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### INTRODUCTION

Metabolic dysfunction-associated steatotic liver disease (MASLD) has a prevalence exceeding 25% in the general population. Non-invasive tests, namely the FIB-4 index, NFS, APRI, and AST/ALT, play a crucial role in differentiating advanced fibrosis stages (F012 vs. F34 in metavir score).

### AIM

Developing and optimizing an interpretable machine learning model that employs the aforementioned non-invasive test parameters as features, surpassing their individual performances in diagnosing advanced fibrosis in MASLD patients.

## METHOD

- Open data from two cohorts, China (train) and Malaysia (test), are used, with 540 participants (early/advanced fibrosis: 391/149) and 147 participants (116/31) having liver biopsyconfirmed hepatic fibrosis
- Features: age, sex, BMI, ALB, PLT, AST, ALT, ALT/AST, AST/PLT, presence of diabetes/impaired fasting glycemia (DM.IFG), FIB-4 score, NFS score, APRI score
- The machine learning model is trained, tuned, and validated using the train dataset, followed by testing on the test dataset
- 10-fold cross-validation (10CV) enhances model robustness
- Feature engineering expands feature space by considering power and product combinations of initial features
- Shapley values from explainable artificial intelligence are utilized to understand feature importance in catboost predictions

## OPEN DATA, EXPLAINABLE AI, DATA SCIENCE AND CONVENTIONAL NITS THE RECIPE FOR NEW MACHINE LEARNING DIAGNOSTIC TESTS ON MASLD

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#### RESULTS

We conducted a comparative analysis between our catboost model (ML) and four non-invasive tests (NITs) FIB-4, NFS, APRI, and AST/ALT using spec threshold values for advanced fibrosis, namely 1.30, -1.455, 0.64, and 0. respectively. The comparison was performed on identical folds within the 10-f cross-validation (10CV) framework, with consistent random seed selection.



### CONCLUSIONS

• By utilizing data science with catboost, feature engineering, and the parameters/scores from four NITs, we achieve superior performance and robustness compared to those NITs

• Explainable artificial intelligence identifies the importance of FIB-4, NFS^6, and PLT^8, potentially leading to a novel noninvasive test

• We propose a paradigm shift for classifying MASLD patients into early and advanced fibrosis stages

• Linear cut-off values alone are inadequate, necessitating data science and explainable artificial intelligence to provide insights

• Combining parameters from all four NITs in an interpretable machine learning framework improves outcomes

	NIT	Spec. 10CV	Sens. 10CV	ROC-AUC 10CV	F1 10CV	Spec. Test	Sens. Test	<b>ROC-AUC Test</b>	F1 Tes
	FIB-4	0.8500+/-0.0744	0.4893+/-0.0855	0.6997+/-0.0692	0.5499+/-0.0797	0.9109	0.4783	0.7514	0.571
he	NFS	0.8764+/-0.0549	0.4559+/-0.0893	0.7076+/-0.0678	0.5673+/-0.0802	0.9271	0.4706	0.7707	0.585
fic	APRI	0.8076+/-0.0600	0.4663+/-0.0945	0.6452+/-0.0538	0.4806+/-0.0792	0.8857	0.4524	0.7073	0.520
old	AST/ALT	0.7483+/-0.0471	0.3521+/-0.1420	0.5445+/-0.0504	0.2994+/-0.1181	0.8140	0.3889	0.5655	0.285
	ML	0.8845+/-0.0601	0.5903+/-0.0877	0.7671+/-0.0632	0.6425+/-0.0708	0.9434	0.6098	0.8343	0.694
				Feature Imp	ortance of LossFun	ctionChange			
		0.016							
		0.016 0.014 0.012 0.010 0.008 0.006 0.004 0.004							

## REFERENCES

**Angelakis A. et al**. Using FIB-4's parameters an explainable black-box machine learning model outperforms FIB-4 index on the diagnosis of advanced fibrosis of non alcohol related fatty liver disease patients in three cohorts from China, Malaysia and India. Journal of Hepatology 22023 vol. 78(S1) | S100–S101, DOI: https://doi.org/10.1016/S0168-8278(23)00435-X





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