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Risk of preeclampsia and small for gestational age according to antioxidants levels throughout pregnancy

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Review question(s)

To conduct a comprehensive systematic literature review to address the question of whether, among pregnant women, the levels of maternal antioxidants at different time points throughout pregnancy are associated with risk of preeclampsia or SGA.

Searches

Our search strategy will initially be developed in Medline and then adapted to the other databases, as described below. A number of other databases will be searched in order to identify relevant studies. We will assess our search strategy by its ability to identify records of known studies and we will further refine it after preliminary searches to include all relevant controlled vocabulary (indexing subject heading including MeSH terms in Medline and Emtree terms in Embase) and free text words in the title or abstract.

The final search strategy will include searches of MEDLINE, EMBASE, ProQuest Dissertations & Theses, CAB Abstracts, CINAHL, FSTA Direct, BIOSIS Previews, and POPLINE databases.

Search terms will include but are not limited to the following: The MeSH terms “antioxidants” [MeSH] (includes “tocopherols” [MeSH], “vitamin E” [MeSH]), “carotenoids” [MeSH], “vitamin A” [MeSH], “ascorbic acid” [MeSH], “preeclampsia” [MeSH], “infant, small for gestational age” [MeSH], and “fetal growth retardation” [MeSH] will be used. Other search terms that will be combined using appropriate Boolean operators include “antioxidant*”, “tocopherol*”, “vitamin E”, “carotenoid*”, “lycopene”, “cryptoxanthin”, lutein, “vitamin A”, “retinol”, “vitamin C”, “ascorbic acid”, “ascorbate”, “preeclampsia”, “pre-eclampsia”, “toxemia”, “small for gestational age”, “SGA”, “fetal growth retardation”, “fetal growth restriction”, “FGR”, “intrauterine growth retardation”, “intra-uterine growth retardation”, “intrauterine growth restriction”, “intra-uterine growth restriction”, and “IUGR.”

Types of study to be included

We will include prospective longitudinal studies (including RCTs where antioxidants levels were measured in the placebo arm), nested case-control studies, case-control studies, and cross-sectional studies. We will include cross-sectional studies of mothers of SGA infants (as SGA can only be reliably diagnosed postpartum) that recruited women at the time of delivery.

Condition or domain being studied

Preeclampsia is a hypertensive disorder that affects 2-8% of pregnancies worldwide, and is most common in first pregnancies. Although it can be diagnosed from the 20th week of pregnancy, most cases occur near term. Preeclampsia is a poorly understood and heterogeneous disorder that is typically diagnosed by the presence of high blood pressure and protein in the urine. It is a multisystem disorder, which may affect the brain, lungs, kidney, and liver. Pregnant women with preeclampsia are at risk for severe complications including seizures (eclampsia), multi-organ failure, stroke, and death. Risks to the fetus include preterm birth, suboptimal growth resulting in small for gestational age (SGA) neonate, and stillbirth.

Small for gestational age (SGA) is defined as the birth of an infant at less than the 10th percentile of gestational-age-

and sex-specific birth weight, according to a given population reference. Hence, SGA is often the result of intrauterine growth restriction (IUGR), also known as fetal growth restriction (FGR), but may also include some constitutionally small babies. The prevalence of SGA is, by definition, approximately 10% among live births. SGA shares a similar risk factor profile to preeclampsia and potentially shared underlying pathology.

Participants/ population

We are interested in pregnant women with preeclampsia or SGA risk factors such as pre-existing disease that increases preeclampsia or SGA risk, personal or family history of preeclampsia, or abnormal ultrasound findings, as well as in women taking part in population-based studies where the outcomes of SGA and preeclampsia were recorded.

Intervention(s), exposure(s)

We are interested in maternal blood levels of non-enzymatic antioxidant levels including tocopherols (vitamin E), carotenes (alpha-carotene, beta-carotene and lycopene), oxy-carotenoids (lutein, zeaxanthin, cryptoxanthin), retinol (vitamin A), and ascorbate (vitamin C). We will include studies of maternal antioxidant levels measured from samples obtained at any point before delivery, or samples taken shortly after delivery. We will not assess cord blood levels or maternal samples taken >72 hours after delivery. However, we believe that our search strategy will capture these studies, which will enable us to briefly summarize them in a section on excluded studies. We will exclude studies published prior to 1970, when high-performance liquid chromatography technology used for precise measurement of many antioxidant biomarker levels was developed.

Comparator(s)/ control

Control subjects will be pregnant women who did not develop preeclampsia or fetal growth restriction in the course of pregnancy or deliver a small for gestational age baby.

Outcome(s)

Primary outcomes

Primary outcomes of interest include preeclampsia and SGA or intrauterine/fetal growth restriction documented by serial ultrasound scans.

Secondary outcomes

Secondary outcomes include severe preeclampsia (including eclampsia), early-onset preeclampsia (onset

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