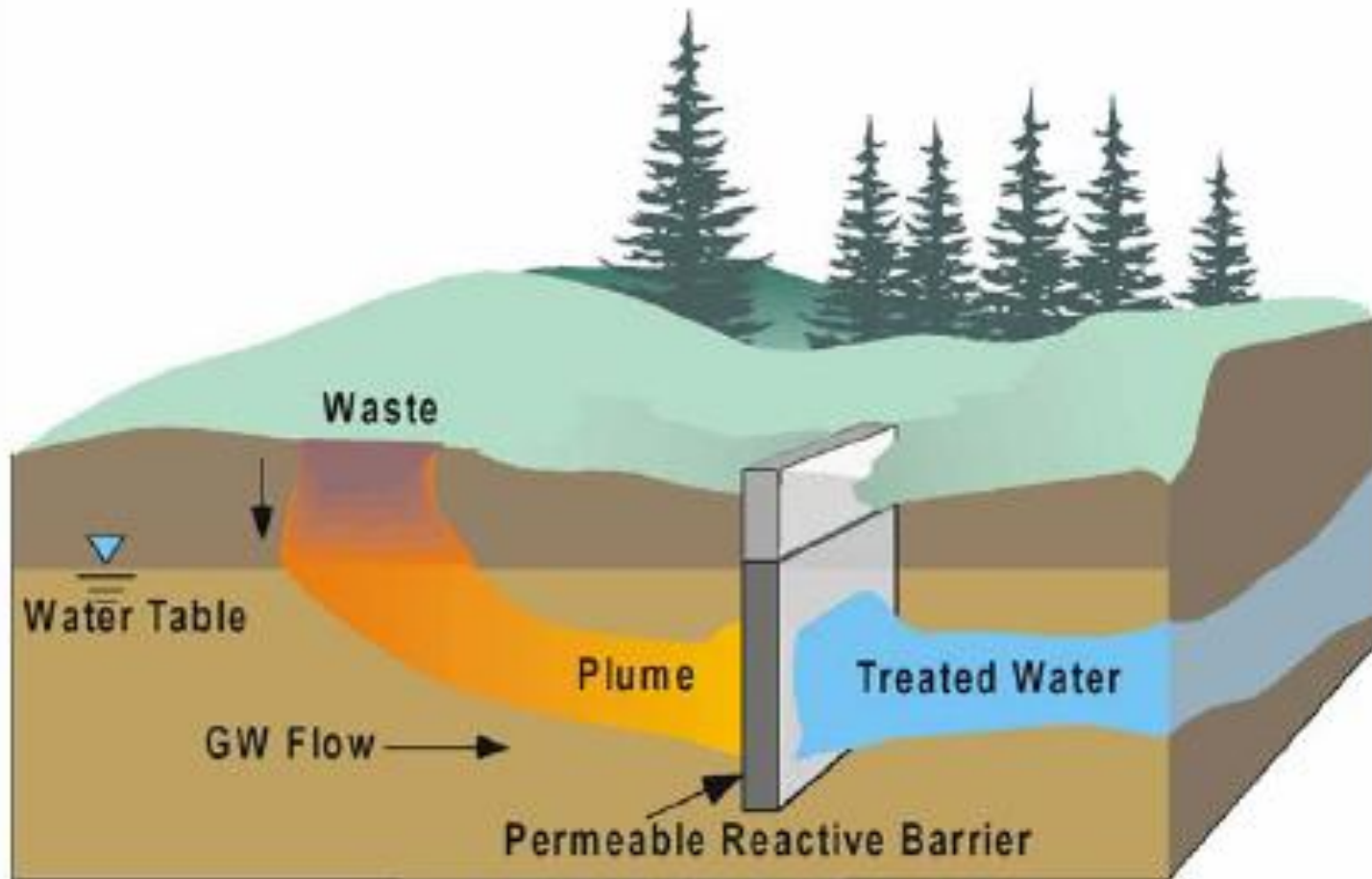


CLEANUP METHODS FOR FIELD CONDITIONS



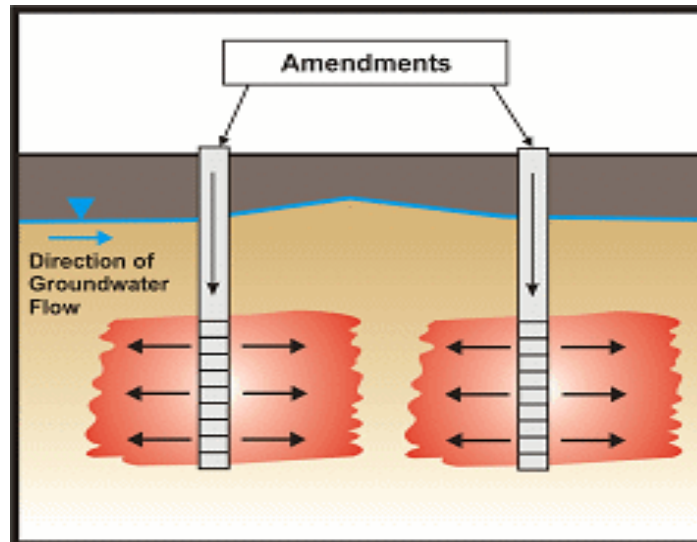
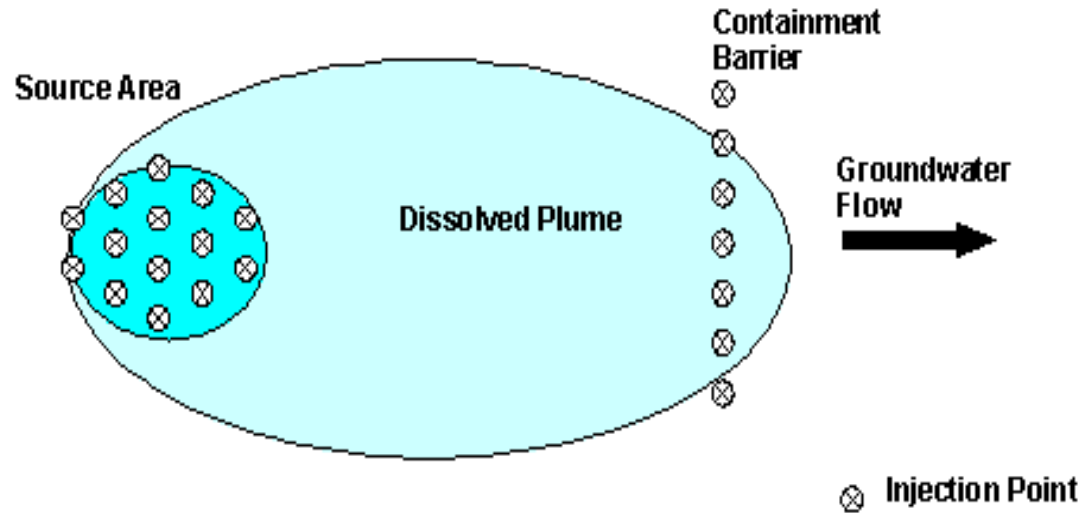
Methods for Remediation of Cr(VI) Contaminated Aquifers

- Pump and Treat systems
- Geochemical fixation
- Permeable Reactive Barriers
- Reactive Zones
- Natural attenuation
- Phyto-remediation



Schematic Representation of a Permeable Reactive Bio-barrier

REACTIVE ZONES



2

BATCH STUDIES

- **Bio-kinetic parameters**
- **Adsorption Parameters**

Cr(VI) Reduction Studies with CRB, SRB and IRB, in Combinations

1. CRB –Aerobic
2. CRB -Anaerobic
2. SRB-Anaerobic
3. IRB- anaerobic
4. CRB+SRB
5. CRB+IRB
6. CRB+SRB+IRB

Adsorption Studies

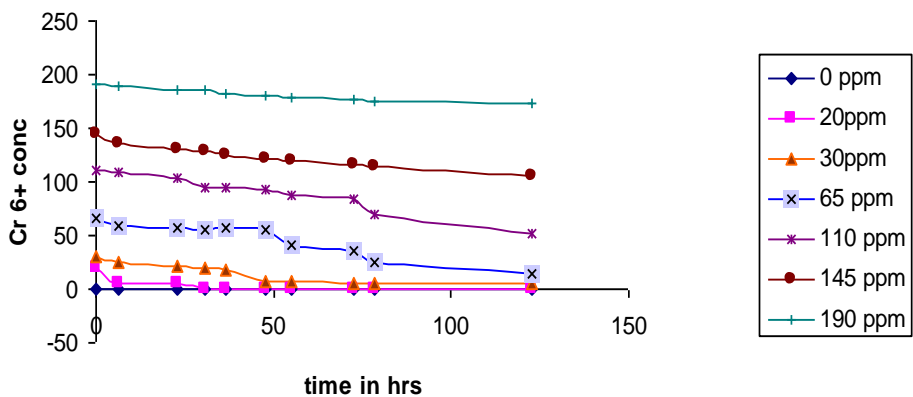
Adsorbents– Soil , Sand

Adsorbates:

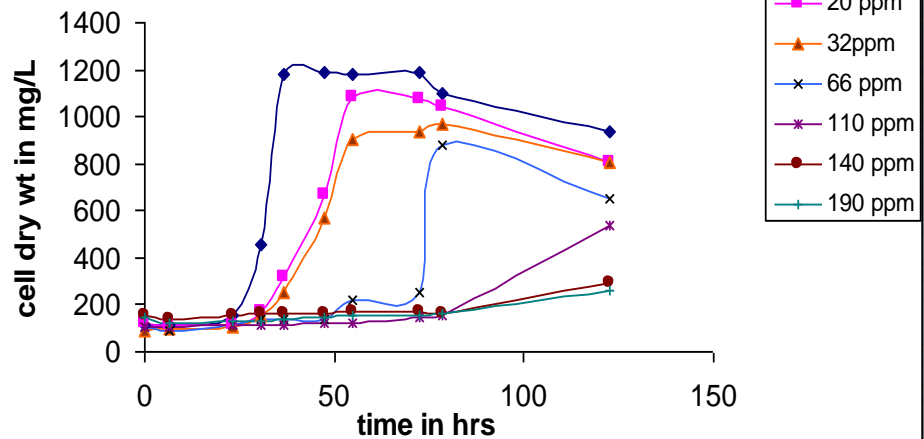
1. Cr(VI)
2. Molasses/Sugar
3. Lithium
4. Cr(III)

Cr(VI) Reduction in Aerobic Conditions

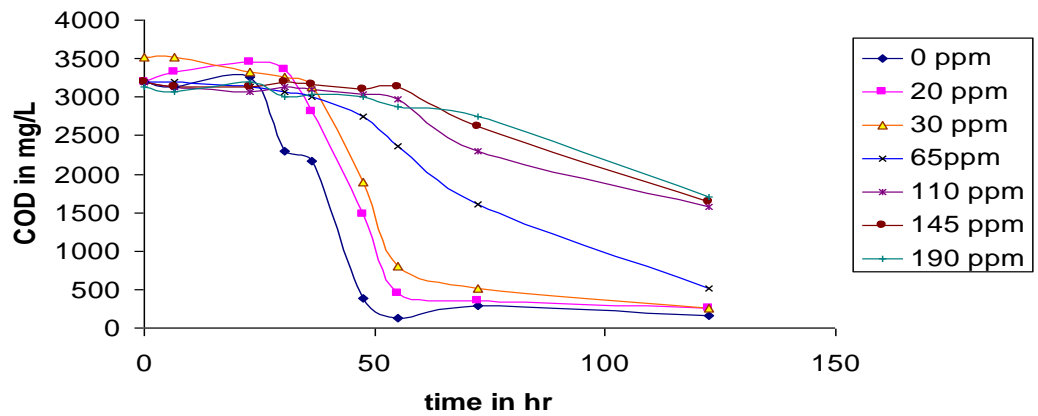
Chromium(VI) reduction



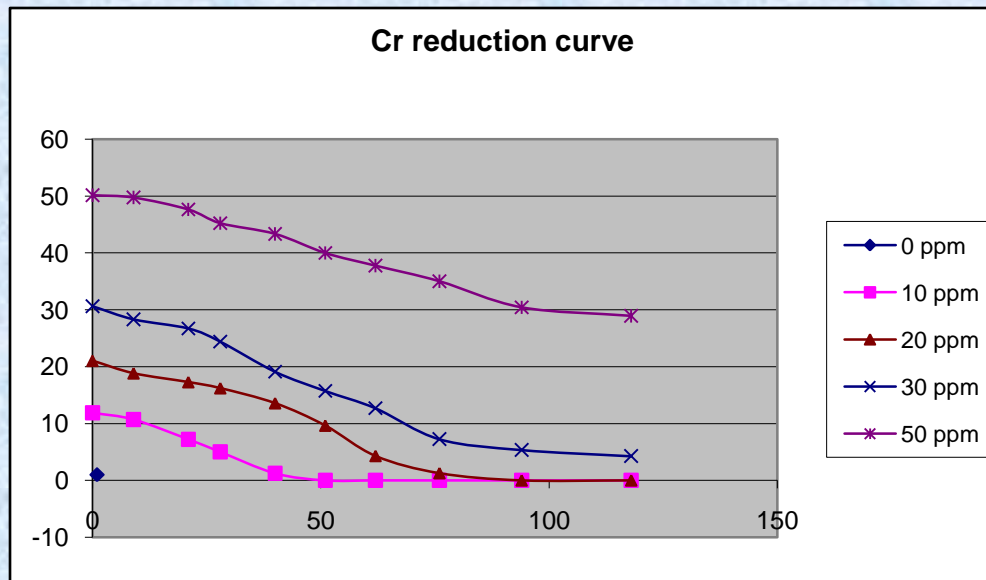
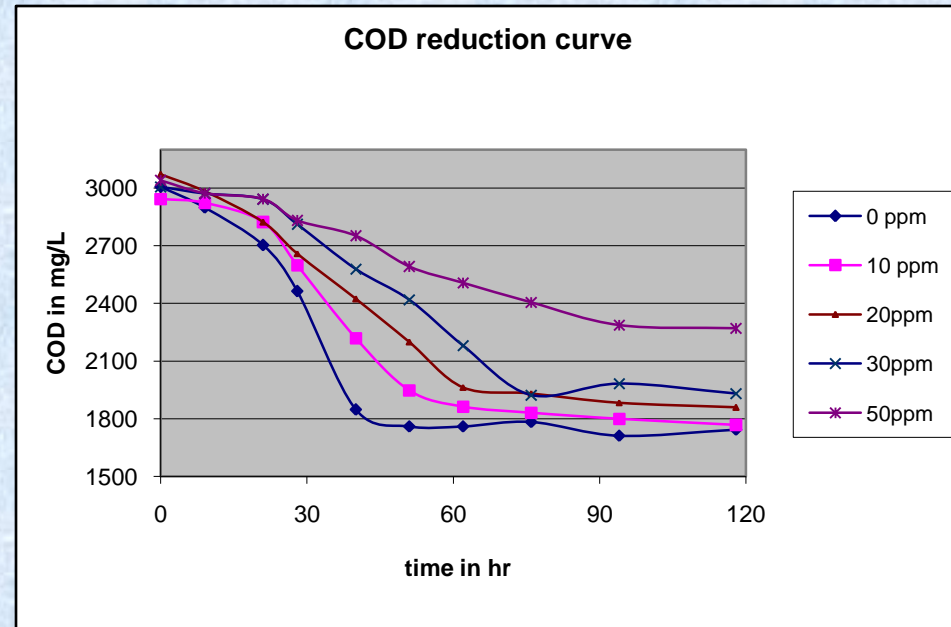
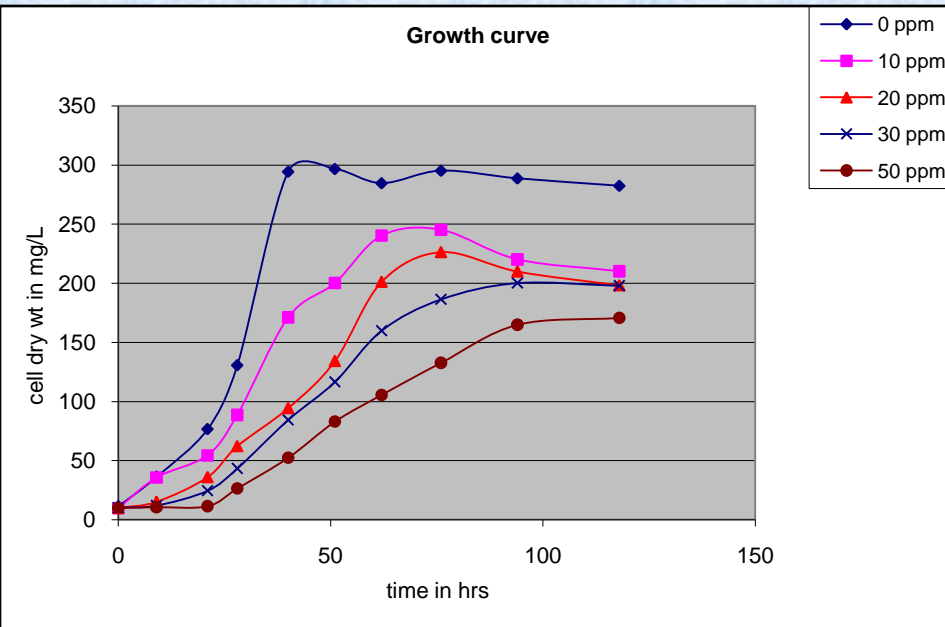
Growth curve



COD removal rate

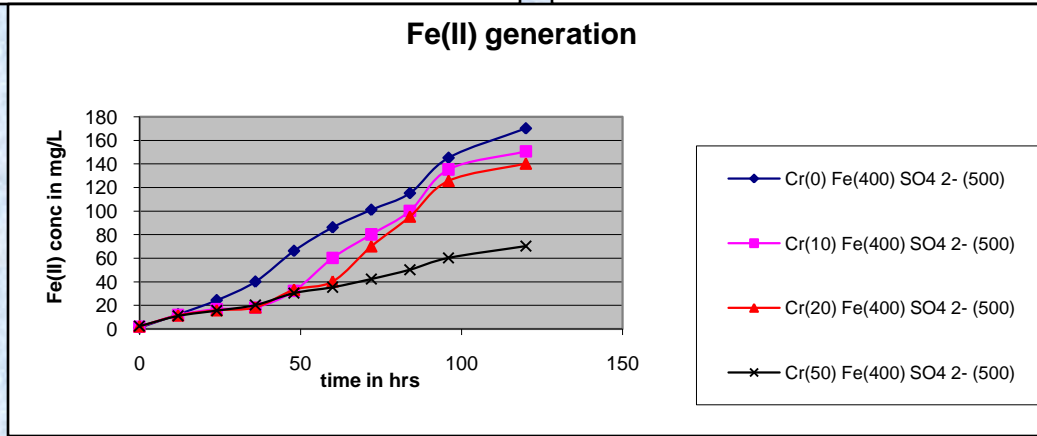
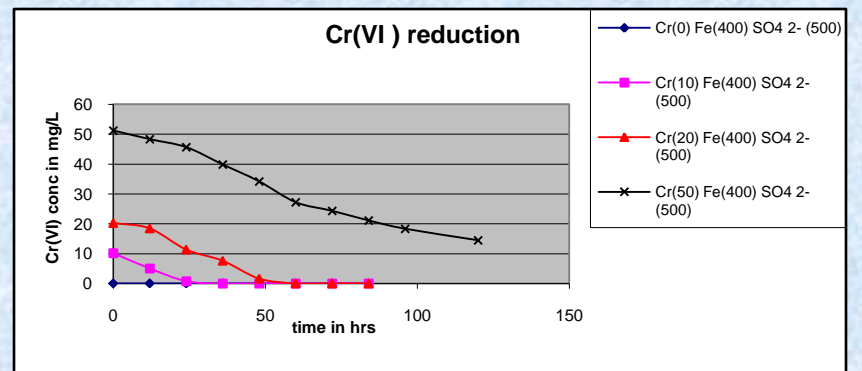
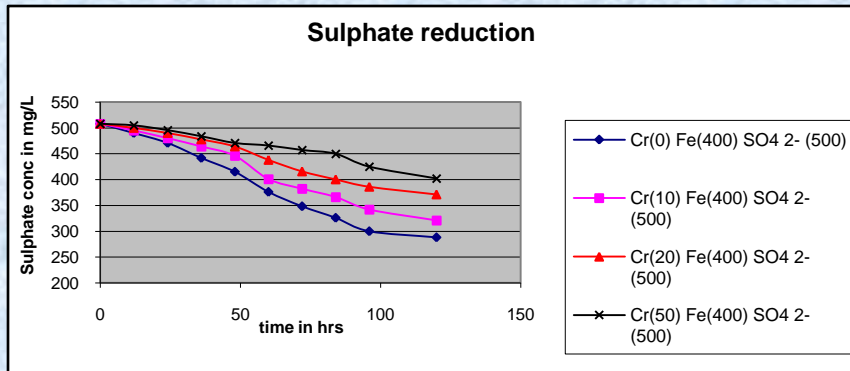
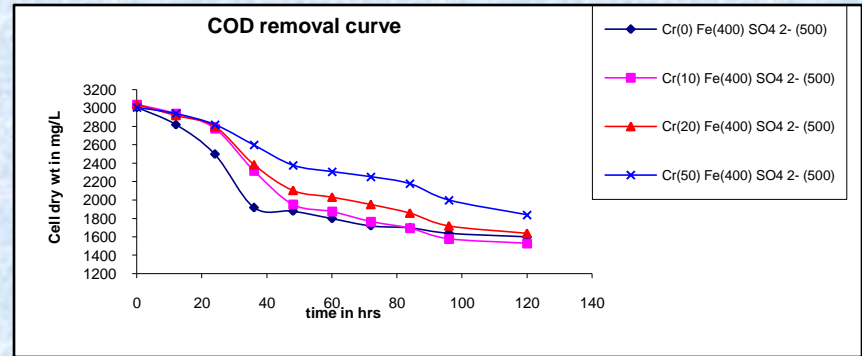
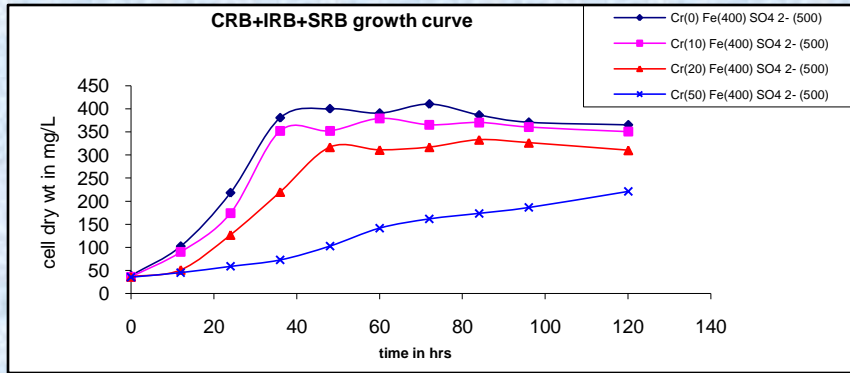


Cr (VI) Reduction by CRB under Anaerobic Condition



Growth of CRB+IRB+SRB under Anaerobic Condition

Fe(400ppm, Sulphate(500ppm))



Model

Suffix 1,2,3 represents CRB,SRB,IRB respectively

$$M = \sum_{i=1}^3 M_i$$

$$S = \sum_{i=1}^3 S_i$$

$$Cr_6 = \sum_{i=1}^3 Cr_{6i}$$

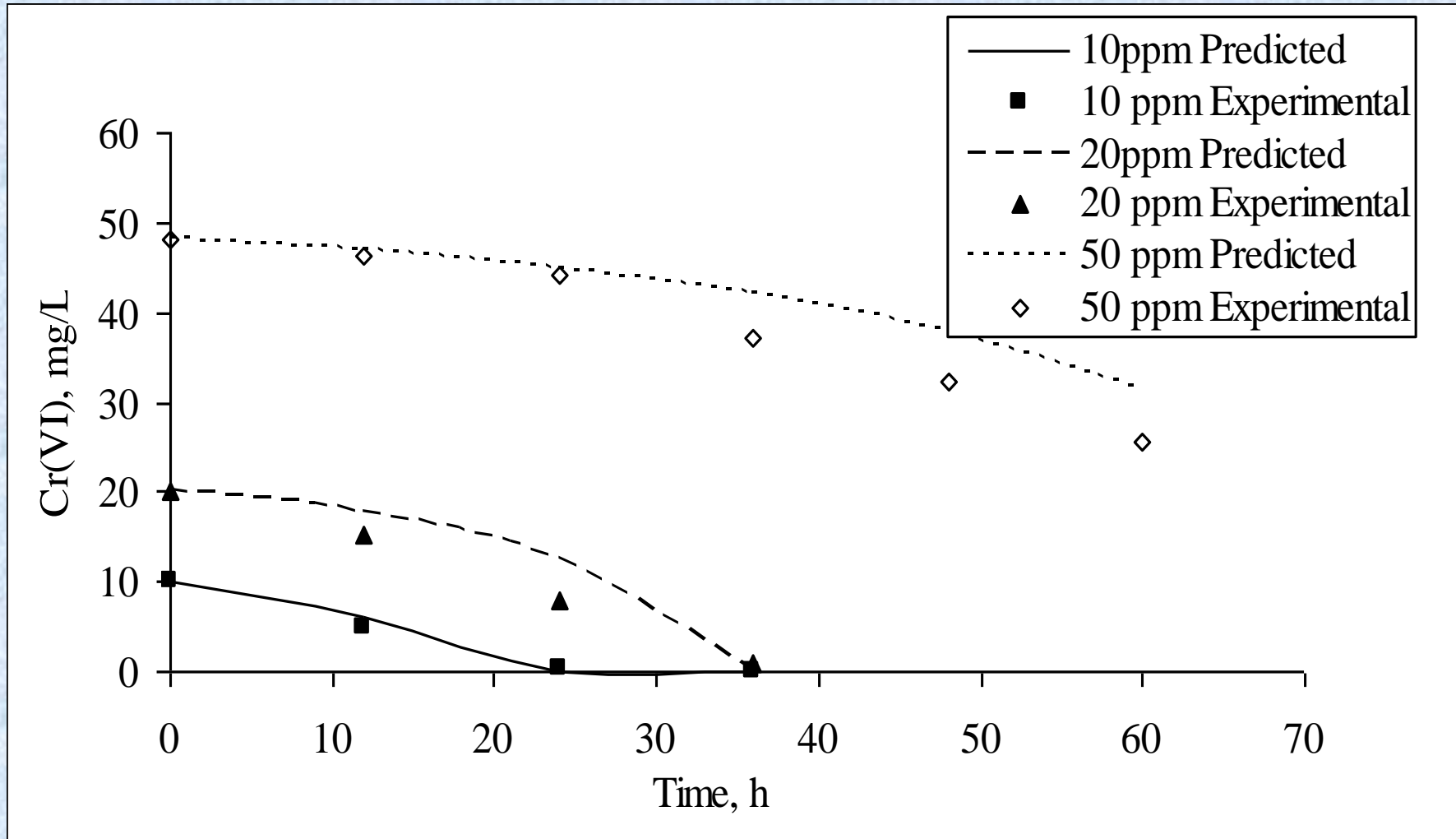
$$S_i = S \left(\frac{M_i}{M} \right)$$

$$Cr_{6,i} = Cr_6 \left(\frac{M_i}{M} \right)$$

$$\frac{dM_{CRB}}{dt} = \frac{M_{CRB} \cdot \mu_{\max, CRB} \cdot S\left(\frac{M_i}{M}\right)}{K_{s, CRB} + S\left(\frac{M_i}{M}\right)} \left(\frac{K_{i, CRB}}{K_{i, CRB} + Cr_6\left(\frac{M_i}{M}\right)} \right)$$

$$\frac{dM_{SRB}}{dt} = \frac{M_{SRB} \cdot \mu_{\max, SRB} \cdot S\left(\frac{M_i}{M}\right)}{K_{s, SRB} + S\left(\frac{M_i}{M}\right)} \left(\frac{K_{i, SRB}}{K_{i, SRB} + Cr_6\left(\frac{M_i}{M}\right)} \right)$$

$$\frac{dM_{IRB}}{dt} = \frac{M_{IRB} \cdot \mu_{\max, IRB} \cdot S\left(\frac{M_i}{M}\right)}{K_{s, IRB} + S\left(\frac{M_i}{M}\right)} \left(\frac{K_{i, IRB}}{K_{i, IRB} + Cr_6\left(\frac{M_i}{M}\right)} \right)$$

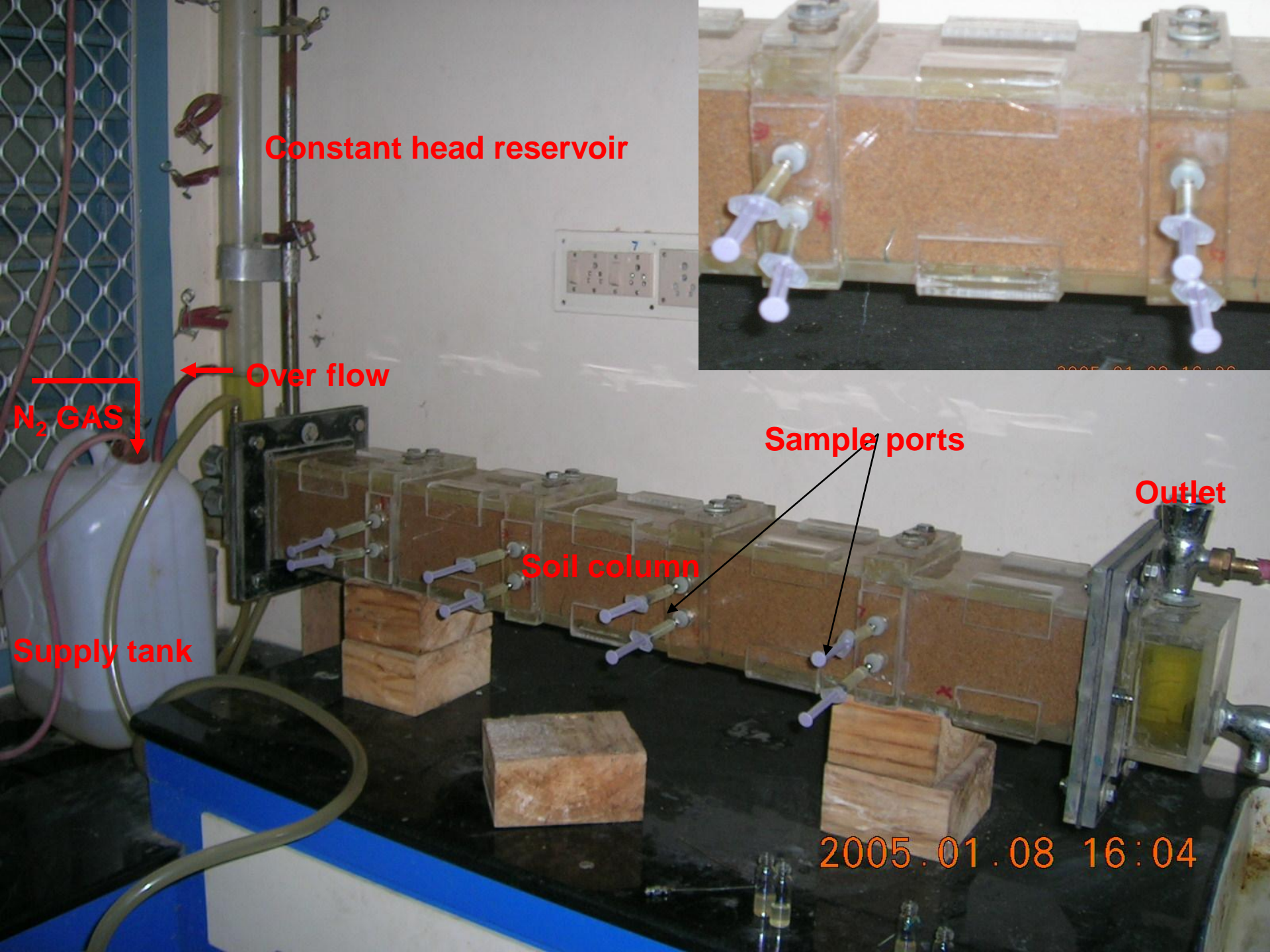


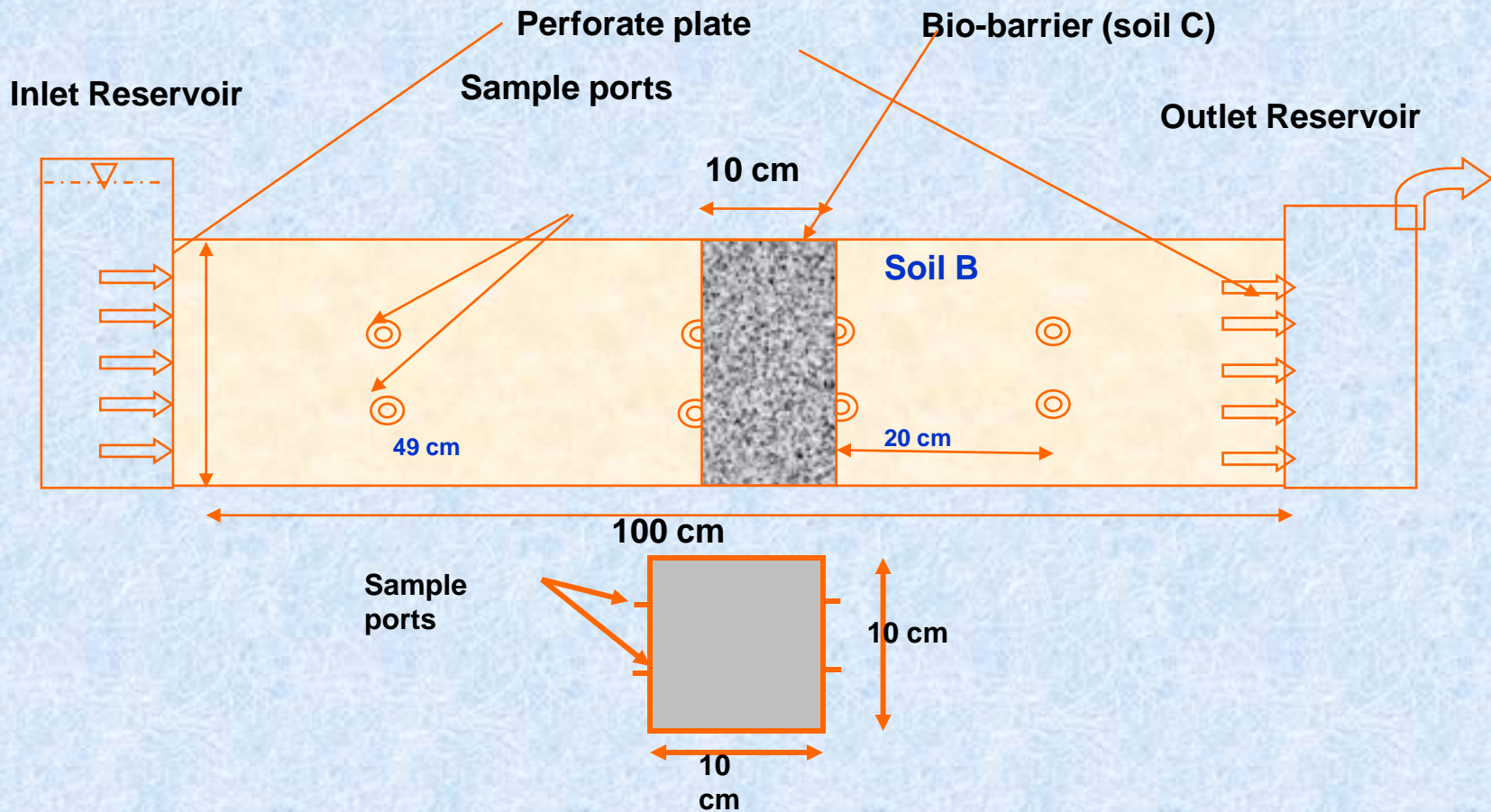
Cr(VI) reduction by CRB, SRB and IRB under anaerobic conditions for different initial Cr(VI) concentrations

Somasundaram et al., Jl. of Hazard. Mater., 2009

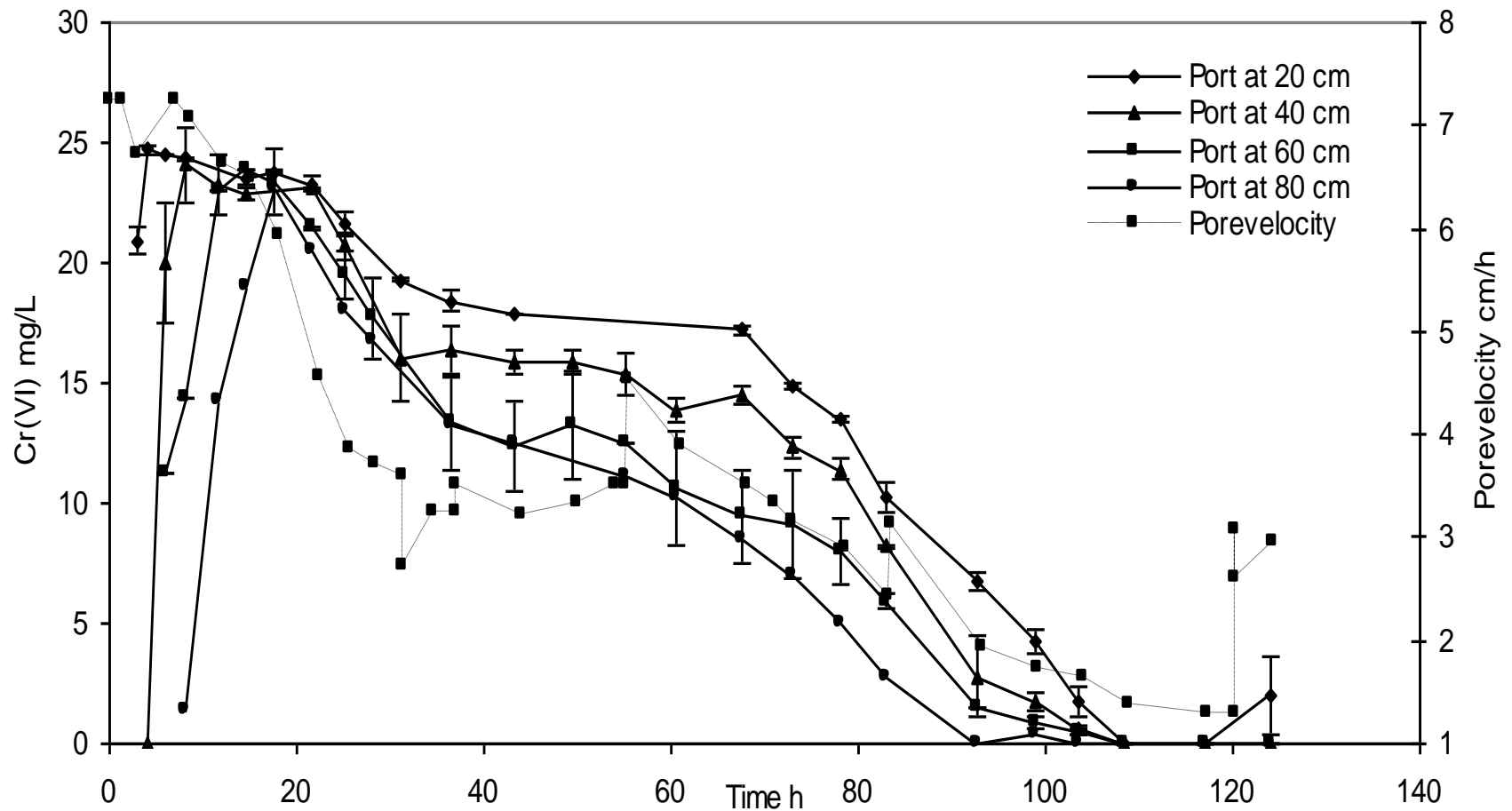
3

BENCH SCALE STUDIES



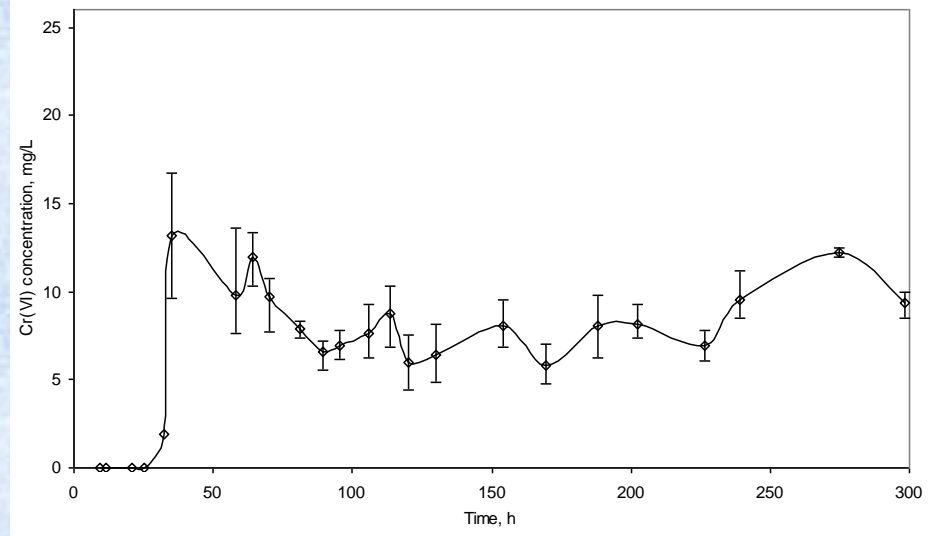
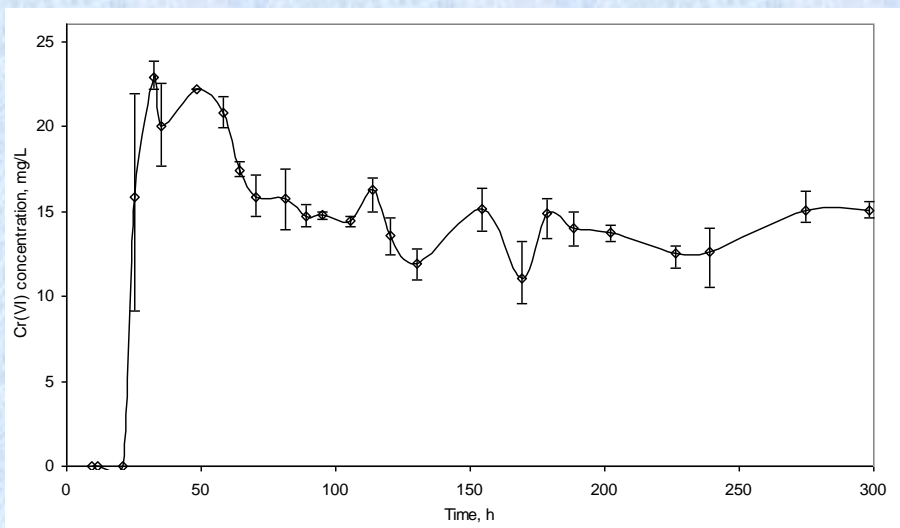


Schematic of experimental setup

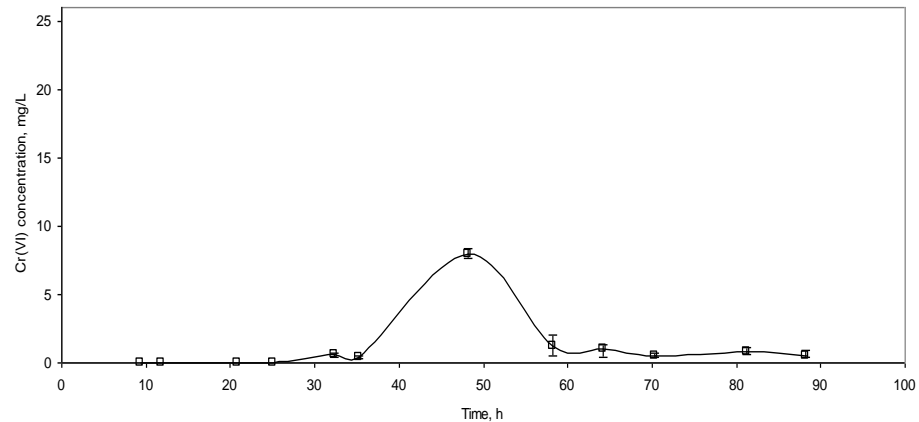
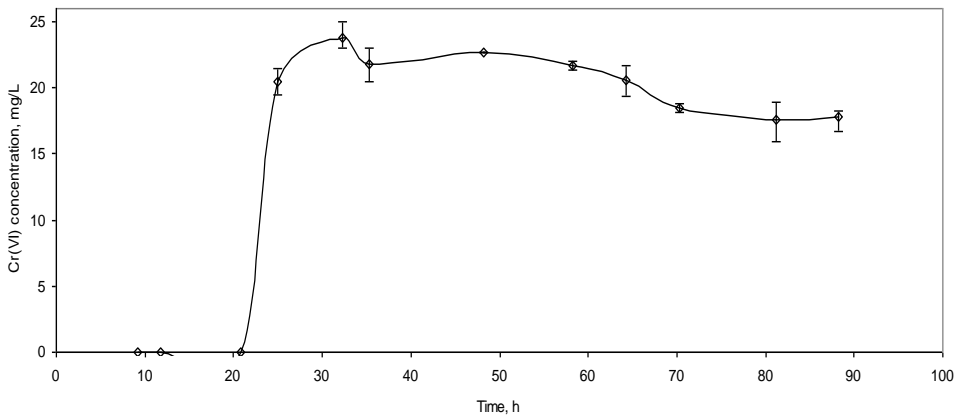


Cr(VI) break-through curve with biotransformation, Soil A

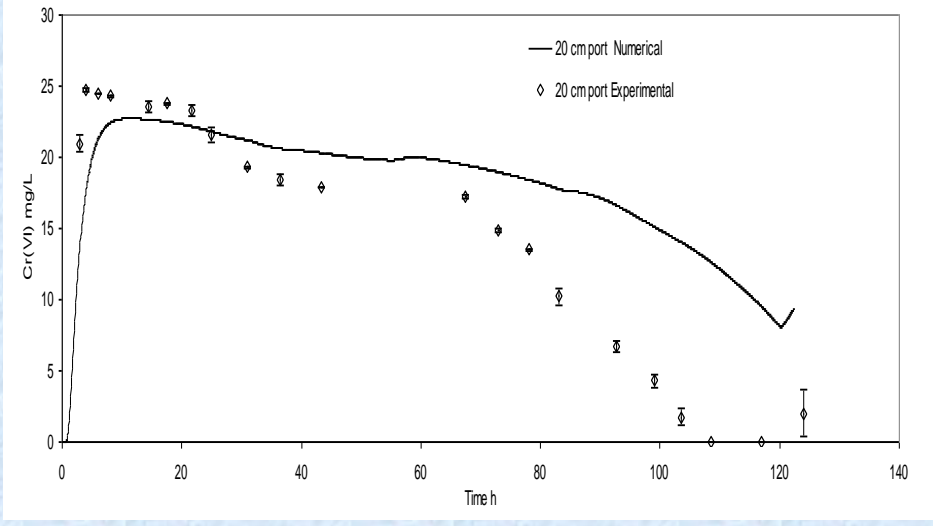
Shashidhar et al., JI. of Hazard. Mater., 2006



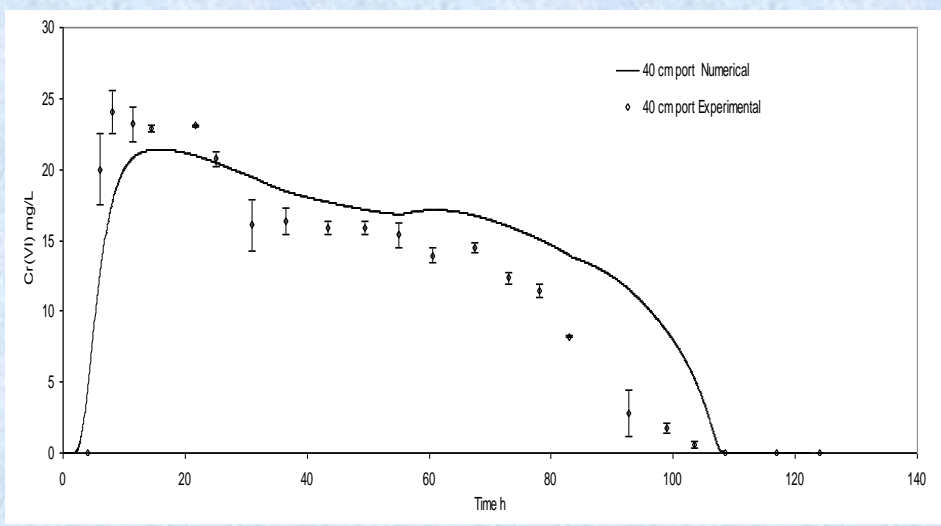
**Cr(VI) breakthrough just before and after Biobarrier BB1
(Bact conc= 0.0205 mg/g of soil)**



**Cr(VI) breakthrough just before and after Biobarrier (BB2)
(Bact conc= 0.205 mg/g of soil)**

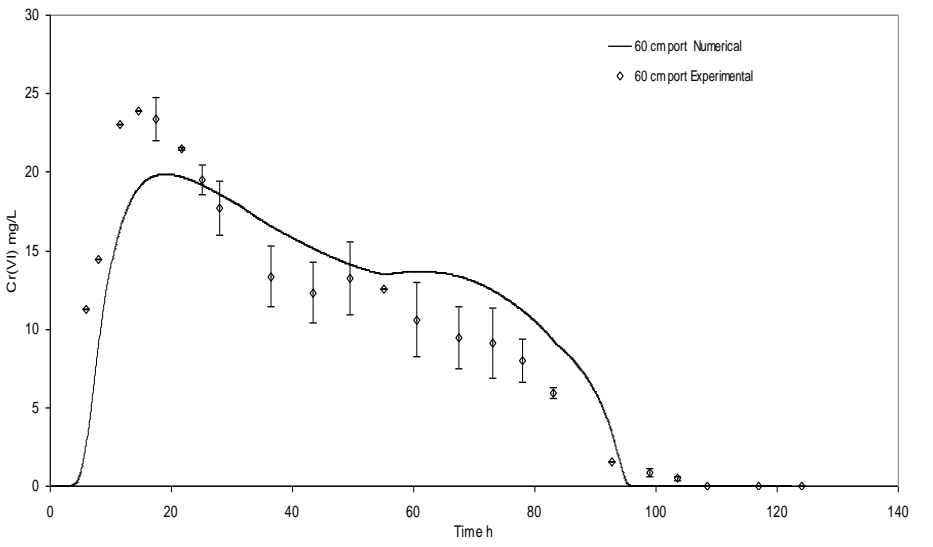


20 cm Port

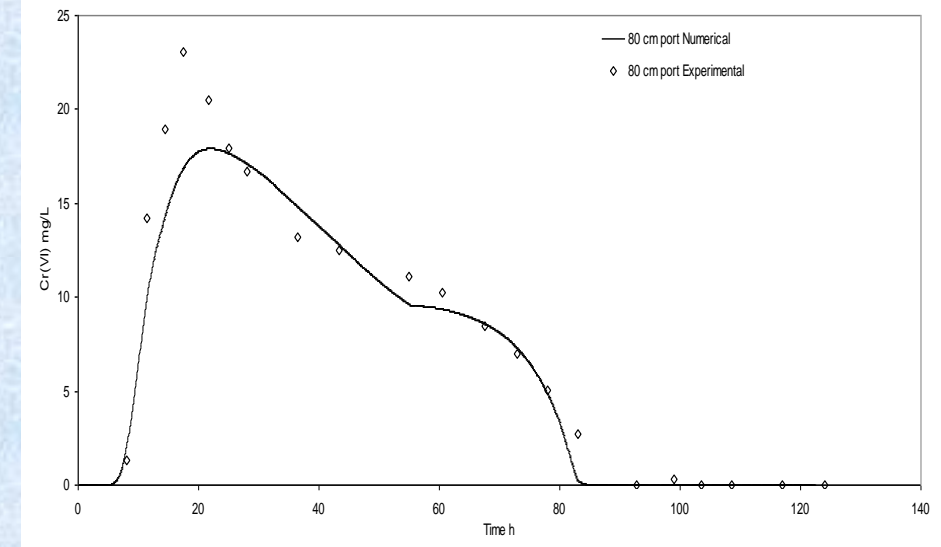


40 cm Port

Initial pore velocity 7.3 cm/h



60 cm Port



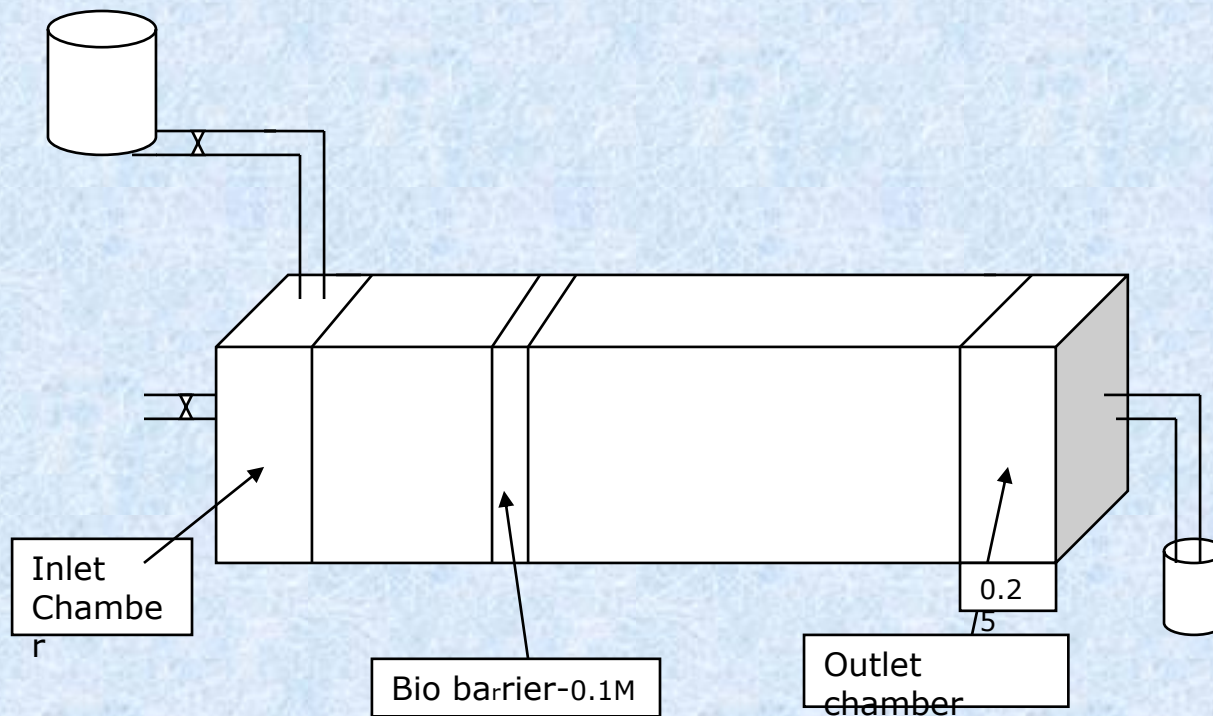
80 cm Port

4

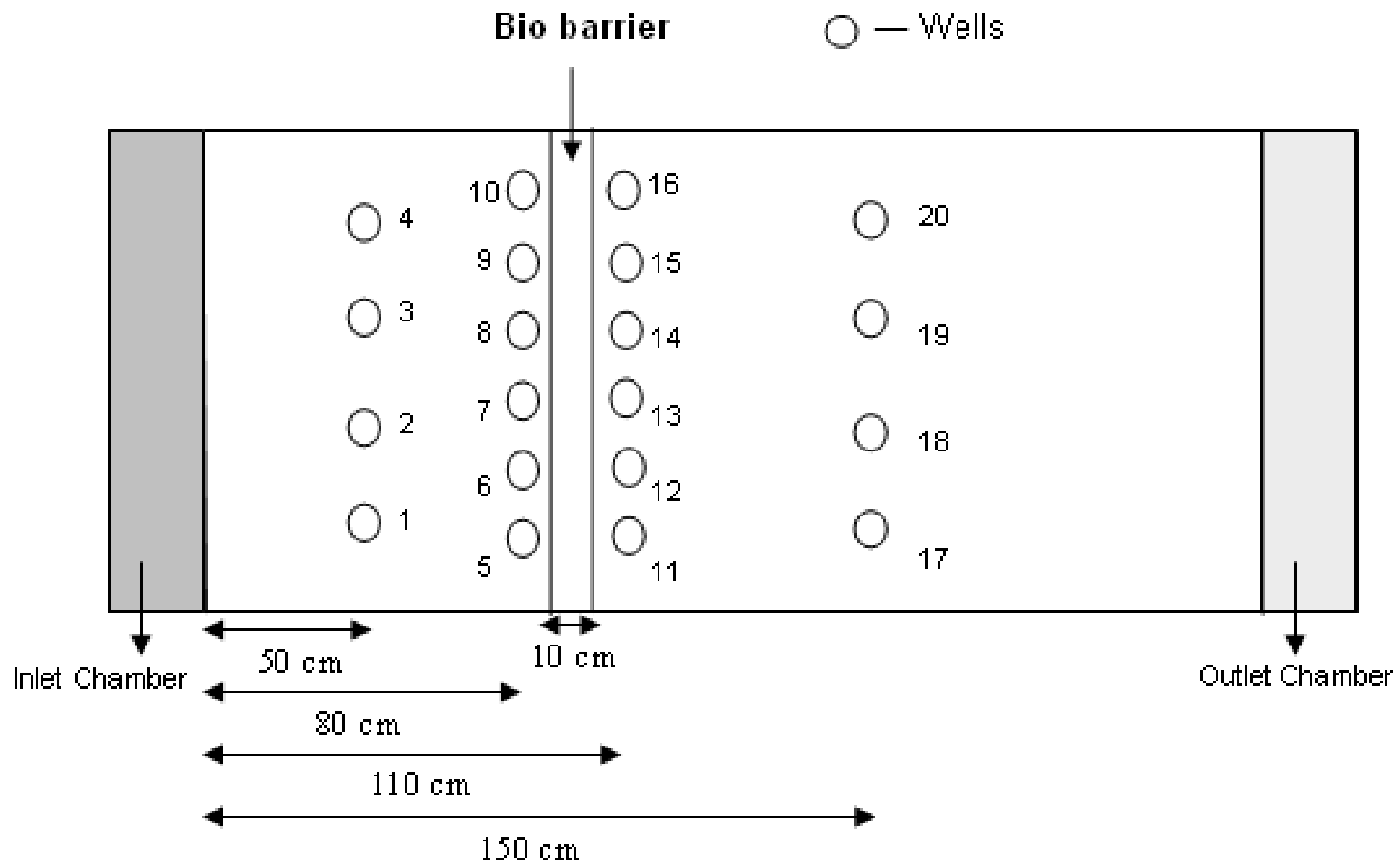
PILOT SCALE STUDIES



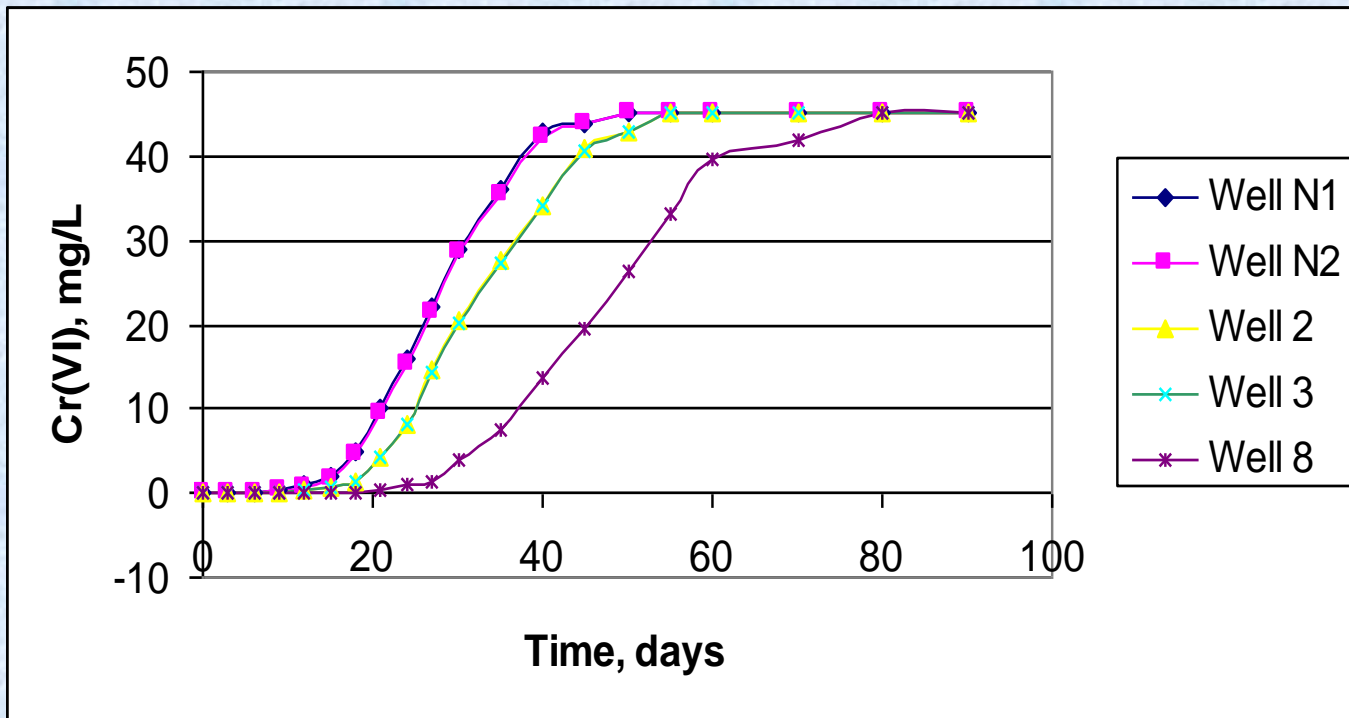
Schematic Diagram of the Reactor



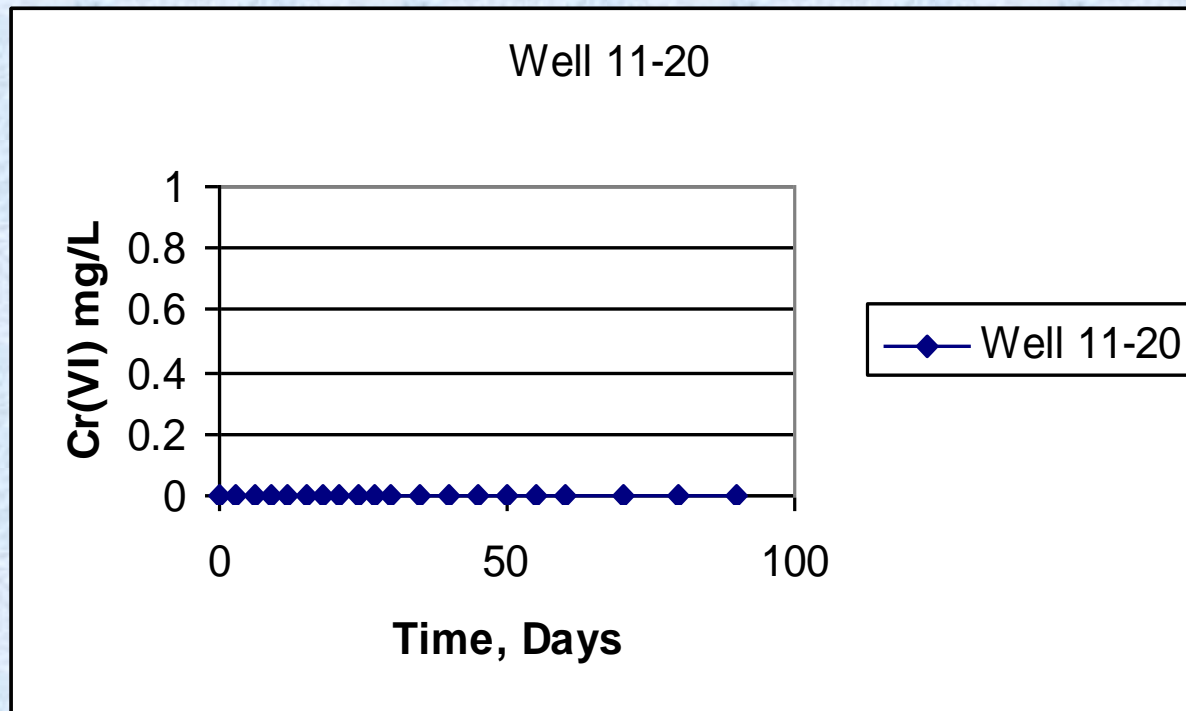
Location of Wells



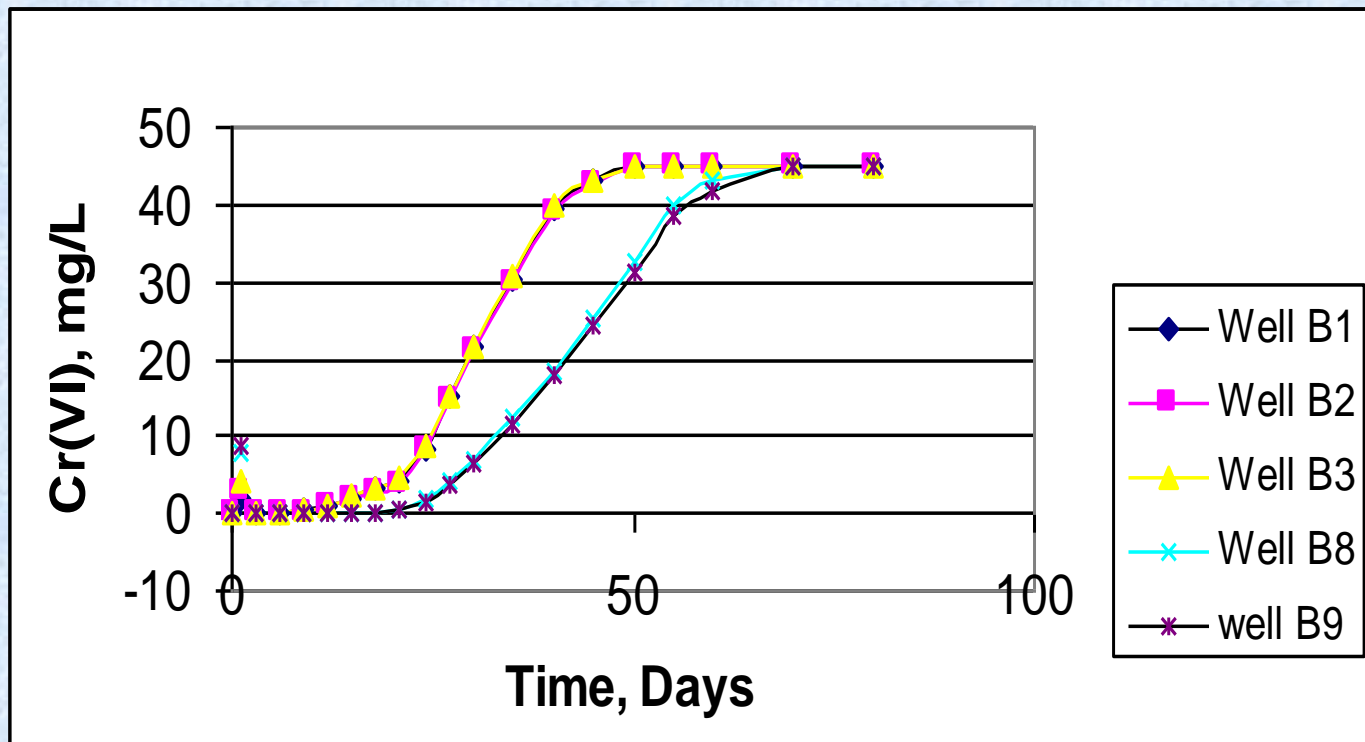
Cr (VI) Concentration before the Bio-barrier in Bioreactor



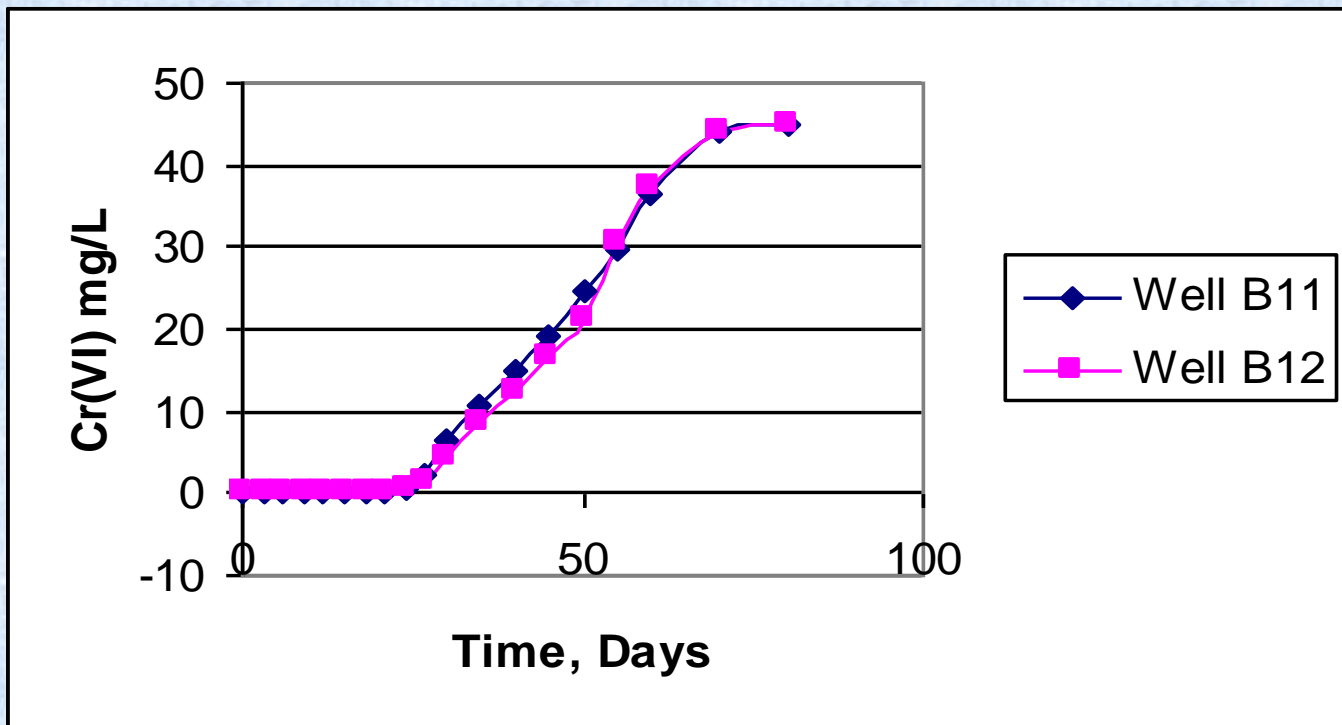
Cr (VI) Concentration after the Bio-barrier in Bioreactor



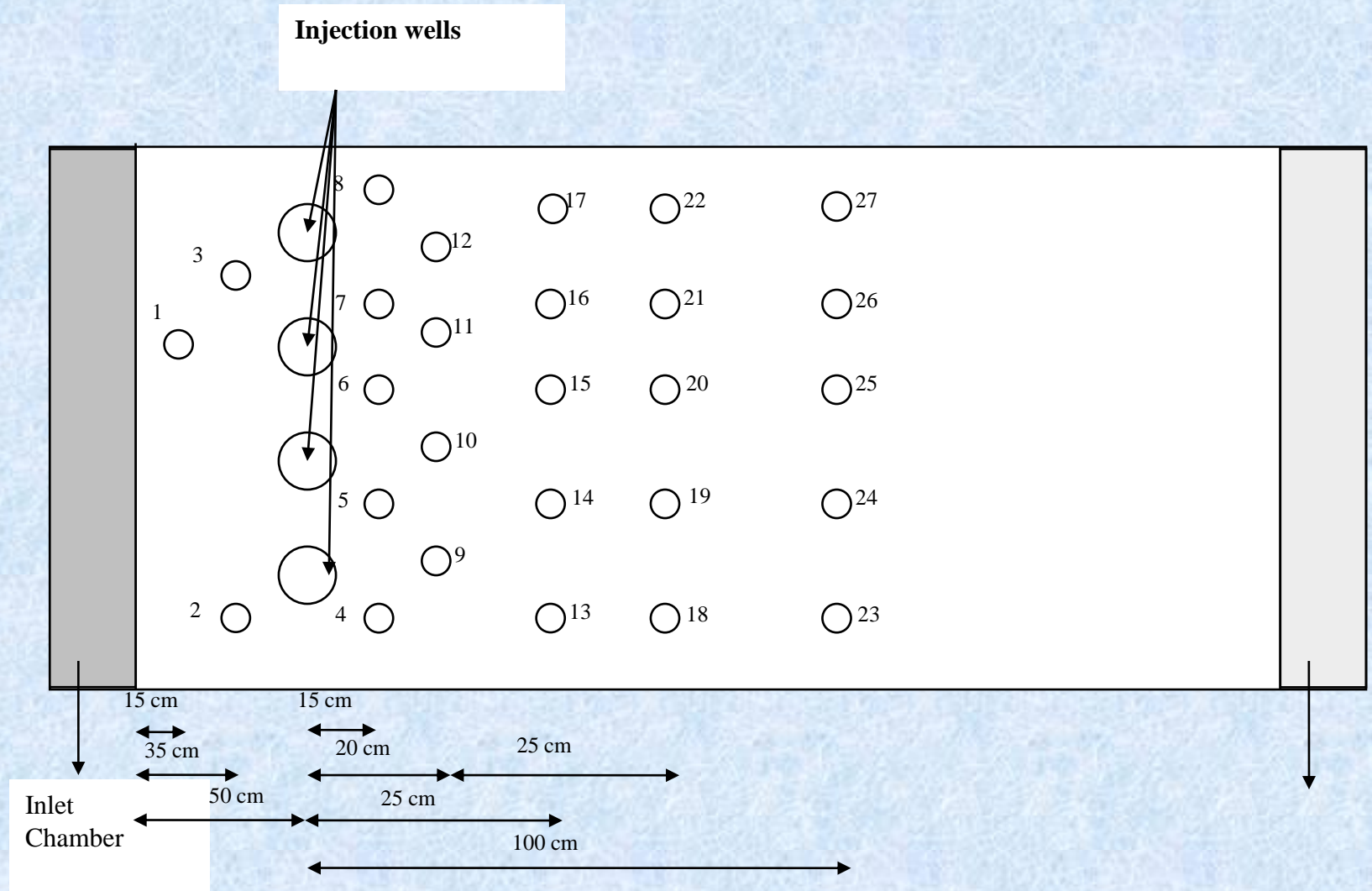
Cr (VI) Concentration in the Blank Reactor before Barrier



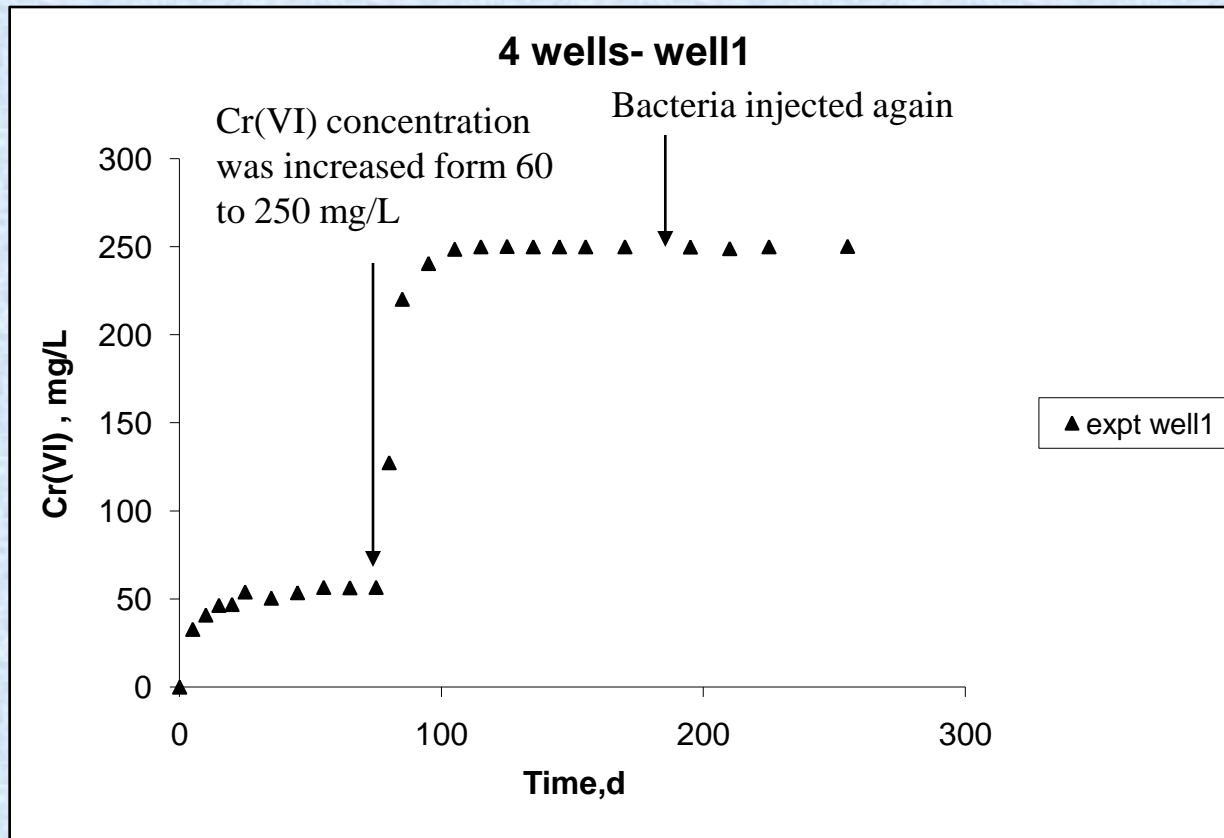
Cr (VI) Concentration in the Blank Reactor after the Barrier



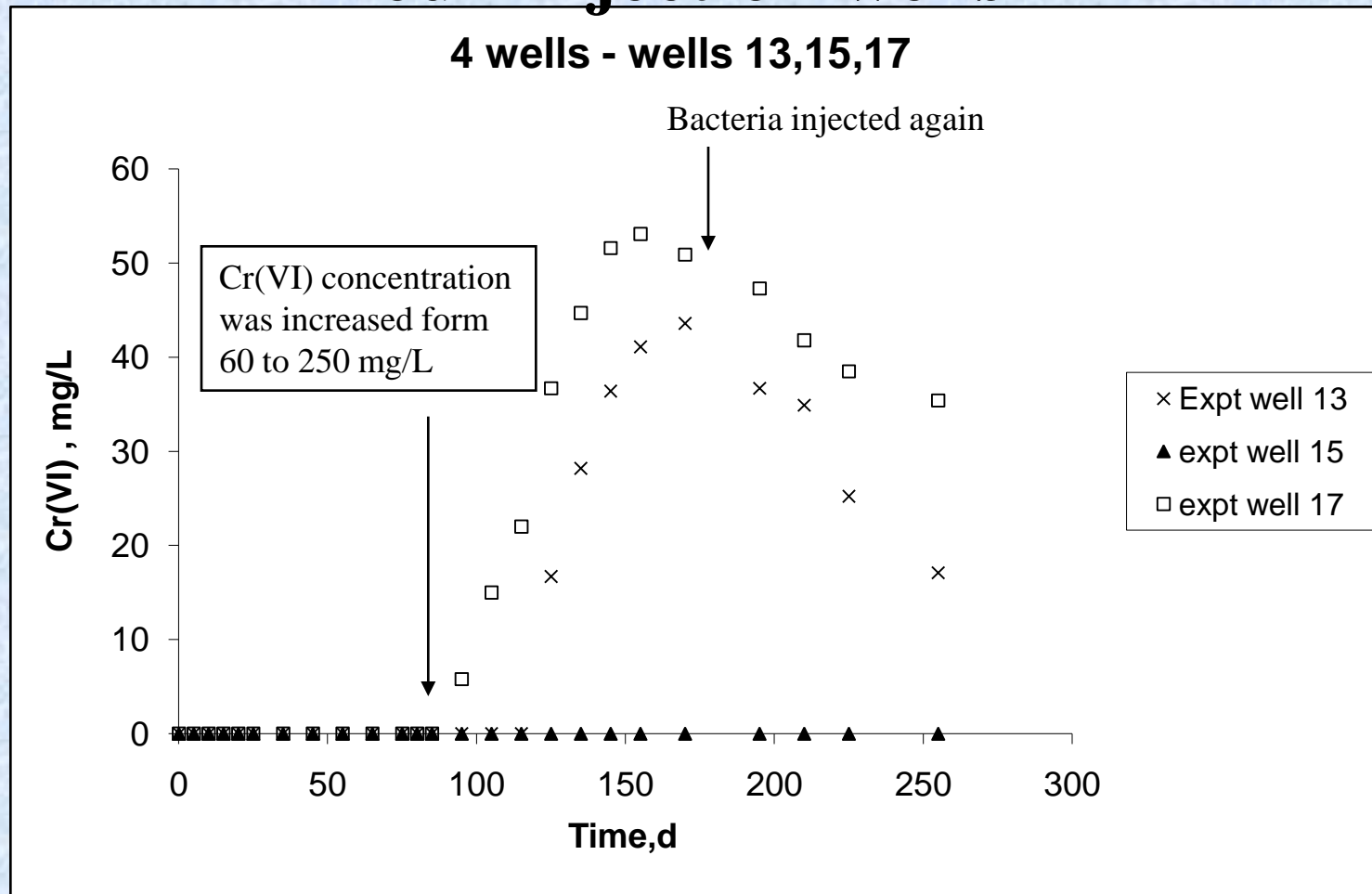
PLAN VIEW OF REACTOR CONTAINING **FOUR** INJECTION WELLS

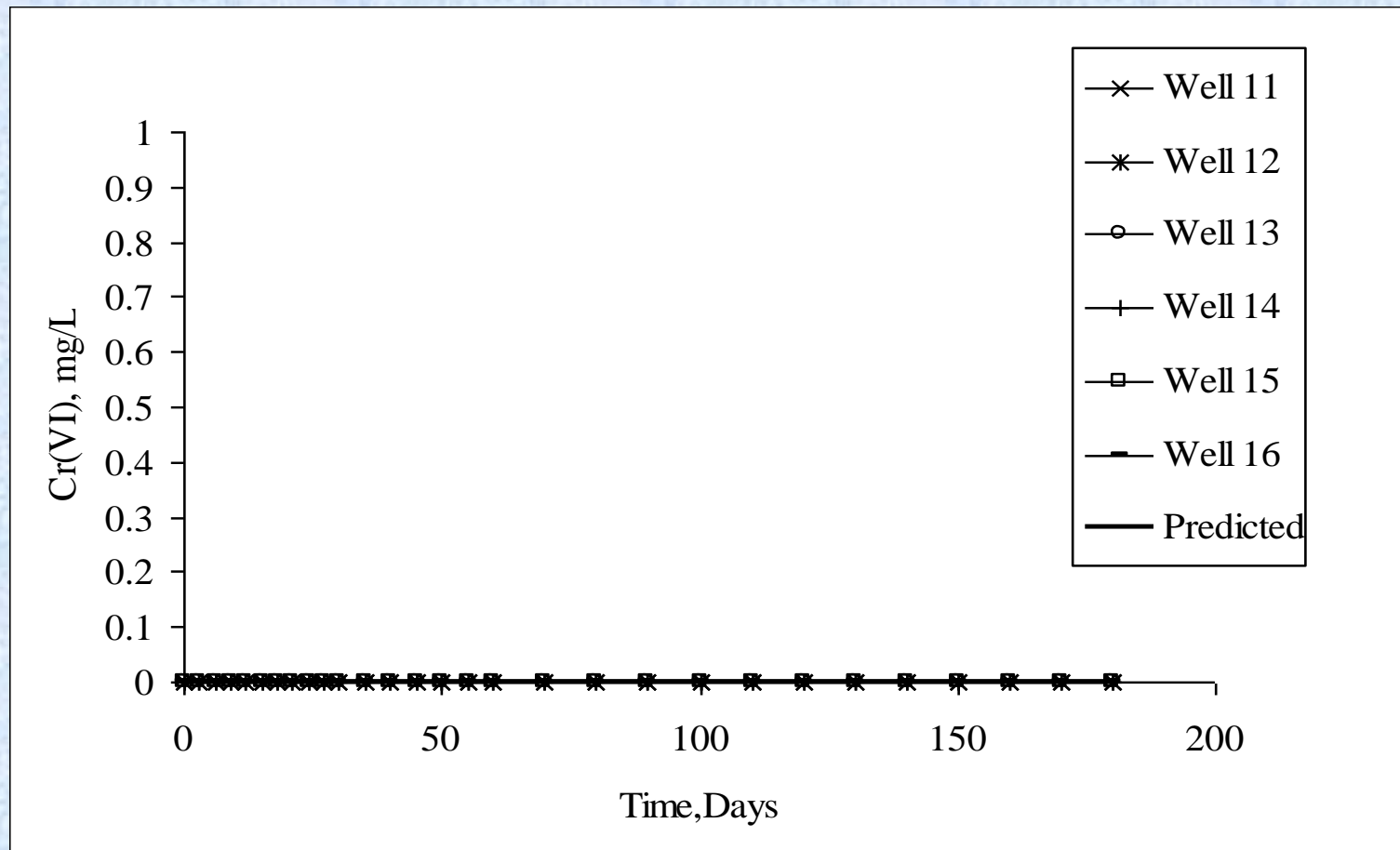


Cr (VI) Concentration in Reactor before four Injection wells



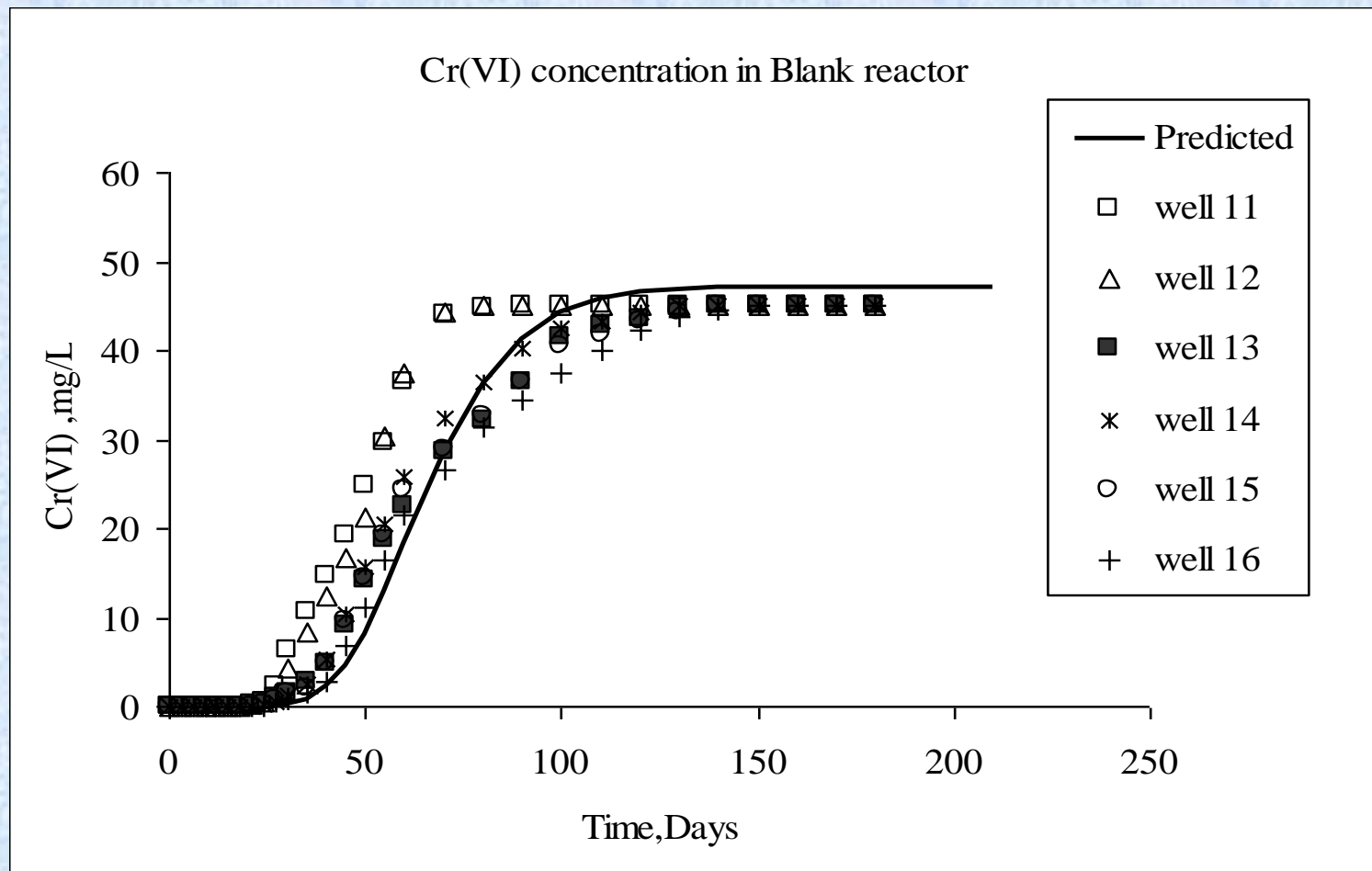
Cr (VI) Concentration in Reactor after four Injection wells





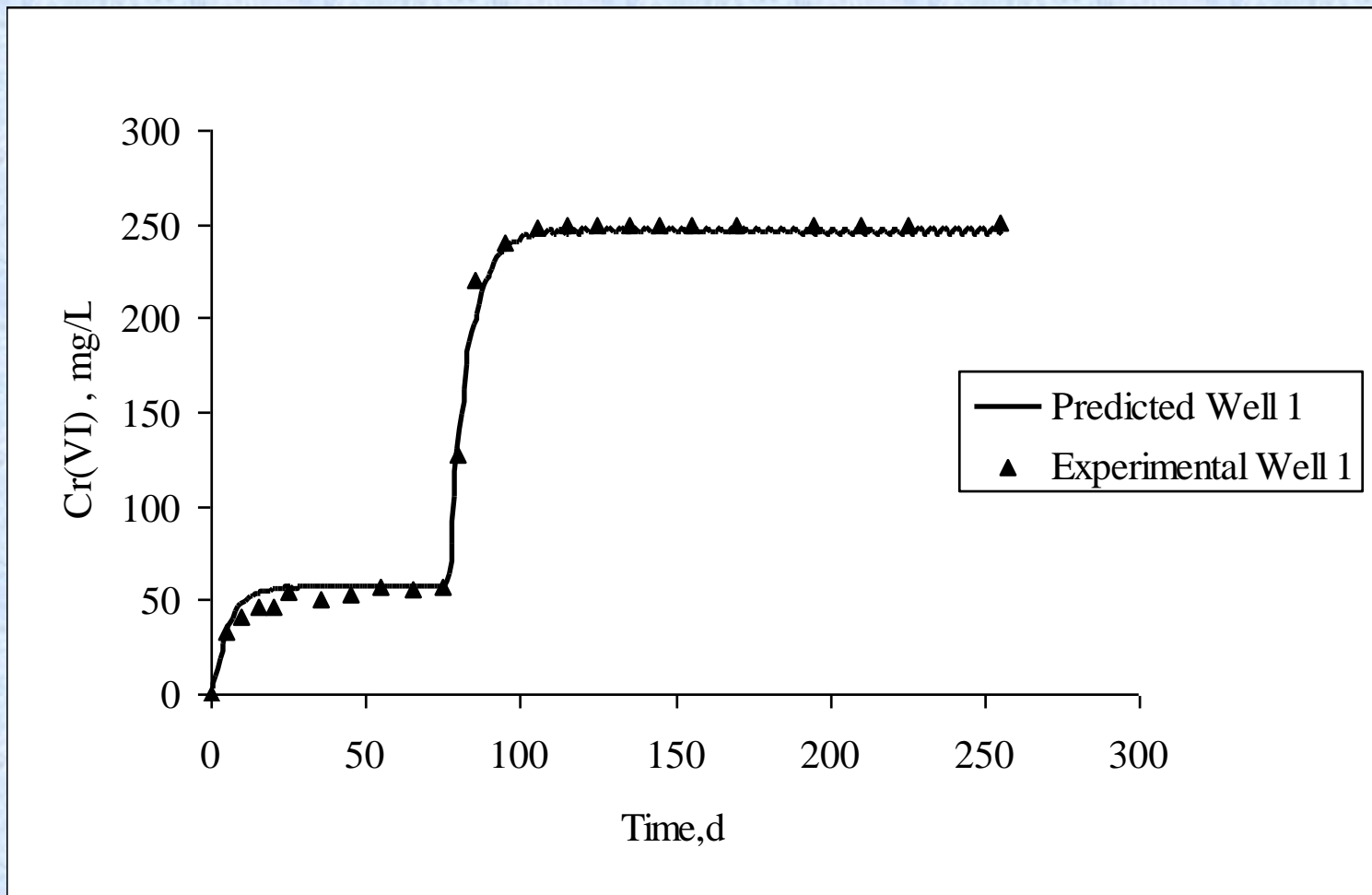
Experimental and modeling results for temporal variation of Cr(VI) concentration in wells 11-16 (at a distance of 110 cm from inlet) in reactor R1

Jeyasigh et al., Chem. Engrg. Jl., 2011



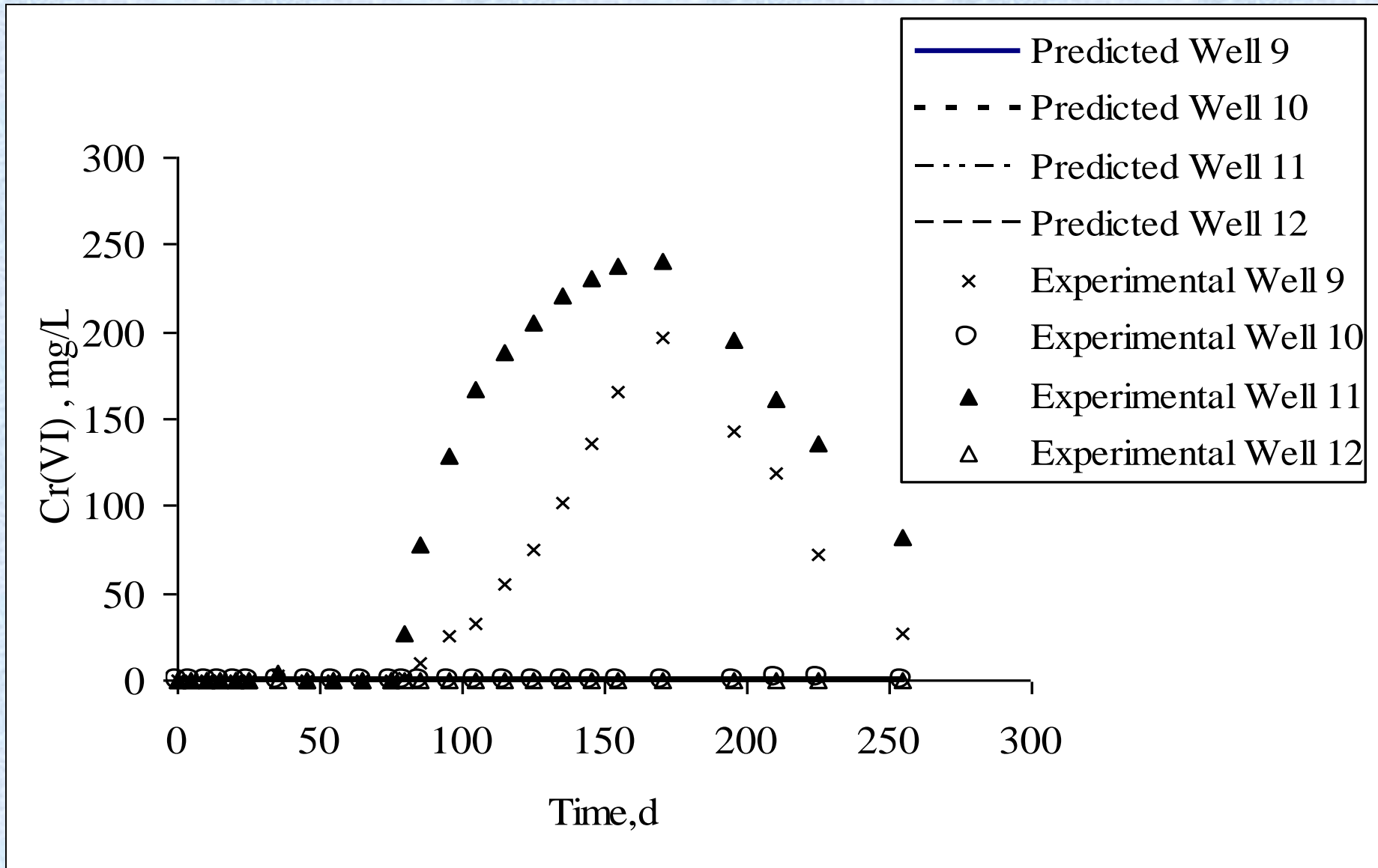
Experimental and modeling results for temporal variation of Cr(VI) concentration in wells 11-16 in reactor R2

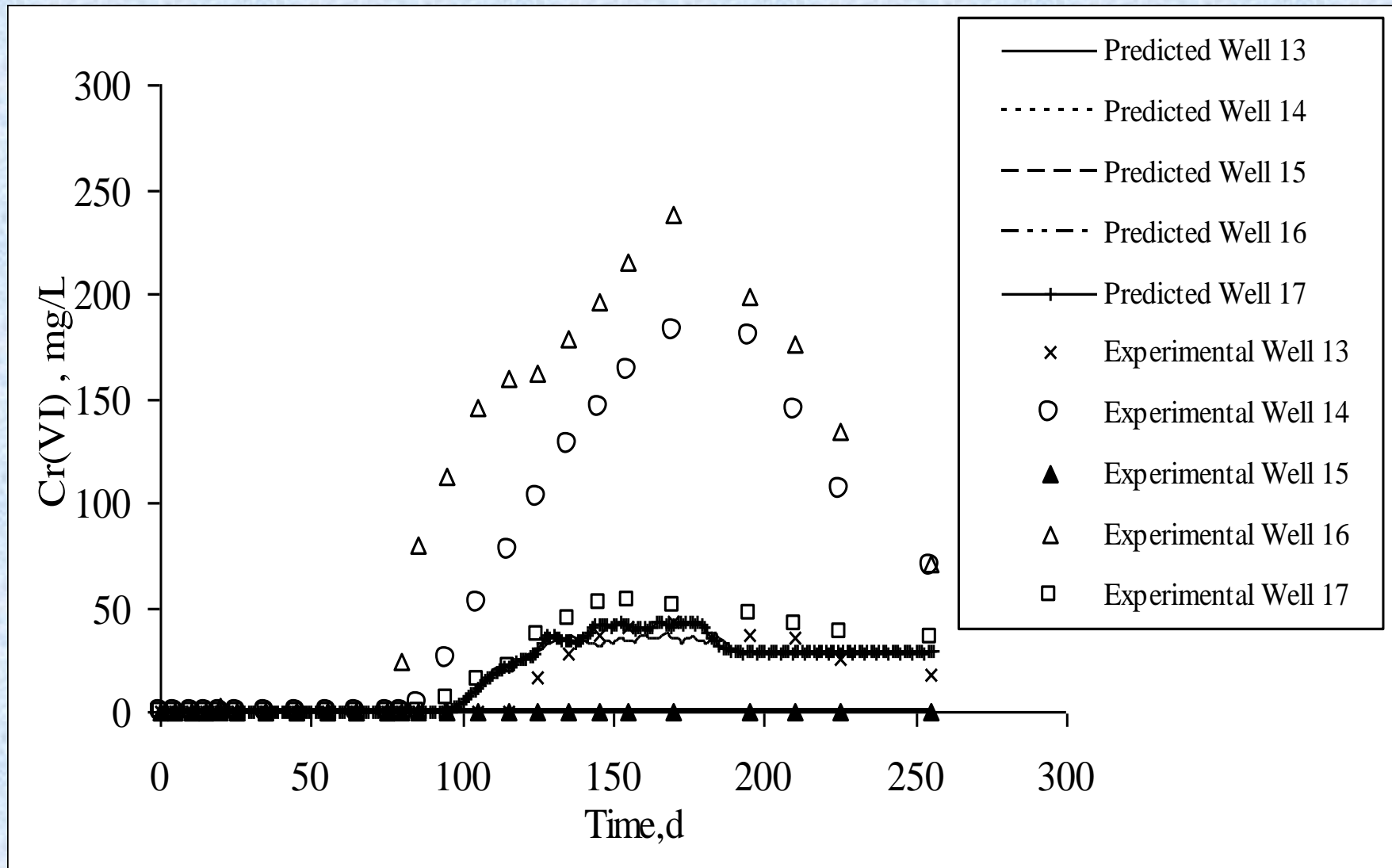
Jeyasigh et al., Chem. Engrg. Jl., 2011



Experimental and modeling results for temporal variation of Cr(VI) concentration at well no 1 in reactor R4 (4 wells system)

Jeyasigh et al., Chem. Engrg. Jl., 2011



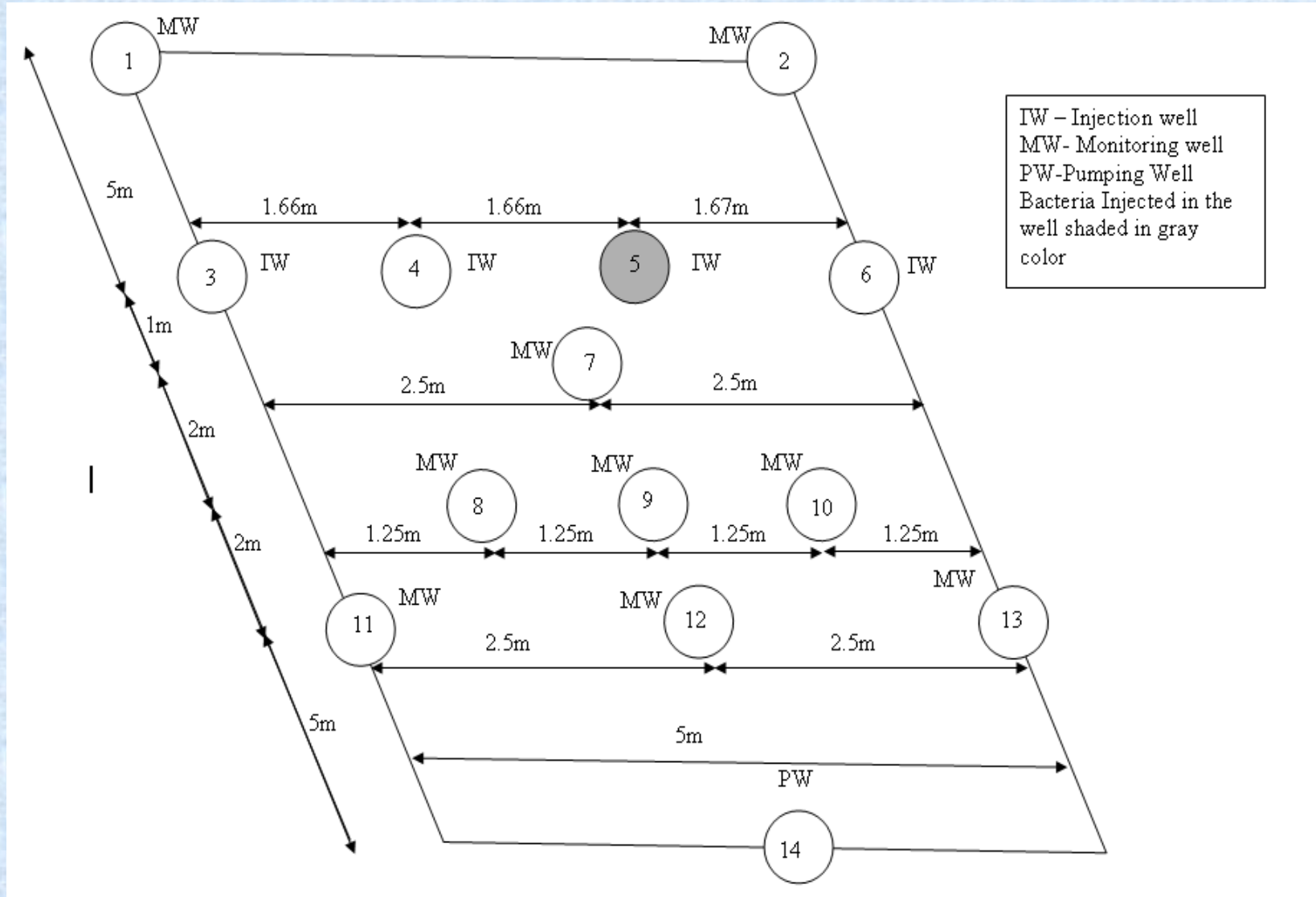


**Field Demonstration of
Bioremediation of Cr(VI)
Contaminated Soil and Aquifer in
Ranipet, Tamilnadu**

SCOPE

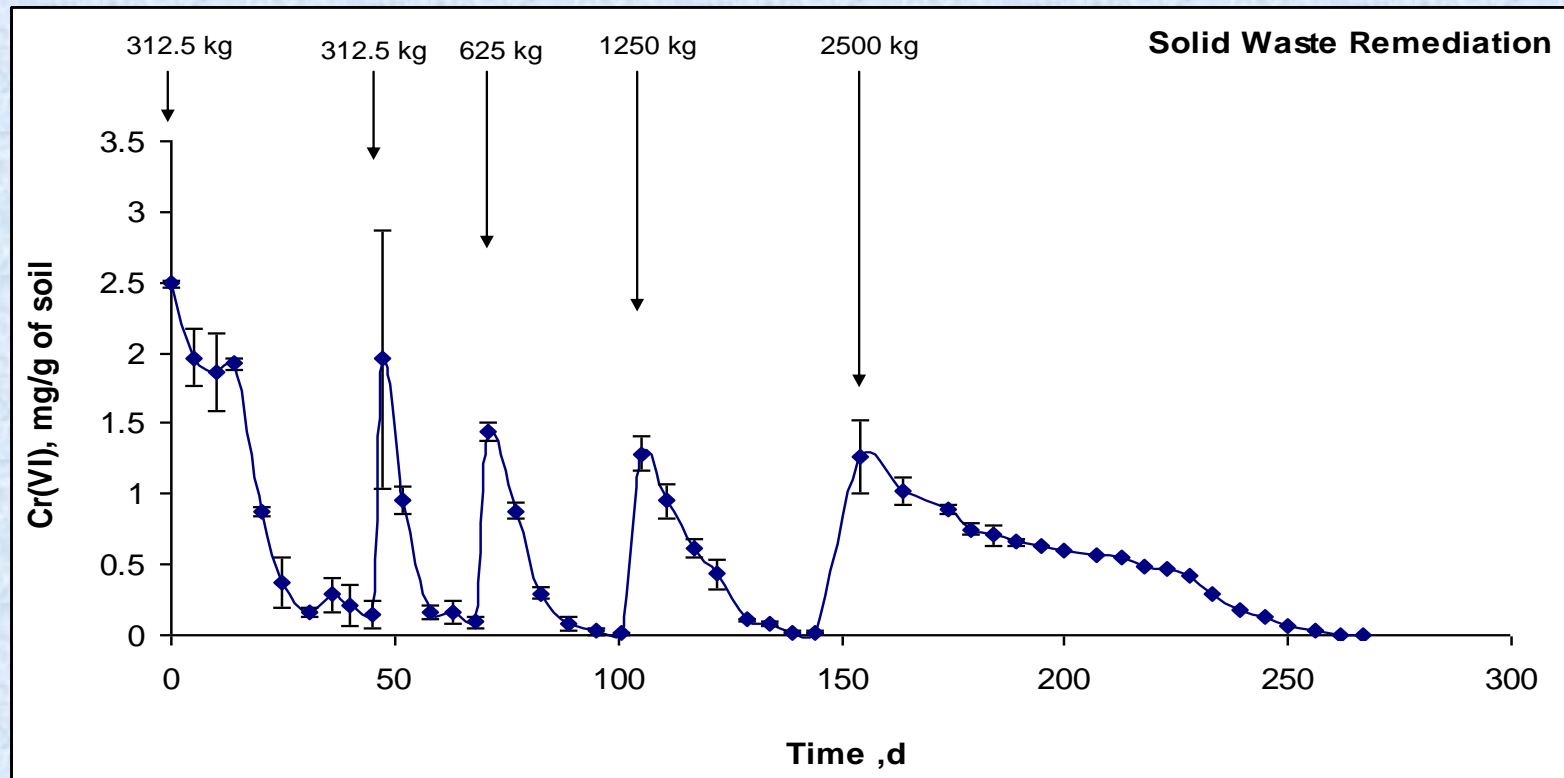
- **Remediation of at least 5 tons of chromium sludge in the vicinity of Tamilnadu Chromates and Chemicals Limited (TCCL) at the site;**
- **Demonstration of in-situ bioremediation of Cr(VI) contaminated aquifer in a 5 m ×5 m area of aquifer in the vicinity of Tamilnadu Chromates and Chemicals Limited (TCCL), Ranipet, by injection well - reactive zone technology;**

Well locations in the experimental plot



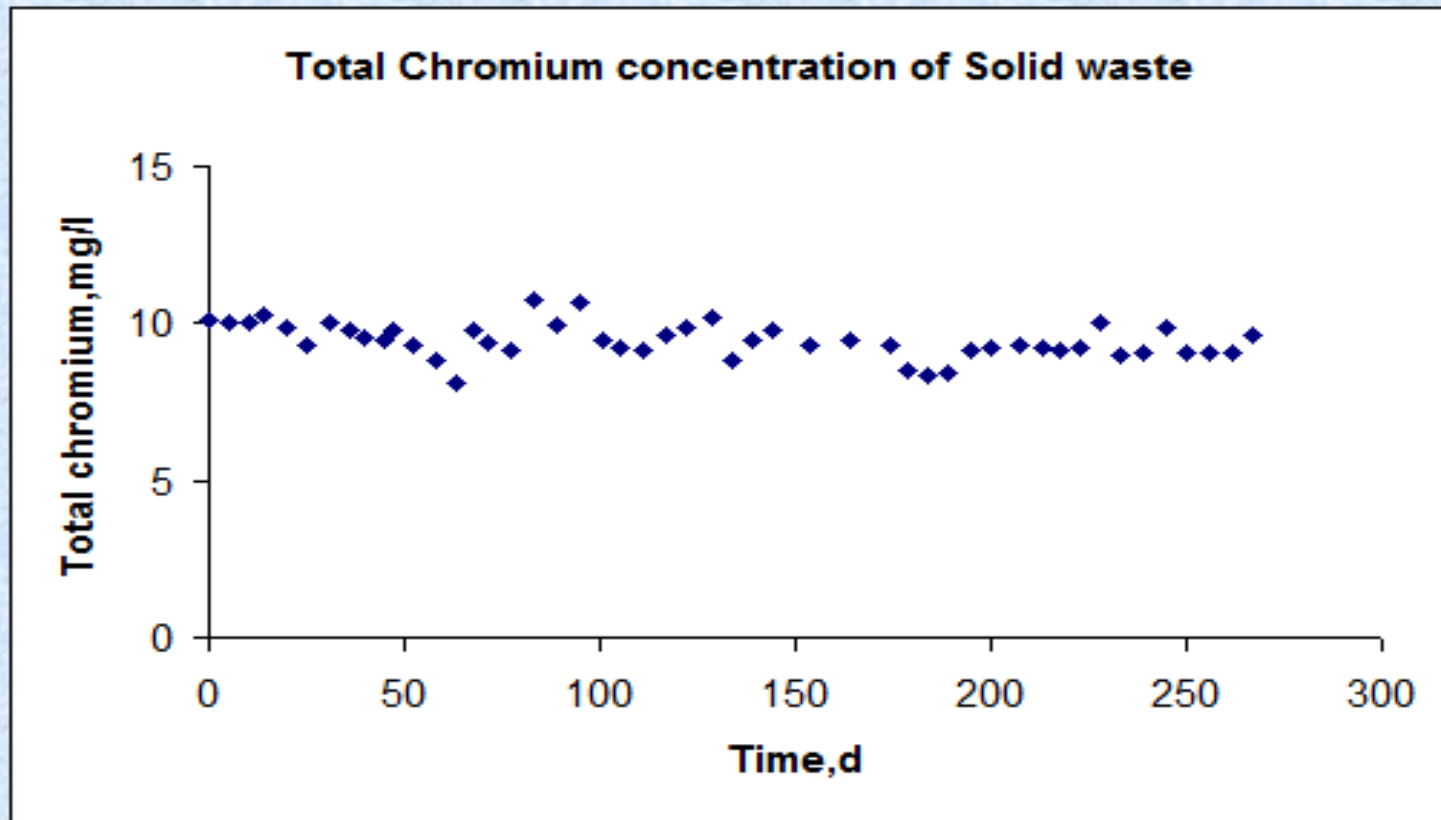
RESULTS

Soil Remediation



Variation of Cr(VI) concentration with respect to time in solid waste remediation (Mass of untreated sludge added at various time is mentioned inside the graph)

Variation of total chromium concentration with respect to time in solid waste remediation



Remediated and un-remediated soils



**Five Tones of Remediated Soil
Leachate from remediated soil**

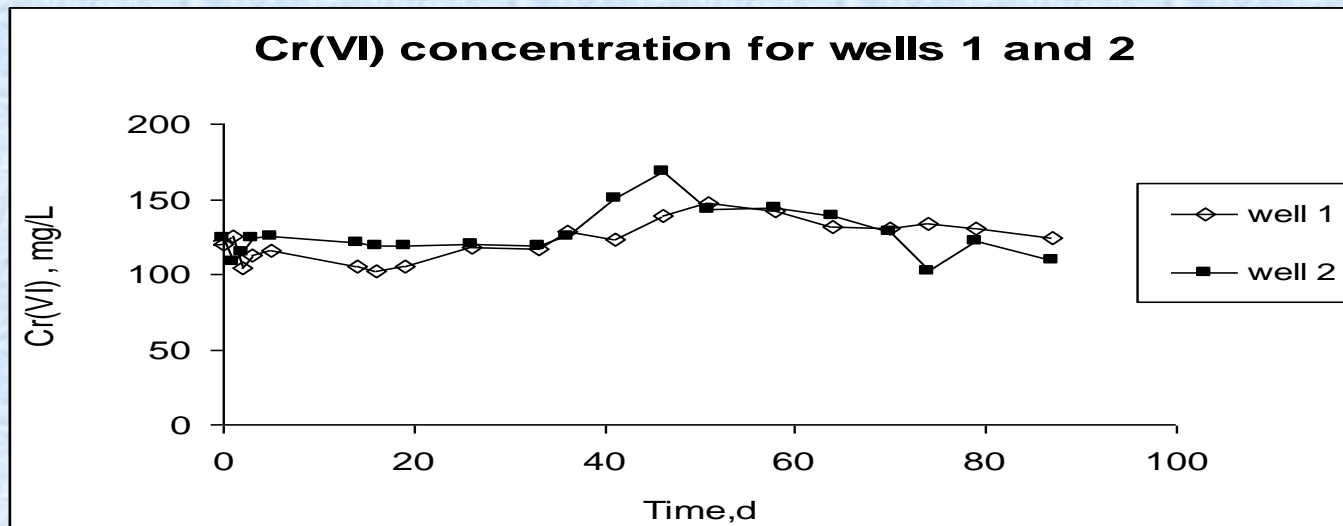


**Un-remediated Soil and
Leachate from un-
remediated soil**



Aquifer Remediation

Bioremediation using Molasses (Jaggery) as the Carbon Source

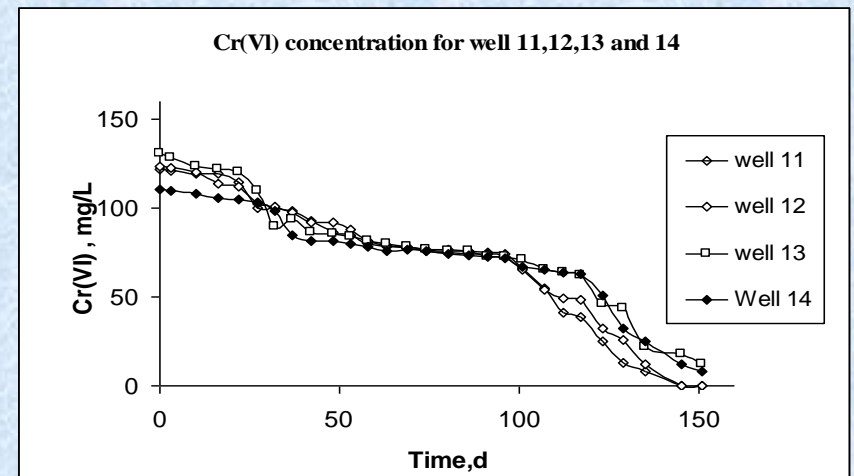
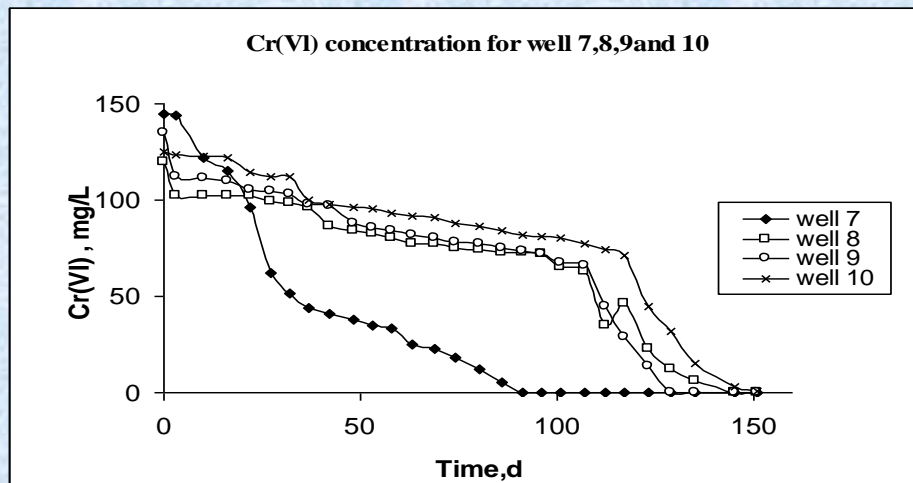
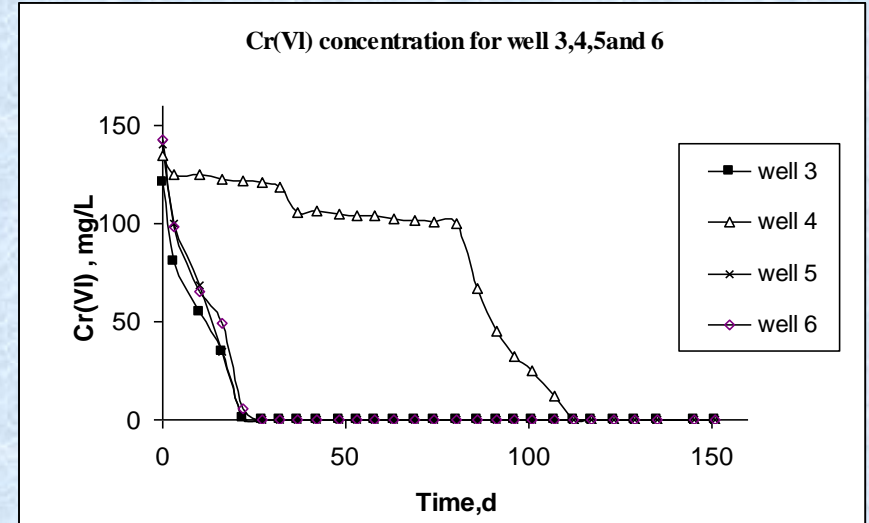
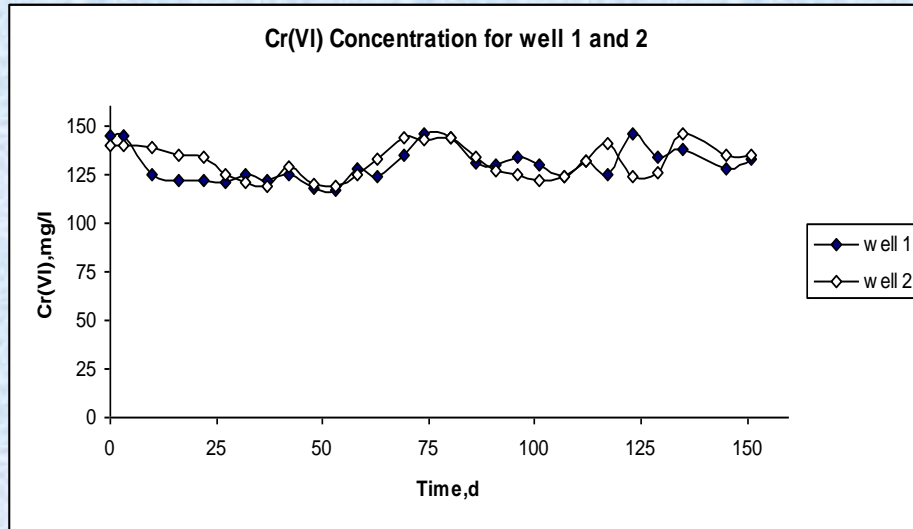


Variation of Cr (VI) concentration with respect to time in wells 1 and 2 (molasses as carbon source)

Bioremediation using Sugar as the Carbon Source

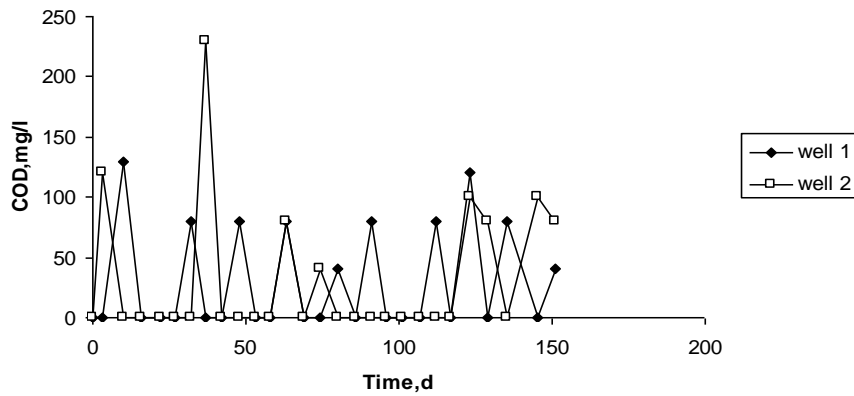
- Remediation of Cr(VI) aquifers were also carried out using sugar as the carbon source.
- For this study the initial biomass concentration was reduced to $1/10^{\text{th}}$ of that used in the previous case.
- Carbon source concentration also was reduced to $1/4^{\text{th}}$ and feeding interval was increased to 7- 10 days.
- The fate and transport of chromium (both Cr(VI) and Cr(III)), molasses and its derivatives, and microbes during the study period was monitored.

Bioremediation using Sugar as the Carbon Source: Cr(VI) concentrations

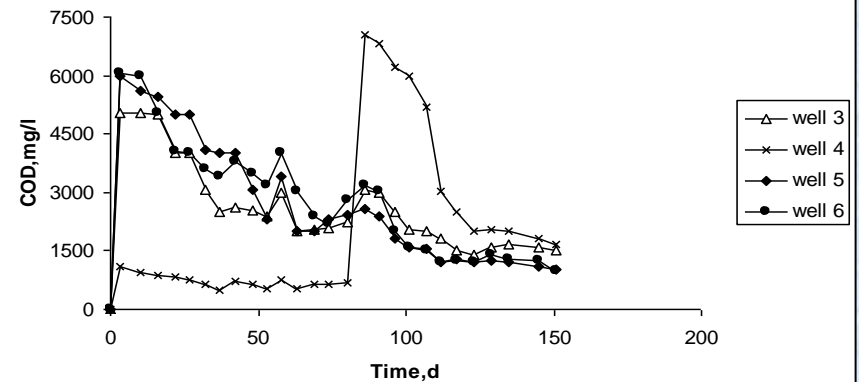


COD concentrations in various wells during bioremediation using sugar as carbon Source

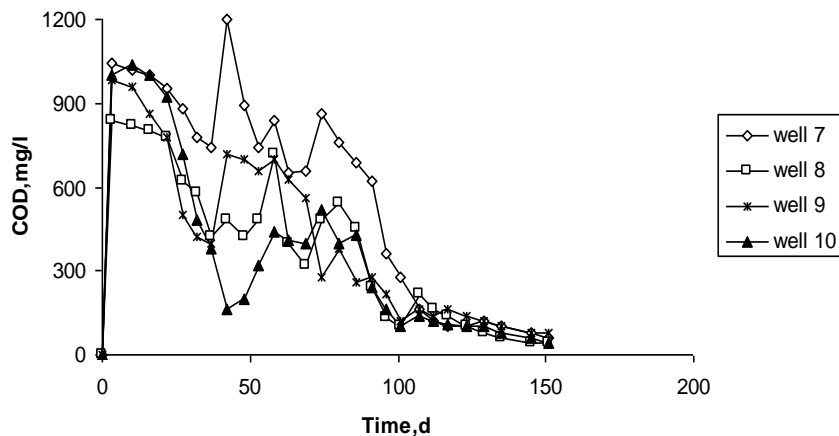
COD for well 1 and 2



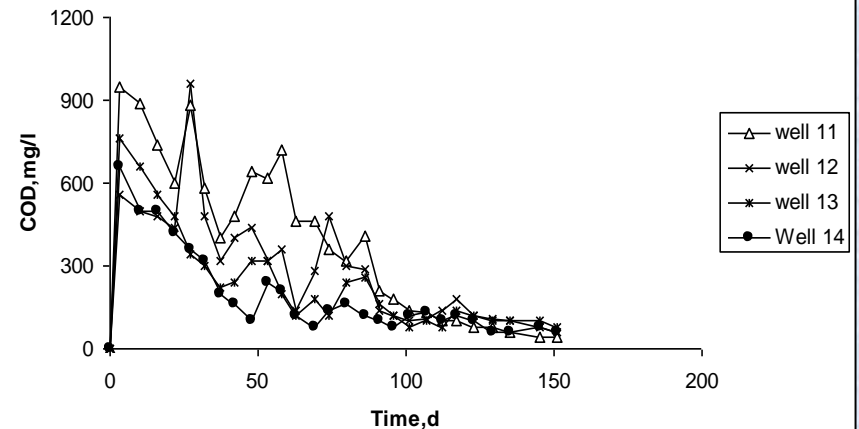
COD for well 3,4,5 and 6



COD for well 7,8,9 and 10

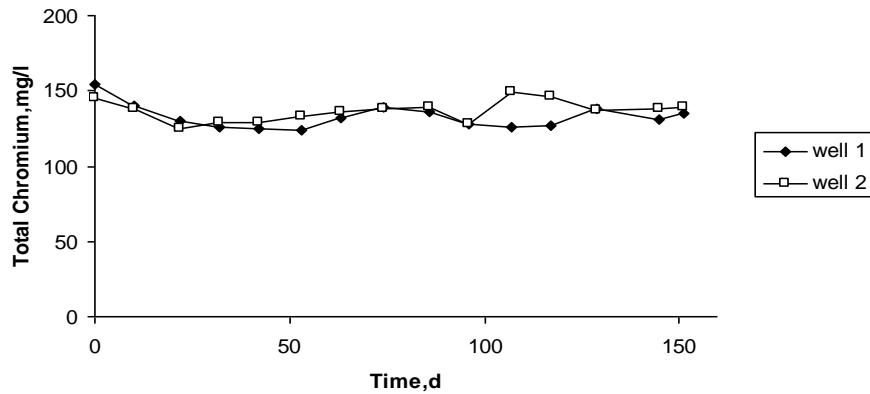


COD for well 11,12,13 and 14

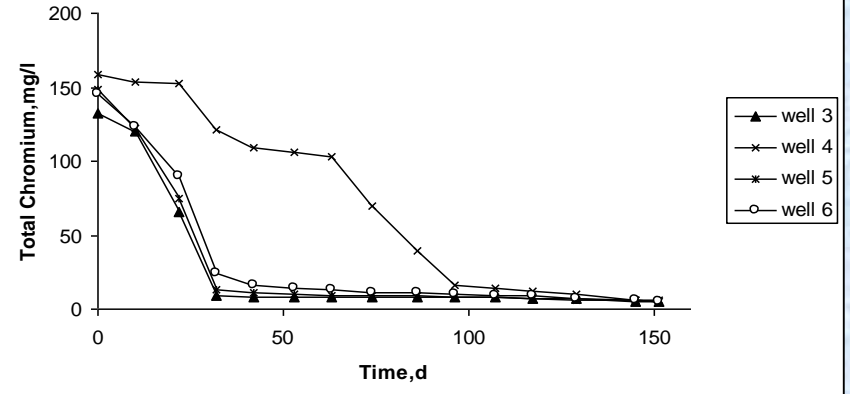


Total Cr concentrations in various wells during bioremediation using sugar as carbon Source

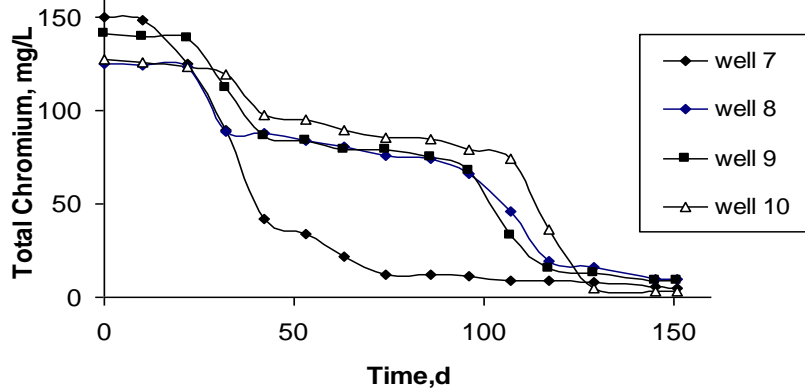
Total Chromium for well 1 and 2



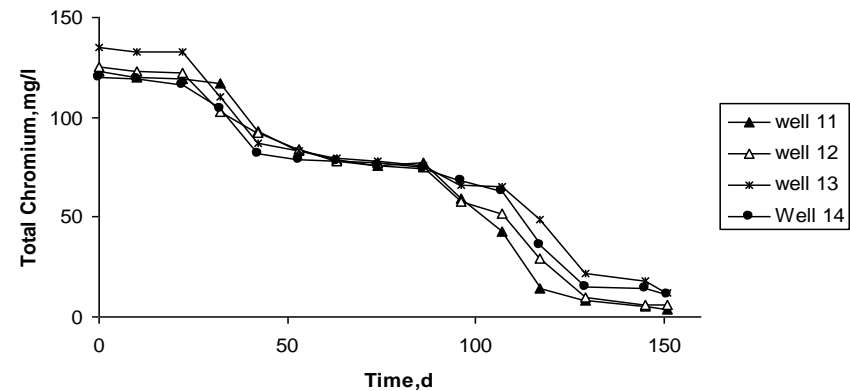
Total Chromium for well 3,4,5 and 6



Total Chromium for well 7,8,9 and 10



Total Chromium for well 11,12,13 and 14



Water samples from various wells after remediation



Water samples from various wells after remediation



Field Applications: Technology Transfer

Radiant Electroplaters

- **MR. ALI AKBAR, Radiant Electroplaters,
32 KMA Garden Road, Kodungaiyur,
Chennai-600118**

NGT Case

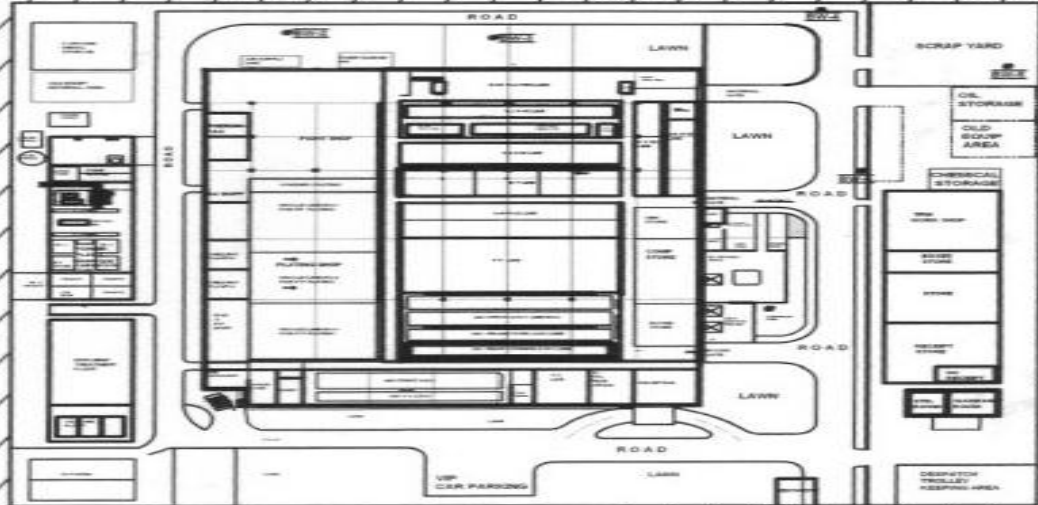
- **The wastes storage tank breached.**
- **Contaminated the neighboring industrial plot
and groundwater**
- **Industry was closed**

Munjal Showa Ltd.,

- Court Order to Close the Industry
- Fine Rs 5 crores.

MUNJAL SHOWA LTD
PLOT NO. 9 TO 11, GURGAON

N
↑



MAINROAD

VILLAGE STREET

DEAD BOREWELL

RESIDENCE OF MR. MAHAVI YADAV

BOREWELL

RESIDENCE OF MR. VIRENDER YADAV

BOREWELL

BOREWELL

Hydro-Geological Conditions

TUBEWELL NO.1

DEPTH: 100 MTRS

STRATA	TABLE
GL-12.5 M	DRY SAND & CLAY
12.5-20 M	FINE SAND
20-30 M	HARD CLAY
30-50 M	FINE SAND
50-60 M	FINE SAND
60-80 M	HARD CLAY
80-100 M	FINE SAND

TUBEWELL NO.2

DEPTH: 150 MTRS

STRATA	TABLE
GL-12.5 M	DRY SAND & CLAY
12.5-20 M	FINE SAND
20-30 M	HARD CLAY
30-55 M	FINE SAND
55-65 M	FINE SAND
65-85 M	HARD CLAY
85-105 M	FINE SAND
105- BELOW 150 M	HARD ROCK

TUBEWELL NO.3

DEPTH: 105 MTRS

STRATA	TABLE
GL-12.5 M	DRY SAND & CLAY
12.5-20 M	FINE SAND
20-35 M	HARD CLAY
35-55 M	FINE SAND
55-65 M	FINE SAND
65-85 M	HARD CLAY
85-105 M	FINE SAND

TUBEWELL NO.4

DEPTH: 105 MTRS

STRATA	TABLE
GL-12.5 M	DRY SAND & CLAY
12.5-20 M	FINE SAND
20-35 M	HARD CLAY
35-55 M	FINE SAND
55-65 M	FINE SAND
65-85 M	HARD CLAY
85-105 M	FINE SAND

TUBEWELL NO.5

DEPTH: 105 MTRS

STRATA	TABLE
GL-12.5 M	DRY SAND & CLAY
12.5-20 M	FINE SAND
20-35 M	HARD CLAY
35-55 M	FINE SAND
55-65 M	FINE SAND
65-85 M	HARD CLAY
85-105 M	FINE SAND

TUBEWELL NO.6

DEPTH: 100 MTRS

STRATA	TABLE
GL-12.5 M	DRY SAND & CLAY
12.5-20 M	FINE SAND
20-30 M	HARD CLAY
30-50 M	FINE SAND
50-60 M	FINE SAND
60-80 M	HARD CLAY
80-100 M	FINE SAND

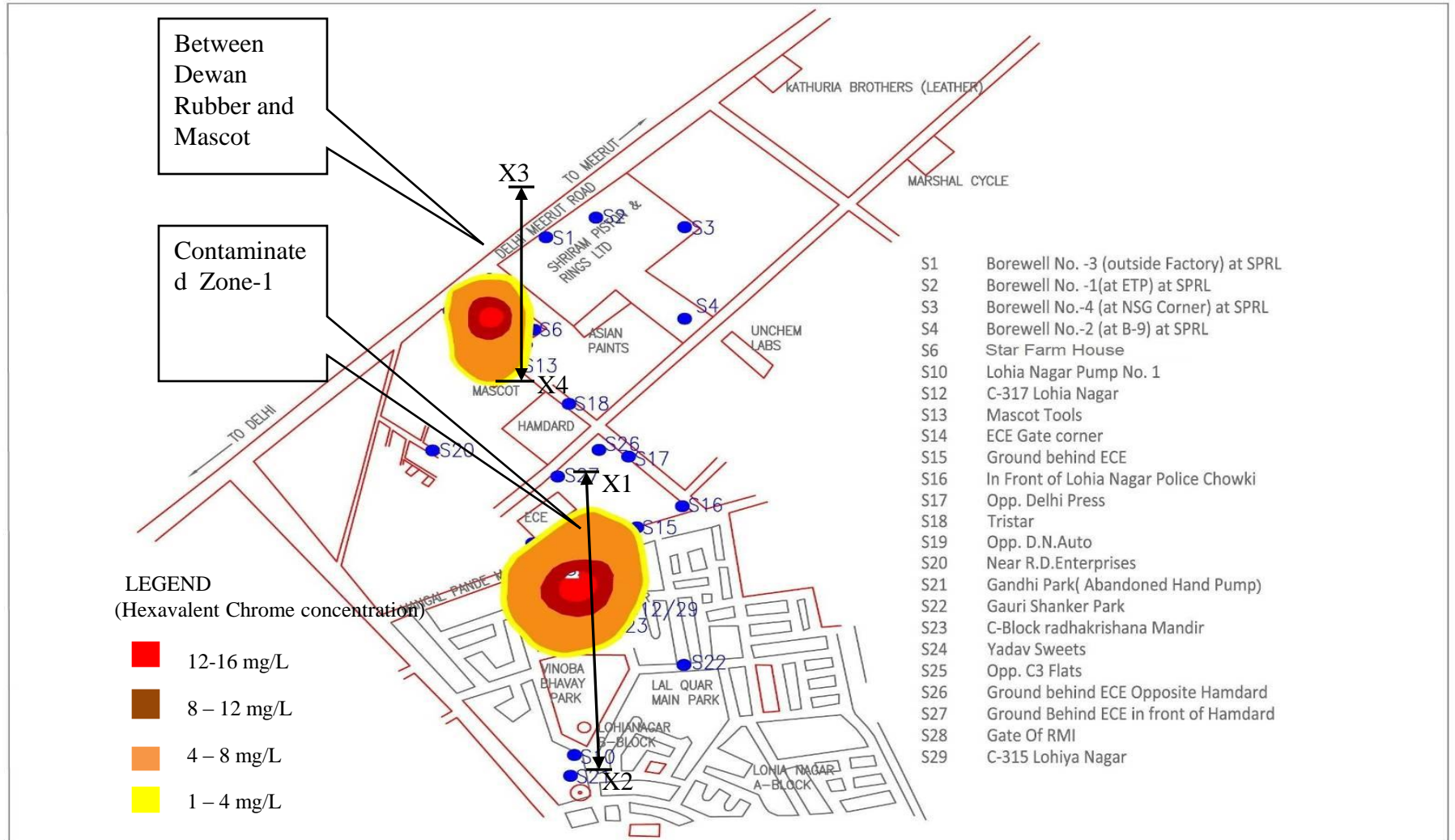
TUBEWELL NO.7

DEPTH: 100 MTRS

STRATA	TABLE
GL-12.5 M	DRY SAND & CLAY
12.5-20 M	FINE SAND
20-30 M	HARD CLAY
30-50 M	FINE SAND
50-60 M	FINE SAND
60-80 M	HARD CLAY
80-100 M	FINE SAND

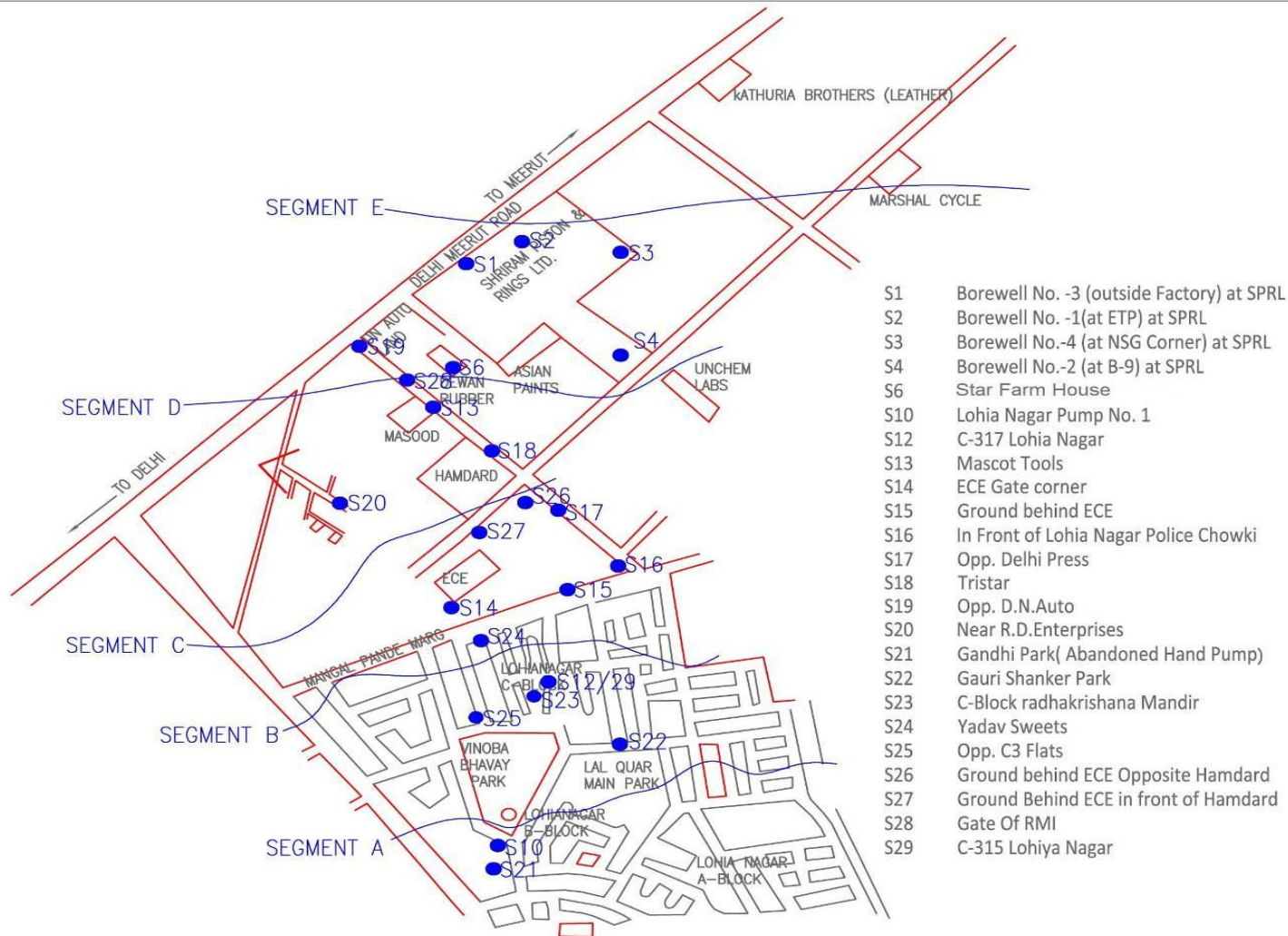
Shriram Pistons and Rings Ltd, Meerut Road, Ghaziabad, INDIA

MAP SHOWING 02 PLUME FORMATIONS IN AND AROUND LOHIA NAGAR



CONTAMINATED ZONE IDENTIFIED FOR SETTING UP ETP

MAP OF LOHIANAGAR AND ADJOINING AREA SHOWING SEGMENTS A – E



Quantification of Contaminated Groundwater

S. No.	Segment	Quantity of Contaminated Groundwater, $Q=A*Wlf*Sp.Y.$	Range of Hexavalent Chromium in Mg/L
1	Segment A	69,600 cu.m./yr.	Nil – 3.4
2	Segment B	2,08,800 cu.m./yr.	0.2 – 16.3
3	Segment C	52,200 cu.m./yr.	0.1 -1.3
4	Segment D	1,74,000 cu.m./yr.	1.3 – 15.4
5	Segment E	1,04,400 cu.m./yr.	Nil – 1.3

Thank you

