



EARLY ONSET PREECLAMPSIA

THE POWER
TO KNOW
SOONER

EARLY ONSET PREECLAMPSIA: A CLINICAL DILEMMA

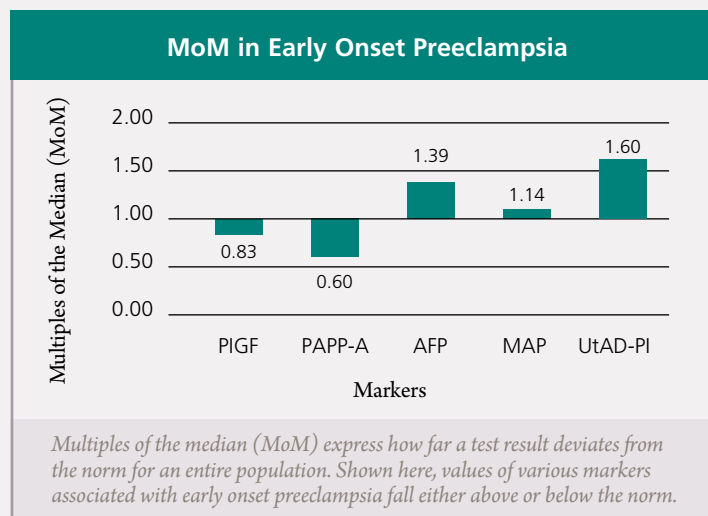
Early onset preeclampsia is defined as preeclampsia resulting in the delivery of the fetus before 34 weeks' gestation. It is estimated to occur in approximately 0.5% of all pregnancies.¹⁻⁵ Early onset preeclampsia is less common than the late form of the disorder but contributes more to the morbidity and mortality of pregnant mothers and babies.¹⁻⁵

Possible early intervention options for preeclampsia include increased monitoring, modified activity, bed rest and medications, including aspirin.⁵⁻¹⁰ To date, there has been no reliable way to detect early onset preeclampsia, particularly for patients experiencing their first pregnancy. Therefore, there has been no satisfactory method with which to stratify patients at high risk of early onset preeclampsia from those at lower risk. Until now.



PreeclampsiaScreen™ | T1. The Power to Know Sooner.

PreeclampsiaScreen™ | T1 is a first-of-its-kind serum screening test that detects three biochemical markers in the mother's blood: PAPP-A (pregnancy-associated plasma protein-A); PIGF (placental growth factor) and AFP (alpha fetoprotein). Together, these three biochemical markers can contribute to accurate prediction of risk for early onset preeclampsia.^{3,5,10-16}



PreeclampsiaScreen™ | T1 enables accurate detection of early onset preeclampsia risk, allowing for earlier intervention and management of the pregnancy.

Earlier Prediction for Earlier Intervention.

Adding PreeclampsiaScreen™ I T1 to your current screening protocol can enhance your ability to identify at-risk patients sooner so that more may be done to prevent or delay onset.

THE BENEFITS:

- **Highest sensitivity in a biochemical screen.** Provides a detection rate of 60% at a 5% false positive rate for early onset preeclampsia when combined with patient history and demographics.¹⁷
- **91% sensitivity when combined with 2 additional biophysical markers.** Detection rate as high as 91% at a 5% false positive rate for early onset preeclampsia when combined with two additional biophysical markers: uterine artery Doppler pulsatility index (UtAD-PI) and mean arterial blood pressure (MAP).¹⁷
- **Practical.** Helps you identify those patients who may require increased vigilance.
- **Convenient.** Can be ordered at the same gestational age/patient visit as first trimester aneuploidy screening.
- **Timely.** Results can be obtained as early as 10 weeks, 2 days' gestation.

PreeclampsiaScreen™ I T1 can enhance the detection rate of your current screening protocol.

60%


Biochemistry
+ History

77%

Biochemistry
+ History
+ MAP

91%

Biochemistry
+ History
+ MAP + UtAD-PI



Widen the Window of Opportunity, for Your Patients at Risk of Early Onset Preeclampsia.

Early detection of early onset preeclampsia allows a valuable window of opportunity for vigilance and intervention.¹⁶ First trimester identification of risk for preeclampsia enables:

- Early identification of asymptomatic high-risk group of patients
- Increased surveillance of high-risk pregnancies
- Wider-ranging intervention possibilities, including increased monitoring, modified activity, bed rest and medications, including aspirin

Don't wait and wonder. Know and act sooner, with PreeclampsiaScreen™ I T1.

For more information, contact your PerkinElmer Labs/NTD sales representative or visit/call us at www.ntdlabs.com/preeclampsia, 1-888-NTD-LABS (683-5227).

Increase Your Vigilance, Guide Your Treatment.

Combined with two biophysical markers, PreeclampsiaScreen™ I T1 can greatly increase your ability to detect patient risk for early onset preeclampsia.

TEST SPECIFICATIONS FOR EARLY ONSET PREECLAMPSIA, COMBINATIONS OF MARKERS + HISTORY

	Biochemistry + History	Biochemistry + History + MAP	Biochemistry + History + UtAD-PI	Biochemistry + History + MAP + UtAD-PI
Markers	PIGF, PAPP-A, AFP	PIGF, PAPP-A, AFP, MAP	PIGF, PAPP-A, AFP, UtAD-PI	PIGF, PAPP-A, AFP, MAP, UtAD-PI
Gestational age (ultrasound dated)	10 weeks, 0 days – 13 weeks, 6 days	11 weeks, 1 day – 13 weeks, 6 days	11 weeks, 1 day – 13 weeks, 6 days	11 weeks, 1 day – 13 weeks, 6 days
Detection rate at 5% FPR*	60%	77%	82%	91%
Requirements	5 ml maternal serum in SST (red/gray speckled or gold) tube or red top tube	5 ml maternal serum in SST tube or red top tube, MAP measurement	5 ml maternal serum in SST tube or red top tube, UtAD-PI measurement	5 ml maternal serum in SST tube or red top tube, MAP and UtAD-PI measurement

*False positive rates are representative of a general screening population. Rates may vary depending on history. Risk cutoff in each case is 1 in 50.

About PerkinElmer Labs/NTD

The PreeclampsiaScreen™ I T1 testing service is offered by PerkinElmer Labs/NTD, a pioneer in screening for aneuploidy and open neural tube defects (ONTDs).

- Known for the highest first trimester detection rates available for biochemical Down syndrome screening¹⁸
- Long recognized as a leading innovator with more than 30 years' experience in prenatal screening

Learn more. For more information, contact your PerkinElmer Labs/NTD sales representative or visit/call us at:

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Pursuant to applicable federal and/or state laboratory requirements, PerkinElmer Labs/NTD has established and verified the accuracy and precision of its testing services.

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