

The Use Of CTPA In Diagnosing Pulmonary Embolism During Pregnancy And Puerperium Period: A 5 years Retrospective Study At A Regional Referral Hospital in Malaysia

ABSTRACT

Introduction: The cumulative incidence of pulmonary embolism (PE) in South Asia is at 1.03 per 10 000 with mortality rate of 11.1%. Therefore in pregnancy the threshold to perform computed tomography pulmonary angiography (CTPA) is low due to the fear of missing the diagnosis of pulmonary embolism (PE). It is often the pre-test clinical score are bypassed especially the modified Well score, which is widely used in the general population yet not validated in pregnant women. The objectives of our studies are:

1. To assess CTPA ordering trend and its positive rate
2. To evaluate the diagnostic accuracy of modified Wells score (MWS) in predicting PE
3. To identify the rate of contrast-induced nephropathy (CIN) post CTPA

Study design: This is a retrospective study which included all pregnant and postpartum patients who underwent CTPA for suspected PE in Hospital Raja Permaisuri Bainun, Malaysia from Jan 2018 to December 2022. A total of 185 CTPA results were retrieved and MWS were calculated retrospectively: ≤ 4 (unlikely PE) or >4 (likely PE).

Results: The positive rate of CTPA was 20/185 (10.5%). From 185 patients, the sensitivity, specificity, positive predicate value (PPV) and negative predictive value (NPV) of MWS were 90%, 14.5%, 11.3%, and 92.3% respectively, in predicting PE. The receiver operating characteristic (ROC) curve analysis showed a non-discriminating value (0.5). 7.1% of patients had CIN post-CTPA.

Conclusion: The trend of using CTPA in diagnosing PE in pregnant women had been steady despite our finding of its low diagnostic yield. Although Modified Wells score is widely used in the general population, its implications for pregnant patients are still a matter of debate.

Keywords: CTPA, modified Wells score, pulmonary embolism, venous thromboembolism

1. INTRODUCTION

Venous thromboembolism is one of the leading causes of death in pregnant women. The incidence of VTE is about 1-2 per 1000, and the incidence of pulmonary embolism is almost 1 in 10,000^{1,2}. A recent study³ showed the cumulative incidence of pulmonary embolism (PE) in South Asia is 1.03 per 10,000 with a mortality rate of 11.1%. With low prevalence but fatal consequences for mother and fetus, pulmonary embolism remains one of the most common diagnoses in pregnancy. The symptoms of pulmonary embolism often overlap with symptoms of physiological changes occurring during pregnancy, such as shortness of breath or tachycardia. Thus, there is a low threshold to suspect pulmonary embolism due to the fear of missing a PE, leading to overuse of CTPA in our setting.

Over the years, with the advancement of technology, CTPA has become the most widely used and reference standard of diagnostic imaging technique for patients suspected of PE. It is easily accessible in our setting, even bypassing the initial clinical probability assessment.

A study from a single tertiary center in Switzerland⁴ showed an annual 4-fold increase in CTPA examinations in 17 years. A similar result from another center in Africa⁵ showed an increase of 25% per year over the past 10 years. In a paper published in 2008, the PE positive rate for CTPA was only 10%, and they commented on the overuse of CTPA as a screening rather than a diagnostic examination⁶. Interestingly, the PE diagnostic yield from CTPA performed differs geographically⁷. The yield of positive CTPA in Europe is about 20–31%, while in the US it is only about 10%^{8,9,10}. In Malaysia, we are using a threshold of 15.4% as recommended by the Royal College of Radiologists, while other studies in the general Malaysian population quoted about 25–33% of a positive rate¹¹.

In view of the concern about the low yield of PE-positive CTPA, C. Rotzinger⁴ also suggested the need for new diagnostic strategies to safely exclude PE with fewer radiological examinations. With the introduction of pre-test clinical score in one of the large interventional studies¹², the yield of CTPA to diagnose PE has shown an increase from 9.2% to 12.6%. Recently, Medson et al.¹³ showed the CTPA PE yield decreased by about 8–21% when the pre-test clinical score was bypassed. In a meta-analysis¹⁴ showing similar accuracy using different pre-test clinical scores in diagnosing PE. More specifically, Wang et al¹⁵ proved that with Wells criteria, there is a modest increase in CTPA yield in diagnosing PE.

Multiple algorithms have been investigated to identify high-risk patient groups; however, none of those have yet to be validated in obstetrics patients. More recently, pregnancy-adapted Geneva score¹⁶ and YEARS¹⁷ algorithm have been investigated in pregnant women. Without the need for imaging, those algorithms are shown to be able to safely exclude a small set of patients from pulmonary embolism in conjunction with D-dimer level. However, in present guidelines as well as in our setting, D-dimer is rarely ordered to exclude pulmonary embolism in view of the normally raised level in pregnancy, especially in maternal conditions such as preeclampsia, which causes more false positive results^{18,19}.

One of the other commonly used pre-test clinical scores is Modified Well's²⁰ score in order to risk stratify high-risk patients during pregnancy and the puerperium period. There are several studies looking into the diagnostic accuracy of MWS. Back in 2011, a study from Hospital Dublin single tertiary referral hospital where a total 125 woman were investigated. The significant result for patients with MWS scores of 6 has 100% sensitivity and 90% specificity, with a 36% positive predictive value for PE on CTPA²¹. This study also showed that there was

no patient with low MWS who had PE (NPV of 100%). It was the first application of MWS to pregnancy, and the article emphasized that low MWS scores in women suggest that PE could avoid unnecessary investigations by at least one third. Similar studies^{22,23} also suggested a sensitivity of 100% and a specificity of 90% with a high MWS. However, there were a few subsequent studies that concluded CTPA was overused in patients with low and intermediate MWS. For example, Hanieh Raji *et al.*²⁴ found that from 120 patients, the positive CT angiography test in patients with low, intermediate, and high clinical probability was 18, 44, and 82%. Crichlow *et al.*²⁵ studied 152 patients with suspected PE in the Hospital of the University of Pennsylvania and concluded that 13.8% of the CTPA procedures could have been avoided by proper use of Wells/D-dimer. In another study with 575 sample sizes, it was also shown that up to 25% of CTPA scans were unnecessary in those patients with a low or intermediate probability of PE⁶.

Overuse of CTPA certainly imposes a financial burden on the hospital. Furthermore there is a risk of administering iodinated contrast material in pregnant women, causing contrast-induced nephropathy. The Clare O Connor *et al.*⁷ study showed that 14% of the patients who underwent CTPA sustained AKI after CTPA. Thus, there is a need for the proper use of pretest clinical score to reduce the use of CPTA in our pregnant population.

In this study, we are determining the imaging rate of CTPA over the past 5 years and its diagnostic yield. Subsequently, we retrospectively assessed the pre-test clinical score based on modified Wells score (MWS) and evaluated the diagnostic accuracy of MWS in predicting PE. The last part of the research will be looking into the rate of contrast-induced nephropathy in these patients after CTPA.

2. METHODOLOGY

This research has received ethical approval from the National Medical Research Registry of Malaysia (NMRR- ID-23-00208-Q6M). Informed consent was waived owing to the cross-sectional nature of this study.

2.1 STUDY SETTING

Hospital Raja Permaisuri Bainun is a tertiary hospital for the Perak state of Malaysia and provides care for about 670,000 people in Perak. The Obstetrics and Gynecology Department has over 3,000 deliveries per year. There are also a variety of services offered by the radiology department, including CT scans and nuclear medicine.

2.2 STUDY POPULATION

This is a cross-sectional analytical of all pregnant and postpartum patients who underwent diagnostic testing CTPA for suspected PE from January 1, 2018 to December 31, 2022. This study was approved by the Medical Research and Ethics Committee (MREC) of Ministry of Health (MOH).

For a total of 199 studies, 14 CTPAs were excluded in view of a non-diagnostic result or a severely degraded image after being carefully reviewed by the radiologist. All other 185 patients who underwent CTPA in the study timeframe were included. Exclusion criteria of this study were all pregnant or postpartum patient with low suspicious of pulmonary embolism not

requiring further definitive imaging. Also, patients with incomplete medical record **will be** excluded from the studies.

2.3 STUDY DESIGN/DATA COLLECTION

All patient were identified via electronic medical record. The number of scans ordered each year **will** were documented. In this retrospective study, all pregnant or postpartum patients with suspected PE who **has** a diagnostic imaging (CTPA) **will** be retrospectively assessed using MWS and correspond to the CTPA result.

A data collection form used to collect data from patients' electronic record. The collected data included patient's age and trimester, clinical signs and symptoms of DVT (=3 points), an alternative diagnosis is less likely than PE (=3 points), heart rate more than 100(=1.5 points), immobilization for ≥ 3 days or surgery within 30 days (=1.5 points), previous diagnosed PE or DVT (=1.5 points), hemoptysis(=1 points), malignancy (on treatment/treatment in last 6 months or palliative) (=1 point), COVID status with CTPA result.

Table 1: Modified Wells Score

CRITERIA	POINTS
Clinical signs and symptoms of DVT	3 points
An alternative diagnosis is less likely than PE	3 points
Heart rate more than 100	1.5 points
Immobilization for ≥ 3 days or surgery within 30 days	1.5 points
Previous diagnosed PE or DVT	1.5 points
Hemoptysis	1 point
Malignancy	1 point
MWS ≤ 4 -> PE unlikely	
MWS > 4 -> PE likely	

The modified well score (Table 1) was then calculated from a manual review of patients' electronic records for each component of the scoring system by a single observer blinded to

the final diagnosis. The patients are stratified into 2 groups: PE unlikely with a score ≤ 4 and PE likely with scores > 4 . The two categories of patients will be compared with the CTPA outcome.

CTPA results were retrieved from Radiology Information System and classified into positive (presence of PE) or negative (absence of PE). Those indeterminate / non-diagnostic CTPA were further reviewed by experienced radiologist before excluded or included in the studies.

Patients' creatinine values will be identified (baseline and within 72 hours after CTPA). A 25% increase in serum creatinine from baseline within 48-72 hours of contrast administration is defined as contrast- induced nephropathy.

2.4 IMAGING TECHNIQUES

CTPA was acquired with 64- and 128- MDCT helical scanners (Canon and GE Healthcare). Two acquisition methods were used: bolus-tracking and timing bolus. With bolus tracking a region of interest (ROI) is placed over the main pulmonary trunk in the axial image and a few dynamic images are obtained in the same position after the injection of contrast material. When a pre-determined threshold is met (e.g., 100 HU), scanning is initiated. In the timing bolus method, a ROI is placed within the pulmonary trunk after a 20-mL timing contrast bolus is given. 100 mL of nonionic low-osmolar contrast medium was administered via IV (iopromide, iohexol, or iopamidol). Caudocranial 1-mm helical images were acquired from the thoracic inlet to the adrenal glands at 100–120 kV with automodulated mAs. Multiplanar reconstruction were performed. Malaysia Ministry of Health (MoH) recognized radiologists who interpreted the CTPA images. All studies were interpreted on a PACS workstation. The scans were classified as positive (presence of PE) or negative (absence of PE). Less than 15 of the scans were excluded due to indeterminate or non-diagnostic CTPA images within this cohort after being reviewed again by another experienced radiologist during our study.

2.5 DATA ANALYSIS

The number of CTPA orders for the past 5 years **is** recorded and compared as a number and frequency (in%). The CTPA diagnostic yield **is** the percentage of CTPA tests **that are** positive for PE.

Using CTPA as the reference diagnostic test, the diagnostic accuracy of MWS **will** be determined using sensitivity, specificity, positive predictive value, and negative predictive value. Sensitivity **is** defined by the proportion of patients with CTPA-confirmed PE who had a PE-likely probability. Specificity **is** the proportion of patients with negative CTPA who had a PE-unlikely probability. The positive predictive value was the proportion of patients with a PE likely score who had CTPA-confirmed PE. The negative predictive value was the proportion of patients with PE-unlikely probability who had a negative CTPA.

The accuracy was determined using receiver operating characteristic (ROC) curve analysis (SPSS software, version 23.0).

3. RESULTS

A total of 199 CTPAs performed and 14 were excluded due to non-diagnostic images. The number of CTPA ordered **increase** about 39-46% form the year of 2018 and steadily ranging about 39-41 scans were ordered from 2020 to 2022. The positive rate of CTPA was 19/185 (10.3%). (Figure 1)

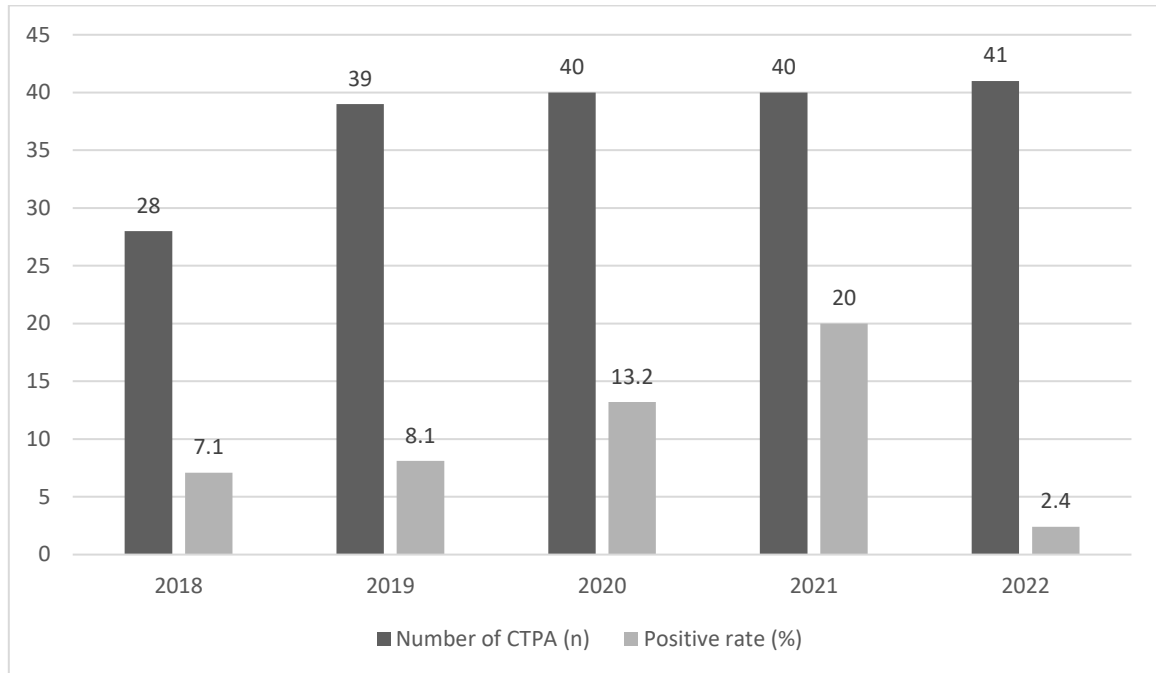


Figure 1: Number of CTPA over the year of 2018-2022 and their positive rate

Among 185 patients, 159 (82%) **was** dichotomized into low probability group and 26 patients (17%) was in high probability category. The percentage of patients diagnosed with PE in these 2 groups **are** 7.6% and 11.3% respectively (Table 2). The positive rate of CTPA is linearly proportional to Well score.

Table 2: Classification of patients and the prevalence of PE in the two probability groups according to the Modified Wells Scores.

	Modified wells score		Total
	Low	High	
All patients, n (%)	26 (17)	159 (82)	185 (100)
Patients with PE, n (%)	2 (7.6)	18 (11.3)	20 (100)

Table 3 Diagnostic accuracy of MWS

Diagnostic accuracy	Sensitivity	Specificity	Positive predictive value	Negative predictive value
Modified Wells score	90%	14.5%	11.3%	92.3%

The sensitivity, specificity, positive predictive value (PPV) and negative predictive value (NPV) of MWS are 90%, 14.5%, 11.3% and 92.3% respectively in predicting PE. The receiver operating characteristic (ROC) curve analysis showed non-discriminating value (0.5) (figure 2)

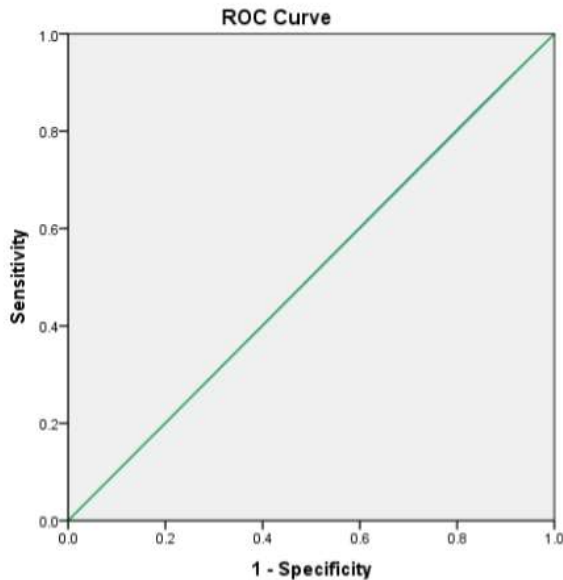


Figure 2: ROC curve analysis of MWS

There is 74% (136 out of 185) has pre-CTPA renal function, and out of 136 of patient only 84 has post-CTPA renal function test done (62%)- About 7.1% of patients had contrast-induced nephropathy post CTPA ($n=84$).

4. DISCUSSION

Despite the low incidence of PE in pregnancy, it remains the second most common cause of maternal death in Malaysia^{26,27}. With the widely available CTPA in tertiary hospitals throughout Malaysia and the significance of timely diagnosis of PE in pregnancy, there is always a low test threshold for diagnostic imaging in this group of patients²⁸. To our knowledge, this is the first study in Malaysia looking into the number of CTPA orders and the diagnostic yield of CTPA in pregnant and puerperium women over 5 years.

We observed a substantial increase in the number of scans ordered from 2019 to 2022. However, the positive rate was in a wide range about 2.4–20%. In the cohort of 2020–2021, the CTPA positive rate was as high as 13.2–20%, with 1/5 of the scans ordered for pregnant

patients with COVID-19. This might be a reflection of the association between thrombotic events and PE in pregnancy during the COVID-19 pandemic²⁹.

Overall, our diagnostic yield was only 10.3%. This diagnostic yield was below the recommendation by Royal College of Radiologists and other studies conducted in the general population. Nonetheless, the finding was in line with a 17-year Swiss study in pregnant population (7%)⁴. This has further validated the challenges of diagnosing PE in pregnancy.

Prior research has indicated that MWS is applicable to pregnancy with notable results demonstrating great sensitivity and specificity with the conclusion of low MWS score could avoid unnecessary imaging by at least one third ^{21,22}. However, the unblinded study with retrospective nature and small sample size rendered the identification of low-risk group less reliable in those studies.

On the other hand, our result revealed a lower sensitivity rate of 90% and specificity of 14.5% which was partially supported by another study³⁰. Low specificity might be contributed by a significant percentage of pregnant women who presented with nonspecific clinical symptoms and signs; in our study, about 82% of patients were classified as high-probability PE group. Pregnancy-related physiological alterations like tachycardia are frequently the most prevalent presenting signs that lead to the suspicion of PE. The utilization of MWS is even more limited during the COVID-19 pandemic, where tachycardia is not uncommon in COVID-19 pregnant women³¹. In addition, one of the score components, "alternative diagnosis less likely than PE, was also debated for its subjectivity causing interobserver variability ^{32,33}. Furthermore, there is also a lack of inclusion of pregnancy-specific variables that may be associated with PE, such as caesarean, delivery, preeclampsia, and infection, in the scoring system. All these highlights highlight the drawbacks of Well's score and the majority of the clinical prediction model scores in this group of patients.

Previous studies also showed that none of the patients with low MWS had PE (NPV 100%)^{21,22}. This is inconsistent with our finding, where the prevalence of PE was 7.6% with an NPV of 92.3% in the low probability group. This outcome can be due to the distinct patient profile, symptoms, and multi-ethnicity population in our study. Out of the 26 patients with low MWS, there are 2 patients diagnosed with PE: one patient has COVID-19, and the other patient's primary symptom was chest pain. The positive rate of CTPA is linearly proportional to the Well score (7.6% for a low score and 11.3% for high risk); however, the receiver operating characteristic (ROC) curve analysis demonstrated poor discriminatory accuracy of MWS in predicting PE in pregnancy. Essentially, it is important to have personalized risk assessment despite low clinical probability.

According to the Malaysia Clinical Practice Guideline of venous embolism, the first step in the algorithm to diagnose PE in pregnancy is by imaging modality³⁴. There is always a risk of contrast-induced nephropathy (CIN) with CTPA. The reported incidence ranged between 6.5 and 19%³⁵. In our studies we discovered that up to 7.1% patient had CIN post-CTPA. All the 6 patients with CIN had either a pre-existing medical condition such as systemic lupus erythematosus, multiple myeloma, or were critically ill. However, in view of the fact that about half of the patients did not have a post-CTPA renal profile as it was not routinely performed, the actual rate might be underestimated.

The strength of the study was the consistent application of MWS by a single reviewer who was blinded by the CTPA result. There were a few limitations in our study: the retrospective assignment of MWS instead of the physician evaluating the patient. This study had a relatively small sample size from a single tertiary center and may not be representative of the population.

In addition, our study did not involve patients with suspected PE who had not undergone CTPA. We did not identify those patients who were suspected to having PE but had venous ultrasonography done instead especially when they had lower limb symptoms. However, we believe that we included those patients in our study as it is one of the components of MWS, so the diagnosis of DVT without CTPA has little effect on our study.

5. CONCLUSION

CTPA has been the first diagnostic modality in diagnosing PE and the diagnostic yield of CTPA performed at our facility has been comparable with larger world studies.

Modified Well's score is widely used in the general population but its implications for pregnant patients are still a matter of debate. With the lack of validated pretest clinical scores in this specific patient category, it is improbable that we are overusing CTPA when a prompt diagnosis is necessary for this fatal disorder. To avoid the unnecessary risk of CTPA to pregnant mothers, future studies on large prospective cohorts are needed to look into the safety and efficacy of other pretest clinical scores to rule out PE in pregnant women.

ETHICAL APPROVAL

This research has received ethical approval from the National Medical Research Registry of Malaysia (NMRR- ID-23-00208-Q6M). Informed consent was waived owing to the cross-sectional nature of this study.

REFERENCES

1. Cantwell, R.; Clutton-Brock, T.; Cooper, G.; Dawson, A.; Drife, J.; Garrod, D.; Harper, A.; Hulbert, D.; Lucas, S.; McClure, J.; *et al.* Saving Mothers' Lives: Reviewing maternal deaths to make motherhood safer: 2006–2008. The Eighth Report of the Confidential Enquiries into Maternal Deaths in the United Kingdom. *BJOG* 2011, 118 (Suppl. 1), 1–203.
2. Heit, J.A.; Kobbervig, C.E.; James, A.H.; Petterson, T.M.; Bailey, K.R.; Melton, L.J., 3rd. Trends in the incidence of venous thromboembolism during pregnancy or postpartum: A 30-year population-based study. *Ann. Intern. Med.* 2005, 143, 697–706.
3. Tan TC, Goh CMY, Tan SSX, Tan LK, Yang Y, Lee LH. Epidemiology of pregnancy-associated pulmonary embolism in South Asian multi-ethnic country: Mortality trends over the last four decades. *J Obstet Gynaecol Res.* 2021;47(1):174-183. doi:10.1111/jog.14450
4. Rotzinger DC, Dunet V, Ilic V, Hugli OW, Meuli RA, Schmidt S. Pulmonary embolism during pregnancy: a 17-year single-center retrospective MDCT pulmonary angiography study. *Eur Radiol.* 2020 Mar;30(3):1780- 1789
5. Lazarus E, Debenedictis C, North D, Spencer PK, Mayo-Smith WW. Utilization of imaging in pregnant patients: 10-year review of 5270 examinations in 3285 patients—1997–2006. *Radiology.* 2009;251:517–24

6. Costantino MM, Randall G, Gosselin M, Brandt M, Spinning K, Vegas CD. CT angiography in the evaluation of acute pulmonary embolus. *AJR Am J Roentgenol*. 2008;191(2):471-474. doi:10.2214/AJR.07.2552
7. Dalen JE, Waterbrook AL. Why are nearly all CT pulmonary angiograms for suspected pulmonary embolism negative? *Am J Medicine*. 2017;130:247–8.
8. Douma RA, le Gal G, Söhne M, et al. Potential of an age-adjusted D-dimer cut-off value to improve the exclusion of pulmonary embolism in older patients: a retrospective analysis of three large cohorts. *BMJ*. 2010;340:c1475.
9. Van Belle A, Büller HR, Huisman MV, et al; Christopher Study Investigators. Effectiveness of managing suspected pulmonary embolism using an algorithm combining clinical probability, D-dimer testing and computed tomography. *JAMA*. 2006;295(2):172-179.
10. Penalzoza A, Verschuren F, Meyer G, et al. Comparison of the unstructured clinical gestalt, the Wells Score, and the revised Geneva score to estimate pretest probability for suspected pulmonary embolism. *Ann Emerg Med*. 2013;62:117-124.
11. Low C, Kow R, Abd Aziz A, et al. (June 15, 2023) Diagnostic Yield of CT Pulmonary Angiogram in the Diagnosis of Pulmonary Embolism and Its Predictive Factors. *Cureus* 15(6): e40484. DOI 10.7759/cureus.40484
12. Prevedello, Luciano M et al. "Does clinical decision support reduce unwarranted variation in yield of CT pulmonary angiogram?." *The American journal of medicine* vol. 126,11 (2013): 975-81. doi:10.1016/j.amjmed.2013.04.018
13. Medson K, Yu J, Liwenborg L, Lindholm P, Westerlund E. Comparing 'clinical hunch' against clinical decision support systems (PERC rule, wells score, revised Geneva score and YEARS criteria) in the diagnosis of acute pulmonary embolism. *BMC Pulm Med*. 2022;22(1):432. doi:10.1186/s12890-022-02242-1
14. Ceriani E, Combescure C, Le Gal G, et al. Clinical prediction rules for pulmonary embolism: a systematic review and meta-analysis. *J Thromb Haemost*. 2010;8(5):957-970. doi:10.1111/j.1538-7836.2010.03801.x
15. Wang RC, Bent S, Weber E, Neilson J, Smith-Bindman R, Fahimi J. The impact of clinical decision rules on computed tomography use and yield for pulmonary embolism: a systemic review and meta-analysis. *Ann Emerg Med*. 2016;67(6):693-701.e3.
16. Robert-Ebadi H, Elias A, Sanchez O, Le Moigne E, Schmidt J, Le Gall C, et al. Assessing the clinical probability of pulmonary embolism during pregnancy: The Pregnancy-Adapted Geneva (PAG) score. *Journal of Thrombosis and Haemostasis*. 19 (12): 3044–3050, 2021
17. Van der Pol LM, Tromeur C, Bistervels IM, Ni Ainle F, van Bommel T, Bertolotti L, et al. Pregnancy- adapted YEARS algorithm for diagnosis of suspected pulmonary embolism. *N Engl J Med* 2019;380:1139– 49.
18. Murphy N, Broadhurst DI, Khashan AS, Gilligan O, Kenny LC, O'Donoghue K. Gestation-specific D-dimer reference ranges: a cross-sectional study. *BJOG*. 2015;122(3):395-400. doi:10.1111/1471-0528.12855

19. Pinheiro Mde B, Junqueira DR, Coelho FF, et al. D-dimer in preeclampsia: systematic review and meta-analysis. *Clin Chim Acta*. 2012;414:166-170. doi:10.1016/j.cca.2012.08.003
20. Wells PS, Anderson DR, Rodger M, et al. Excluding pulmonary embolism at the bedside without diagnostic imaging: management of patients with suspected pulmonary embolism presenting to the emergency department by using a simple clinical model and d-dimer. *Ann Intern Med*. 2001;135(2):98-107. doi:10.7326/0003-4819-135-2-200107170-00010
21. O'Connor C, Moriarty J, Walsh J, Murray J, Coulter-Smith S, Boyd W. The application of a clinical risk stratification score may reduce unnecessary investigations for pulmonary embolism in pregnancy. *J Matern Fetal Neonatal Med* 2011;24:1461–4.
22. Cutts BA, Tran HA, Merriman E, Nandurkar D, Soo G, DasGupta D, et al. The utility of the Wells clinical prediction model and ventilation-perfusion scanning for pulmonary embolism diagnosis in pregnancy. *Blood Coagul Fibrinolysis* 2014;25(4):375–8.
23. Parilla BV, Fournogerakis R, Archer A, Sulo S, Laurent L, Lee P, et al. Diagnosing Pulmonary Embolism in Pregnancy: Are Biomarkers and Clinical Predictive Models Useful? *AJP Rep* 2016;6(2):e160–4.
24. Raji, Hanieh et al. "Overuse and underuse of pulmonary CT angiography in patients with suspected pulmonary embolism." *Medical journal of the Islamic Republic of Iran* vol. 32 3. 4 Feb. 2018, doi:10.14196/mjiri.32.3
25. Crichlow A, Cuker A, Mills AM. Overuse of computed tomography pulmonary angiography in the evaluation of patients with suspected pulmonary embolism in the emergency department. *Acad Emerg Med*. 2012;19(11):1219-1226. doi:10.1111/acem.12012
26. Kaur J, Singh H. Maternal Health in Malaysia: A review. *Webmed Central Public Health* 2011; 2: WMC002598. [https:// doi.org/10.9754/journal.wmc.2011.002598](https://doi.org/10.9754/journal.wmc.2011.002598)
27. Ministry of Health Malaysia , Prevention & treatment of thromboembolism in pregnancy and puerperium A training Manual, 2018
28. Kline JA, Richardson DM, Than MP, Penaloza A, Roy PM. Systematic review and meta-analysis of pregnant patients investigated for suspected pulmonary embolism in the emergency department. *Acad Emerg Med*. 2014;21(9):949-959. doi:10.1111/acem.12471
29. Servante J, Swallow G, Thornton JG, et al. Haemostatic and thrombo-embolic complications in pregnant women with COVID-19: a systematic review and critical analysis. *BMC Pregnancy Childbirth*. 2021;21(1):108. Published 2021 Feb 5. doi:10.1186/s12884-021-03568-0
30. Touhami O, Marzouk SB, Bennisr L, et al. Are the Wells Score and the Revised Geneva Score valuable for the diagnosis of pulmonary embolism in pregnancy?. *Eur J Obstet Gynecol Reprod Biol*. 2018;221:166-171. doi:10.1016/j.ejogrb.2017.12.049
31. Askary E, Poordast T, Shiravani Z, et al. Coronavirus disease 2019 (COVID-19) manifestations during pregnancy in all three trimesters: A case series. *Int J Reprod Biomed*. 2021;19(2):191-204. Published 2021 Feb 21. doi:10.18502/ijrm.v19i2.8477

32. Leclercq MG, Kruip MJ, Mac Gillavry MR, Van Marwijk Kooy M, Büller HR. Observer variability in the assessment of clinical probability in patients with suspected pulmonary embolism. *J Thromb Haemost.* 2004;2(7):1204-1206.
33. Testuz A, Le Gal G, Righini M, Bounameaux H, Perrier A. Influence of specific alternative diagnoses on the probability of pulmonary embolism. *Thromb Haemost.* 2006;95(6):958-962. doi:10.1160/TH06-02-0114
34. Ministry of Health, Clinical Practice Guideline: Prevention and Treatment of Venous Thromboembolism, 2013
35. Kooiman J, Klok FA, Mos IC, et al. Incidence and predictors of contrast-induced nephropathy following CT-angiography for clinically suspected acute pulmonary embolism. *J Thromb Haemost.* 2010;8(2):409-411. doi:10.1111/j.1538-7836.2009.03698.x

UNDER PEER REVIEW