





BabySEQ, a Non-Invasive Prenatal Testing method, is a new generation screening that screens microdeletion diseases as well as many chromosomal diseases such as Trisomy 21 (Down Syndrome), Trisomy 18 (Edward Syndrome), Trisomy 13 (Patau Syndrome) in the fetus without any invasive procedures that may cause risk. Currently, NIPT-based tests such as BabySEQ are the most successful tests known for screening these diseases. BabySEQ, which is performed in our center, is an increasingly preferred screening method in recent years since it is easy and it does not contain any invasive risks. BabySEQ includes two separate panels according to your preference.

## **All Chromosomes Panel**

- Trisomy 21 (Down Syndrome)
- Trisomy 18 (Edward Syndrome)
- Trisomy 13 (Patau Syndrome)
- Gonadal Chromosome
- Investigation of All Other Chromosomal Anomalies.

## **Extended Panel**

It screens microdeletion diseases in addition to the All Chromosomes panel.

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