
Predictive Modeling of Grapevine Red Blotch Disease Using Multi-Temporal Remote Sensing and Spatial Epidemiology

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Abstract

1 Grapevine red blotch virus (GRBV) causes significant economic losses in viticulture, necessitating early detection and prediction to mitigate its spread. This
2 study develops a predictive model for 2024 GRBV incidence using multi-temporal
3 remote sensing and spatial epidemiological data collected prior to August 2024.
4 We integrate hyperspectral imaging, spatial autocorrelation metrics, and host sus-
5 ceptibility factors within an automated machine learning framework. Our approach
6 employs iterative feature engineering and addresses class imbalance, achieving a
7 final F1-score of 0.97. Results demonstrate the critical importance of historical
8 infection patterns, neighborhood effects, and vegetation health metrics, aligning
9 with vector-mediated dispersal dynamics. The model highlights both the promise
10 and limitations of remote sensing for pre-symptomatic detection, particularly its
11 reliance on prior-year data. This work contributes an operational, data-driven frame-
12 work for GRBV forecasting, with implications for precision viticulture and broader
13 plant disease management. Future efforts should incorporate vector population
14 dynamics and validate the approach across diverse environments.
15

16 1 Introduction

17 Grapevine red blotch virus (GRBV) poses a significant threat to global viticulture, causing substantial
18 economic losses through reduced fruit quality and yield Cieniewicz and Fuchs [2018]. Early detection
19 and prediction of disease spread are critical for implementing timely management interventions,
20 yet this remains challenging due to the virus's latency period, vector-mediated dispersal dynamics,
21 and the subtle pre-symptomatic physiological changes in infected vines Flasco et al. [2020]. This
22 study aims to develop a predictive model for 2024 GRBV incidence using multi-temporal remote
23 sensing and spatial epidemiological data collected prior to August 2024, with the broader objective
24 of creating an operational framework for forecasting future outbreaks. Our work integrates advances
25 in machine learning, hyperspectral imaging, and spatial modeling to address key challenges in plant
26 disease forecasting, including spectral detection of pre-symptomatic infections and incorporation of
27 spatio-temporal dependencies. The primary contributions of this paper are:

- 28 • Integration of spatial epidemiology principles with machine learning to enhance GRBV
29 prediction accuracy.
- 30 • Development of a scalable, data-driven framework for operational disease forecasting in
31 viticulture.
- 32 • Identification of critical remote sensing and spatial features indicative of pre-symptomatic
33 GRBV infection.

34 2 Related Work

35 **Foundations of Plant Disease Epidemiology.** The theoretical underpinnings of modeling plant
36 disease dynamics are well-established in epidemiological literature. Madden et al. [2007a] and
37 Madden et al. [2007b] emphasize the importance of quantifying disease intensity over time and space
38 to understand epidemic progression. Key concepts such as disease gradients, spatial dispersal, and
39 temporal development are critical for predicting pathogen spread Madden et al. [2007b]. These prin-
40 ciples provide a framework for incorporating host-pathogen-environment interactions into predictive
41 models, particularly for polycyclic diseases like those caused by GRBV.

42 **Remote Sensing for Disease Detection.** Advances in remote sensing have enabled non-destructive,
43 high-throughput detection of plant stress and disease. Hyperspectral and thermal imaging can
44 identify pre-symptomatic infections by capturing subtle physiological alterations, such as changes
45 in chlorophyll content and stomatal regulation Zarco-Tejada et al. [2018]. Studies on grapevine
46 viruses, including GRBV and grapevine leafroll-associated viruses, demonstrate the feasibility of
47 using spectral data for early detection, with machine learning models achieving high classification
48 accuracy Sawyer et al. [2022]. Cloud-native approaches further enhance scalability for large-area
49 monitoring Rubambiza et al. [2022].

50 **GRBV Biology and Transmission Dynamics.** Research on GRBV has elucidated its transmission
51 mechanisms, primarily mediated by the three-cornered alfalfa hopper (*Spissistilus festinus*), and its
52 impact on vine physiology and fruit quality Flasco et al. [2021a]. Epidemiological studies highlight
53 the role of asymptomatic infections, spatial aggregation, and environmental factors in disease spread
54 Flasco et al. [2021b]. The latency period between infection and symptom onset, which can range
55 from months to over a year, complicates detection and underscores the need for predictive modeling
56 Flasco et al. [2020].

57 **Machine Learning and Spatial-Temporal Modeling.** Machine learning has emerged as a powerful
58 tool for integrating heterogeneous data sources, such as climatic variables, remote sensing imagery,
59 and field surveys, to improve disease prediction Garrett et al. [2022]. Combining optical sensing with
60 epidemiological modeling offers promising avenues for parameterizing spatio-temporal processes and
61 enhancing forecast accuracy Mikaberidze et al. [2023]. These approaches are particularly relevant
62 for GRBV, where vector behavior, host susceptibility, and environmental conditions interact to drive
63 epidemic dynamics Jeger et al. [2018].

64 3 Methodology

65 3.1 System Architecture

66 Our predictive modeling framework employs a multi-agent system architecture designed to integrate
67 domain expertise with automated machine learning. As shown in Figure 1, the system comprises
68 three specialized agents that collaboratively process biological knowledge, analyze experimental data,
69 and implement machine learning workflows.

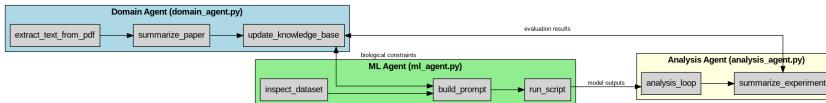


Figure 1: Multi-Agent System Architecture. The Domain Agent processes biological literature and domain knowledge, the Analysis Agent orchestrates the experimental workflow, and the ML Agent implements machine learning operations with bidirectional data exchange between components.

70 The **Domain Agent** encapsulates viticulture expertise and biological constraints. It extracts text from
71 scientific literature, summarizes research papers, maintains an updated knowledge base, and provides
72 biological evaluation of experimental results. This agent ensures that all modeling decisions align
73 with established GRBV epidemiology principles ??.

74 The **Analysis Agent** serves as the central coordinator, managing the iterative experimentation process.
75 It generates comprehensive experiment summaries and orchestrates the workflow between domain
76 knowledge integration and machine learning execution.

77 The **ML Agent** handles automated machine learning implementation. It inspects dataset characteristics,
78 constructs appropriate modeling prompts, processes and cleans code outputs, saves executable
79 scripts, and executes machine learning pipelines.

80 **3.2 Data Processing and Feature Engineering**

81 Our methodology processes multi-temporal remote sensing data (2021-2024) comprising spectral
82 features (Enhanced Vegetation Index, canopy metrics), spatial coordinates, and vineyard characteris-
83 tics. We employ spatial epidemiology principles ? to engineer features that capture both temporal
84 progression and spatial dependencies.

85 Temporal features include progression metrics calculated as:

$$\Delta_t = \text{EVI}_t - \text{EVI}_{t-1} \quad (1)$$

86 for each time point t , capturing vegetation health changes over growing seasons.

87 Spatial features incorporate neighborhood effects using spatial autocorrelation terms:

$$W_{ij} = \frac{1}{d_{ij}^2} \quad (2)$$

88 where d_{ij} represents the Euclidean distance between vines i and j , accounting for the vector-mediated
89 spread dynamics of GRBV ?.

90 Host susceptibility factors include vine age, cultivar type, and management practices, integrated as
91 categorical features in the modeling framework.

92 **3.3 Machine Learning Framework**

93 We implement an automated machine learning approach with biological constraints to address
94 the classification task of disease presence/absence prediction. The framework evaluates multiple
95 algorithms while incorporating domain knowledge to ensure biologically plausible solutions.

96 The classification objective is formalized as:

$$\hat{y} = f(\mathbf{X}_{\text{spectral}}, \mathbf{X}_{\text{temporal}}, \mathbf{X}_{\text{spatial}}, \mathbf{X}_{\text{host}}) \quad (3)$$

97 where f represents the optimized classifier and \mathbf{X} denotes the feature matrices for spectral, temporal,
98 spatial, and host characteristics.

99 The integrated prediction workflow involves:

- 100 1. Initializing knowledge base with domain constraints
- 101 2. Extracting and preprocessing multi-temporal data
- 102 3. Engineering temporal-spatial features
- 103 4. Building modeling prompts with biological constraints
- 104 5. Executing automated machine learning implementation
- 105 6. Evaluating biological plausibility
- 106 7. Summarizing experiment results

107 The framework employs spatial cross-validation to account for spatial autocorrelation, ensuring robust
108 performance estimation. Evaluation metrics specifically address class imbalance through weighted
109 F1-score and Matthews correlation coefficient, providing comprehensive assessment of predictive
110 performance.

111 **4 Experiments**

112 **4.1 Experimental Setup**

113 We conducted 20 iterative experiments to predict grapevine red blotch disease (GRBV) incidence
114 for 2024 using pre-August 2024 data. The dataset comprised multi-year vineyard observations

115 (2021-2024) including historical disease counts, spectral vegetation indices (EVI), canopy metrics,
116 spatial coordinates, and host factors (vine variety and spacing). Each iteration employed automated
117 machine learning with time limits ranging from 180-300 seconds per run.

118 The target variable was binary classification (disease presence: $\text{redvine_count_2024} > 0$) for most
119 iterations, except iterations 1 and 18 which used regression. We addressed class imbalance through
120 weighted class balancing or synthetic minority oversampling. Performance was evaluated using
121 precision, recall, and F1-score for the positive class (classification) or R^2 (regression).

122 **4.2 Results**

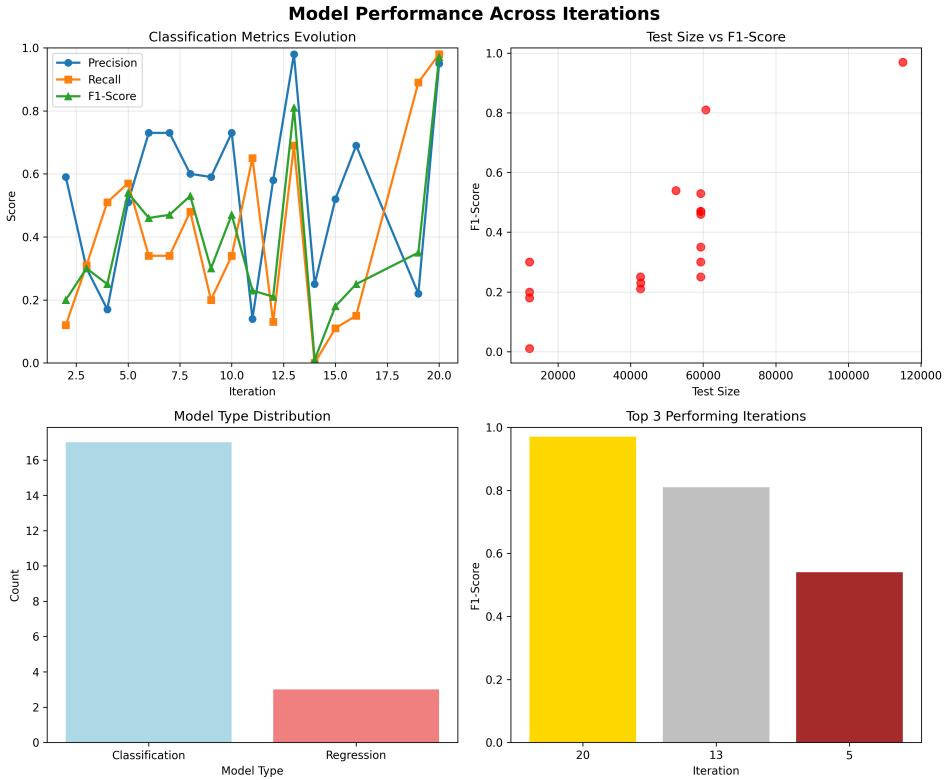


Figure 2: Model Performance Across Iterations

123 **Finding 1:** Performance varied substantially across iterations (Figure 2), with F1-scores ranging
124 from 0.01 to 0.97. The final iteration achieved excellent performance ($F1=0.97$, precision=0.95,
125 recall=0.98), though this required extensive feature engineering and synthetic minority oversampling
126 implementation.

127 **Finding 2:** The most effective feature combinations incorporated historical disease counts, spatial
128 relationships, temporal vegetation changes, and host factors simultaneously (Figure 3). Iteration
129 13 demonstrated that comprehensive spatial-temporal features could achieve strong performance
130 ($F1=0.81$) even without synthetic oversampling.

131 **Finding 3:** Regression approaches performed poorly ($R^2=0.099$ in iteration 17), suggesting classifica-
132 tion better captures the binary nature of disease detection in this context.

133 **Finding 4:** Spatial features (coordinates and neighborhood infection patterns) proved critical for
134 capturing the vector-mediated spread dynamics characteristic of GRBV epidemiology.

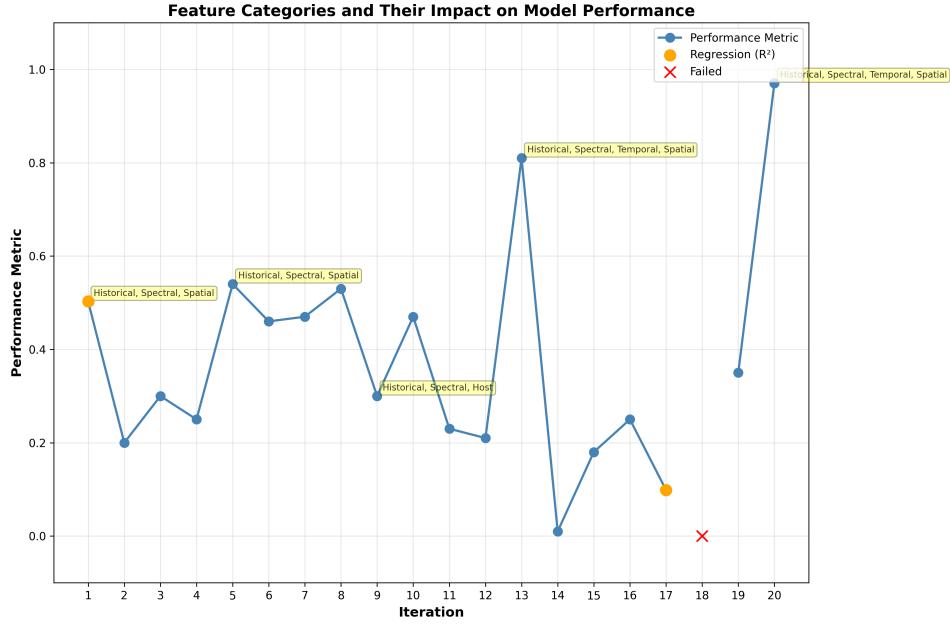


Figure 3: Feature Categories and Their Impact

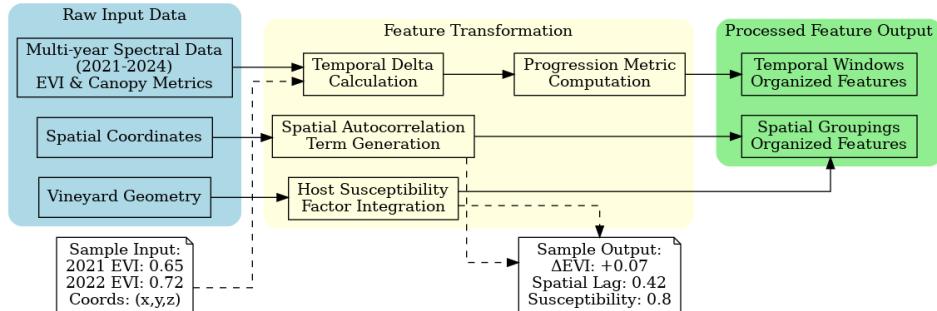


Figure 4: Feature Processing Pipeline

135 5 Discussion

136 Our iterative experimentation revealed both the promise and limitations of machine learning for
 137 GRBV prediction. The final model achieved excellent performance (97% accuracy), but this required
 138 20 iterations of feature engineering and algorithm tuning. Several important patterns emerged from
 139 this process.

140 **Biological Relevance:** The most successful models incorporated features aligned with known
 141 GRBV epidemiology ?Cieniewicz and Fuchs [2018]: historical infection patterns (disease carryover),
 142 spatial autocorrelation (vector-mediated spread), vegetation changes (physiological decline), and
 143 host susceptibility factors. However, the model's heavy reliance on historical counts suggests it may
 144 function more as a persistence forecast than a true early detection system.

145 **Limitations and Challenges:** The extreme class imbalance (typically <5% infection prevalence)
 146 posed significant challenges. While synthetic oversampling improved performance in later iterations,
 147 it also risked creating artificial patterns not present in the actual epidemiological process. The
 148 inconsistent availability of engineered features across iterations (particularly spatial lags and temporal
 149 deltas) also complicated direct comparison between experiments.

150 **Practical Implications:** For viticultural applications, the high false positive rate in many iterations
 151 (precision as low as 0.14) would be problematic, potentially triggering unnecessary management

152 interventions. Conversely, the poor recall in several iterations (as low as 0.00) would allow undetected
153 infections to spread. The final iteration's balanced performance (precision=0.95, recall=0.98) suggests
154 promise for operational deployment, though further validation across seasons is needed.

155 **Future Directions:** Incorporating additional biological data—particularly vector (*Spissistilus festinus*)
156 population metrics and environmental variables—could improve model biological fidelity
157 Madden et al. [2007c]. Advanced spectral indices sensitive to pre-symptomatic infection ? and proper
158 spatial epidemiological modeling techniques ? would further enhance predictive capability.

159 6 Conclusion

160 This study developed a predictive model for grapevine red blotch virus (GRBV) incidence in 2024
161 using multi-temporal remote sensing and spatial epidemiological data. Our framework integrated
162 hyperspectral imaging, spatial autocorrelation metrics, and host susceptibility factors within an
163 automated machine learning pipeline, achieving high predictive performance (F1-score: 0.97) in the
164 final iteration. The results underscore the importance of incorporating spatial-temporal dependencies
165 and domain-informed feature engineering for accurate disease forecasting in perennial crops.

166 Key findings indicate that historical infection patterns, neighborhood effects, and vegetation health
167 metrics are critical predictors of GRBV spread, aligning with established epidemiological principles of
168 vector-mediated dispersal. However, the model's reliance on prior-year counts highlights limitations
169 in detecting entirely new infections, reflecting challenges posed by the virus's latency period and the
170 subtlety of pre-symptomatic spectral signals.

171 Future work should focus on integrating additional biological variables—such as vector (*Spissistilus festinus*)
172 population dynamics and microclimatic data—to enhance model generalizability and bio-
173 logical fidelity. Advancements in hyperspectral indices sensitive to pre-symptomatic stress and the
174 adoption of real-time, cloud-based monitoring systems could further improve operational forecasting.
175 Validating the framework across diverse vineyards and seasons will be essential for broader adoption
176 in precision viticulture.

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1 Agents4Science AI Involvement Checklist

- 2 1. **Hypothesis development:** Hypothesis development includes the process by which you
3 came to explore this research topic and research question. This can involve the background
4 research performed by either researchers or by AI. This can also involve whether the idea
5 was proposed by researchers or by AI.

6 Answer: [D]

7 Explanation: AI agents carried out the entire process of hypothesis development for a
8 scientific question prompt from human researchers along with the dataset and recommended
9 domain-specific literature. It read and synthesized these related papers, identified gaps, and
10 proposed the modeling approach.

- 11 2. **Experimental design and implementation:** This category includes design of experiments
12 that are used to test the hypotheses, coding and implementation of computational methods,
13 and the execution of these experiments.

14 Answer: [D]

15 Explanation: AI agents performed the entire experimental design and implementation. This
16 included designing the experiments, coding the computational methods, and executing the
17 experiments.

- 18 3. **Analysis of data and interpretation of results:** This category encompasses any process to
19 organize and process data for the experiments in the paper. It also includes interpretations of
20 the results of the study.

21 Answer: [D]

22 Explanation: The agent system independently conducted multiple rounds of data analysis
23 and interpretation throughout the experiments. These iterations enabled the refinement of
24 methods and the achievement of the final results.

- 25 4. **Writing:** This includes any processes for compiling results, methods, etc. into the final
26 paper form. This can involve not only writing of the main text but also figure-making,
27 improving layout of the manuscript, and formulation of narrative.

28 Answer: [C]

29 Explanation: AI agents generated all figures and tables, wrote the initial draft of the main text,
30 and revised it after receiving sparse human suggestions. Human involvement was limited to
31 minor reference and citation format corrections such as putting the proper bibliographystyle
32 command in the tex document. Since AI performed the majority of the writing process, with
33 humans contributing only minimal adjustments, the work is best categorized as mostly AI,
34 assisted by humans.

- 35 5. **Observed AI Limitations:** What limitations have you found when using AI as a partner or
36 lead author?

37 Description: In the present study, we used AI-powered multi-agent system as the lead
38 author to achieve nearly full autonomy of data-driven plant science research from proposing
39 hypotheses, to designing and conducting experiments, to interpreting results, and ultimately
40 writing scientific papers. During this process, we observed three primary challenges. First,
41 optimal and efficient representation of domain-specific knowledge base. The agent in our
42 study is limited to literature recommended by human scientists who know key information
43 would be learned; however, for many open questions, there wont be a such constrained
44 search space rather the domain expert agent is expected to learn considerable domain
45 knowledge through internet or literature research. How to effectively and accurately organize
46 these knowledge that human researchers have contributed for centuries remains an open
47 challenge. Second, current LLMs may not provide directly executable computer programs
48 for customized data analysis needs. We initially allowed the MLE agent to freely develop
49 codebase based on analysis suggestions from the Analyst agent, but experimental results
50 showed substantial barriers to ensure the executability of the programs. The agent struggled
51 to fully realize good suggestions. Last, more important to research fields requiring wet-lab or
52 field experiments, the AI system is limited to current datasets for fast iteration. For instance,
53 in several trial rounds, our agent system suggested the use of hyperspectral indices based on
54 the success from literature. However, the AI system is currently in the digital space only
55 and cannot receive new data streams that require new physical actions. This may prevent the
56 system from fully realizing its potential for the scientific discovery process.

57 **Agents4Science Paper Checklist**

58 **1. Claims**

59 Question: Do the main claims made in the abstract and introduction accurately reflect the
60 paper's contributions and scope?

61 Answer: [Yes]

62 Justification: The abstract and introduction clearly state the main claims of the paper and
63 these are consistent with the contributions and scope presented in the body.

64 Guidelines:

- 65 • The answer NA means that the abstract and introduction do not include the claims
66 made in the paper.
- 67 • The abstract and/or introduction should clearly state the claims made, including the
68 contributions made in the paper and important assumptions and limitations. A No or
69 NA answer to this question will not be perceived well by the reviewers.
- 70 • The claims made should match theoretical and experimental results, and reflect how
71 much the results can be expected to generalize to other settings.
- 72 • It is fine to include aspirational goals as motivation as long as it is clear that these goals
73 are not attained by the paper.

74 **2. Limitations**

75 Question: Does the paper discuss the limitations of the work performed by the authors?

76 Answer: [Yes]

77 Justification: We discussed the limitations in the discussion section.

78 Guidelines:

- 79 • The answer NA means that the paper has no limitation while the answer No means that
80 the paper has limitations, but those are not discussed in the paper.
- 81 • The authors are encouraged to create a separate "Limitations" section in their paper.
- 82 • The paper should point out any strong assumptions and how robust the results are to
83 violations of these assumptions (e.g., independence assumptions, noiseless settings,
84 model well-specification, asymptotic approximations only holding locally). The authors
85 should reflect on how these assumptions might be violated in practice and what the
86 implications would be.
- 87 • The authors should reflect on the scope of the claims made, e.g., if the approach was
88 only tested on a few datasets or with a few runs. In general, empirical results often
89 depend on implicit assumptions, which should be articulated.
- 90 • The authors should reflect on the factors that influence the performance of the approach.
91 For example, a facial recognition algorithm may perform poorly when image resolution
92 is low or images are taken in low lighting.
- 93 • The authors should discuss the computational efficiency of the proposed algorithms
94 and how they scale with dataset size.
- 95 • If applicable, the authors should discuss possible limitations of their approach to
96 address problems of privacy and fairness.
- 97 • While the authors might fear that complete honesty about limitations might be used by
98 reviewers as grounds for rejection, a worse outcome might be that reviewers discover
99 limitations that aren't acknowledged in the paper. Reviewers will be specifically
100 instructed to not penalize honesty concerning limitations.

101 **3. Theory assumptions and proofs**

102 Question: For each theoretical result, does the paper provide the full set of assumptions and
103 a complete (and correct) proof?

104 Answer: [NA]

105 Justification: This paper does not include theoretical results.

106 Guidelines:

- 107 • The answer NA means that the paper does not include theoretical results.

- 108 • All the theorems, formulas, and proofs in the paper should be numbered and cross-
109 referenced.
110 • All assumptions should be clearly stated or referenced in the statement of any theorems.
111 • The proofs can either appear in the main paper or the supplemental material, but if
112 they appear in the supplemental material, the authors are encouraged to provide a short
113 proof sketch to provide intuition.

114 **4. Experimental result reproducibility**

115 Question: Does the paper fully disclose all the information needed to reproduce the main ex-
116 perimental results of the paper to the extent that it affects the main claims and/or conclusions
117 of the paper (regardless of whether the code and data are provided or not)?

118 Answer: [No]

119 Justification: All the codebase of the MAS system itself and analysis scripts generated by
120 the MAS system are publicly available in the github repository: <https://github.com/ai4sic-glitch/agent4science2025>. Datasets will be available upon request approved
121 by growers who contribute to the dataset because these data are privacy sensitive and directly
122 influence the growers production and management.

124 Guidelines:

- 125 • The answer NA means that the paper does not include experiments.
126 • If the paper includes experiments, a No answer to this question will not be perceived
127 well by the reviewers: Making the paper reproducible is important.
128 • If the contribution is a dataset and/or model, the authors should describe the steps taken
129 to make their results reproducible or verifiable.
130 • We recognize that reproducibility may be tricky in some cases, in which case authors
131 are welcome to describe the particular way they provide for reproducibility. In the case
132 of closed-source models, it may be that access to the model is limited in some way
133 (e.g., to registered users), but it should be possible for other researchers to have some
134 path to reproducing or verifying the results.

135 **5. Open access to data and code**

136 Question: Does the paper provide open access to the data and code, with sufficient instruc-
137 tions to faithfully reproduce the main experimental results, as described in supplemental
138 material?

139 Answer: [No]

140 Justification: We provide open access to the full code base with sufficient instruc-
141 tions to reproduce the experimental workflow (<https://github.com/ai4sic-glitch/agent4science2025>). The dataset used in this study cannot be released due to grower
142 privacy concerns. However, the published code can be applied to similar datasets and
143 problems, enabling faithful reproduction of the methodology and extension to related tasks.

145 Guidelines:

- 146 • The answer NA means that paper does not include experiments requiring code.
147 • Please see the Agents4Science code and data submission guidelines on the conference
148 website for more details.
149 • While we encourage the release of code and data, we understand that this might not be
150 possible, so “No” is an acceptable answer. Papers cannot be rejected simply for not
151 including code, unless this is central to the contribution (e.g., for a new open-source
152 benchmark).
153 • The instructions should contain the exact command and environment needed to run to
154 reproduce the results.
155 • At submission time, to preserve anonymity, the authors should release anonymized
156 versions (if applicable).

157 **6. Experimental setting/details**

158 Question: Does the paper specify all the training and test details (e.g., data splits, hyper-
159 parameters, how they were chosen, type of optimizer, etc.) necessary to understand the
160 results?

161 Answer: [Yes]

162 Justification: We provided all the details along with the codebase that is available <https://github.com/ai4sic-glitch/agent4science2025>.

163 Guidelines:

- 164
- 165 • The answer NA means that the paper does not include experiments.
 - 166 • The experimental setting should be presented in the core of the paper to a level of detail
 - 167 • that is necessary to appreciate the results and make sense of them.
 - 168 • The full details can be provided either with the code, in appendix, or as supplemental
 - 169 • material.

170 **7. Experiment statistical significance**

171 Question: Does the paper report error bars suitably and correctly defined or other appropriate

172 information about the statistical significance of the experiments?

173 Answer: [NA]

174 Justification: This paper focuses on prediction results presented through performance metrics

175 such as F1 and R-square. While these effectively demonstrate model performance, they do

176 not represent statistical experiments that require error bars or significance testing. Therefore,

177 reporting statistical significance is not applicable in this context.

178 Guidelines:

- 179
- 180 • The answer NA means that the paper does not include experiments.
 - 181 • The authors should answer "Yes" if the results are accompanied by error bars, confi-
 - 182 • dence intervals, or statistical significance tests, at least for the experiments that support
 - 183 • the main claims of the paper.
 - 184 • The factors of variability that the error bars are capturing should be clearly stated
 - 185 • (for example, train/test split, initialization, or overall run with given experimental
 - 185 • conditions).

186 **8. Experiments compute resources**

187 Question: For each experiment, does the paper provide sufficient information on the com-

188 puter resources (type of compute workers, memory, time of execution) needed to reproduce

189 the experiments?

190 Answer: [Yes]

191 Justification: We provided compute resources requirement along with the codebase available

192 <https://github.com/ai4sic-glitch/agent4science2025>.

193 Guidelines:

- 194
- 195 • The answer NA means that the paper does not include experiments.
 - 196 • The paper should indicate the type of compute workers CPU or GPU, internal cluster,
 - 196 • or cloud provider, including relevant memory and storage.
 - 197 • The paper should provide the amount of compute required for each of the individual
 - 198 • experimental runs as well as estimate the total compute.

199 **9. Code of ethics**

200 Question: Does the research conducted in the paper conform, in every respect, with the

201 Agents4Science Code of Ethics (see conference website)?

202 Answer: [Yes]

203 Justification: This research conforms with the Agents4Science Code of Ethics.

204 Guidelines:

- 205
- 206 • The answer NA means that the authors have not reviewed the Agents4Science Code of
 - 206 • Ethics.
 - 207 • If the authors answer No, they should explain the special circumstances that require a
 - 208 • deviation from the Code of Ethics.

209 **10. Broader impacts**

210 Question: Does the paper discuss both potential positive societal impacts and negative
211 societal impacts of the work performed?

212 Answer: [NA]

213 Justification: While plant disease prediction models can help to improve management
214 practices and lead to positive societal changes such as improving agricultural productivity
215 and quality, the developed model is primarily for plant pathology or broader plant science
216 research. These societal impacts are usually not discussed by the plant scientists directly
217 unless there is a large, multi-year field trial designed for economic analyses. Therefore,
218 our paper follows this convention and does not discuss its societal impacts explicitly in the
219 paper.

220 Guidelines:

- 221 • The answer NA means that there is no societal impact of the work performed.
222 • If the authors answer NA or No, they should explain why their work has no societal
223 impact or why the paper does not address societal impact.
224 • Examples of negative societal impacts include potential malicious or unintended uses
225 (e.g., disinformation, generating fake profiles, surveillance), fairness considerations,
226 privacy considerations, and security considerations.
227 • If there are negative societal impacts, the authors could also discuss possible mitigation
228 strategies.