

Machine Learning-based Models for Disease Prediction

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Source: iconape.com



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- 1. Introduction
- 2. Machine Learning
- 3. Machine Learning-based Disease Prediction Models
 - 4. Machine Learning-based Model for Disease Prediction Applications
 - 5. Conclusion



Introduction



- According to World Health Organization report [1], in 2019, the top 10 causes of death accounted for 55% of the 55.4 million deaths worldwide.
- These top 10 serious diseases take an 'immense and increasing toll on lives' [1].

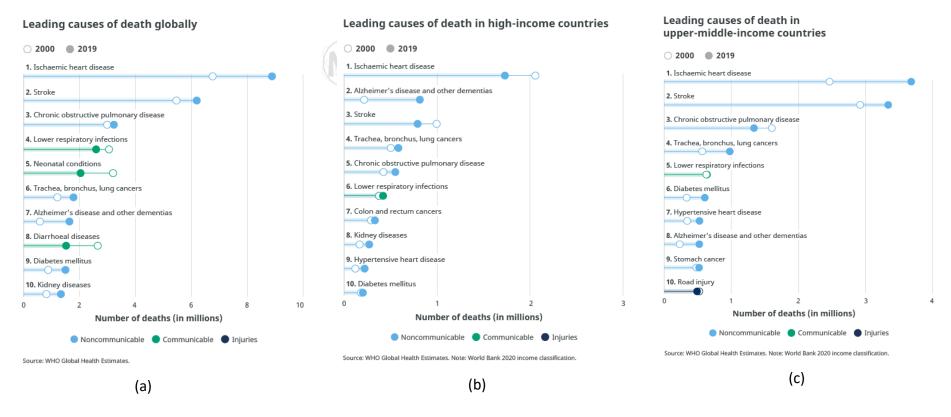


Fig 1. Worldwide leading cause death (a), in high-income countries (b), in upper-middle-income (c) [1]

Introduction



- As the number of deaths due to chronic diseases rose annually, the cost of medical diagnosis, tests, and treatment also followed rising [2, 3].
- Several methods and strategies should be developed as a solution to help individuals more easily and cost effectively check their health status, thus could help early detect the diseases and prevent from the occurrence of the worst-case scenario.

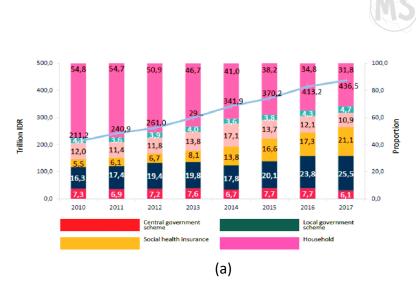
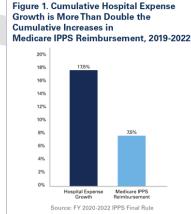
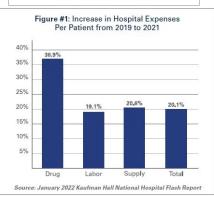
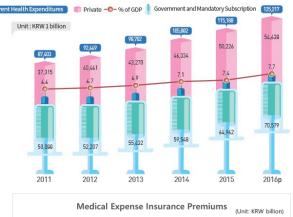


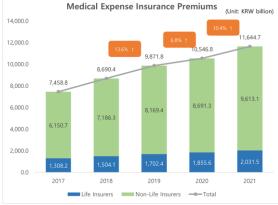
Fig 2. Total Health Expenditure of Indonesia (a), USA (b), Korea (c). [4, 5, 6, 7]





(b)





(c)

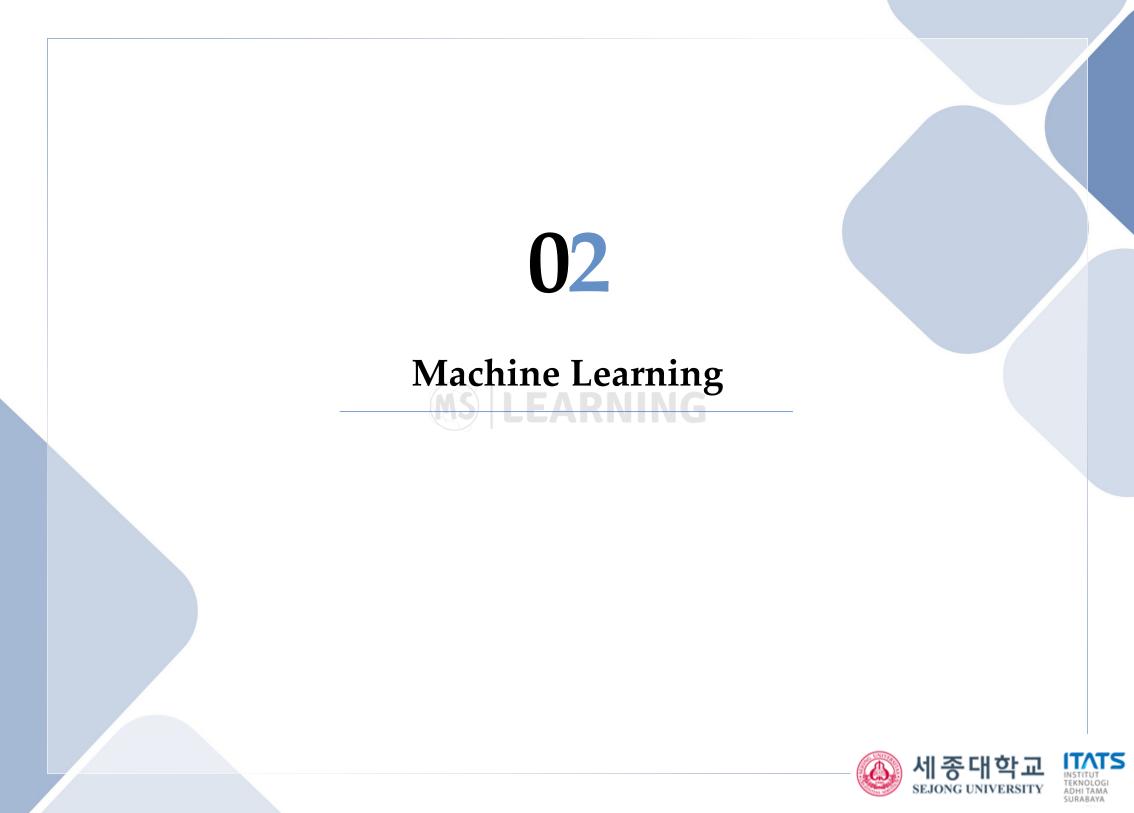
Introduction

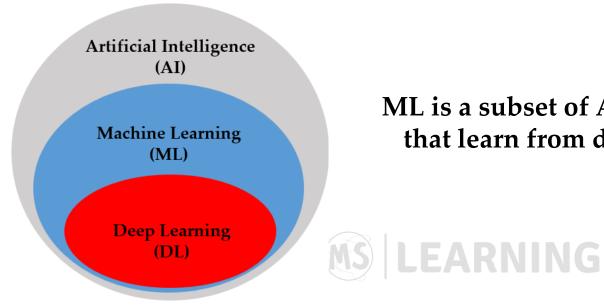


- One of the solutions that could be used to early detect the disease is machine learning-based prediction models development and utilization.
- Recent studies have utilized machine learning algorithms as decision-making tools to diagnose various diseases at an early stage, so that preventive action can be taken by individuals.
- The machine learning algorithms have showed high performance on predicting the diabetes [3, 8, 9, 10, 11], heart disease [12, 13, 14, 15], lung cancer [16, 17], and other diseases based on current conditions of individuals.

Disease	Author	Algorithm	Dataset	Accuracy (%)
Diabetes	Fitriyani et. al [3]	Forward Logistic Regression and MLP	NAGALA	92.11
	Patil et. al [8]	Decision Tree C4.5	The Pima Indians	92.38
	Wu et. al [9]	K-Mean and Logistic Regression	The Pima Indians	93.50
	ljaz et. al [10]	DBSCAN+SMOTE+Random Forest	Dr John Schorling	92.55
	Fitriyani et. al [11]	iForest+SMOTETomek+Ensemble Learning	Dr John Schorling	100.00
Heart Disease	Bhatt et. al [12]	GridSearchCV+MLP	CVD (Kaggle)	87.28
	Fitriyani et. al [13]	DBSCAN+SMOTE-ENN+XGBOOST	Statlog Cleveland	95.90 98.40
	Ali et. al [14]	Stacked SVMs	Cleveland	92.22
	Gupta et. al [15]	FAMD + Random Forest	Cleveland	93.44
Lung Cancer	Dritsas and Trigka [16]	SMOTE + Rotation Forest	Lung Cancer (Kaggle)	97.10
	Alam et. al [17]	Watershed Transform + GLCM + SVM	Lung Cancer (UCI ML Rep)	97.00

Adopted from "Introduction to machine learning by Delta Analytics"





ML is a subset of AI, which uses algorithms that learn from data to make predictions

Fig 3. Hierarchy of AI, ML, and DL

Like a human would, the ML algorithm learns from historical data or past experiences and is thus able to make predictions about the future.

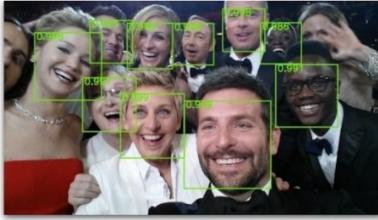


Fig 4. Machine learning concept



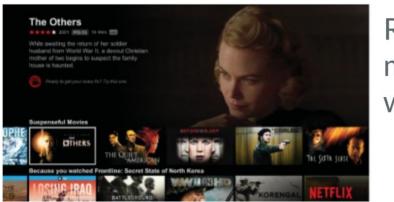
Age (Years)* 52	ah	Predictions	
D-Dimer (mg/L FEU)* 2.52	ıth	Al Severity Score Likelihood of	Severe Clinical Events
LDH (IU/L)* 1764	ah	0	97% 🛉 86%
Lymphocyte (%)* 8	ith	3.6	Ventilator use End Organ Damage
Eosinophil (%)* 0.1	ıtlı	Severity 4 - Mortality	ііііііііііііііііііііііііііііііііііііі
Creatinine (mg/dL)* 0.2	ıth	Sevency 4 - Mortancy	In Hospital 30 Day Mortality
CRP (mg/L)* 63.9	ıth		
Ferritin (ng/mL)* 20310	ıtlı		
INR* 1.15	ıtlı	Cohort Severity Distribution Patients in Cohort: 8427	Percentage Distribution of Similar Patients by Severity
Tropanin-I (pg/mL)* 59	ith	Construction for a construction of the co	Seenty 2 Seenty 1 Seenty 2 Seenty 3 Seenty 4
		727 Mill 228	179 199 244
		· * * * * * * * * * * * * * * * * * * *	0 10 20 30 43 50 60 70 50 90 109

Machine learning is a powerful tool; it can...



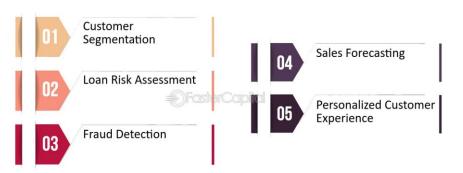
Recognize your face in a photo.

Identify potential disease progression and predict disease

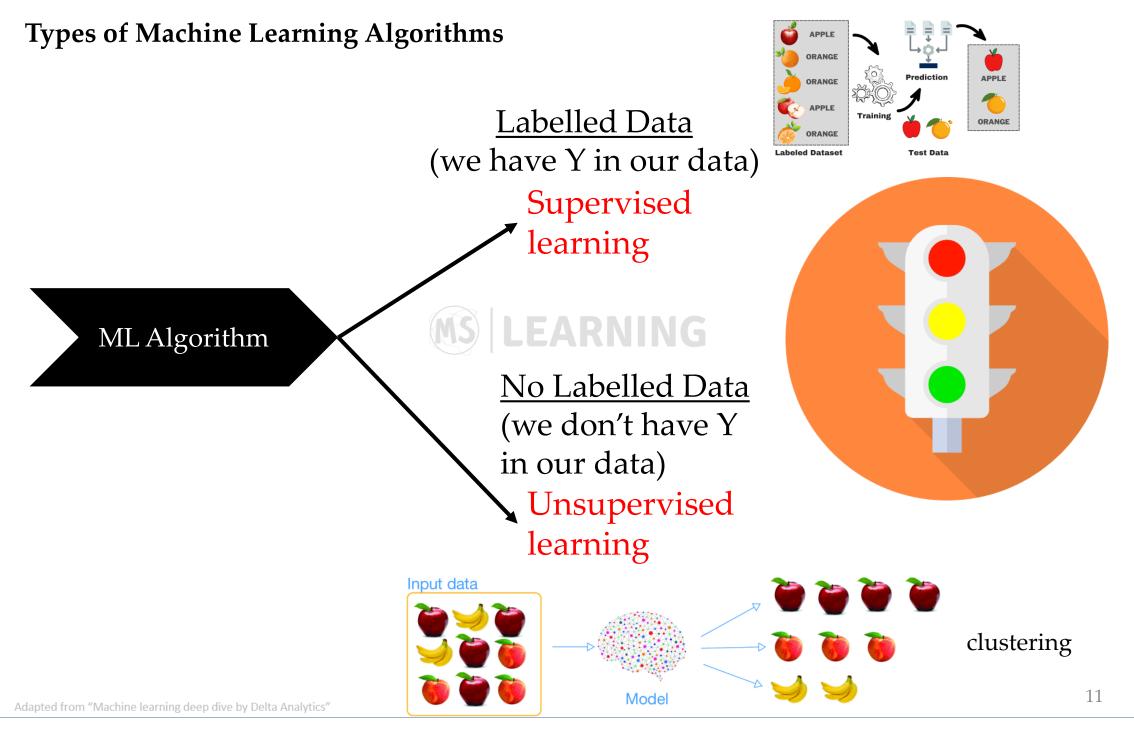


Recommend movies you will like.

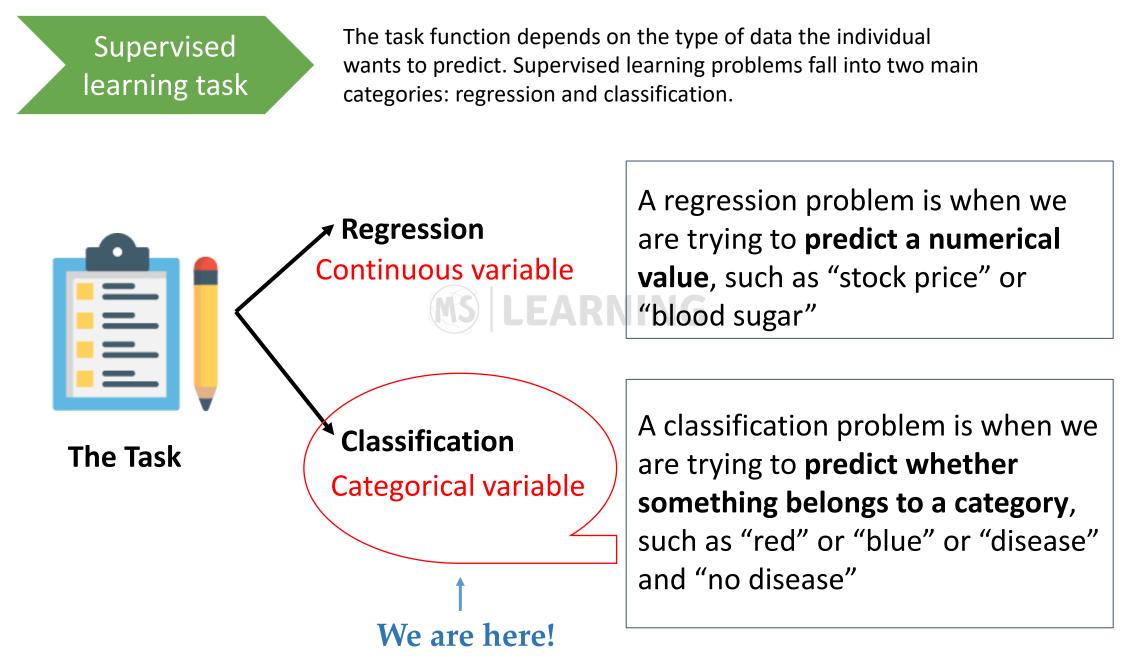
Predict potential bank customer











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Machine Learning

Supervised Machine Learning

- Logistic Regression
- Decision Tree
- K-Nearest Neighbor (KNN)
- Support Vector Machine (SVM)
- Neural Network : Multilayer Perceptron (MLP)
- Naïve Bayes
- Random Forest
- AdaBoost
- Extreme Gradient Boosting (XGB)
- LightGBM
- CatBoost

03

Machine Learning-based Disease

Prediction Models





- 1. Data Collection
- 2. Data Preparation
- 3. Choose the Model/Algorithm

Data Preprocessing

Missing value

elimination

Learning and Classification

K-fold Cross

Validation

Multi Layer Perceptron Classifier

With Weight Sensiting ALT Sensiting ALT Shatter Photos Pho Testing

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(a) [3]

- 4. Training the Model
- 5. Evaluate the Model

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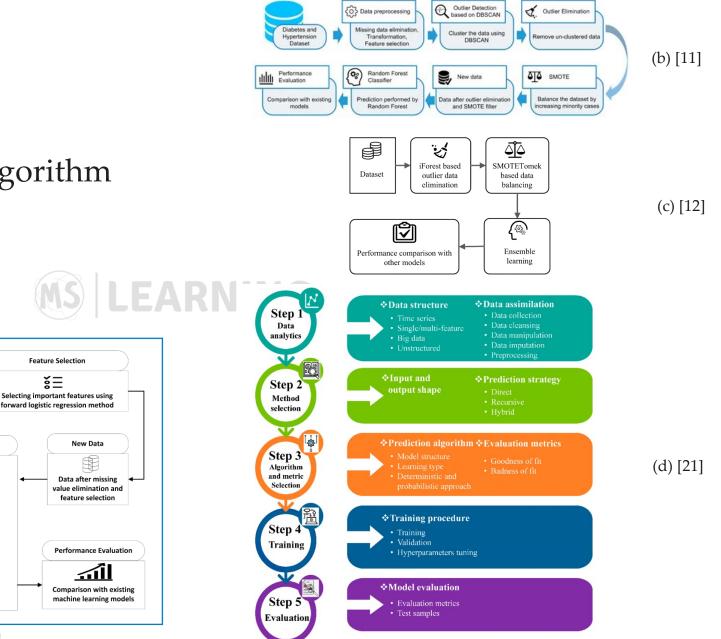
Training

π

T2D-NAFLD

DATASETS

Ø



Adapted from "Machine learning deep dive by Delta Analytics" Fig 6. Machine learning-based model deployed in real-case health applications



1. Data Collection

There are two main categories of data [22]:

- Primary data is newly collected data;

Primary Data

- It can be gathered directly from people's responses (surveys), or
 - from their biometrics (blood pressure, weight, blood tests, etc.).
- The data is collected for other (medical) purposes by extracting the data from medical records.
- Secondary data is data that already exists;
- It has already been published or complied.

Secondary Data

 There are extant local, regional, national and international databases such as Public Health Data, government statistics, and WHO data.
 Public health data sources : UCI ML Repository, Kaggle, data.world, KHNES, NHIS, etc.



1. Data Collection

Public health dataset example:

	- · · ·										-				-													
Age	Gender(I	l Body M	a Obesity	Hyperter	Hyperlip Me	taboli Smok	king_S[AST	ALT	r alf				glycei HDL					y Fibros	sis NAS	5 sco NAS s	co Fibrosis			e Cirrhosis Dia	agnosi Ty	pe of Disease (Mild illne	s: Clas:	÷
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32	2	26.09) 0	0	0	2	23	31	100	23	141	72	40	87	119	2	2	1	4	1	1 (ו כ	0 0	1		2	1
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39	1	32.		1 0	1	1	1	18	12	60	9	182	210	35	105	110	2	2	0	4	1 () (ו ו	0 0	1		2	0
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35	2	37.8		1 0	1	1	1	16	26	53	19	226	170	40	152	99	2	2	1	4	1	1 (ו ו	0 0	1		2	0
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565	44		0.35	1	0 1		1	30	39	77	29	183	106	60	110	133	1	4	3	6	0	1	1	1 0	1	2	5	1
566	64		9.74	0	1 1		3	20	20	65	176	170	262	49	58	112	1	4	3	6	0	1	1	1 0	1	2		1
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565 rows × 29 columns



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Development of machine learning-based disease prediction model

2. Data Preparation

Ove Und Hyb

- Data preparation in machine learning is the process of cleaning, transforming, and organi zing raw data into a format that machine learning algorithms can understand.
- Processing the data into good quality data [22] due to lack quality of data.

Missing value elimination		Microsoft Excel, Python, etc.
	DR	MING
Duplicate data elimination		Microsoft Excel, Python, etc.
	_	
Noise or outlier elimination	1	Isolation Forest, DBSCAN, Local Outlier Factor,
		standard deviation, interquartile range

• Processing the data due to the high quantity of data (size or dimension) that could affect the performance of the model.

Imbalance class distribution (high difference number of '+' and '–' class)	Too large data (number of features or dimension)
er sampling Technique : SMOTE, ADSYN, Random Over Sampling (ROS)	Feature selection : Information Gain, Chi Square Test, Fisher Score,
der sampling Technique : Tomek Link, Random Under Sampling, NearMiss	Correlation Coefficient, Forward Selection, Backward Selection,
brid Technique : SMOTE-ENN, SMOTE-Tomek	Recursive Feature Elimination, Tree-based (RF, XGB)
"Machine learning deep dive by Delta Analytics"	Feature Extraction : Principal Component Analysis



2. Data Preparation

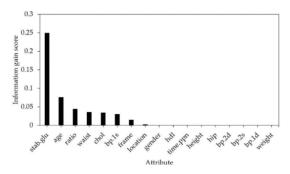


Fig 7. Feature selection based on Information Gain [11]

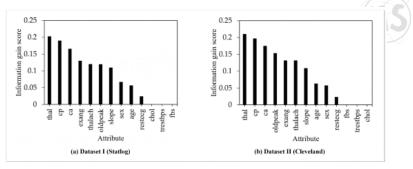


Fig 10. Feature selection based on Information Gain [13]

Dataset	MinPts	eps	# Outlier Data
Dataset I (Statlog)	5	9	3
Dataset II (Cleveland)	5	8	6

Fig 10. Outlier elimination based on DBSCAN [13]

	Before SM	OTE-ENN	After SM	OTE-ENN
Dataset	Minority class (%)	Majority class (%)	Minority class (%)	Majority class (%)
Ι	44.19	55.81	50.79	49.21
II	46.05	53.95	49.5	50.5

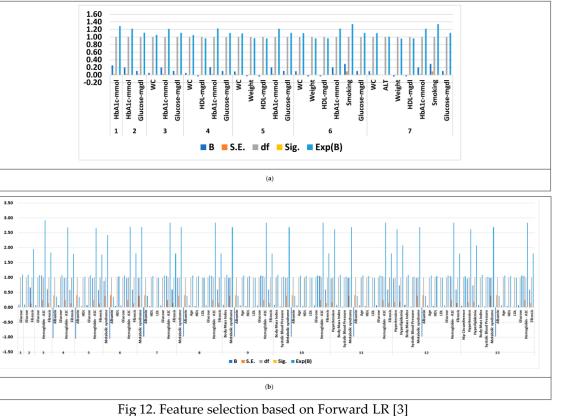
Fig 11. Data balancing based on SMOTE-ENN[13]

Dataset	MaxSample	NumTree	Number of Outliers	Number of subjects before outlier removal	Number of subjects after outlier removal
Ι	41	100	94	403	309
Π	18	100	36	175	139
III	23	100	38	224	186
IV	40	100	156	398	242

Fig 8. Outlier elimination based on Isolation Forest [11]

Datasat	Before SM	OTETomek	After SMC)TETomek				
Dataset	Minority (%)	Majority (%)	Minority (%)	Majority (%)				
Ι	15 (4.85%)	294 (95.15%)	293 (50%)	293 (50%)				
П	31 (22.30%)	108 (77.70%)	99 (50%)	99 (50%)				
III	73 (39.25%)	113 (60.75%)	94 (50%)	94 (50%)				
IV	51 (21.07%)	191 (78.93%)	191 (50%)	191 (50%)				

Fig 9. Data balancing based on SMOTETomek [11]





- 3. Choose the Model/Algorithm
 - In machine learning, choosing the right model is one of the most important steps in building a successful predictive model.
 - Choosing the wrong model can lead to poor performance, wasted time and resources, and inaccurate results.





- 3. Choose the Model/Algorithm
 - Steps to choose the right machine learning model:
 - Define the problem : the researcher needs to understand what kind of problem he/she is dealing with.
 Is it a classification problem or a regression problem? is he/she trying to predict a categorical or continuous outcome?
 - **Consider the data** : the researcher should know the *feature types* (numerical or categorical, text or image), different models may be better suited for different feature types. *Feature importance:* are all features equally important, or are some more important than others? If some features are more important, and want to use a model that can perform feature selection or feature weighting, such as random forests [23]. *Data size*: how much data does the researcher have? If the dataset is small, simpler models may be more appropriate to avoid overfitting [23]. If dataset is large, more complex models may be able to capture the patterns. *Data distribution*: Is the data distribution balanced or imbalanced?
 - Evaluate different models or conducting model comparison: each type of model has its own strengths and weaknesses, and it's important to evaluate each one carefully to determine which is best suited for the researcher's problem.



4. Training the Model

- Model training is the stage where the ML algorithm is trained by feeding datasets.
- Model training in machine language is the process of feeding an ML algorithm with data to help identify and demonstrate correlation between the input data and outcomes.

iteration 10 MLP 0.33333333333333333 - youden: 0.6666666666666666 LR - Precision: 46.73913043478261 - Recall: 50.0 - F-1 Score: 48.31460674157303 - Accuracy: 93.47826086956522 - AUC: 0.5 - sensitivity: 0.0 - specificity: 1.0 - ppv: nan - npv: 0.9347826086956522 - lrp: nan - lrm: 1.0 - youden: 0.0 KNN - Precision: 97.77777777777777 - Recall: 66.6666666666666 - F-1 Score: 73.863636363636 - Accuracy: 95.65217391304348 - AUC: 0.666666666666666 - sensitivity: 0.33333333333 - specificity: 1.0 - ppv: 1.0 - ppv: 1.0 - ppv: 1.0 - ppv: 1.0 - pv: 1.0 - DT 0.33333333333333333 - youden: 0.666666666666666666 SVM NB 0.33333333333333333 - youden: 0.6666666666666666 0.333333333333333337 - youden: 0.66666666666666666 - Precision: 100.0 - Recall: 100.0 - F-1 Score: 100.0 - Accuracy: 100.0 - AUC: 1.0 - sensitivity: 1.0 - specificity: 1.0 - ppv: 1.0 - npv: 1.0 - lrp: inf - lrm: 0.0 - youden: 1.0 XGB - Precision: 100.0 - Recall: 100.0 - F-1 Score: 100.0 - Accuracy: 100.0 - AUC: 1.0 - sensitivity: 1.0 - specificity: 1.0 - ppv: 1.0 - npv: 1.0 - lrp: inf - lrm: 0.0 - youden: 1.0 [LightGBM] [Info] Number of positive: 28, number of negative: 391 [LightGBM] [Info] Auto-choosing col-wise multi-threading, the overhead of testing was 0.000245 seconds. You can set 'force_col_wise=true' to remove the overhead. [LightGBM] [Info] Total Bins 520 [LightGBM] [Info] Number of data points in the train set: 419, number of used features: 10 [LightGBM] [Info] [binary:BoostFromScore]: pavg=0.066826 -> initscore=-2.636503 [LightGBM] [Info] Start training from score -2.636503 [LightGBM] [Warning] No further splits with positive gain, best gain: -inf [LightGBM] [Warning] No further splits with positive gain, best gain: -inf [LightGBM] [Warning] No further splits with positive gain, best gain: -inf [LightGBM] [Warning] No further splits with positive gain, best gain: -inf [LightGBM] [Warning] No further splits with positive gain, best gain: -inf [LightGBM] [Warning] No further splits with positive gain, best gain: -inf [LightGBM] [Warning] No further splits with positive gain, best gain: -inf [LightGBM] [Warning] No further splits with positive gain, best gain: -inf [LightGBM] [Warning] No further splits with positive gain, best gain: -inf [LightGBM] [Warning] No further splits with positive gain, best gain: -inf LGB

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- 5. Evaluating the Model
 - The model evaluation is a step where the performance of the model is tested on previously unseen data.
 - The unseen data used is the testing set that is split from the master dataset before model selection.
 - The performance of the model is evaluated used numerous evaluation metrics in machine learning such as accuracy, precision or positive predictive value (ppv), recall or sensitivity or true positive rate (tpr), negative predictive value (npv), f1 score, specificity, area under the curve (AUC), etc.

						80
	MLP Final Acc: 96.559	LR Final Acc: 94.408	KNN Final Acc: 93.978	DT Final Acc: 96.785	SVM Final Acc: 96.341	60
	Final Prec: 93.234	Final Prec: 67.185	Final Prec: 61.975	Final Prec: 88.800	Final Prec: 93.125	40
	Final Spec: 74.583 Final Rec: 74.583	Final Spec: 58.333 Final Rec: 58.333	Final Spec: 55.000 Final Rec: 55.000	Final Spec: 88.247 Final Rec: 88.247	Final Spec: 72.917 Final Rec: 72.917	20
	Final F1: 80.097 Final AUC: 0.746	Final F1: 60.048 Final AUC: 0.583	Final F1: 55.938 Final AUC: 0.550	Final F1: 86.591 Final AUC: 0.882	Final F1: 78.541 Final AUC: 0.729	MLP LR KNN DT SVM NB RF AB XGB ROC Curve Analysis
	NB Final Acc: 87.405 Final Prec: 83.009 Final Spec: 83.239 Final Rec: 83.239 Final F1: 78.524 Final AUC: 0.832	RF Final Acc: 96.989 Final Prec: 93.964 Final Spec: 83.704 Final Rec: 83.704 Final F1: 86.400 Final AUC: 0.837	ADA Final Acc: 96.998 Final Prec: 90.437 Final Spec: 86.810 Final Rec: 86.810 Final F1: 87.186 Final AUC: 0.868	XGB Final Acc: 97.428 Final Prec: 87.000 Final Spec: 90.144 Final Rec: 81.667 Final F1: 79.810 Final AUC: 0.901	LightGBM Final Acc: 96.984 Final Prec: 91.667 Final Spec: 82.154 Final Rec: 65.000 Final F1: 71.476 Final AUC: 0.822	0.8 0.6 0.6 0.4 0.2 0.2 0.2 0.6 0.4 0.4 0.4 0.4 0.4 0.4 0.4 0.4
ro	m "Machine learning deen dive	hy Delta Analytics"	Fig 14. Mo	del Performance		0.0 0.2 0.4 0.6 0.8 1.0 False Positive Rate

Adapted from "Machine learning deep dive by Delta Analytics"



5. Evaluating the Model

MLP

69.99

80.69

53.75

53.78

53.97

				Performance	evaluation			
NB 84 LR 84 MLP 85 SVM 69 DT 74 RF 82 Promosed 82	acc (%)	pre (%)	rec/sen/TPR (%)	f(%)	MCC	FPR (%)	FNR (%)	TNR (%)
NB	84.07 ± 4.70	84.36 ± 7.85	80.00 ± 9.28	81.61 ± 5.46	0.68 ± 0.10	12.67 ± 7.57	20.00 ± 9.28	87.33 ± 7.57
LR	84.81 ± 4.21	85.49 ± 7.38	80.83 ± 11.81	82.21 ± 5.71	0.70 ± 0.08	12.00 ± 7.18	19.17 ± 11.81	88.00 ± 7.18
MLP	85.56 ± 4.21	86.12 ± 6.52	81.67 ± 12.25	82.99 ± 6.03	0.67 ± 0.12	11.33 ± 6.00	18.33 ± 12.25	88.67 ± 6.00
SVM	69.63 ± 7.37	72.90 ± 11.29	50.83 ± 10.83	59.52 ± 10.17	0.38 ± 0.16	15.33 ± 7.33	49.17 ± 10.83	84.67 ± 7.33
DT	74.81 ± 8.57	74.28 ± 13.61	70.83 ± 12.50	71.39 ± 9.27	0.49 ± 0.17	3.33 ± 12.74	28.33 ± 11.90	76.67 ± 12.74
RF	82.96 ± 8.15	85.15 ± 10.71	75.83 ± 12.05	79.64 ± 9.70	0.68 ± 0.14	12.00 ± 8.33	23.33 ± 11.06	88.00 ± 8.33
	95.90 ± 5.55	97.14 ± 5.71	94.67 ± 11.08	95.35 ± 6.52	0.92 ± 0.10	4.52 ± 6.94	3.33 ± 6.67	95.48 ± 6.94

recall/ sensitivity/ true positive rate, f = f-measure, MCC = Matthews correlation coefficient, FPR accuracy, pre = Precision, rec/sen/TPR -FNR = false negative rate, TNR = true negative rate.

(a)

Model rec/sen/TPR pre (%) f(%)MCC FPR (%) FNR (%) TNR (%) acc (%) (%) NB 83.17 ± 7.64 84.18 ± 9.75 78.79 ± 8.29 81.25 ± 8.29 0.66 ± 0.15 13.12 ± 8.59 21.21 ± 8.29 86.88 ± 8.59 LR 84.85 ± 6.91 86.12 ± 7.85 80.22 ± 9.30 82.90 ± 7.80 0.70 ± 0.14 11.25 ± 6.73 19.78 ± 9.30 88.75 ± 6.73 MLP 84.15 ± 7.76 85.01 ± 9.74 80.22 ± 9.83 82.28 ± 8.66 0.68 ± 0.12 14.37 ± 9.29 18.41 ± 9.22 85.62 ± 9.29 SVM 71.06 ± 6.16 74.65 ± 9.51 59.23 ± 14.88 64.53 ± 9.43 0.43 ± 0.12 18.75 ± 10.46 40.77 ± 14.88 81.25 ± 10.46 DT 74.21 ± 7.29 75.16 ± 8.00 76.09 ± 4.86 74.31 ± 5.18 0.52 ± 0.09 24.38 ± 8.12 27.80 ± 7.31 75.62 ± 8.12 RF 82.14 ± 6.84 83.69 ± 8.63 76.54 ± 10.09 12.50 ± 8.39 22.03 ± 10.27 87.50 ± 8.39 79.63 ± 8.36 0.66 ± 0.13 Proposed 98.40 ± 3.21 98.57 ± 4.29 98.33 ± 5.00 98.32 ± 3.37 1.67 ± 5.00 0.00 ± 0.00 98.33 ± 5.00 0.97 ± 0.06 HDPM

Performance evaluation

(b)

Fig 15. Performance of the machine learning models for predicting heart disease in combination with the Information Gain-based feature selection, DBSCAN-based outlier removal, SMOTE-ENN-based data balancing methods in Cleveland (a) and Statlog (b) datasets [13]

Classification			ormance m			Classification		Perf	ormance m	etric		Classification		Per	formance m	etric		Classification		Perfo	ormance m	etric	
model	p (%)	r (%)	f(%)	acc (%)	AUC	model	p (%)	r (%)	f(%)	acc (%)	AUC	model	p(%)	r(%)	f(%)	acc (%)	AUC	model	n(%)	r (%)	f(%)	acc (%)	AUC
MLP	52.59	48.57	45.08	84.9	0.85	MLP	73.23	97.56	83.56	72.02	0.5	MLP	57.31	69.81	57	54.92	0.53	MLP	67.78	70.49	67.02	80.84	0.89
SVM	88.235	41.096	56.075	81.9	0.62	SVM	73.04	99.17	84.1	72.58	0.45				57								
DT	36.37	42.68	32.18	69.69	0.59	DT	71.35	69.42	70.13	56.96	0.46	SVM	56.38	85.26	67.77	53.48	0.44	SVM	64.49	57.69	56.66	75.85	0.77
LR	52.5	47.14	43.77	84.9	0.91	LR	73.67	95.06	82.77	71.33	0.58	DT	57.33	54.1	54.69	49.96	0.49	DT	61.48	56.43	56.13	72.48	0.75
K-means + LR	91.6	96.4		90.7	0.957	CART [19]	-	58.38	-	-	0.68	LR	61.53	82.76	69.52	59.41	0.62	LR	67.98	77.14	69.67	81.57	0.89
[16]	91.0	90.4	-	90.7	0.937	DBSCAN +						CART [19]	_	45.65	-	-	0.566	DBSCAN +	83.665	84.677	84.168	83.644	
DBSCAN +						SMOTE + RF	78.788	70.270	74.286	76.419	-	Proposed DPM	75.6	81.78	77.12	75.78	0.76	SMOTE + RF [40]	85.005	04.077	04.100	05.044	-
SMOTE + RF	91.497	93.403	92.440	92.555	-	[40]						Proposed DPM	/5.0	01./0	//.12	/5./8	0.76	Proposed DPM	100	100	100	100	1
[40]						Proposed	93.57	84.89	88.8	85.73	0.87							Tioposed DT M	100	100	100	100	
Proposed DPM	94.49	98.62	96.32	96.74	0.99	DPM	95.57	04.07	00.0	05.75	0.07												
(2) (C)												(d))										

(C)(a) (b) 100 100 90 90 80 80 70 70 60 60 50 40 50 30 40 20 30 10 20 0 Before After Before After Before After Before After 10 Precision (%) Recall (%) F1 (%) Accuracy (%) After LR 54.62 59.74 52.81 50.81 51.72 49.51 91.34 91.85 Precision (%) Recall (%) KNN 57.26 68.01 51.46 53.13 50.83 53.39 90.43 90.43 79.13 84.06 75.68 80.01 DT 52.06 52.42 51.77 51.64 50.83 51.17 80.04 80.12 KNN 66.68 78.30 64.68 73.91 DT 71.57 74.22 70.45 73.26 XGB 50.73 52.04 47.92 50.02 48.08 49.20 83.48 88.12 XGB 80.73 82.25 72.54 80.97 (a) (b) 50.00 SVM 45.93 45.93 50.00 47.88 47.88 91.85 91.85 SVM

54.84

90.46

92.11

F1 (%) Accuracy (%) 75.56 80.80 77.47 82.87 64.45 74.27 66.59 77.66 69.81 73.23 71.19 74.96 79.71 81.22 80.88 82.48 80.35 77.28 82.12 72.58 76.98 72.92 77.64 75.31 MLP 79.19 84.12 76.23 80.74 76.20 81.37 77.71 83.05

Fig 17. Performance of the machine learning models for predicting T2D in patient with NAFLD in combination with forward logistic regression in NAGALA (a) and NAFLD (b) datasets [3]

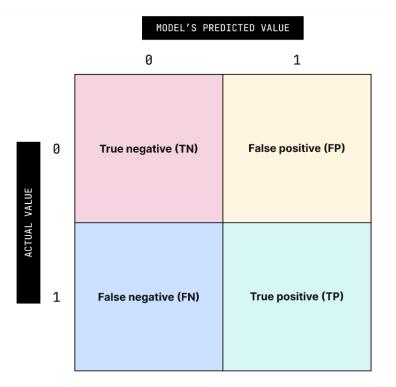
Fig 16. Performance of the machine learning models for predicting T2D and hypertension in combination with the Information Gain-based feature selection, iForest-based outlier removal, SMOTETomekbased data balancing methods [11] in Dr John Schorling (a), Golino et al male hypertension (b), Golino et al female prehypertension (c), and Dr. P. Soundarapandian, M.D., D.M CKD datasets [11]



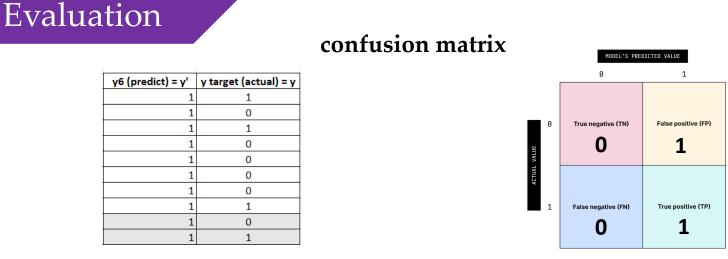
Important

What should be concerned when evaluating the disease prediction model's performance?

- In order to evaluate the classification model's performance, a summarized table called the confusion matrix is used.
- The confusion matrix consists of four categories:
 - **1. True Negative (TN)** represents the number of samples correctly classified or predicted as belonging to the negative class. For example, the actual class is negative (0), and the predicted class is also negative (0).
 - 2. **True Positive (TP)** represents the number of samples correctly classified or predicted as belonging to the positive class. For example, the actual class is positive (1), and the predicted class is also positive (1).
 - **3. False Negative (FN)** represents the number of samples incorrectly predicted as the negative class. For example, the actual class is positive (1), but the predicted class is negative (0).
 - 4. **False Positive (FP)** represents the number of samples incorrectly predicted as the positive class. For example, the actual class is negative (0), but the predicted class is positive (1).
- According to Hicks et al [24], the most commonly used for evaluating the performance of the MLbased disease prediction model are accuracy, recall or sensitivity or true positive rate (tpr), precision or positive predictive value (ppv), negative predictive value (npv), f1 score, Matthew's correlation coefficient (MCC), and threat score (TS).







• Accuracy : percentage of correctly classified samples over the total number of samples. Accuracy measures the overall correctness of the model's predictions.

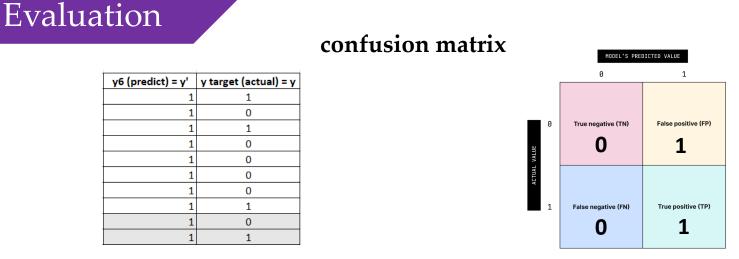
Accuracy=
$$\frac{TN+TP}{TN+TP+FP+FN}$$
 Testing Accuracy = $\frac{0+1}{0+1+1+0}$ = 0.5 = 50 %

 Recall/Sensitivity/TPR : the ratio between correctly classified positive samples and all samples assigned to the positive class [23]. When it's actually yes, how often does it predict yes?

$$\operatorname{Recall} = \frac{TP}{Actual Yes (1)} = \frac{TP}{TP + FN}$$
 Testing $\operatorname{Recall} = \frac{1}{1+0} = 1 = 100 \%$

Performance





Precision/Positive Predictive Value (PPV) : the ratio between correctly classified samples and all samples assigned to that class. When it predicts yes, how often is it correct? Τe

 $Precision = \frac{TP}{Predicted Yes (1)} = \frac{TP}{TP + FP}$

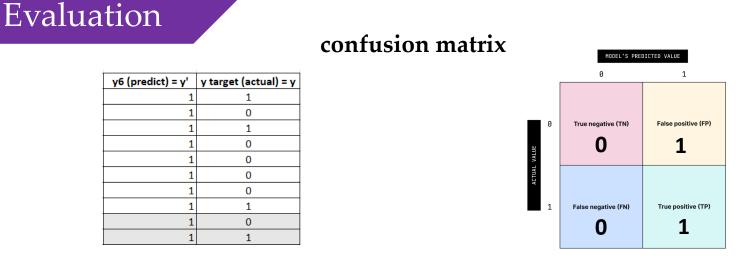
esting Precision =
$$\frac{1}{1+1}$$
 = 0.5 = 50 %

Negative Predictive Value (NPV) : the ratio between correctly classified negative samples and all samples classified as negative. When it predicts no, how often is it correct?

$$NVP = \frac{TN}{Predicted No(0)} = \frac{TN}{TN+FN}$$
 Testing NPV = $\frac{0}{0+0} = \frac{0}{0} = 0 = 0 \%$

Performance





• **F1 score :** represents the harmonic mean or weighted average of precision and recall. A large F1 score of 1 indicates excellent precision and recall, while a low score indicates poor model performance.

F1 score =
$$\frac{2 x \ precision x \ recall}{precision+recall}$$
 Testing F1 = $\frac{2 x \ 0.5 \ x \ 1}{0.5+1}$ = $\frac{1}{1.5}$ = 0.6667 = 66.67 %

• **Specificity/True Negative Rate :** how often the model predicts a negative for a value that is actually negative.

Specificity=
$$\frac{TN}{Actual No(0)} = \frac{TN}{TN+FP}$$
 Testing Specificity = $\frac{0}{0+1} = 0 = 0$ %

Performance

04

Machine Learning-based Model for Disease Prediction Applications



Machine Learning-based Model for Disease





30

Prediction Applications

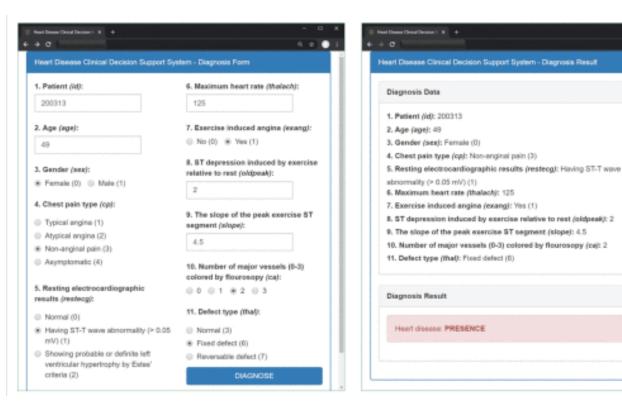
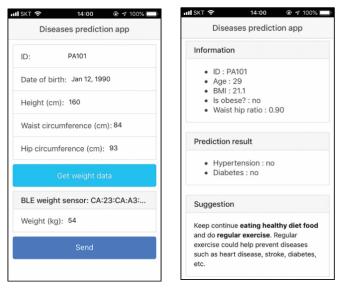


Fig 20. Web-based heart disease clinical support system [3]



8. ú 🕕

Fig 19. Web-based disease prediction application [11]



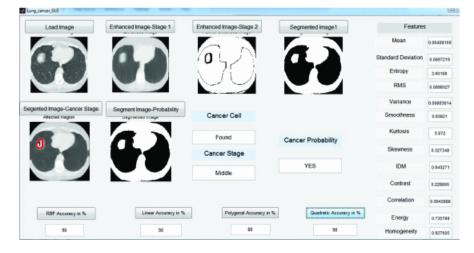
Machine Learning-based Model for Disease



Prediction Applications



(a) Siemens Healthineers - Prediction and early identification of disease: identify potential disease progression in COVID-19 patient [18]



(b) Siemens Healthineers - Prediction and early identification of lung disease [19]



(c) and (d) Health system – detection and prediction lung cancer utilized by medical institutions [17, 20]

Adopted from "Introduction to machine learning by Delta Fig 22 Real-case application of machine learning-based model in healthcare or medical



05 Conclusion



- Machine learning is a powerful tool that can be utilized as one of the alternatives to early detection of the disease.
- By utilizing ML as a disease prediction tool, it could help individuals know their current health status, thus preventing the occurrence of the worst-case scenario.
- Not only in the healthcare or medical domain, but machine learning has also been widely utilized in many other domains.

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Thank you for your attention! Any questions?

Feel free to send any research/project collaboration proposals via norma@sejong.ac.kr

