

## The Mab013 data product

<b>Original number of samples</b>	2,994
<b>Number of samples (per 27.11.2023)</b>	2,987
<b>Number of unique participants</b>	2,968
<b>Biological sample type</b>	Plasma
<b>Participant type(s)</b>	MoBa mothers
<b>Collection timepoint</b>	Gestational week ~17
<b>Case-control selection criteria</b>	None
<b>Biomarker type(s)</b>	Carotenoids
<b>Original reference article</b>	<a href="#">Kelsey et al. 2022</a>
<b>Analytical method(s)</b>	HPLC
<b>Related MoBaBIO product(s)</b>	Mab011, Mab012, Mab014, Pro003, Pro004
<b>FHI Project number(s)</b>	PDB1440

## The project that generated these data

### **Norwegian Environmental Biobank, part I: The importance of nutritional status for the effect of heavy metals on the health of mothers and their children (MoBa-ETox)**

*Project lead: Line Småstuen Haug*

This project formed the first part of the establishment of a Norwegian environmental biobank. The overarching goal of the Norwegian environmental biobank is to monitor levels of nutrients, environmental toxicants, and other unwanted substances in the body over time and examine how these substances affect our health. MoBa-ETox aims to obtain knowledge about nutritional and heavy metal status during pregnancy in the Norwegian Mother, Father and Child Cohort Study (MoBa), and to investigate what significance this may have for subsequent health outcomes in mothers and children. There will be a special focus on whether nutritional status can protect against the negative effects of unwanted environmental substances. The project uses biological samples and questionnaire data from the MoBa to analyze the amount of a selection of nutrients, essential elements and heavy metals in existing MoBa samples from the 2nd trimester of pregnancy, describe the results and assess these in relation to established recommendations and acceptable intakes, and investigate the importance of specific nutrients (vitamins and essential elements) and heavy metals for the risk of developing health problems in later life.

### **Study population**

The original Mab013 biomarker data source is based on plasma samples from **2,975 mothers** in MoBa who were pregnant in 2002-2008. Mothers were eligible for inclusion if they had completed questionnaires 1–6, if data were available from the father's questionnaire, if they had available blood and urine samples collected in pregnancy, and if they had genetic data available in MoBa. Mothers were ineligible for inclusion based on exclusion criteria applied for genotyping, which included participants who were not registered in the Medical Birth Registry, plural pregnancies, and pregnancies with children with autism, suspected autism, or symptoms of severe language delay. For a more detailed overview of the participant selection procedure in this study, refer to [Caspersen \*et al.\* 2019](#).

### **Available biomarker measures (variable names in bold)**

$\alpha$ -Carotene (**AlphaCarotene**)  
 $\alpha$ -Cryptoxanthin (**AlphaCryptoxanthin**)  
 $\beta$ -Carotene (**BetaCarotene**)  
 $\beta$ -Cryptoxanthin (**BetaCryptoxanthin**)  
Unidentified Carotene (**UnidentifiedCaro**)  
Lycopene (**Lycopene**)  
 $\gamma$ -Carotene (**GammaCarotene**)  
Total Lutein (**TotalLutein**)

Retinol (**Retinol**)  
 $\alpha$ -Tocopherol (**AlphaTocopherol**)  
 $\gamma$ -Tocopherol (**GammaTocopherol**)

## Biological sampling and processing

Non-fasting blood samples were collected from mothers at 17-18 weeks' gestation into ethylenediaminetetraacetic acid (EDTA) tubes, centrifuged within 30 minutes, and temporarily placed in a refrigerator at 4 °C. They were shipped from the collecting hospital overnight to MoBa's biobank at the Norwegian Institute of Public Health (NIPH). The samples most often arrived at the biobank within 1–2 days of blood donation, where EDTA plasma were aliquoted onto polypropylene microtiter plates (96-well format, 300  $\mu$ L per well), sealed with the use of heat-sealing foil sheets, and placed in long-term storage at  $-80$  °C.

For more information on biological sampling, processing and storage, please refer to the original reference articles for NIPH's biobank by [Rønningen \*et al.\* 2006](#) and [Paltiel \*et al.\* 2014](#).

## Analytical methodology

Retinol (vitamin A), carotenoids, and tocopherols (vitamin E) were analyzed by **high-performance liquid chromatography (HPLC)** using an Agilent HPLC 1260 system with a diode array detector (Agilent Technologies Inc., Santa Clara, CA, USA). This system detects carotenoids at 450 nm, except  $\beta$ -lycopene which was detected at 472 nm. Tocopherols were detected at 292 nm and retinol at 326 nm. For more information, refer to the article by [Kelsey \*et al.\* 2022](#).

For more detailed information of the methods used in this study, you may refer to the specific methods description documentation developed by the project study group in MoBa-ETox. This will be provided to approved studies in accompaniment of biological datasets.

### Measurement units:

Concentration in mg/L for all measures.

### Limit of quantification (LOQ):

$\alpha$ -Carotene (AlphaCarotene\_Conc): 0.020 mg/L  
 $\alpha$ -Cryptoxanthin (AlphaCryptoxanthin\_Conc): 0.009 mg/L  
 $\beta$ -Carotene (BetaCarotene\_Conc): 0.030 mg/L  
 $\beta$ -Cryptoxanthin (BetaCryptoxanthin\_Conc): 0.018 mg/L  
Unidentified Carotene (UnidentifiedCaro\_Conc): 0.009 mg/L  
Lycopene (Lycopene\_Conc): 0.029 mg/L  
 $\gamma$ -Carotene (GammaCarotene\_Conc): 0.008 mg/L  
Total Lutein (TotalLutein\_Conc): 0.048 mg/L  
Retinol (Retinol\_Conc): 0.165 mg/L

$\alpha$ -Tocopherol (AlphaTocopherol\_Conc): 5.647 mg/L  
 $\gamma$ -Tocopherol (GammaTocopherol\_Conc): 0.167 mg/L

## Published articles using Mab013

*This section also includes articles related to study design, sampling, and data collection.*

- ❖ Kelsey PT, Papadopoulou E, Borge TC, et al. Ultra-processed food consumption and associations with biomarkers of nutrition and inflammation in pregnancy: The Norwegian Environmental Biobank. *Front Nutr.* 2022 Dec 8;9:1052001.
- ❖ Caspersen IH, Thomsen C, Haug LS, et al. Patterns and dietary determinants of essential and toxic elements in blood measured in mid-pregnancy: The Norwegian Environmental Biobank. *Sci Total Environ.* 2019 Jun 25;671:299-308.

## Restrictions for use

None currently known.

## Acknowledgements recommended for use

We recommend that any use of these data in analyses that are presented in peer-review publications acknowledges the original articles describing sampling and data collection:

Kelsey PT, Papadopoulou E, Borge TC, et al. Ultra-processed food consumption and associations with biomarkers of nutrition and inflammation in pregnancy: The Norwegian Environmental Biobank. *Front Nutr.* 2022 Dec 8;9:1052001.

## Disclaimer

The data in Mab013 that are available for use are provided by MoBa on an *as is* basis as they were received from the generating laboratory and have not been curated or quality controlled prior to release. FHI does not provide any guarantees related to data quality and assurance of the original dataset. We reserve the right to periodically remove samples from the dataset belonging to participants who have retracted their consent to participate in this cohort study, and may alter the contents of the associated documentation accordingly.