Economic Evaluation of Transient Elastography for Significant Fibrosis Detection in Patients with Diabetes Mellitus in Thailand



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OBJECTIVES

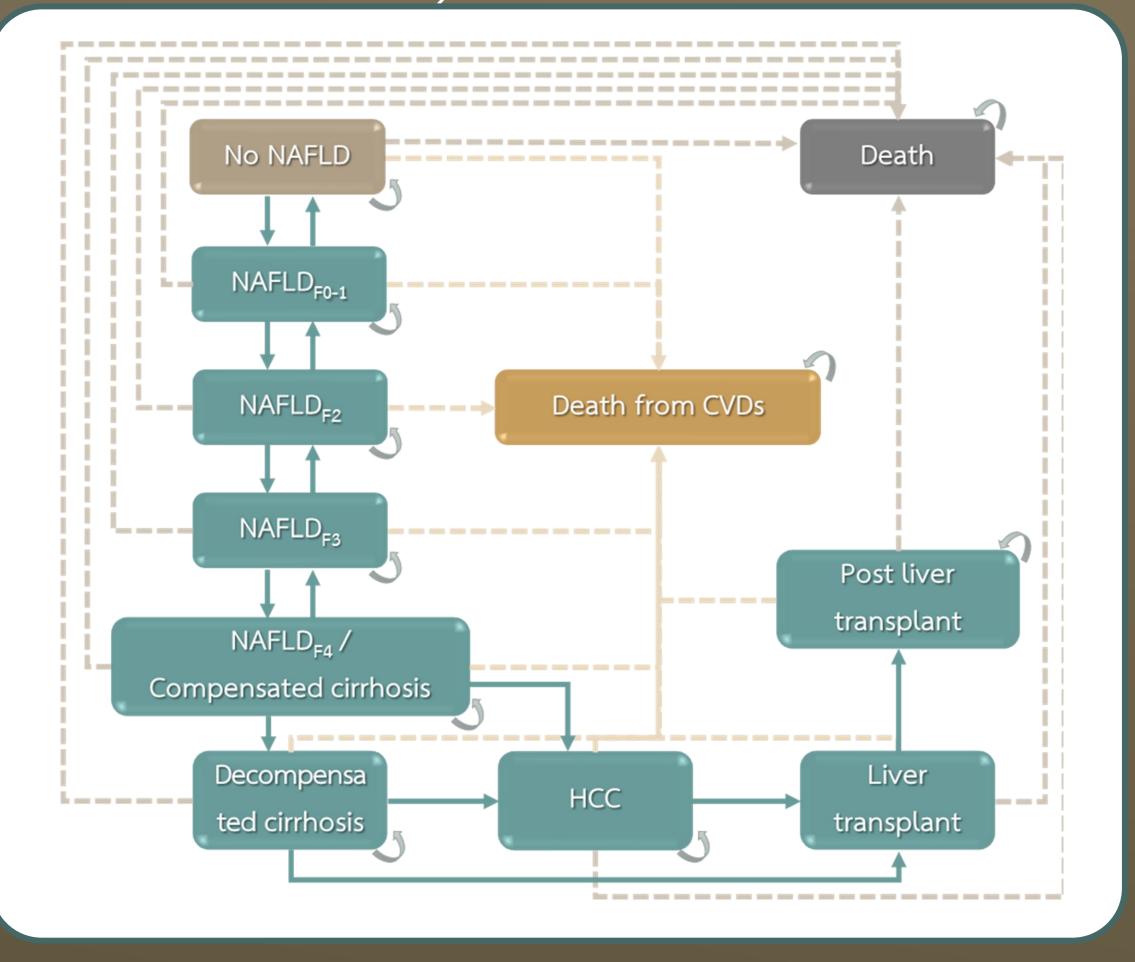
The association between diabetes mellitus (DM) and non-alcoholic fatty liver disease (NAFLD) is well-documented, with DM heightens risk of developing NAFLD and liver fibrosis. This study aimed to evaluate the cost-effectiveness and budget impact of transient elastography (TE) for detecting significant fibrosis in patients with DM in Thailand.

METHODS (1 USD = 34.64 THB)

A cost-utility analysis was conducted using a decision tree and Markov model over a lifetime horizon with a one-year cycle length, adopting a societal perspective. The model included 11 health states related to NAFLD (**Figure 1**), comparing no screening against one-time screening strategies involving fibrosis-4 index then TE (FIB-4+TE), steatosis-associated fibrosis estimator score then TE (SAFE+TE), and TE alone. Key parameters are shown in the **Table**. The analysis applied a discount rate of 3% per year on costs and outcomes. Incremental cost-effectiveness ratios (ICERs) were calculated and compared against a willingness-to-pay threshold of 4619 USD per quality-adjusted life-year (QALY) gained. A 5-year budget impact was estimated from a payer perspective.

RESULTS

Among the screening methods evaluated, TE alone yielded the highest total lifetime costs (5,785 USD) and QALYs (12.81 QALYs). Compared to no screening, all strategies demonstrated cost-effectiveness with ICERs of 2193, 2321, and 2857 USD per QALY gained for FIB-4+TE, SAFE+TE, and TE alone, respectively. SAFE+TE emerged as the best-buy option when compared to the other strategies. However, the probabilistic sensitivity analysis illustrated FIB-4+TE had the highest chance of being cost-effective (**Figure 2**). Estimated annual budget impacts were substantial, amounting to 13.6, 21.8, and 19.9 million USD for FIB-4+TE, SAFE+TE, and TE alone, respectively.



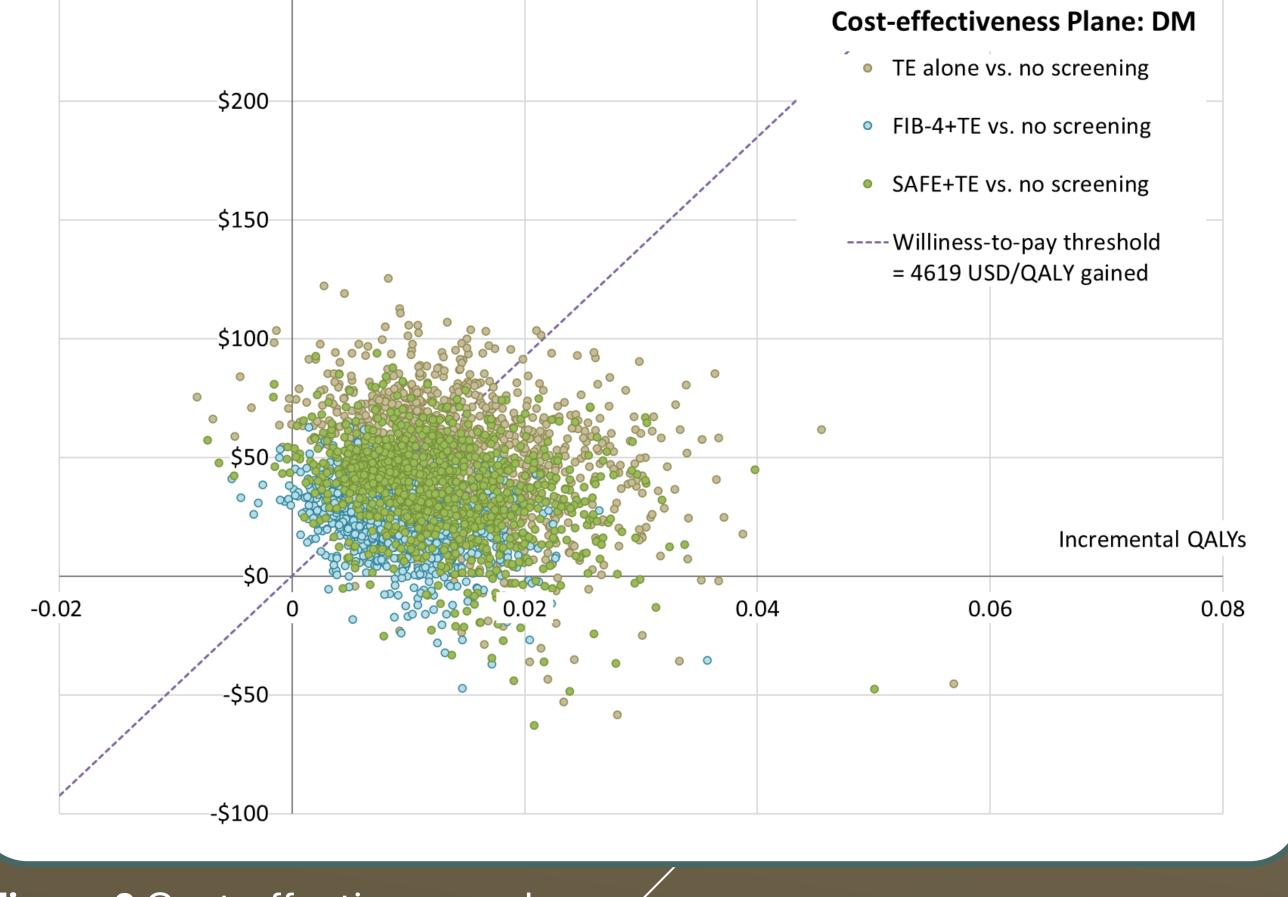


Figure 2 Cost-effectiveness plane /

Incremental cost (USD)

Table Key input parameters

Input parameters	Values (ranges)	References
Performance of tests (%) $FIB-4 \ge 1.3$ $SAFE \ge 0$ $TE \ge 7.0 \text{ kPa}$	Sens: 66 (64, 78); Spec: 65 (65, 68) Sens: 87 (86, 89); Spec: 35 (32, 38) Sens: 80 (76, 83); Spec: 73 (68, 77)	Alkhouri, 2025 Alkhouri, 2025 Xiao, 2017; Selvaraj, 2021
Prevalence (%) DM NAFLD NAFLD NAFLD _{F2} (1 st detection)	9.9 (9.4, 10.4) 34.8 (34.4, 35.2) 30.4 (23.3, 38.2)	Aekplakorn, 2018 Phisalprapa, 2021 Primary data
Incidence (%) DM NAFLD NAFLD NAFLD F2	0.9 (0.7, 1.1) 4.3 (3.9, 4.7) 6.7 (1.7, 11.6)	Papier, 2016 Park, 2021 Le, 2023
Treatment effectiveness	Risk reduction: 0.20 (0.16, 0.45)	Vilar-Gomez, 2015
Utilities DM, NAFLD _{F0-F3} NAFLD _{F4} DC HCC LT Post-LT	0.753 (0.526, 0.980) 0.748 (0.666, 0.830) 0.603 (0.590, 0.754) 0.380 (0.360, 0.410) 0.570 (0.540, 0.600) 0.683 (0.640, 0.690)	Deerochanawong, 2023 Chongmalaxme, 2019 Prakongsai, 2014 Levy, 2008 Levy, 2008 Prakongsai, 2014
Costs (USD per year) Screening tests DM treatment Lifestyle modification NAFLD DC HCC LT Post-LT Food Transportation	FIB-4: 7.8 SAFE: 10.3 TE: 57.7 (± 14.4) 277 (± 69.1) 41.2 (± 10.3) 356.3 (± 89.1) – 1,118.6 (± 279.6) 4,363.9 (± 1,091.0) 5,335.6 (± 1,333.9) 19,729.9 (± 4,932.5) 3,183.9 (± 796.0) 1.9 (± 0.2), 5.2 (± 0.4)	Riewpaiboon, 2009 Riewpaiboon, 2009 Primary data, Siriraj Hospita Primary data, Siriraj Hospita Riewpaiboon, 2009 Primary data Chongmalaxme, 2019 Thongsawat, 2014
Food, Transportation	1.9 (± 0.2), 0.2 (± 0.4)	Riewpaiboon, 2009

DC, decompensated cirrhoxis; F, fibrosis; FIB-4, fibrosis-4 index; HCC; hepatocellular carcinoma; LT, liver transplant; NAFLD, non-alcoholic fatty liver disease; SAFE, steatosis-associated fibrosis estimator score; sens, sensitivity; spec, specificity; TE, transient elastography

CONCLUSIONS

Implementing screening for significant fibrosis in patients with DM is deemed cost-effective. However, considerations regarding the budget impact and accessibility of TE are critical for practical implementation.

Acknowledgement

We would like to extend our heartfelt thanks to Euarat Mepramoon, our research assistant, for her exceptional coordination of the project.

We are also deeply grateful to Assistant Professor Surasak Saokaew and Assistant Professor Pimsiri Sripongpun for sharing their expert perspectives, which have significantly enhanced our understanding and contributed to the depth of this study.

Ethics approval

This study was approved by ethical committee of Faculty of Medicine Siriraj Hospital, Mahidol University (certificate of approval number: Si 990/2023) and the Human Research Ethics Committee of Thammasat University (Medicine) (certificate of approval number: 124/2024).