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Developing a Smart and Clean Technology for Bioremediation of Antibiotic Contamination in Arable Lands

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1 Developing a Smart and Clean Technology for Bioremediation of

2 Antibiotic Contamination in Arable Lands

4 **Abstract**

5 This study presents a smart technological framework to efficiently remove azithromycin
6 from natural soil resources using bioremediation techniques. The framework consists of several
7 modules, each with different models such as *Penicillium Simplicissimum* (PS) bioactivity, soft
8 computing models, statistical optimisation, Machine Learning (ML) algorithms, and Decision Tree
9 (DT) control system based on Removal Percentage (RP). The first module involves designing
10 experiments using a literature review and the Taguchi Orthogonal design method for cultural
11 conditions. The RP is predicted as a function of cultural parameters using Response Surface
12 Methodology (RSM) and three ML algorithms: Instance-Based K (IBK), KStar, and Locally
13 Weighted Learning (LWL). The sensitivity analysis shows that pH is the most important factor
14 among all parameters, including pH, Aeration Intensity (AI), Temperature, Microbial/Food (M/F)
15 ratio, and Retention Time (RT), with a p-value of < 0.0001 . AI is the next most significant
16 parameter, also with a p-value of < 0.0001 . The optimal biological conditions for removing
17 azithromycin from soil resources are a temperature of 32°C, pH of 5.5, M/F ratio of 1.59 mg/g,
18 and AI of 8.59 m³/h. During the 100-day bioremediation process, RP was found to be an
19 insignificant factor for more than 25 days, which simplifies the conditions. Among the ML
20 algorithms, the IBK model provided the most accurate prediction of RT, with a correlation
21 coefficient of over 95%.

22 **Keywords:** Azithromycin; bioremediation; machine learning; *penicillium simplicissimum*;
23 Taguchi design.

24 1. Introduction

25 Managing soil health is a significant challenge in agriculture. Moreover, it is a complex
26 issue that is difficult to address using traditional methods (Mohammad et al., 2021). With the
27 increasing prevalence of new diseases worldwide, people especially in low-income countries are
28 resorting to antibiotics to treat infections (Klein et al., 2021). This trend can have adverse health
29 effects due to the carcinogenic properties of antibiotics (Llor and Bjerrum, 2014). Recent studies
30 have also shown that the widespread use of antibiotics in natural resources such as air, soil, water,
31 and humans can lead to health risks as microorganisms develop drug resistance (Zhang et al.,
32 2015).

33 Azithromycin (AZ) is one of the most widely used antibiotics for treating bacterial
34 infections. According to the Food and Drug Administration (USFDA), AZ (Zithromax or Zmax)
35 can disrupt the electrical processes of the heart, leading to a potentially fatal irregular heartbeat
36 (Patel et al., 2020). Due to the persistent nature of antibiotics, their recalcitrance, and the
37 emergence of resistance genes, their widespread presence in natural resources can cause a global
38 environmental problem, making their remediation essential (Cycoń et al., 2019).

39 Azithromycin is classified as an Endocrine Disrupting Compound (EDC), which can pose
40 unprecedented health risks due to pollutant emissions in soil and water (Lau et al., 2020). Soil
41 pollution by EDCs can occur when pharmaceutical and industrial solid wastes are released into the
42 environment without adhering to regulations and landfill standards (Hu et al., 2010). According to
43 a survey conducted by SOM-institute¹ in Sweden in 2020, environmental pollution and antibiotic
44 resistance are considered significant concerns by the public. The release of azithromycin in soil

¹ Source(s): SOM-institute; ID 909223

45 exacerbates both issues and increases public concerns. Therefore, reducing the concentration of
46 azithromycin in soil would alleviate these problems and increase public satisfaction.

47 Addressing antimicrobial resistance (AMR) requires clear global guidelines and regulatory
48 options, but this has been challenging due to the diversity and dynamic nature of healthcare and
49 regulatory systems across different countries ([Chokshi et al., 2019](#)). As a result, the World Health
50 Organisation (WHO) has made policies and recommendations, but it is up to each nation to
51 regulate antibiotic resistance. To tackle AMR, individuals, healthcare professionals, policymakers,
52 the healthcare industry, and the agricultural sector all need to act.

53 The WHO recommends that individuals use antibiotics only with a prescription, follow
54 infection prevention guidelines, and practice safe food preparation according to the WHO Five
55 Keys to Safer Food (FKSF). Policymakers can develop a national action plan to reduce antibiotic
56 resistance, improve surveillance systems of antibiotic use, regulate the proper use and disposal of
57 medicine, and educate the public on antibiotic resistance. Healthcare professionals can prevent
58 infections by maintaining a clean and sterile working environment, prescribing antibiotics only
59 when necessary, reporting antibiotic-resistant infections to surveillance systems, and educating
60 patients on the risks of antibiotic misuse and how to prevent infections. The healthcare industry
61 should invest in research and development of new antibiotics, diagnostics, and vaccines. The
62 agricultural sector must use antibiotics in animals only under the supervision of a veterinarian,
63 avoid using antibiotics to prevent diseases, and use vaccines instead. Fine practices can help reduce
64 infections and improve biosecurity on farms ([WHO, 2019](#)).

65 In 2015, the World Health Assembly established a global action plan on AMR, consisting of 5 key
66 policies: 1) enhancing public awareness about AMR, 2) promoting stewardship of antibiotics, 3)
67 reducing and preventing infection, 4) improving surveillance systems and research, and 5)

68 ensuring sustainable investment in combating AMR. The Assembly also urged each country to
 69 develop its own national action plan (NAP) to combat AMR (Anderson, 2020). In response, Iran
 70 created its own NAP (IRI-NAP) in 2016 in five sections (summarised in Table 1) aligned with the
 71 main policies outlined in the global action plan (Moradi et al., 2018).

72 Among the most common types of antibiotics (e.g., *β-lactams*, *Macrolides*, *Fluoroquinolones*,
 73 *Tetracyclines*, *Sulfonamides*, *Diaminopyrimidines*, *Lincosamides*, and their degradation products),
 74 according to the reports, Fluoroquinolones, Tetracyclines, and Sulfonamides have the highest
 75 concentration in soil samples mostly caused by manure and wastewater irrigation (Yang et al.,
 76 2021). Table 2 shows the concentration data of some of the frequently found antibiotics in soil.

77 **Table 1. Selected policies of the national action plan in Iran against Antimicrobial resistance**

Goal	Policy
Enhance public awareness of AMR	<ul style="list-style-type: none"> -Conduct education courses for specific groups such as children and elders -Run awareness campaigns for people working in related fields -Initiate targeted activities
Optimise the use of antibiotics	<ul style="list-style-type: none"> -Strategic purchases of antimicrobial medicines to improve the quality -Empowering medical institutions to create guidelines and manuals for antimicrobial stewardship of their own
Prevent and control infections	<ul style="list-style-type: none"> -Promotion of vaccines -Support NGO activities in coordination with hospitals to prevent and control infection
Improve surveillance system	<ul style="list-style-type: none"> -Control antimicrobial residue in food productions -Sending AMR experts all over the country to respond quickly while outbreaks happen -Create a monitoring system of prescriptions and antibiotics -Update and monitor prescription criteria for antibiotics -Increase the capacity of laboratories dealing with AMR
Guarantee sustainable investment and research in combating AMR	<ul style="list-style-type: none"> -Research the AMR surveillance systems -Promote research to clarify the necessity of AMR -Create a database of resistant genes -Conduct more research to deeply investigate the impact of AMR on health -Reconsidering the microbial diagnosis, treatment, and prevention approaches -Promote research and industry with the international collaboration

78

Table 2. The highest concentrations of some major antibiotics reported in the soil environment.

Type	Antibiotic	Concentration (ng/g)	Reference
<i>β-lactams</i>	<i>Amoxicillin</i>	200	Braschi et al., 2013
<i>Fluoroquinolones</i>	<i>Ciprofloxacin</i>	350	Al Masud et al., 2023; Martínez-Carballo et al., 2007; Karci and Balcioglu, 2009; Hu et al., 2010; Van Doorslaer et al., 2014; Pan and Chu, 2017b
	<i>Difloxacin</i>	21.5	
	<i>Enrofloxacin</i>	1,347.60	
	<i>Norfloxacin</i>	5,610	
	<i>Ofloxacin</i>	898	
<i>Quinolone</i>	<i>Sarafloxacin</i>	5.92	Rashid et al., 2023
<i>Macrolides</i>	<i>Enrofloxacin</i>	22.93	Thiele-Bruhn, 2003; Leal et al., 2012; Tasho and Cho, 2016; Pan and Chu, 2017, 55
	<i>Erythromycin</i>	100	
	<i>Tylosin</i>	1,250	
	<i>Azithromycin</i>	1,000	Topp et al., 2016
	Total macrolides	1.471	Li et al., 2023
<i>Sulfonamides</i>	<i>Sulfachloropyridazine</i>	52.9	Thiele-Bruhn, 2003; Dolliver et al., 2007; Karci and Balcioglu, 2009; Hu et al., 2010; Carter et al., 2014; Pan and Chu, 2017
	<i>Sulfadiazine</i>	85.5	
	<i>Sulfadimethoxine</i>	40.4	
	<i>Sulfadoxine</i>	9.1	
	<i>Sulfamethoxazole</i>	54.5	
	<i>Sulfamethazine</i>	200–25,000	
	<i>Sulfamonomethoxine</i>	5.37	
	<i>Sulfapyridine</i>	5.11	
Total sulfonamides	18.497	Li et al., 2023	
<i>Tetracyclines</i>	<i>Chlortetracycline</i>	12,900	Hamscher et al., 2002; Thiele-Bruhn, 2003; Karci and Balcioglu, 2009; Hu et al., 2010; Liu et al., 2016; Tasho and Cho, 2016; Pan and Chu, 2017; Łukaszewicz et al., 2018
	<i>Doxycycline</i>	728	
	<i>Oxytetracycline</i>	50,000	
	<i>Minocycline</i>	32	
	<i>Tetracycline</i>	2,683	

80

81 Various techniques are available for degrading antibiotics in different environments such as water,
82 wastewater, soil, and solid waste. These techniques include adsorption (Gheibi et al., 2023),
83 integrated biological treatment with membranes (Zhao et al., 2021), permeable reactive barriers

84 (Zhao et al., 2018), fungus-based bioremediation (Mohammadi et al., 2021), and electrochemical
85 systems (Bicudo et al., 2017). However, each technique has its strengths and weaknesses and is
86 applicable in specific real field situations. For example, permeable reactive barriers, membranes,
87 biofilm membranes, and adsorption processes provide an acceptable efficiency of more than 95%,
88 but they have limited capacity for decontamination and may require regeneration or recovery of
89 the system in a short time (Zhang et al., 2020c). On the other hand, coagulation and
90 electrocoagulation procedures have high efficiency but involve chemical material addition and
91 unusual energy consumption (Bicudo et al., 2021). Bioremediation, which involves living
92 organisms such as fungi, algae, bacteria, plants, and animals, is a process that removes or detoxifies
93 pollutants in the environment (Jagtap, 2020). It is environmentally friendly, requires low capital
94 investment, and has minimal energy consumption, making it a popular method for degradation
95 (Irshad et al., 2021). Bioremediation is particularly suitable for removing antibiotic compounds
96 that are sensitive to pH and temperature (Liu et al., 2017).

97 Mycoremediation is a type of bioremediation that involves the use of fungi as the
98 decomposer of pollutants (Schmit and Mueller, 2007). Fungi are a diverse group of
99 microorganisms with unique characteristics, including the ability to form extensive mycelial
100 networks, low specificity of their catabolic enzymes, and their independence from using pollutants
101 as a growth substrate (Harms et al., 2011). Fungi are known to be capable of degrading and
102 mineralising recalcitrant antibiotics due to their non-specific, non-stereoselective enzymatic
103 systems based on free-radical levels (Čvančarová et al., 2015). Over the past two decades, fungi
104 have been widely used for the treatment of waste and wastewater as well as for the degradation of
105 hazardous compounds (Khatoon et al., 2021). Fungi have multiple strategies to cope with toxic
106 compounds, such as antibiotics, including *bioadsorption*, *biomineralisation* (bio-precipitation) as

107 well as biotransformation and biodegradation mediated by enzymatic systems (Olicón-Hernández
108 et al., 2017).

109 Fungi have been found to play a critical role in removing heavy metals and mineralising
110 various types of pollutants, including phenols, halogenated phenolic compounds, petroleum
111 hydrocarbons, and polycyclic aromatic compounds (Sing, 2006). They are known to excrete
112 enzymes for the decomposition of carbohydrates without prior hydrolysis, making them highly
113 effective in degrading a vast number of pollutants (Bellaouchi et al., 2021). Fungi have several
114 advantages such as being easy to grow in fermenters and having a filamentous structure that allows
115 for easy separation of fungal biomass (Akhtar and Abdullah, 2014). *Penicillium* strains are popular
116 among all other fungi species as they can live in saline environments and have been reported to
117 treat heavy metals, polycyclic aromatic hydrocarbons, phenol and its derivatives, wastewater, and
118 wastes (Leitão et al., 2007). Among these species, *Penicillium simplicissimum* (PS) has been
119 selected as the decomposer in this study to learn the effectiveness of this specie in the removal of
120 AZ in soil (Sowmya et al., 2015).

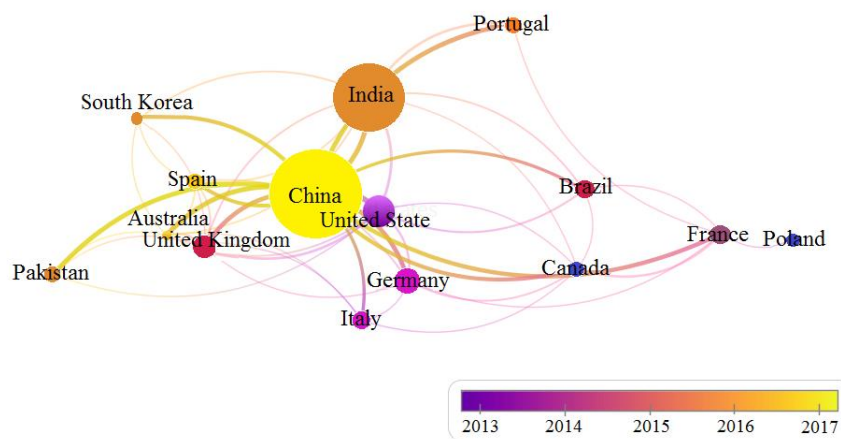
121 The performance of microorganisms is mainly dependent on the cultural conditions
122 (Khayati and Barati, 2017). The efficiency of biodegradation processes relies on several factors
123 such as pH, temperature, soil properties and substrate (Oliveira et al., 2020). Conventional
124 optimisation methods take a lot of time and cost due to numerous variables, which can be overcome
125 by using multi-factor methods such as Taguchi orthogonal design (Carraro et al., 2022). This
126 approach helps to investigate parameters, obtain more data, and reduce time and cost by suggesting
127 effective adjustments to control factors. Taguchi has been used for myco-synthesis of nano-silver,
128 wastewater treatment processes, and bioremediation and biodegradation process optimisation.

129 The use of Machine Learning (ML) approaches has significantly helped in modelling
130 biodegradation procedures with high accuracy and non-limited applicability (Sodhi and Singh,
131 2022). Several studies have been conducted on the bioremediation of various pollutants by fungi,
132 and ML algorithms such as Random Forest (RF), Adaptive Neuro-Fuzzy Inference System
133 (ANFIS), Random Tree (Mohammadi et al., 2021), Support Vector Machine (SVM) (Liu et al.,
134 2022), M5 Pruned model tree, Gaussian Processes (GP), and Sequential Minimal Optimisation
135 (SMOreg) (Akbarian et al., 2022) have been applied to predict the biodegradation efficiency and
136 cultural conditions.

137 The fate and remediation of antibiotic pollutants in the environment have been extensively
138 researched. Vermillion and Tjeerdema (2017) studied the degradation kinetics of AZ under aerobic
139 and anaerobic conditions. Sidhu et al. (2019) used Continuous Stirred-Tank Reactors (CSTRs) to
140 remediate *ciprofloxacin* and AZ with soil-based biomaterials, while Li et al. (2019) used
141 electrokinetic remediation for tetracycline-polluted soil. Zhan et al. (2021) evaluated heat
142 treatment for the decontamination of *tetracycline* and *roxarsone*. Mohammadi et al. (2021)
143 proposed a fungus-based purification method for *amoxicillin*, while Sidhu et al. (2021) assessed
144 AZ resistance in biosolids. Zelt et al. (2021) emphasised the importance of purifying antibiotics
145 from agricultural soil to ensure food chain health.

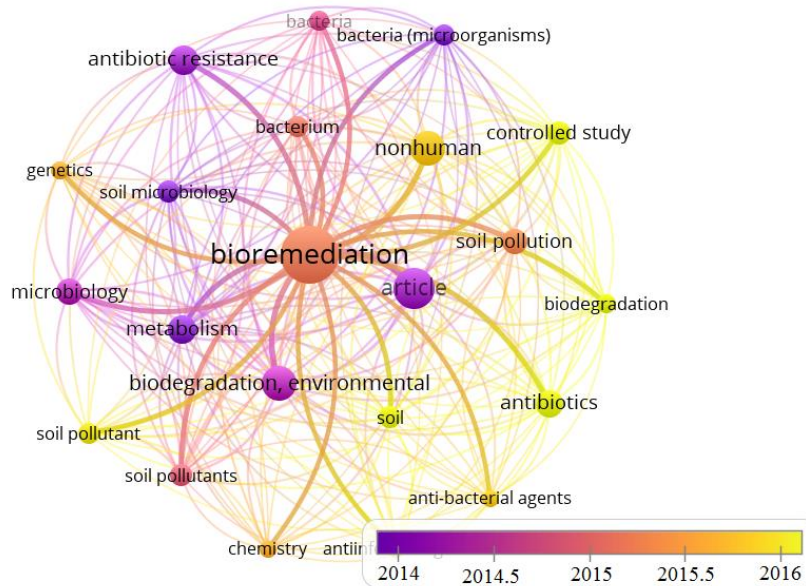
146 To evaluate the research on bioremediation process for decontamination of antibiotics from
147 soil, the library assessment was carried out using the VOSviewer software, where more than ten
148 documents (Fig. 1a) and over 100 repetitions of keywords (Fig. 1b) related to soil, bioremediation,
149 and antibiotics were filtered (Fig. 1). The results showed that China, India, and the United States
150 have conducted most of the investigations on this topic. The research on this subject in Iran was
151 challenging due to the limitation of facilities and equipment. The bioremediation subject is

152 associated with soil pollution, antibiotics, soil microbiology, and biodegradation issues, and it is
 153 regarded as a pressing issue, essential for scientific communities, which is the aim of this study.
 154 The figures generated by the software represent different ranges of time series and show the
 155 accumulated published documents based on different subjects and the focus of the country's
 156 contributions.



157
 158

(a)



159
 160
 161
 162

(b)

Fig. 1. Scientometry analysis of antibiotic bioremediation from soil resources through (a) country, and (b) occurrence of keywords aspects

163 [Table 3](#) provides an overview of research that has used fungi for the removal of antibiotics.
 164 The authors argue that a sustainable system for soil quality management is necessary to reduce the
 165 impact of biomagnification, epidemiological disease, and immunological issues ([Fasihi et al.,](#)
 166 [2021](#)), but previous studies have not considered a smart system for controlling bioremediation
 167 structures. Therefore, this study aims to develop a smart framework for optimal control and
 168 degradation of AZ in soil using *Penicillium Simplicissimum* (PS) bioactivities, the Taguchi design
 169 method for optimisation, and three lazy machine learning models to predict bioremediation
 170 behaviour. The study also aims to design a control system for the bioremediation system based on
 171 decision tree modelling and evaluate a sustainable industry using conceptual models. The authors
 172 argue that this study will address a knowledge gap related to optimising decontamination of AZ
 173 by considering optimisation, system performance prediction, and process control through
 174 conceptual modeling at the same time.

175 **Table 3. Bioremediation studies using fungus for antibiotic removal.**

Fungal name	Antibiotic contamination	Mechanism	Reference
<i>Trametes versicolor</i>	Azithromycin	Bio-oxidation	Del Álamo et al., (2022)
<i>Ganoderma lucidum</i>	20 different antibiotics	Biodegradation	Salandez et al., (2022)
<i>Penicillium oxalicum RJJ-2</i>	Erythromycin	Biodegradation	Ren et al., (2021)
<i>Penicillium commune</i>			
<i>Epicoccum nigrum</i>			
<i>Trichoderma harzianum</i>	Oxytetracycline	Biodegradation	Ahumada-Rudolph et al., (2021)
<i>Aspergillus terreus</i>			
<i>Beauveria bassiana</i>			

Fungal name	Antibiotic contamination	Mechanism	Reference
<i>Penicillium restrictum</i>	Sulfamethoxazole, Erythromycin, Tetracycline	Biodegradation	Fakhri et al., (2021)
<i>Aspergillus flavus</i>	Amoxicillin	Biodegradation	Mohammadi et al., (2021)
<i>Trametes versicolor</i>	Azithromycin	Biodegradation	Tormo-Budowski et al., (2021)
<i>Pleurotus ostreatus</i>	Sulfonamides Tetracyclines	Biodegradation	Camacho-Arévalo et al., (2021)
<i>Trametes polyzona</i>	Amoxicillin	Biodegradation	Lueangjaroenkit et al., (2019)
<i>Pycnoporus sanguineus,</i> <i>Phanerochaete</i> <i>chrysosporium</i>	Ciprofloxacin	Biodegradation	Gao et al., (2018)
<i>Pleurotus ostreatus</i>	Ciprofloxacin	Biodegradation	Singh et al., (2017)
<i>Aspergillus terreus</i> FZC3	Gentamicin	Biosorption and biodegradation	Liu et al., (2016)
<i>Trichoderma harzianum</i>	Clarithromycin	Biodegradation	Buchicchio et al., (2016)
<i>Trametes versicolor</i>	<i>Cefalexin, Ciprofloxacin,</i> <i>Etracycline</i>	Biodegradation	Badia-Fabregat et al., (2016)
<i>Irpex lacteus</i> ^b , <i>Trametes</i> <i>versicolor</i>	<i>Ciprofloxacin,</i> <i>Norfloxacin, Ofloxacin</i>	Biodegradation	Čvančarová et al., (2015)
<i>Trametes versicolor</i>	<i>Ofloxacin</i>	Biodegradation	Gros et al., (2014)
<i>Trametes versicolor</i>	<i>Erythromycin,</i> <i>Ciprofloxacin,</i> <i>Azithromycin, Cefalexine</i>	Biodegradation	Cruz-Morató et al., (2013)
<i>Trametes versicolor</i>	<i>Norfloxacin</i> <i>Ciprofloxacin</i>	Biodegradation	Prieto et al., (2011)
<i>Cunninghamella elegans</i>	<i>Flumequine</i>	Biotransformation	Williams et al., (2007)

Fungal name	Antibiotic contamination	Mechanism	Reference
<i>Trichoderma viride</i>	<i>Ciprofloxacin</i>	Identification of degraded products	Parshikov et al., (2002)
	<i>Norfloxacin</i>		
<i>Pestalotiopsis guepini</i>	<i>Ciprofloxacin</i>	Biotransformation	Parshikov et al., (2001)
	<i>Norfloxacin</i>		
<i>Mucor ramannianus</i>	<i>Enrofloxacin</i>	Biotransformation	Parshikov et al., (2000)
<i>Gloeophyllum striatum</i>	<i>Ciprofloxacin</i> ,	Biodegradation and metabolite identification	Wetzstein et al., (1997, 1999)
	<i>Enrofloxacin</i>		

176

177 2. Materials and Methods

178 2.1. Methodology

179 This investigation proposes a new smart framework for the bioremediation of AZ in soil,
180 which includes lab-scale tests and a prediction system. The study highlights the need for
181 sustainable and nature-based approaches to manage Emerging pollutants (EPs) such as AZ. The
182 proposed framework involves setting up a lab-scale bioremediation system and optimising control
183 factors and designing a prediction system for AZ bioremediation. The study also presents a flow
184 cycle of AZ in the environment as illustrated in [Fig. 2](#) that emphasises the carcinogenic effects of
185 EPs on human health, highlighting the urgency of managing them with sustainable and nature-
186 based approaches ([Abubakr et al., 2020](#)). The study emphasises the importance of smart,
187 sustainable systems for environmental purification and highlights the use of bioremediation
188 techniques with a decision support system as a no-chemical and sustainable method for the
189 decontamination of soil from AZ. The soil samples were collected in the Industrial Centre of

190 Mashhad, Iran for the purpose of investigating the soil properties and preparing the experimental
191 setup.



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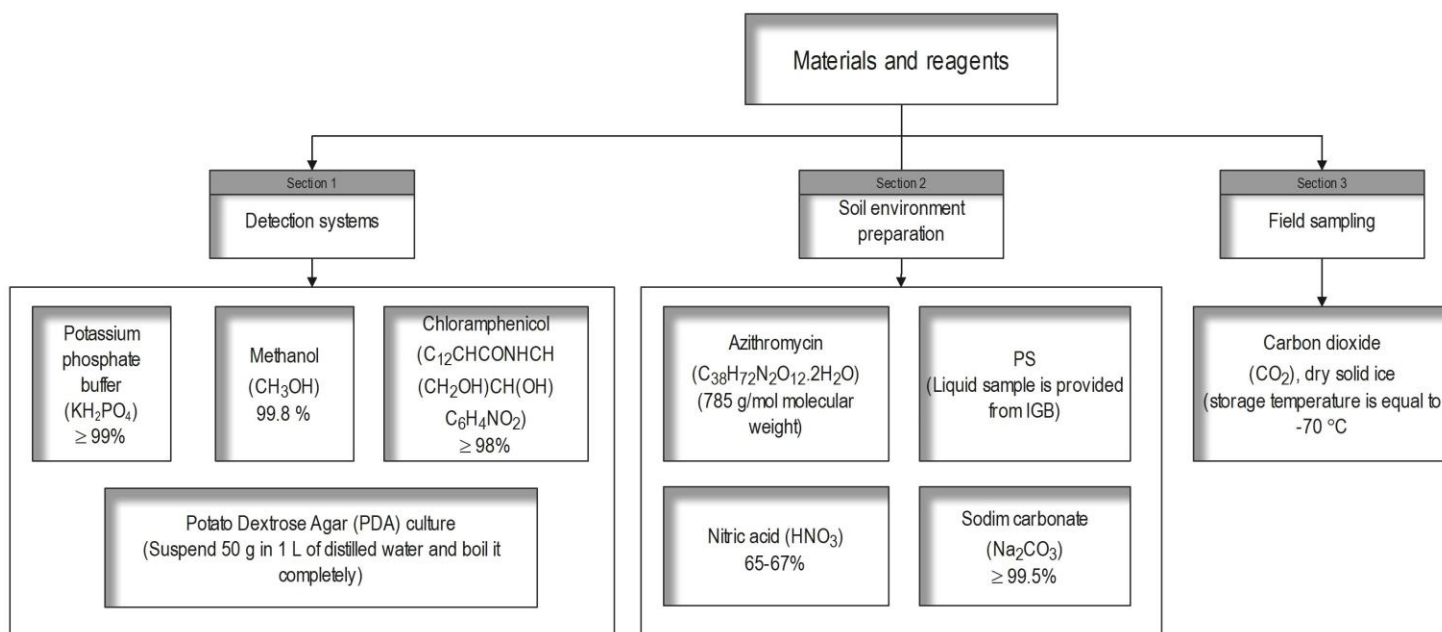
193 **Fig. 2. Schematic plan of sustainable antibiotic bioremediation cycle in the study**
194 ***PS: Penicillium Simplicissimum***

195

196 2.2. Experimental techniques

197 **Fig. 3** provides a list of materials and reagents used in the experiments, including
198 measurement procedures, soil evaluation, and field practices. The PS seed was obtained from the
199 Iranian Genetic Bank (IGB), while all other materials were purchased from Merck, Darmstadt,
200 Germany. Distilled water was used throughout the experiments, and the soil sample was collected
201 from the Quchan road region in Mashhad, Iran, where a pharmaceutical complex is located. The
202 purpose of the experiments was to mitigate the adverse effects of polluted effluent on the
203 surrounding soil. As the AZ concentration was synthesised in the laboratory, all soil samples were
204 planted deep to ensure the absence of AZ.

205 The study collected soil samples from three points in a polluted district for lab-scale
 206 experiments and field performance assessment. AZ solutions were prepared using a 1000 mg/g
 207 stock standard solution from Merck, Germany, and the concentrations of AZ in the samples were
 208 measured at five points (12, 9, 3, 7, and 1.5 mg/g) with a mean amount of 6.5 mg/g, representing
 209 the simulated concentrations of the real accumulated pollution condition in the field.



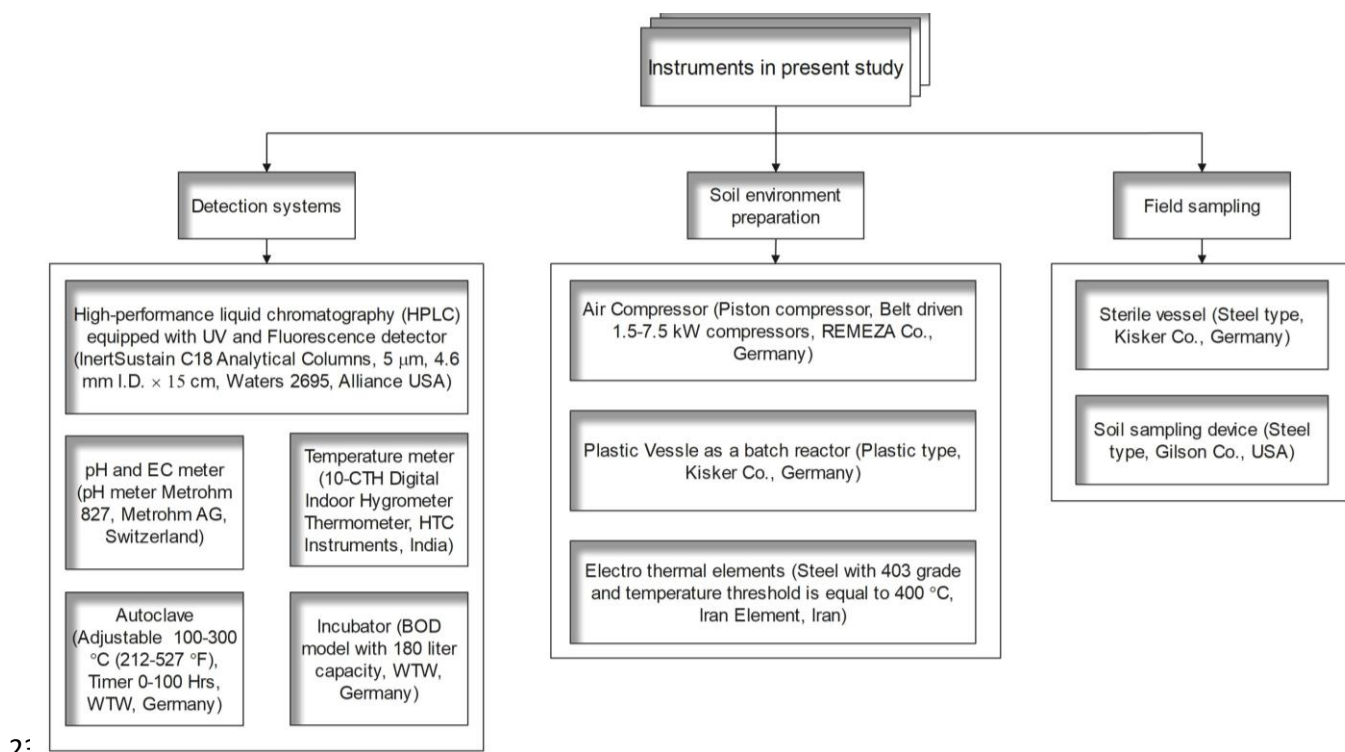
211 **Fig. 3. Materials and reagents used in the experiments.**

212

213 The detection system used in the experiment includes several instruments from different
 214 manufacturers, such as the Waters Alliance 2695 HPLC from the USA, a pH and EC meter from
 215 Switzerland, an autoclave, and an incubator by WTW from Germany, and a temperature meter by
 216 HTC Instrument from India. Other equipment used includes a belt-driven air compressor 1.5-7.5
 217 kW made by REMEZA Co. from Germany, a plastic batch reactor by Kisker Co. from Germany,
 218 steel electrothermal elements by Element Co. from Iran, a sterile steel vessel by Kisker Co., from
 219 Germany, and a steel soil sampling device by Gilson Co. from USA. The study's controlling system

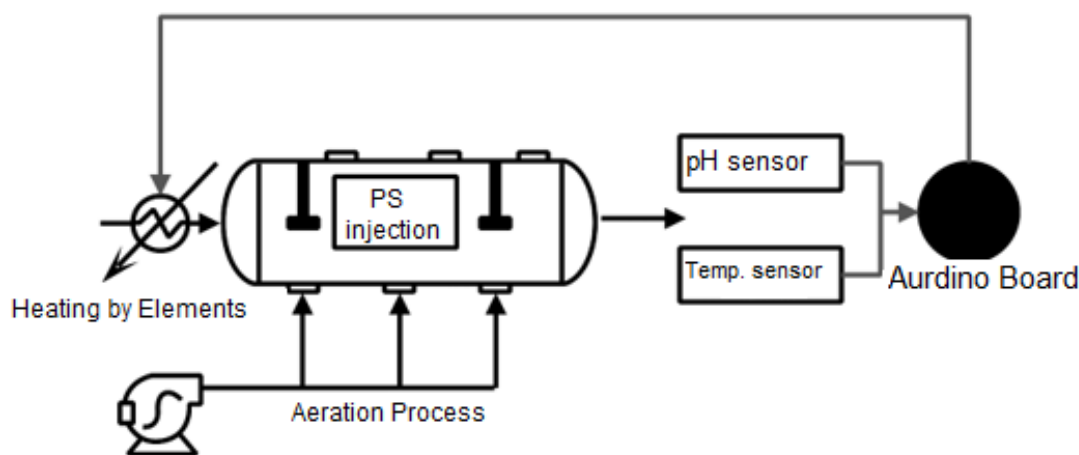
220 uses Arduino hardware and adjusts temperature by coordinating thermal sensors and elements. The
 221 mobile phase for AZ measurement is adjusted with a concentration on isocratic flow, and the AZ
 222 detector is set on UV (210 nm) and fluorescence with corresponding emission (435nm) and
 223 excitation wavelengths (365nm). Fig. 4 elaborates on the information (including model numbers)
 224 of the instruments used to detect AZ, preparation of the soil medium, and samples from different
 225 points of the soil (Hussain et al., 2021).

226 Selective isolates of PS in Petri dishes with three repetitions for each sample containing
 227 Potato Dextrose Agar (PDA) were placed in a growth chamber at 28°C and under light cycle
 228 conditions (12:12) for two weeks for insemination and transfer. After inoculation, spore
 229 suspension using distilled water sterile containing 0.1% Tween 80 was prepared. Purification of
 230 isolates was done by single spore method on a water-agar culture medium (WA) (Babaahmadi et
 231 al., 2018).



232 **Fig. 4. Instruments and their specifications used in the experiments.**

234 Fig. 5 provides a diagram of the key elements and sensors utilised in the laboratory-scale
235 experiments, which were based on bioreactor design principles and environmental regulations. The
236 lab-scale setup consists of two parts: online and offline systems. The offline system includes a
237 fungal seedling, a pH meter (Metrohm 827, Metrohm AG, Switzerland), and manually adjusted
238 air pressure (using piston compressor, Belt driven 1.5-7.5 kW, REMEZA Co., Germany). The soil
239 layers used in the experiment are uniform and have the same characteristics. Heating is controlled
240 by solid elements and the temperature is monitored using an Arduino controller. The experiment
241 package size 10×15×30 cm with a soil mass of 6.750 kg.



242

243

Fig. 5. Schematic layout of the lab-scale set up in the study.

244

245 This study involves measuring AZ and culturing PS. The stages for the determination of
246 AZ and PS culturing are described by Mohammadi et al. (2021). To measure AZ, the HPLC
247 instrument (InterSustain C1 Analytical Columns, 5µm, 4.6mm I.D. × 15cm, Waters 2695,
248 Alliance, USA) must be adjusted with specific settings for the detector wavelengths, oven column
249 temperature, and flow rate equal to 210 nm, 40 °C, and 0.8 mL/min, respectively and adding 50
250 mm Methanol-phosphate buffer 0.02 M (90:10, v/v).

251 To reach pH 8, phosphate acid was used, and the injection volume was set to 50 μ L. A
252 calibration curve was plotted with 5 data points resulting in an R^2 greater than 0.96. By 1 mg/g
253 spiked concentration, the value of AZ is equal to 0.005 mg/g (the limit of AZ is equal to
254 0.005mg/g). To culture the PS for seeding into the bio-engine, several steps are required. These
255 include obtaining PS seeds from the IGB, creating a suspension of initial seeds with sterilised
256 water under laboratory conditions, mixing the provided suspension with WA culturing
257 environment in a sterilised hood, transferring the created seedlings to normal laboratory conditions
258 after 12-18 hours, and charging the seedlings onto PDA Petri dishes with suitable slops for
259 complete growth in the incubator for 4-5 days. Finally, the cultivated PS can be stored in the
260 laboratory's refrigerator at 4°C for future experimental tests.

261 **2.3. Optimisation and numerical models**

262 **2.3.1. Taguchi design**

263 Taguchi is a useful tool for designing complex systems and finding the best set of designs
264 for quality parameters, despite noise factors. Experiments are conducted using an experimental
265 matrix known as the orthogonal array, and quality loss values are calculated for each quality
266 characteristic. The quality loss function is categorised into three types, and the values are
267 transformed into a signal-to-noise (S/N) ratio that shows the dispersion around the desired value.
268 The larger S/N ratio indicates better quality, and it is calculated using [Eq. 1](#) as:

$$269 \quad S / N = -10 \log \left(\frac{1}{n} \sum_{i=1}^n \frac{1}{y_i^2} \right) \quad \text{Eq. 1}$$

270 where $y_i = i^{th}$ observed response value, n = the number of observations in a trial. The S/N ratio is a
271 measure of the effect of control factor levels on the response quality. A higher S/N ratio corresponds to

272 better performance of the response. Therefore, the optimal levels of parameters can be obtained by
273 identifying the levels that result in the highest *S/N* ratios.

274 The Taguchi model is used to design experiments by determining parameter characteristics,
275 defining levels, designing the orthogonal matrix, and inspecting results by desirability and *S/N*
276 ratio (Koilaraj et al., 2012). This study's experimental parameters include Temperature (*T*) (°C),
277 Retention Time (*RT*) (day), pH, Aeration Intensity (*AI*) (m³/h), and Microorganism to Food weight
278 ratio (*M/F*) (mg/g) as influential parameters according to the literature. The Taguchi design method
279 in MiniTab16 software is used to obtain the most efficient range of parameters, and the primary
280 levels are set in Table 4. All experiments are initially conducted for 20 days, and AZ concentration
281 is measured in both influent and effluent every five days from day five to day 100. The PS used in
282 the bio-engine for AZ decontamination is fed by local agricultural waste from the municipal waste
283 centre in Mashhad.

284

285

Table 4. The primary data of Taguchi design for being optimised.

Parameter	1	2	3
Temperature (°C)	24	28	32
Retention time (day)	20	40	60
pH	3	6	9
Aeration intensity (m³/h)	8	14	20
Microbial/Food ratio (mg/g)	1	4	7

286

287

288

289 2.3.2. Response Surface Methodology (RSM)

290 RSM is a statistical tool used to predict the interactions between factors that may be
291 difficult to observe or too complex to test experimentally. RSM provides a way to investigate the
292 relationship between variables and a response variable or performance characteristic of a system
293 under control. The relationship between the controlling variables (X_1, X_2, \dots, X_n) and the response
294 (Z) is represented by a function f , as shown in Eq. 2. In this equation, ε represents other sources of
295 variables that are not included in the function f , which is assumed to have a normal distribution
296 with a mean of zero and variance of σ^2 . Therefore, the expected value of the response function is
297 the f function.

$$298 Z = f(X_1, X_2, \dots, X_n) + \varepsilon \quad \text{Eq. 2}$$

299 To simplify the estimation of the complicated f function, a polynomial equation based on
300 experimental data can be used. This polynomial equation can estimate the response of the variables
301 in points where there is insufficient experimental data. A second-order polynomial function,
302 represented by Eq. (3) is commonly used as an estimation of the actual response surface around a
303 desired point. This approach is known as response surface methodology and can help researchers
304 understand the relationship between variables and the response (Sarabia and Ortiz, 2009).

$$305 Z = a_0 + a_1X_1 + a_2X_2 + \dots + a_nX_n + a_{12}X_1X_2 + a_{13}X_1X_3 + \dots + a_{1n}X_1X_n + a_{23}X_2X_3 + a_{24}X_2X_4 \\ 306 + \dots + a_{2n}X_2X_n + \dots + a_{(n-1)n}X_{n-1}X_n + a_{11}X_1^2 + \dots + a_{nn}X_n^2 \quad \text{Eq. 3}$$

307 The ANOVA test is used to determine the significance of the regression (Sarabia and Ortiz
308 2009). Using RSM, researchers can predict the values of culture parameters at points where
309 experimental data is lacking. In this study, the efficiency of the reactor in Fig. 5 is evaluated on a
310 lab-scale setup using optimal conditions, and the resulting data is used in Design Expert 7 software

311 to predict data for other points. These predicted values can be considered as the desired set of
312 control variables based on the operating situation.

313 **2.3.3. Machine learning algorithms**

314 Prediction models based on ML are used in the study to estimate AZ degradation in soil.
315 The performance of ML models can vary depending on the dataset and the specific problem being
316 addressed. Therefore, it's recommended to evaluate multiple algorithms and compare their
317 performance before selecting the most appropriate one for a specific application. As such, several
318 factors were considered for selecting the most effective ML algorithms for this study, including
319 the nature of the problem, available data, and prior experience. After careful consideration, IBK,
320 Kstar, and LWL were selected for their suitability for classification and regression tasks with
321 complex decision boundaries and non-linear relationships. The prediction models of these
322 algorithms are developed using WEKA 3.9 software by training based on 70% of the experimental
323 data and then validating based on the remaining 30% of the data. The main objective is to compare
324 the performance of these methods for application to bioremediation processes. More details of
325 modelling with these algorithms are outlined below.

326 **2.3.3.1 Instance-Based K (IBK) algorithm**

327 The IBK algorithm with the K parameter falls into the category of regression and
328 classification lazy algorithms. The IBK algorithm works by identifying similarities between
329 instances and specifying the number of nearest neighbours to use when classifying a test instance.
330 It can select the most suitable value of K by using cross-validation and distance weighting,
331 (Moayedi et al., 2019). Note that in WEKA software, IBK is based on cross-validation, which
332 helps find the best value for K's nearest neighbour.

333

334 **2.3.3.2 K-Star algorithm**

335 Kstar, another lazy learning algorithm, selects the most relevant features for classification
336 using statistical methods based on the K nearest neighbour method and is suitable for datasets with
337 many features. Unlike other instance-based learners, K-Star attempts to divide n data points into k
338 clusters using an entropic distance measure. This involves computing the probability of
339 transforming one instance into another, which requires measuring the distance between instances.
340 To achieve this, the algorithm determines a finite set of transformations that map an instance into
341 another and transforms an instance using a limited series of transformations starting at point ‘a’
342 and ending at ‘b’. The K-Star computation is as follows:

343
$$K(y_i, n) = -\ln \hat{P}(y_i, n) \tag{Eq. 4}$$

344 where n = new data points attached to the most expected class y_i and P' = the probability of the
345 point i reaching point j through a random path.

346 **2.3.3.2 Locally Weighted Learning algorithm**

347 LWL, a non-parametric regression algorithm, assigns weights to training instances based
348 on their distance from the query point and is suitable for regression tasks with non-linear
349 relationships. prediction in LWL is based on local functions using a subset of data to replace a
350 global function that result in faster predictions. More specifically, a local model is created for each
351 point of interest based on the neighbouring data of the query point instead of a global model for
352 the entire dataset. To satisfy this, each data point becomes a weighting factor that represents its
353 influence on the prediction. This means that the closer a point is to the query point, the more weight
354 it receives, making this method very accurate and allowing new training points to be added easily.
355 If there is a continuous function f with noise ε , then the LWL cost function is as follows:

356
$$y = f(x) + \varepsilon \tag{Eq. 5}$$

$$G = \frac{1}{2} \sum w_i x_q (y_i - x_i \beta_q)^2 \quad \text{Eq. 6}$$

where x_q is the point of interest (or query point) which is the point where we want the prediction y_q . Labelled training data $D = \{(x_i, y_i) | i = 1, 2, \dots, n\}$ where each data point of x_i belongs to a corresponding y_i . w_i represents the weight of the corresponding set of (x_i, y_i) for the current prediction which is computed through a weighting function. β describes the regression coefficient of the linear model.

The algorithm aims to find the β in a way that minimises the G function for the current point of interest x_q . It is one of the main differences of this method to global least functions where β is dependent on the x_q and finding w_i through two these steps: (1) The distance function $d(x_i, x_q)$; $d = \sqrt{(x - q)D(x - q)}$, measures the relevance of the training points in the current prediction. This function receives two inputs and gives one number. D describes the distance metric which is an important parameter expressing the size and shape of the receptive field; and (2) The Kernel function $K(d)$; $K(d) = \exp(-d^2)$ gives out a weight for each distance value.

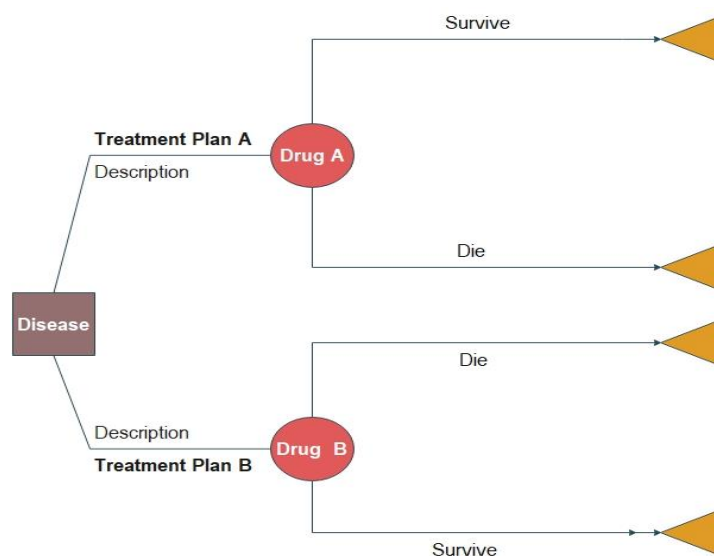
These three algorithms (IBK, Kstar, and LWL) are also specifically suitable for the AZ decontamination process due to a range of factors that are difficult to capture in a simple model. For example, the process may have many factors influencing the performance of fungi for which K-star is a suitable algorithm while other algorithms such as SVM and RF may be inappropriate for problems with complex decision boundaries or many input features.

2.3.4. Decision Tree

After identifying the optimal conditions for the experiments, a decision tree (DT) is created to develop a smart control system (Amini et al., 2021; Gheibi et al., 2019). In this context, the DT

378 serves as a dashboard for the decision support system. DTs are a simple modelling technique that
379 represents a sequence of interventions over a period as a graphical tree-like structure. The tree has
380 branches and leaves representing options between alternatives and outcomes, respectively. It
381 consists of three parts (Fig. 6): the root node, which is the starting point of the tree; branches,
382 which represent possible answers; and leaf nodes, where each branch ends and are shown in three
383 types: decision nodes, chance nodes, and terminal nodes.

384 The DT structure produces a series of rules that explain the path from the root to a leaf of
385 the DT. Each path represents a rule, and the leaf is labelled with the class in which the correct
386 value of records is assigned. DTs have limitations in modelling decision problems but provide
387 several advantages, such as visual aid that requires no further explanation and is easy to
388 understand. They can also cover both quantitative and qualitative data and consider data sets that
389 may contain errors or missing values. DTs are also convenient to draw and free of complicated
390 computations.



391

392

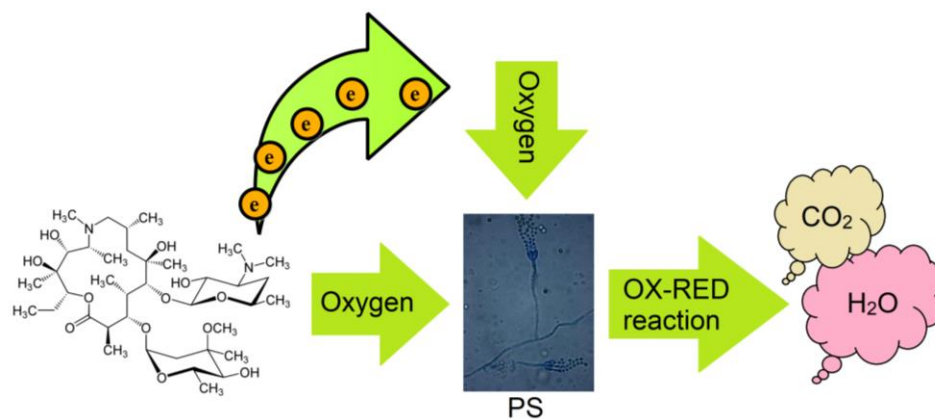
Fig. 6. A simple example of a decision tree

393

394 3. Results and discussion

395 3.1. Mechanism of the bioremediation process

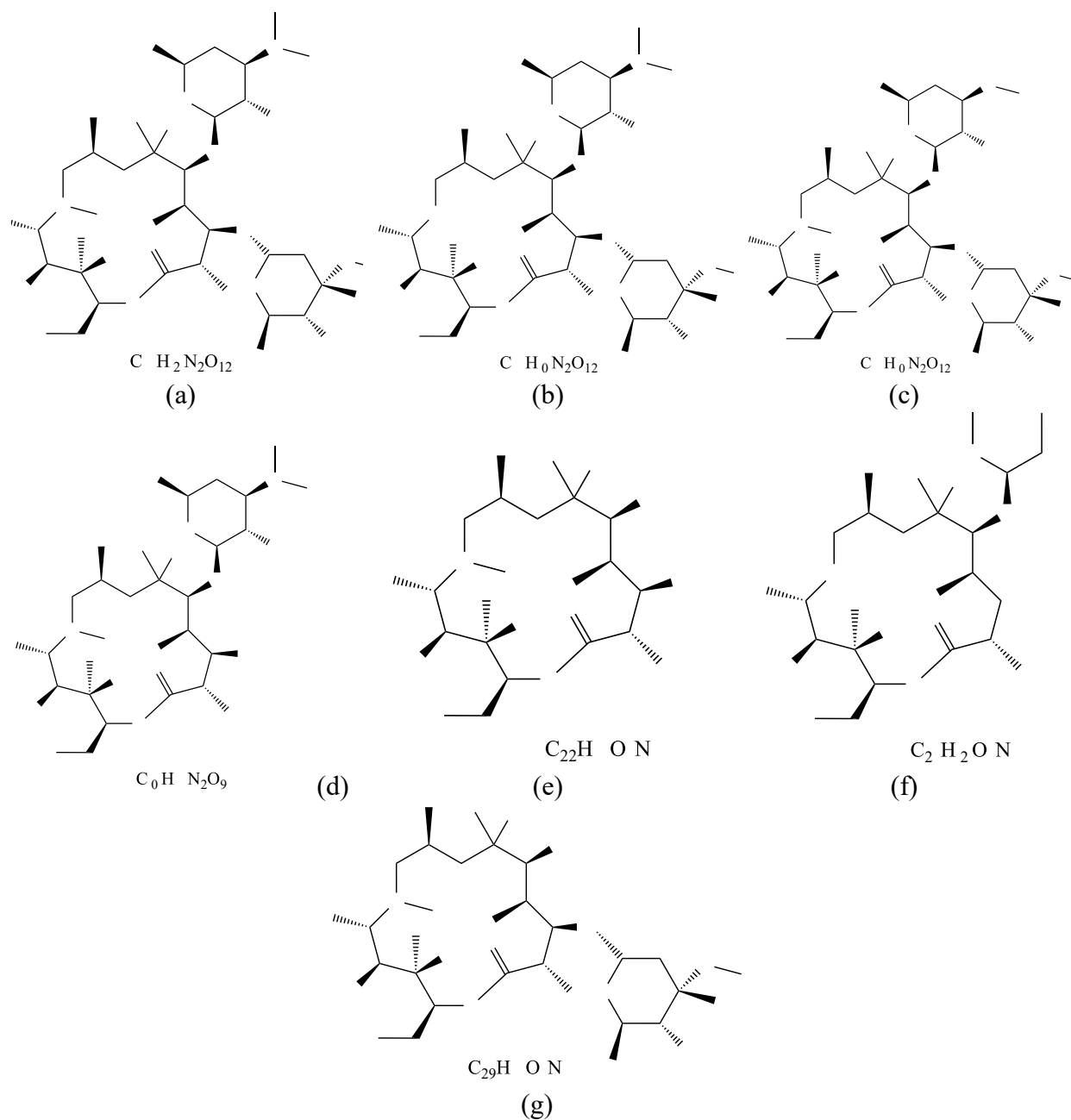
396 Fig. 7 illustrates the biodegradation mechanism of AZ through the *Penicillium*
397 *Simplicissimum* (PS) bioremediation process and other fundamental concepts. The PS fungal cell
398 walls are negatively charged due to the presence of various functional groups, such as carboxylic,
399 phosphate, amine, or sulfhydryl, in different wall components such as hemicelluloses, pectin, and
400 lignin (Fomina et al., 2007). The decomposition process is carried out through the transfer of
401 electrons between O₂ and organic matter by fungus activities (Bell et al., 2011). The PS
402 decomposes AZ compounds using hydrolase and free radical enzymes via the oxidation-reduction
403 process (Dias et al., 2021).



404
405 **Fig. 7. The mechanism of azithromycin biodegradation by fungus activities in this study**
406

407 The degradation process of AZ involves a reduction half-reaction and an oxidation half-
408 reaction catalysed by enzymes secreted by PS. The reduction half-reaction involves the transfer of
409 a hydride from the substrate to the reactant, resulting in a binary complex between the two-electron
410 reduced enzyme and the p-quinone methide of the substrate. In the oxidation half-reaction, the
411 reduced AZ is oxidised by molecular oxygen with the concomitant hydration of the quinone
412 methide intermediate. This process continues until CO₂ and H₂O are the final products of the

413 reaction, which is called mineralisation. The main compounds generated during the degradation
 414 process of AZ are safe intermediate materials containing CO₂ and H₂O (Deblonde et al., 2011)
 415 while some degradation products are also potential to be created under aerobic conditions (Fig. 8).
 416



417 **Fig. 8. (a) Azithromycin and the potential degradation products, (b) 9a-N-desmethyl azithromycin,**
 418 **(c) Desclandose azithromycin, (d, e, f) resulted from the removal of some other groups from azithromycin, (g)**
 419 **N-desmethyl azithromycin**

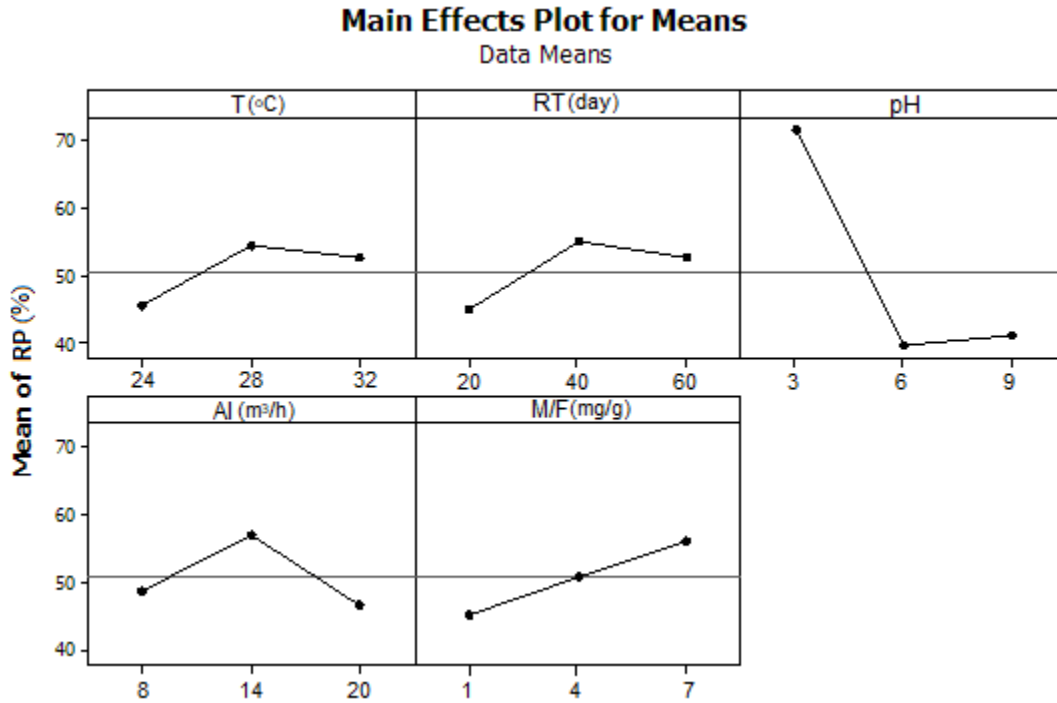
420

421 The bioremediation of AZ by PS involves the formation of a binary complex between the
422 enzyme and the substrate, which could be an important step in breaking down the antibiotic into
423 less harmful forms (Fleming, 1946). Enzymes play a critical role in bioremediation, and PS may
424 use them to break down AZ which is an antibiotic that can accumulate in the environment and
425 potentially lead to negative effects on ecosystems (Clarke, 2015). Therefore, the input of the clean
426 technology was AZ as a hazardous material in nature, and the outputs were CO₂ and H₂O as safe
427 materials for the environment. Hence, the bioremediation process of AZ by PS can be assumed as
428 a clean and green technology for soil protection.

429 **3.2. Optimisation and mathematical modelling**

430 **3.3.2. Taguchi design analysis**

431 The Taguchi model was used to determine the optimal parameters for bioremediation of
432 AZ by PS. The result of the experiments designed by Taguchi method as illustrated in Fig. 9 and
433 Table 5 shows that the best temperature range was 28°C, and lower temperatures resulted in a
434 steeper reduction in RP due to the increase in the hydrolysis rate of AZ. Retention time also
435 affected RP, with a RT of 40 days resulting in the highest RP. The pH of the soil was critical for
436 both antibiotic stability and fungus activity, with an acidic environment being better for removing
437 AZ from the soil. Aeration intensity performed optimally at around 14 m³/h, while the degradation
438 rate improved as the M/F ratio increased.



439

440 **Fig 9. Results of Taguchi design optimisation for Temperature (T), Retention time (RT), pH, Aeration**
 441 **intensity (AI) and Microbial/Food (M/F) ratio**

442

443 The Taguchi Orthogonal Array approach is designed to isolate the effects of selected
 444 variables, and adding extraneous variables outside the orthogonal array can introduce confounding
 445 effects that bias the results. Careful selection of variables is important, and if additional variables
 446 are included, a larger orthogonal array can be used for comparison. In such cases, a comparison
 447 can be made between the original orthogonal array and the expanded orthogonal array to assess
 448 the impact of the additional variables on the prediction model. However, adding variables outside
 449 the orthogonal array can make the process more complex and may not lead to better results. In this
 450 study, important features were extracted from the literature review (Mohammadi et al., 2021), and
 451 the optimisation process is done as per the operational optimum condition.

452

Table 5. Results of experimental trials based on the Taguchi design method.

T (°C)	RT (day)	pH	AI (m ³ /h)	M/F (mg/g)	Performance RP (%)
24	20	3	8	1	48
24	20	3	8	4	57
24	20	3	8	7	69
24	40	6	14	1	38
24	40	6	14	4	45
24	40	6	14	7	51
24	60	9	20	1	27
24	60	9	20	4	34
24	60	9	20	7	39
28	20	6	20	1	31
28	20	6	20	4	32
28	20	6	20	7	36
28	40	9	8	1	42
28	40	9	8	4	48
28	40	9	8	7	50
28	60	3	14	1	76
28	60	3	14	4	84
28	60	3	14	7	89
32	20	9	14	1	40
32	20	9	14	4	44
32	20	9	14	7	45
32	40	3	20	1	66
32	40	3	20	4	73
32	40	3	20	7	81
32	60	6	8	1	38
32	60	6	8	4	41
32	60	6	8	7	45

455 3.3.3. Response surface methodology

456 The RSM used the historical data analysis to determine the maximum degradation rate of
457 AZ and the impact of various cultural conditions on the rate of degradation. As such, the Design-
458 Expert 7.0.0 software was used to predict the response of AZ degradation rate to cultural conditions
459 including temperature, retention time, aeration intensity, pH, and M/F ratio. Therefore, a quadratic
460 polynomial equation was obtained based on the above parameters to predict the response of the

461 AZ degradation rate. This equation considers the interactions between the different cultural
462 conditions to provide a more accurate prediction of the degradation rate as follows:

$$463 \quad P = 54.70 + 4.08A - 2.39B - 31.83C - 23.94D + 5.5E - 33.22AB - 12.61AC - 1.58AE + IBC + \\ 464 \quad 0.083BE - 2CE - 0.33DE - 5.81A^2 - 0.22E^2 \quad \text{Eq. 7}$$

465 Where P = the predicted value of AZ removal percentage and A , B , C , D , and E = coded factors of
466 temperature, retention time, pH, aeration intensity, and M/F ratio, respectively. The statistical data
467 of the quadratic polynomial equation indicates this is a significant equation ($p < 0.0001$ and $R^2 =$
468 0.9887). The regression data also demonstrate that temperature (A), pH (C), aeration intensity (D),
469 M/F ratio (E), and the square term of temperature (A^2) were significant ($p < 0.0002$), whereas the
470 square term of M/F and retention time (B) were insignificant. It is also evident that pH with the
471 $p < 0.0001$ and the highest F-value is the most influential factor followed by aeration intensity.
472 [Table 6](#) also shows the result of the ANOVA analysis including interactions and the squared
473 effects.

474 In other words, [Eq. 7](#) represents a mathematical formula that can be used to predict the
475 removal percentage (RP) of AZ in a bioremediation process using PS, based on certain operational
476 factors. These factors include the initial concentration of AZ, the size of the fungi inoculum, and
477 the duration of the process. This equation can be particularly useful for optimising bioremediation
478 processes for AZ or similar pollutants. By incorporating the variables described in the equation, it
479 can help predict the removal percentage of AZ without having to conduct time-consuming and
480 expensive experimental trials. Moreover, this equation can be applied in various settings beyond
481 the bioremediation process discussed in this study. It can be adapted to fit different experimental
482 designs or operational factors, or it can be used as a foundation for developing more sophisticated
483 models of bioremediation processes.

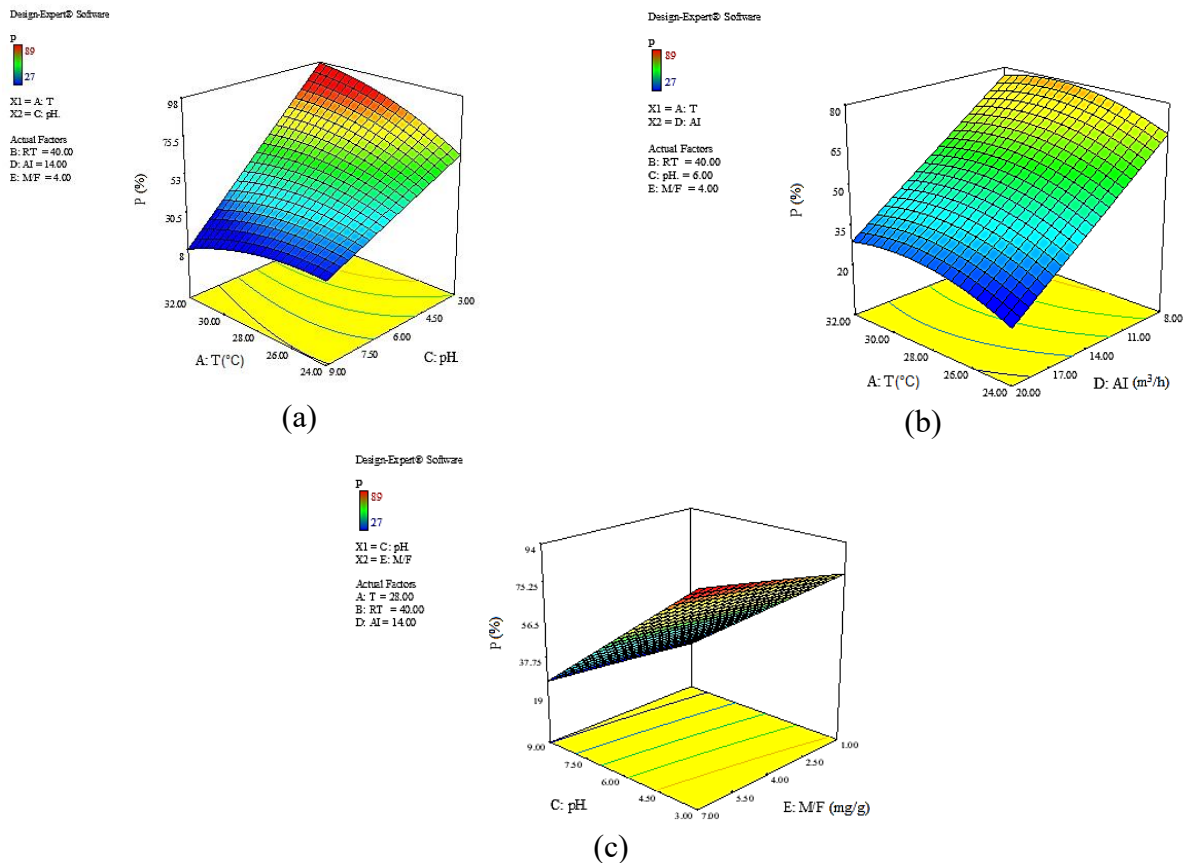
Table 6. Results of ANOVA analysis in this study

Source	<i>Sum of Squares</i>	<i>Mean Square</i>	<i>F-Value</i>	<i>P-value Prob > F</i>
Model	7904.03	564.57	163.10	< 0.0001
Temperature (T)	92.34	92.34	26.67	0.0002
Retention time (RT)	25.68	25.68	7.42	0.0185
pH	4560.12	4560.12	1317.41	< 0.0001
Aeration intensity (AI)	1474.29	1474.29	425.92	< 0.0001
Microbial/Food ratio (M/F)	544.50	544.5	157.30	< 0.0001
T × RT	1655.57	1655.57	478.29	< 0.0001
T × pH	238.56	238.56	68.92	< 0.0001
T × AI	0.00			
T × M/F	30.08	30.08	8.69	0.0122
RT × pH	2.00	2.00	0.58	0.4619
RT × AI	0.00			
RT × M/F	0.08	0.08	0.02	0.8793
pH × AI	0.00			
pH × M/F	48.00	48.00	13.87	0.0029
AI × M/F	1.33	1.33	0.39	0.5464
T²	115.56	115.56	33.38	< 0.0001
(RT)²	0.00			
(pH)²	0.00			
(AI)²	0.00			
(M/F)²	0.30	0.30	0.09	0.7748

486 [Fig. 10](#) depicts the results of some statistical tests according to the ANOVA computation

487 as variance analysis for effective factors used in the prediction model. The sensitivity analysis

488 reveals that the maximum efficiency occurs at a pH of approximately 3 when the retention time
 489 (RT), aeration intensity (AI), and M/F ratio are fixed. The plots show that the slope of pH and AI
 490 are higher than the other parameters, indicating that the removal percentage (RP) is more sensitive
 491 to changes in these factors. These results are consistent with the outcome predicted by the RSM
 492 analysis. Table 7 provides some of the data resulting from the prediction model by the RSM
 493 analysis. Based on Fig. 10, it is evident that the aeration volume and pH control should receive
 494 constant attention during the operation of the system, and the operator should adjust them to
 495 optimal conditions with high precision under different conditions.



496 **Fig 10. The dual sensitive analysis of AZ bioremediation factors in this study a) Temperature (T) against pH,**
 497 **b) T against Aeration Intensity (AI) c) pH against Microbial/Food (M/F) ratio**

498
 499

500

Table 7. The optimisation results obtained from the RSM analysis.

No.	T (°C)	RT (day)	pH	AI (m ³ /h)	M/F (mg/g)	RP (%)
1	29.44	25.88	3.03	13.64	4.63	100
2	31.53	25.83	4.02	14.47	2.13	99.79
3	28.98	45.1	3.42	8.73	1.74	97.96
4	30.95	41.78	3.1	13.76	5.91	96.67
5	31.1	28.21	5.52	8.55	1.59	95.39
6	31.98	25.71	6.39	8.04	1.09	92.28

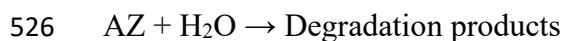
501

502 The data in [Table 7](#) indicates that the best performance for AZ removal is achieved at a pH
503 lower than 4. However, it is also possible to achieve high efficiency at higher pH levels, which are
504 more practical to achieve in real-world settings due to the complexity of the test for reaching the
505 pH of 3. Studies on AZ biodegradation in soil environments are limited, but [Li et al., \(2012\)](#) found
506 that microbiomes in soil are most efficient at pH levels between 6.5-8.5 and temperatures between
507 25-45 °C. They also noted that the optimum pH for tetracycline degradation was around 6.5. [Ding](#)
508 [et al., \(2016\)](#) investigated the simultaneous removal of 14 different antibiotics from soil using
509 laccase oxidation and soil adsorption processes. They achieved over 70% efficiency in the first 15
510 minutes at pH 6 and a temperature of approximately 25 °C.

511 The optimum temperature range for AZ bioremediation under fungal activity is between
512 28 and 32 °C, which is consistent with expectations from Taguchi modelling. Retention time (RT)
513 does not significantly affect the optimum cultural conditions and is effective for periods of more
514 than 25 days. Anaerobic degradation of AZ is ineffective, according to [Vermillion and Tjeerdema](#)
515 [\(2017\)](#), who found that biotic degradation in aerobic environments leads to higher levels of AZ
516 removal.

517 Based on the research findings, the suggested operating conditions for AZ bioremediation
518 under fungal activity are outlined in the fifth row of [Table 7](#). While the removal efficiency may
519 not be the highest, it is at a desirable level, and the recommended cultural conditions, such as pH,
520 are more practical to achieve. The pH level is closer to the natural pH of the soil, and AI is at a
521 low level, reducing power usage and associated costs including the energy cost and
522 depreciation/operating costs. The low M/F ratio means that cheap and abundant agricultural waste
523 can be used as a food source, reducing waste and costs ([El-Ramady et al., 2022](#)). Finally, the low
524 RT range reduces waiting time and allows for early response of the system.

525 The degradation of AZ by fungi in the soil can be represented by the following:



527 It should be mentioned that the moisture percentage of the examined soil in this study is kept
528 between 30%-45% which is enough for completing the reactions. The range of moisture
529 percentage is related to the type of agriculture and land curing process in the case study. To
530 accurately evaluate the effect of increasing the concentration of AZ on the rate of degradation, a
531 comprehensive experimental study should be conducted that considers all the relevant factors and
532 monitors the degradation process over time. This would involve measuring the concentration of
533 AZ and its degradation products at different time intervals and under different operational
534 conditions. The results of such a study could be used to optimise the bioremediation process and
535 develop more efficient strategies for treating contaminated soils.

536 If the reaction is first order concerning AZ, the rate of the reaction is given by $rate = k[AZ]$,
537 where k is the rate constant, and $[AZ]$ is the concentration of AZ. If the concentration of AZ is
538 increased, the rate of the reaction will also increase proportionally, because the new $rate = k[2AZ]$
539 $= 2k[AZ] = 2(old\ rate)$. This shows that the reaction rate is directly proportional to the

540 concentration of AZ when the reaction is first ordered with respect to AZ. If the reaction is second
541 order concerning water, the rate of the reaction is given by $rate = k [H_2O]^2$. If water concentration
542 is increased, the rate of the reaction will increase proportionally to the square of the concentration
543 of water, because the new $rate = k [(2H_2O)^2] = 4k [H_2O]^2 = 4(old\ rate)$. This shows that the
544 reaction rate is proportional to the square of the concentration of water when the reaction is second
545 order concerning water. Changes in the concentration of azithromycin and water can have a
546 significant effect on the rate of degradation of azithromycin by fungi in soil. In this reaction, the
547 rate of degradation is directly proportional to the concentration of azithromycin and proportional
548 to the square of the concentration of water. However, it should be noted that the degradation of
549 AZ by fungi is a complex process that may involve multiple reactions and intermediates.
550 Therefore, the kinetics of the degradation process may not always follow a simple first-order or
551 second-order rate law. The actual rate of degradation may depend on various factors, including the
552 type of fungus used, the initial concentration of AZ, the pH, temperature, moisture content, and
553 the presence of other pollutants or organic matter in the soil. It is important to note that the specific
554 reaction mechanism and the environmental conditions in the soil may also play a role in
555 determining the rate of the reaction. Therefore, changing in concentrations of different elements
556 in the reaction, the rate is changed, and operational features are considered constant approximately
557 because they are related to the origin of the applied fungi and its interactions with AZ.

558 **3.3. Machine learning prediction modelling**

559 The results of the ML models (IBK, KStar, and LWL algorithms) for predicting the
560 bioremediation of AZ under PS degradation are presented in [Table 8](#). All the ML models have
561 correlation coefficients above the acceptance range. However, the IBK with a correlation
562 coefficient of 0.95 outperforms the other two models with high accuracy and more confidence.
563 Furthermore, a correlation coefficient of 0.94 has been achieved through the KStar model which

564 means a close accuracy to the IBK simulation. This suggests that IBK and KStar algorithms may
 565 be more suitable for predicting the bioremediation of AZ under PS degradation, while LWL may
 566 not be as accurate with a correlation coefficient of 0.89. The value of these prediction models is
 567 better understood by looking at the complexity, cost, and time of conducting the experiments.
 568 However, it is important to conduct further testing and validation to confirm the reliability and
 569 generalisability of these models as well as considering other factors such as interpretability,
 570 scalability, and computational efficiency when selecting an ML algorithm for a specific
 571 application.

572 **Table 8. The outcomes of prediction models through IBK, KStar, and LWL algorithms**

Parameters of ML algorithm	IBK	KStar	LWL
<i>Correlation coefficient</i>	0.95	0.94	0.89
<i>Mean absolute error</i>	4.07	4.45	6.28
<i>Root mean squared error</i>	5.099	5.55	8.04
<i>Relative absolute error (%)</i>	28.23	30.83	43.50
<i>Root relative squared error (%)</i>	28.85	31.40	45.46

573

574 Several studies have used various ML algorithms to predict the behaviour of bio-engine
 575 systems. For example, [Mohammadi et al., \(2021\)](#) used RF, ANFIS, and RT algorithms to predict
 576 amoxicillin removal efficiency from soil, achieving correlation coefficients of 0.97, 0.95, and 0.99,
 577 respectively. [Amiri et al., \(2022\)](#) employed an M5 tree model to predict AZ removal efficiency
 578 from aqueous solutions, achieving a correlation coefficient of 0.946 and RMSE of 9.89%. [Mojiri
 579 et al., \(2020\)](#) used an artificial neural network to predict the removal efficiency of ciprofloxacin in
 580 wastewater, achieving $R^2 > 0.99$. [Zhu et al., \(2021\)](#) investigated the adsorption capacity of
 581 antibiotics on carbon-based adsorbents, finding that random forest-based algorithms performed

582 better than other models, with the specific surface area of adsorbents being highly important.
583 Lastly, Arab et al., (2022) used ANN and ANFIS approaches to predict the experimental data for
584 removing cephalexin from the water and found these models with high accuracy, achieving
585 accuracy of 88.21% and 93.87%, respectively, at a pH of 6.14.

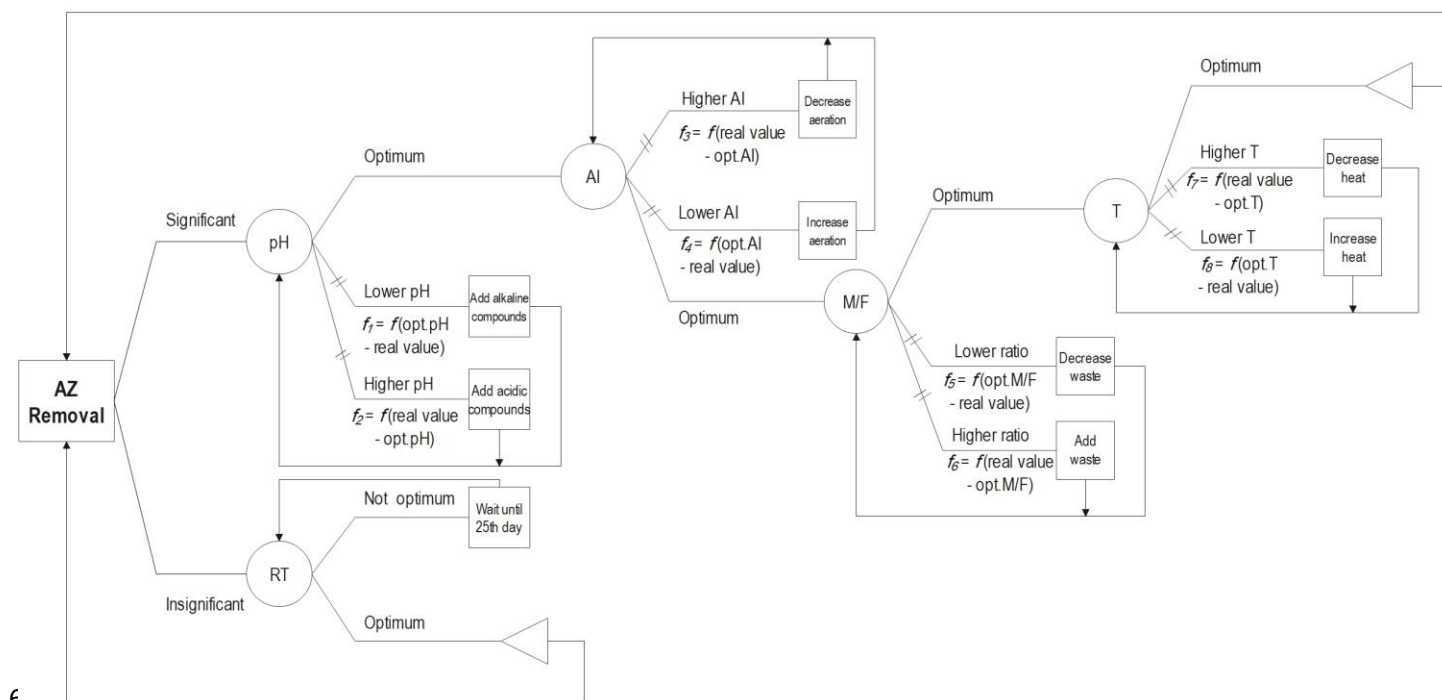
586 The study highlights the benefits of using AI in predicting the performance of a system
587 accurately without conducting costly and time-consuming tests. In other words, if this system
588 undergoes changes in operation for any reason; Using the methods of *AI* and the performance of
589 the operator to adjust the parameters, the soft ML-based system forecasted the removal efficiency
590 of AZ. The ML algorithms used in this study collected data via the results of optimised solutions
591 from Taguchi runs. The size of the data entered in ML modelling is the same as the data derived
592 from Taguchi simulations. However, the ML algorithms may not perform well when used to make
593 predictions outside of that range mainly because the model may not be able to accurately capture
594 the underlying relationships between the variables being modelled when applied to data within the
595 training range. This should be considered as a constraint of *ML* models. Note that as experiments
596 are based on Taguchi's design, the optimal situation is necessarily among the experiments that
597 have taken place.

598 Furthermore, RSM prediction modelling may outperform ML algorithms due to its ability to model
599 complex relationships between variables and response variables, and its capacity to optimise the
600 system being studied. RSM can also provide a clear understanding of the underlying mechanisms
601 driving the system, making it easier to identify the most important variables and interactions. In
602 contrast, ML algorithms may struggle to capture complex interactions and can be opaque, making
603 it difficult to understand why a particular prediction is made.

604

605 3.4. Bioremediation control system

606 The DT modelling in Fig. 11 shows the influential parameters for the bioremediation
 607 process and how they should be controlled. The bioremediation process can be maintained at a
 608 high-efficiency level by adjusting the parameters to their optimised values using the dashboard.
 609 The four most important parameters are M/F, pH, T, and AI, which are adjusted in parallel based
 610 on the ANOVA analysis. Reaction time is then controlled at the optimum value of 25 days,
 611 followed by other effective factors organised in the DT. The DT is based on a linear control system
 612 and can help maintain the process at high efficiency.



614 **Fig 11. The decision tree based on RSM analysis.**

615 The process of AZ bioremediation involves two types of factors: significant and insignificant. The
 616 insignificant factor, which is RT, does not require specific optimisation and only needs to be more than 25
 617 days. On the other hand, the significant factors need to be adjusted accurately. According to ANOVA
 618 analysis, pH is the most influential factor that needs to be monitored consistently. The optimised range for
 619 pH is 5.5, and if the pH is below this level, alkaline compounds should be added to the soil. Conversely,

620 acidic compounds should be added to raise the pH if it is too high. The modification can be calculated
621 through the error functions f_1 and f_2 . Once the optimum pH is achieved, the system moves on to
622 adjust AI. Functions f_3 and f_4 are used to calculate the difference in current aeration intensity and
623 adjust AI to achieve optimum degradation. The condition is being constantly checking and further
624 passes to the next cultural parameter which is the M/F ratio. The same process is repeated for M/F
625 ratio and temperature, and functions f_5 , f_6 , f_7 , and f_8 are used to amend the specified parameters
626 respectively. This whole process is constantly repeated to ensure that the optimum condition is
627 maintained throughout the project.

628 **3.5. Economic evaluation of the system**

629 To implement the bioremediation process for the degradation of AZ in soil in real-world
630 scenarios, it is important to consider its economic evaluation. The process can be assessed using
631 frameworks for smart sustainable operation of systems, although there is no specific framework
632 for the exact economic assessment of biological methods. Scientific knowledge and experiences
633 can be employed to help with the real field operation of the process.

634 The economic evaluation of the bioremediation process for AZ degradation in soil as
635 illustrated in [Fig. 12](#) indicates that operational costs (37%) are more significant than investment
636 costs (63%). Therefore, the bioremediation approach is appropriate for treating antibiotic
637 contamination in soil, with a focus on operational costs. It should be noted that the main challenge
638 of biological decontamination is related to operational costs, and this challenge is addressed by the
639 investigated bioremediation process, as shown by the outcomes of other studies ([Ghadami et al.,](#)
640 [2021](#); [Gheibi et al., 2021](#); [Gheibi et al., 2021](#); [Mirabi et al., 2019](#)). The investment costs for the
641 bioremediation process include a fee for transference, cost of experimental tests, setup preparation,
642 and organisational costs. On the other hand, the operational costs consist of the cost of human
643 resources, sampling practices, energy price, and material consumption.

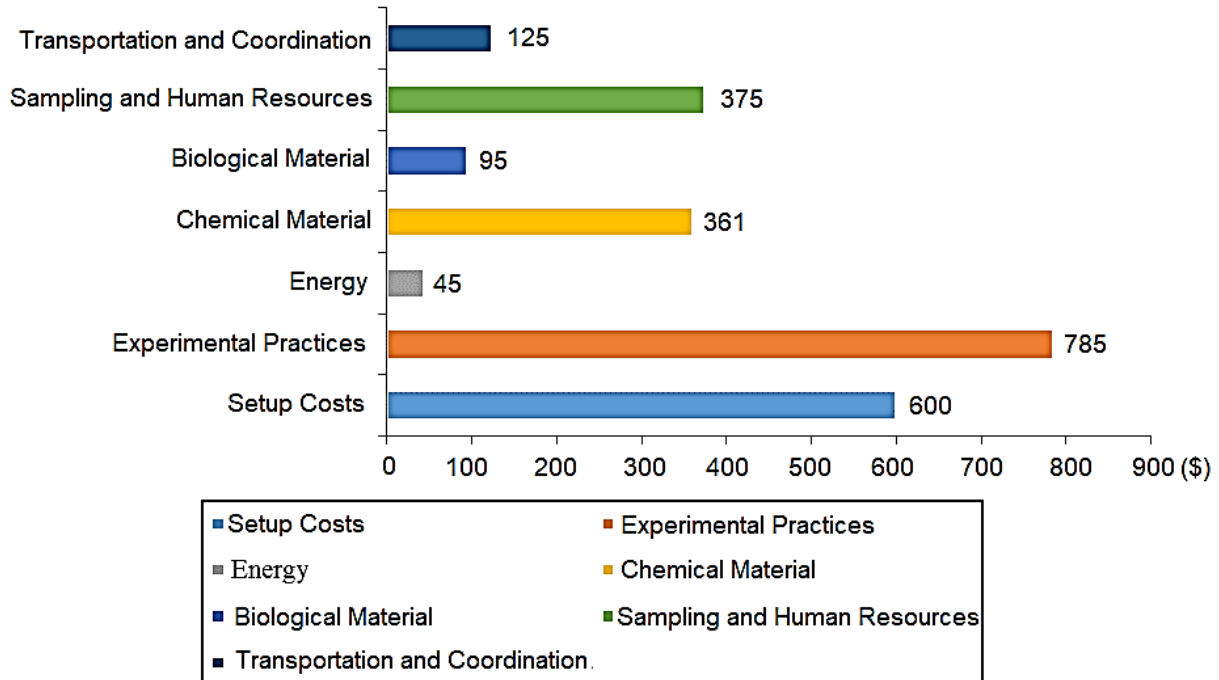


Fig. 12. Cost-effective analysis Economic value in \$ for different stages of the experiment in this study

By using the obtained costs provided for lab scale and assuming a scale-up factor of 10, an estimation of the costs involved in the full-scale implementation of the bioremediation technology can be obtained based on calculations given in the appendix. Based on these calculations, the total estimated cost for full-scale implementation is \$23,865. However, it is important to note that these costs are based on the financial conditions of the case study in Iran, where human resource and energy costs are much lower than in similar cases in Western countries.

According to fundamental equations and models, the bioremediation project is expected to generate an annual profit of \$237,163 and a total revenue of \$250,000 per year. The net present value (NPV) and internal rate of return (IRR) of the project will depend on the discount rate used and the actual costs and revenues incurred during the project's lifetime. It is worth noting that the total annual cost of the project is higher than the annual operational costs alone, which supports the notion that investment costs are higher than operational costs in environmental bioprocesses.

659 Finally, it should be mentioned that a comparison with another research on the
660 decontamination of *amoxicillin* by *Aspergillus Flavus* fungi in the soil environment (Mohammadi
661 et al., 2021) reveals differences in energy values (AI difference), experimental practices
662 (difference in protocols), and chemical materials (difference in roadmap of both studies), which
663 contribute to cost differences.

664 Maier and Tjeerdema, (2018) evaluated AZ biodegradation and sorption by considering
665 the effects of operational parameters. The study conveyed that aerobic bioremediation is the best
666 option between aerobic degradation, anaerobic decomposition, and sorption procedures. However,
667 the bioremediation reaction time in their study is much slower (about 150 days) than PS fungi (at
668 least 25 days) as per the achievements of this study. Another study by Hanamoto and Ogawa
669 (2019) presented a smart system for predicting the sorption of AZ onto organic and inorganic
670 compounds in sediments. They evaluated both ion exchange and adsorption processes. However,
671 the main disadvantage of their achievements is related to regeneration essentially after completing
672 the surface capacity. However, this study continuously conducts antibiotic decontamination with
673 the application of bioremediation.

674 The bioactivities of various strains of *Penicillium* in the process of decontamination are
675 similar, and their characteristics and decontamination mechanisms summarised in Table 9. It is
676 worth noting that all strains can produce enzymes such as proteases and lipases, which break down
677 the complex organic molecules in azithromycin, and they also produce extracellular enzymes that
678 can degrade the molecule. However, the biodegradation mechanism of *Penicillium chrysogenum*
679 differs slightly. Therefore, based on theoretical evaluations, the ML computations, and optimum
680 operational conditions developed in this study, it is possible to apply them to other strains.
681 However, experimental practices and verification are necessary to confirm this hypothesis.

682 **Table 9. The mechanism of azithromycin contamination decomposition by different strains of *Penicillium***

Strain type	Description	Decontamination mechanism	Reference
<i>Penicillium chrysogenum</i>	It is commonly found in soil, and it is applied as a source of penicillin.	Breaking down the β -lactam ring structure of azithromycin by β -lactamases enzyme. It also decomposes molecules by extracellular enzymes, such as proteases and lipases.	Leitão et al., 2007
<i>Penicillium roqueforti</i>	It is used to produce different types of cheese widely.	Breaking down the complex organic molecules in azithromycin	Chang et al., 1996
<i>Penicillium notatum</i>	It is the main source of penicillin and, it is used for the production	Breaking down the complex organic molecules in azithromycin by enzymes	Bujacz et al., 1995
<i>Penicillium camemberti</i>	It is used to produce soft cheeses.	Changing the structure of azithromycin by using proteases and lipases enzymes	Lessard et al., 2014
<i>Penicillium glaucum</i>	It is utilised due to the product's hard cheeses.	Applying proteases and lipases enzymes due to the degraded structure of azithromycin	Hugo, 1991
<i>Penicillium candidum</i>	It is applied to produce blue cheese	Using both proteases and lipases and extracellular enzymes for changing the formulation of azithromycin	Lessard et al., 2014

683

684 **4. Practical applications and prospects**

685 To improve the efficiency of decontaminating antibiotics from soil, the study recommends a

686 combination of adsorption and bioremediation techniques, with bio-adsorption being considered

687 as a potential alternative to conventional adsorption methods. Further research could be conducted

688 to determine the individual contributions of biodegradation, adsorption, aeration, and heating in

689 the decontamination process. Furthermore, studying the kinetics of fungal activity could provide

690 more precise insights into the bioremediation mechanism. Although the study focused on the

691 decontamination of azithromycin by *Penicillium* fungus, future research should also evaluate the

692 competitive effects of co-existing antibiotics such as *amoxicillin* and *azithromycin*, to determine

693 which antibiotic exhibits greater affinity for degradation in soil. In other words, these measures

694 would improve the sustainability of decontaminating antibiotics from soil resources.

695 **5. Conclusions**

696 This study focused on developing a sustainable and eco-friendly approach for removing
697 antibiotics from soil samples using bioactivities, without the need for any hazardous chemicals.
698 The study also explored the potential of ISWM as a novel approach for purifying hazardous
699 materials in the environment. The study found that the M/F ratio was the most significant factor in
700 removing AZ from soil samples and identified the optimal temperature, pH, AI, and M/F ratio for
701 AZ removal. The study found that the IBK model had the highest accuracy in predicting optimal
702 conditions for AZ removal. The study also conducted an economic evaluation of the system and
703 found that 63% of the cost was associated with the investment, while 37% was associated with the
704 operation. The study recommended integrating adsorption and bioremediation techniques for the
705 purification of antibiotics from soil resources, with bio-adsorption being evaluated as an alternative
706 to simple adsorption processes.

707

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