

To develop intelligent, personalized technologies

Dr. Tom Hueting - Evidencio

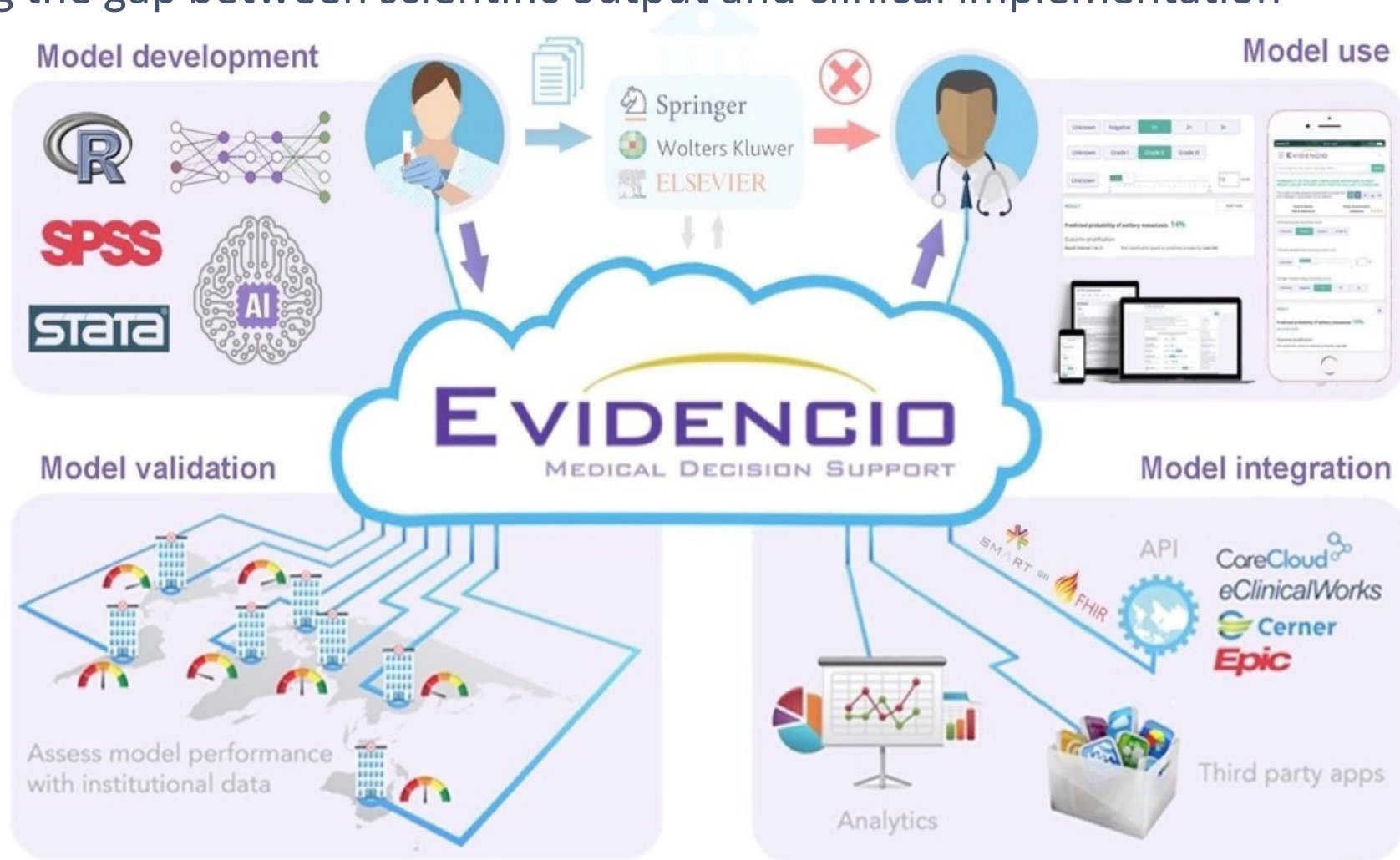
DISCLOSURE SLIDE

Employee at Evidencio B.V.



The Evidencio platform

Bridging the gap between scientific output and clinical implementation



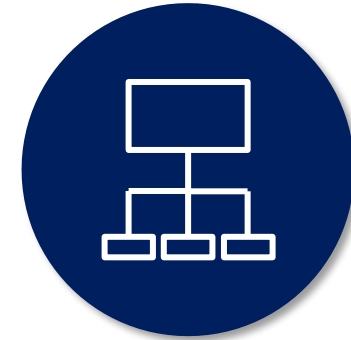
What is a medical algorithm?



Any **calculation, formula,** statistical **survey,** **nomogram** or **look-up table** useful in healthcare.



Can contain **heterogeneous** and **multimodal** data



Include **decision trees** and **tools** for reducing or defining **uncertainty**.

Prediction model lifecycle



Problem definition

Data collection

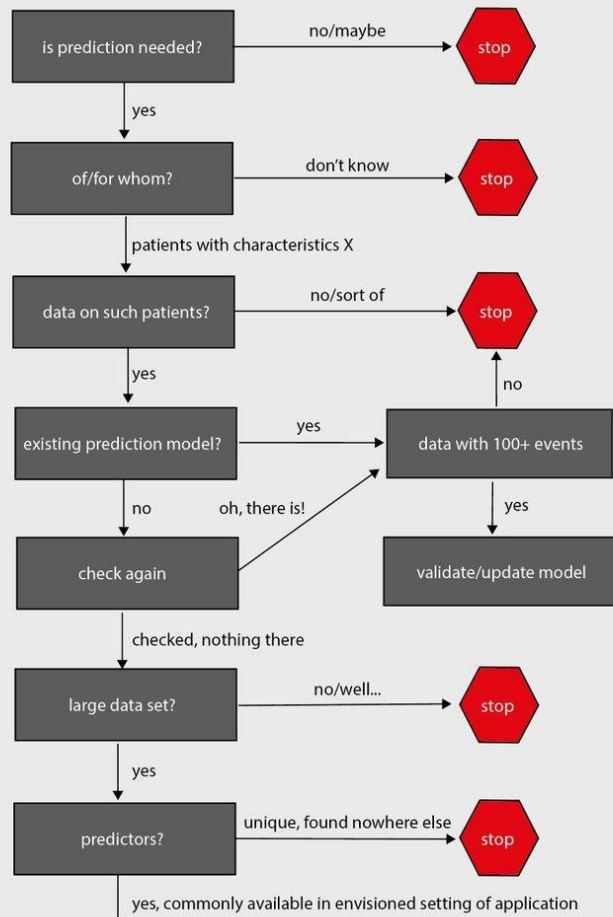
Development

Validation & update

Impact assessment

Clinical adoption

Should a risk prediction model be developed?



proceed: avoid dichotomizing, penalize where possible, do rigorous internal/external validation, study model calibration, think hard about dealing with missing data /andimperfect outcome measurements, don't forget to report everything including your intercept (just follow TRIPOD guideline).



Prediction model

[Advanced](#) [Create alert](#) [Create RSS](#)

Save

Email

Send to

Sort by:

Best match

MY NCBI FILTERS

709,888 results

Page 1

RESULTS BY YEAR



A continual prediction model for inpatient acute kidney injury.

1 Kate RJ, Pearce N, Mazumdar D, Nilakantan V.

Cite Comput Biol Med. 2020 Jan;116:103580. doi: 10.1016/j.combiomed.2019.103580
PMID: 32001013

Share The continual **model predicts** AKI every time a patient's AKI-relevant variable cha
method to comprehensively evaluate the overall performance of a continual **pred**
introduced, and we experimentally show usin ...



New article: **“Development and validation of a prediction model for outcome X in patients with disease Y”**

Development and Validation of a Prediction Model for Hepatitis B Virus-Related Hepatocellular Carcinoma Patients Receiving Postoperative Adjuvant Transarterial Chemoembolization.

Tu X, Zhang J, Li M, Lu F, Wang T, Gong W, Xiang B.

J Hepatocell Carcinoma. 2023 Oct 24;10:1881-1895. doi: 10.2147/JHC.S422565. eCollection 2023.

PMID: 37901717 [Free PMC article.](#)

Development and Validation of a Nomogram for Renal Survival Prediction in Patients with Autosomal Dominant Polycystic Kidney Disease.

Wang X, Zheng R, Liu Z, Qi L, Gu L, Wang X, Zhu S, Zhang M, Jia D, Su Z.

Kidney Dis (Basel). 2023 Jun 6;9(5):398-407. doi: 10.1159/000531329. eCollection 2023 Oct.

PMID: 37901714 [Free PMC article.](#)

Development and validation of a nomogram for predicting pulmonary complications after video-assisted thoracoscopic surgery in elderly patients with lung cancer.

Zhao D, Ma A, Li S, Fan J, Li T, Wang G.

Front Oncol. 2023 Oct 13;13:1265204. doi: 10.3389/fonc.2023.1265204. eCollection 2023.

PMID: 37901337 [Free PMC article.](#)

Development and validation of a nomogram for preoperative prediction of

Box 1: Full PREP prognostic models to calculate the risk of adverse maternal outcomes in women with early onset pre-eclampsia

a. Risk at various time points from diagnosis until 34 weeks' gestation using the survival model (PREP-S) $S_{(t)} = S_0(t)^5 \wedge \exp((\beta_1 * X_1 + \dots + \beta_n * X_n))$ $S_{(t)} = S_0(t) \wedge \exp(-0.031 * \text{maternal age} + 1.514 * (\text{Log}(\text{GA at diagnosis}/10))^{-2} - 0.8345136) + 5.707 * (\text{Log}(\text{GA at diagnosis}/10))^{-2} * \ln(\text{log}(\text{GA at diagnosis}/10)) - 0.0652155) + 0.122 (\text{exaggerated tendon reflexes}) - 0.169 (\text{one pre-existing medical condition}) - 0.384 (\text{two or more pre-existing medical conditions}) + 0.016 * \text{systolic blood pressure} + 0.797 (\text{oxygen saturation} < 94\% \text{ on air}) - 0.002 * \text{platelet count} + 0.126 * \text{log}(\text{alanine amino transferase}) + 0.605 * \text{log}(\text{serum urea})^2 - 0.144 * \text{log}(\text{serum urea})^3 + 0.265 * \text{log}(\text{serum creatinine}) + 0.080 * \text{log}(\text{protein creatinine ratio}) + 0.176 (\text{baseline treatment with any antihypertensive}) + 1.066 (\text{baseline treatment with magnesium sulfate})$ § $S_0(t) - \text{baseline survival adjusted for optimism at time } t$ $S_0(48 \text{ hrs}) = 0.99142, S_0(72 \text{ hrs}) = 0.98542, S_0(1 \text{ week}) = 0.96492, S_0(1 \text{ month}) = 0.87377$ **b. Overall risk by postnatal discharge using the logistic model (PREP-L)** Probability (maternal adverse outcome) = $\exp(X)/(1 + \exp(X))$, Where $X = -1.507 - 0.020 * \text{maternal age} + 12.052 * (\text{log}(\text{gestational age}))^3 - 39.90241) - 7.930 * (\text{log}(\text{gestational age}))^3 * \text{log}(\text{log}(\text{gestational age}) - 49.08188) - 0.330 (\text{if one pre-existing medical condition}) - 0.579 (\text{if two or more pre-existing medical conditions}) + 0.146 * \text{log}(\text{urine protein creatinine ratio}) - 0.951 * (\text{log}(\text{serum urea})^{-1}) - 0.004 * \text{platelet count} + 0.024 * \text{systolic blood pressure} + 0.409 (\text{baseline treatment with antihypertensive}) + 1.252 (\text{baseline treatment with magnesium sulfate})$ Predictor value is 1 when present and 0 when absent



Maternal age 28 Years

Gestational age (weeks) 24 Weeks

Gestational age (days) 1 days

Exaggerated tendon reflexes No Yes

Pre-existing medical condition 0 1 ≥2

PREP-L: Risk of complications in Early-onset Pre-eclampsia ▼
 Predicted risk of adverse event by the time of discharge is: **87.8%**

PREP-S: Risk of complications in Early-onset Pre-eclampsia ▼
 Predicted risk of adverse event by the timepoint indicated is: **39.2%**

Name	Conversion	Option (value)			
Maternal age	Full range	<input checked="" type="checkbox"/> 16 - 45 Years	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Gestational age (weeks)	Full range	<input checked="" type="checkbox"/> 20 - 34 Weeks	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Gestational age (days)	None	<input checked="" type="checkbox"/> 0 - 6 days	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Exaggerated tendon reflexes	None	<input checked="" type="checkbox"/> No (0) Yes (0.122)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Pre-existing medical condition	None	<input checked="" type="checkbox"/> 0 (0) 1 (-0.169) ≥2 (-0.385)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Protein to creatinine ratio (PCR)	Full range	<input checked="" type="checkbox"/> 0 - 5000 mg/mmol	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Serum urea concentration	Full range	<input checked="" type="checkbox"/> 0 - 25 mmol/L	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Platelet count	Full range	<input checked="" type="checkbox"/> 50 - 400 x10 ⁹ /L	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Systolic blood pressure	Full range	<input checked="" type="checkbox"/> 50 - 250 mmHg	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Treatment with antihypertensive drugs	None	<input checked="" type="checkbox"/> No (0) Yes (0.176)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Treatment with magnesium sulphate (MgSO4)	None	<input checked="" type="checkbox"/> No (0) Yes (1.066)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

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b. Overall risk by postnatal discharge using the logistic model (PREP-L) Probability (maternal adverse outcome) = $\exp(X) / (1 + \exp(X))$, Where $X = -1.507 - 0.020 * \text{maternal age} + 12.052 * (\log(\text{gestational age}))^3 - 39.90241) - 7.930 * (\log(\text{gestational age}))^3 * \log(\log(\text{gestational age}) - 49.08188) - 0.330 * (\text{if one pre-existing medical condition}) - 0.579 * (\text{if two or more pre-existing medical conditions}) + 0.146 * \log(\text{urine protein creatinine ratio}) - 0.951 * (\log(\text{serum urea})^{-1}) - 0.004 * \text{platelet count} + 0.024 * \text{systolic blood pressure} + 0.409 * (\text{baseline treatment with antihypertensive}) + 1.252 * (\text{baseline treatment with magnesium sulfate})$ Predictor value is 1 when present and 0 when absent



Presence Of DCIS

Please indicate if any ductal carcinoma in situ (DCIS) is present as assessed on core needle biopsy.

Estimated risk of positive surgical margins:
15%

Histological type

Please indicate the histological type of the primary tumor as assessed on core needle biopsy.

Histological grade

Please indicate the histological grade of the primary tumor as assessed on core needle biopsy.

Estimated risk of positive surgical margins: **15%**

Are the risk predictions accurate?



Show validation data as:

Validation Values Model Labels Clear data

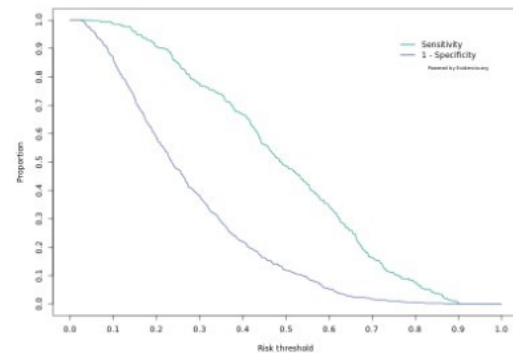
	Preoperative M...	Microcalcifica...	Preoperative N...	Preoperative T...	Density	Palpability	Suspicion of m...	Estrogen recep...	Presence Of DC...	Histological t...	Histological g...	Outcome	Probability
1	No	Present	Negati...	T2	50-75%	Palpa...	Yes	Positive	Present	Ductal	Elston ...	Positive	0.6880
2	Yes	Absent	Negati...	T1	0-25%	Non-P...	No	Negati...	Absent	Other	Elston ...	Negati...	0.0452
3	No	Present	Positive	T1	0-25%	Non-P...	No	Negati...	Absent	Ductal	Elston ...	Negati...	0.1562
4	Yes	Present	Negati...	T1	50-75%	Non-P...	No	Positive	Absent	Ductal	Elston II	Positive	0.1000
5	Yes	Present	Positive	T1	50-75%	Palpa...	No	Negati...	Present	Ductal	Elston II	Negati...	0.1801
6	Yes	Absent	Positive	T1	0-25%	Palpa...	No	Negati...	Absent	Other	Elston II	Negati...	0.0364
7	No	Present	Positive	T1	50-75%	Non-P...	No	Positive	Absent	Ductal	Elston ...	Negati...	0.3318
8	Yes	Absent	Positive	T1	25-50%	Non-P...	No	Positive	Present	Ductal	Elston II	Negati...	0.2800
9	Yes	Absent	Negati...	T1	50-75%	Palpa...	No	Positive	Absent	Ductal	Elston ...	Negati...	0.0774
10	Yes	Absent	Negati...	T1	50-75%	Non-P...	Yes	Negati...	Present	Lobular	Elston I	Positive	0.5532
11	No	Present	Positive	T2	75-10...	Palpa...	Yes	Positive	Absent	Lobular	Elston II	Positive	0.7511
12	Yes	Present	Negati...	T1	50-75%	Non-P...	No	Positive	Present	Ductal	Elston II	Negati...	0.2561
13	Yes	Absent	Positive	T2	50-75%	Non-P...	No	Negati...	Absent	Ductal	Elston ...	Negati...	0.1397
14	Yes	Present	Negati...	T2	50-75%	Non-P...	No	Positive	Present	Ductal	Elston II	Negati...	0.3142
15	No	Absent	Negati...	T2	0-25%	Non-P...	No	Positive	Absent	Other	Elston I	Positive	0.1242
16	No	Present	Negati...	T1	25-50%	Non-P...	Yes	Positive	Absent	Lobular	Elston II	Positive	0.5711
17	Yes	Present	Positive	T1	50-75%	Non-P...	No	Positive	Present	Ductal	Elston ...	Positive	0.4618
18	Yes	Absent	Negati...	T2	75-10...	Palpa...	Yes	Positive	Present	Other	Elston I	Positive	0.4522
19	No	Absent	Positive	T1	25-50%	Palpa...	No	Negati...	Present	Other	Elston I	Negati...	0.2042



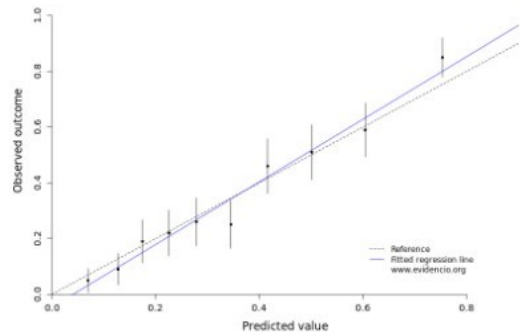
Results

Rows included 1000
 C-index 0.792 | 95% CI: 0.7625 - 0.8204 **ROC**
 Brier score 0.172
 Scaled Brier score 0.24
 Slope 1.123
 Intercept -0.046
 Hosmer-Lemeshow GOF 0.153

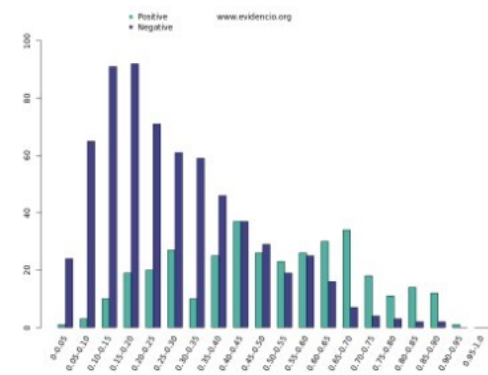
Classification plot



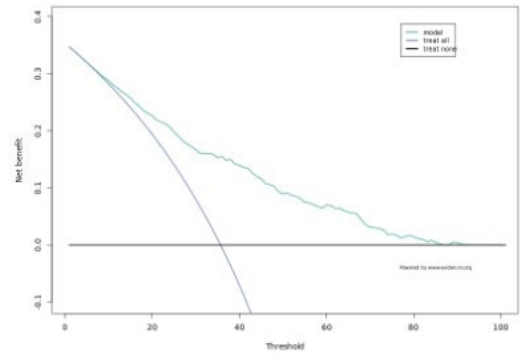
Calibration plot

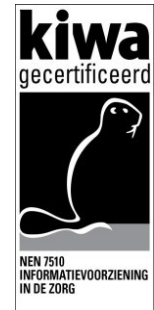


Histogram



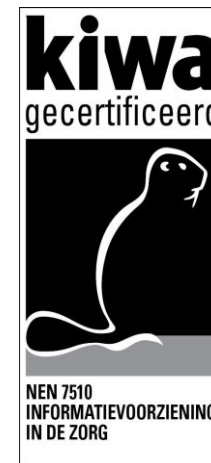
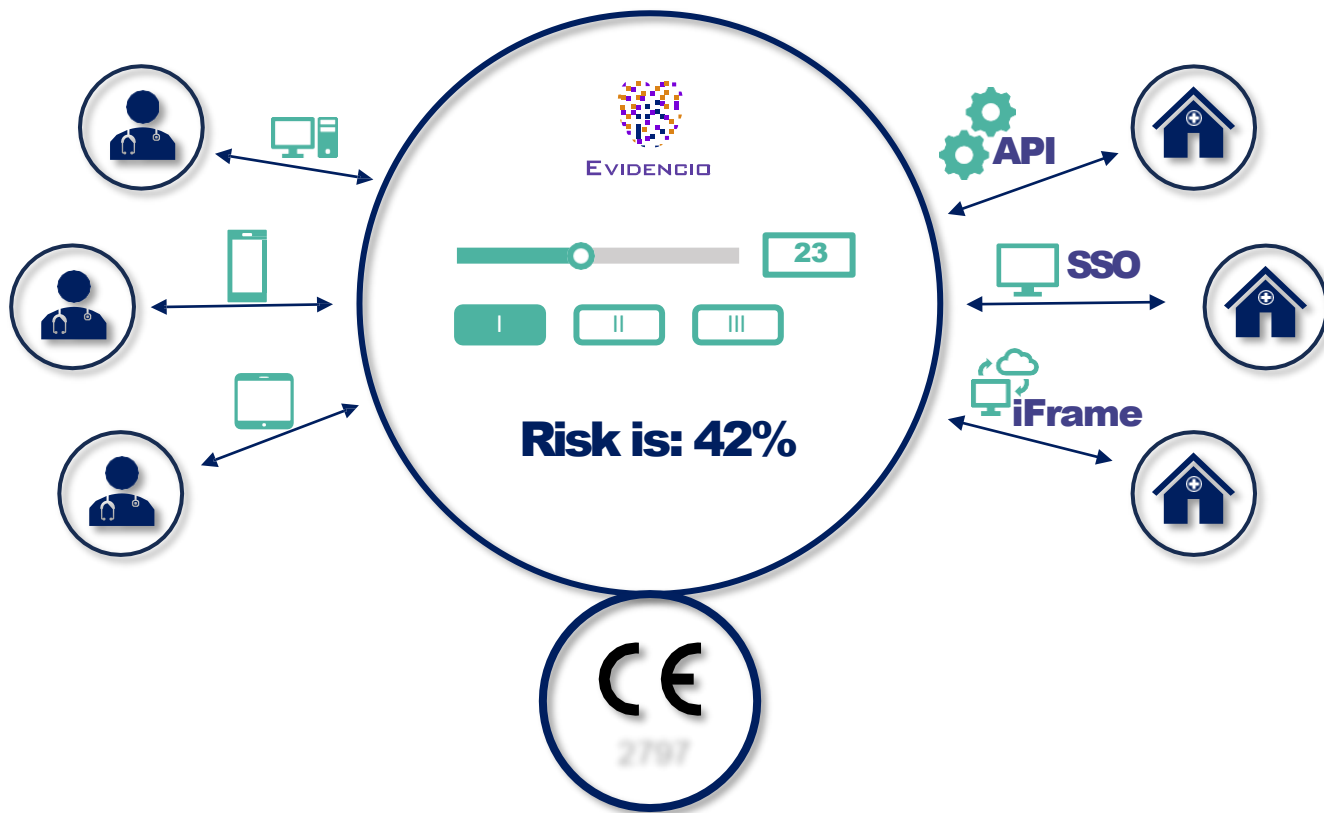
Decision curve





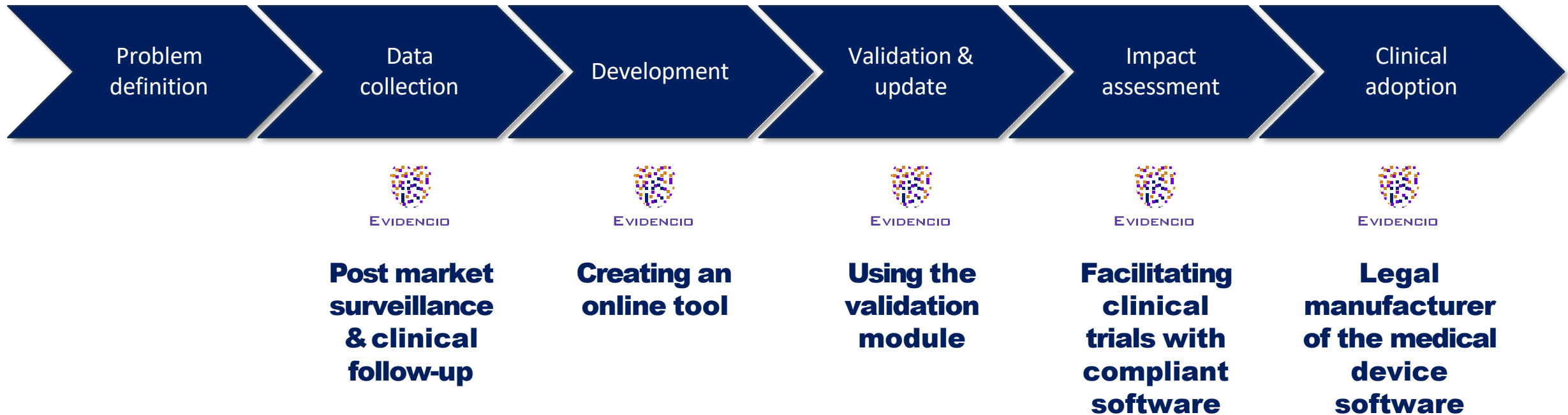
Use the Evidencio platform to assess the impact of your prediction model

Clinical prediction algorithm lifecycle

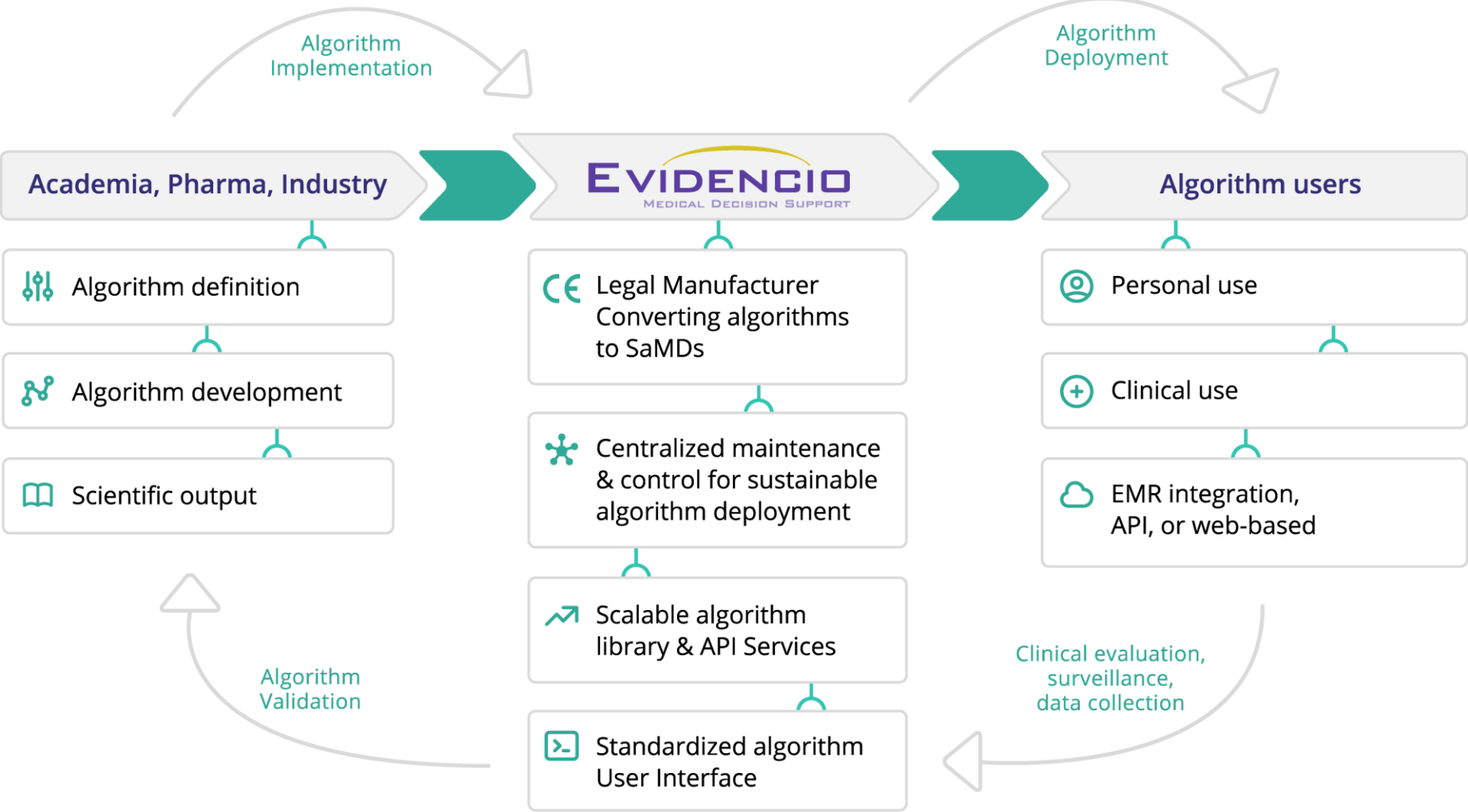


Evidencio can act as a legal manufacturer for your medical device software

Involvement in the prediction model lifecycle



Evidencio's Algorithm library



**A showcase example of the collaboration
with the University of Twente:**

The NABOR project

INFLUENCE 2.0 -> INFLUENCE 3.0

Breast Cancer Research and Treatment (2021) 189:817–826
<https://doi.org/10.1007/s10549-021-06335-z>

EPIDEMIOLOGY



Improved risk estimation of locoregional recurrence, secondary contralateral tumors and distant metastases in early breast cancer: the INFLUENCE 2.0 model

Vinzenz Völkel¹ · Tom A. Hueting^{2,3} · Teresa Draeger¹ · Marissa C. van Maaren^{3,4} · Linda de Munck⁴ · Luc J. A. Strobbe⁵ · Gabe S. Sonke⁶ · Marjanka K. Schmidt⁷ · Marjan van Hezewijk⁸ · Catharina G. M. Groothuis-Oudshoorn³ · Sabine Siesling^{3,4}

the risk for DM. Table 3 gives an overview of the underlying coefficients.

Online calculator

The first DM based on the sex predictions

es; an easy-to-use online risk calculator is a

<https://www.evidencio.com/models/show/2238>.

calculator estimates the risks and the 95% conf

calculator estimates the risks and the 95% conf

Comparison to the original INFLUENCE nomogram and other related prediction models

Compared to the original INFLUENCE nomogram, the INFLUENCE 2.0 model comes with a variety of updates leading to improved flexibility and a broader application range regarding predictable events. Concerning clinical decision-making, discrimination is arguably the most relevant indicator for model performance. The AUC of the five annual prediction models of the original INFLUENCE nomogram which is exclusively concentrating on the endpoint LRR starts with 0.84 for the first year and decreases to 0.62 in the 5th year. A direct comparison to the AUC

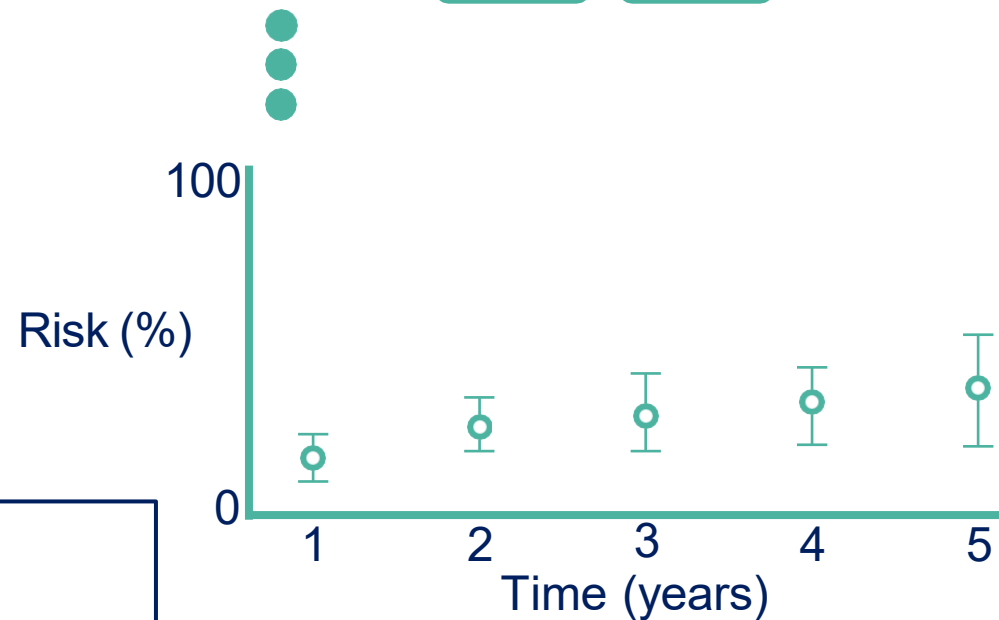
Age **60**

Grade I II III

Tumor stage pT1 pT2 pT3

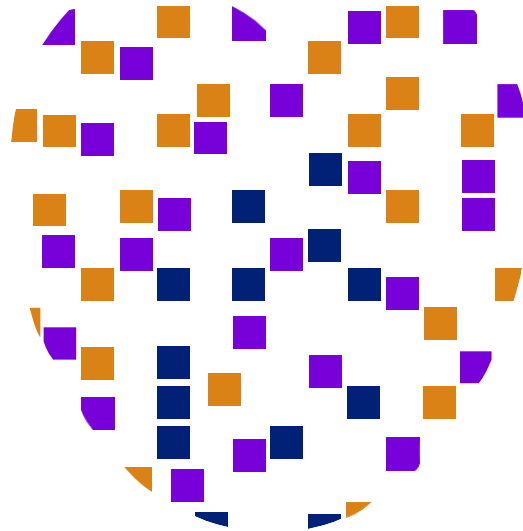
Nodal stage pN0 pN1 pN2 pN3

Multifocality No Yes



LOT V-2.0-2238.21.05.14 **CE**

UDI (01)08720299526440(8012)v2.0(4326)210514(240)2238



EVIDENCIO

Tom@evidencio.com

Dr. Tom Hueting
01 November 2023
TECHMED Event