Isolated Ectopia Cordis in a twin fetus: a case report
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-Introduction
Ectopia cordis is a rare congenital abnormality with a sporadic incidence of 6 per one million live births and may be confused with Cantrell pentad which include other malformations.

-Case Report
We report a case of a twin fetus with a complete isolated ectopia cordis in a dichorionic diamniotic twin gestation. There is no consanguinity, family history of abnormalities, IVF or first trimester medication. Diagnosis of this isolated anomaly was done at 23 weeks in one fetus while the other had normal morphology; The pregnancy was complicated by preterm labor at 28 w and cesarean delivery followed at 33 w. The affected baby weighed 1800g and had an outcome complicated by episodes of desaturation and deceased at day 8. Parents did not wish in-utero termination of pregnancy nor postnatal surgery for reintegration in the thorax and chose palliative care. Videos available.

-Conclusion:
Isolated ectopia cordis can occur in twin pregnancy with a variable outcome.