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Generalised Anasarca due to Pre-eclampsia: A Rare Case

J. B. Sharma, M. Malhotra, Shalini Jain, R. Arora

Department of Obstetries & Sannaccology, Maniana Azad Medical College & Associated Lok Nanak Hospital, New Delb., 1999, June.

I reclampsia is an important cause of maternal and perinatal mortality and morbidity and can manifest in many ways involving many organs of the body like kidney liver fungs, brain, heart, uterus and placenta. Oedema is an important component of pre-eclampsia. We are presenting a case of gross and generalised anasarca in a patient of pre-eclampsia.

Case Report

Mrs 1.B a 25 years old third gravida presented at 13 weeks of pregnancy with gross swelling all over the body that had been gradually worsening for the last 2 months. She was also complaining of decreased urmary output. There was history of feeling sleepy and mild breathnessness on exertion. No history of headache, blurring of vision or epigastric pain. No history of urinary symptoms, tever or flank pain. She was an unbooked case but had tetanus immunisation.

In obstetric history, she had two normal vaginal deliveries at home 6 and 4 years back. No history of pre-ecoampsia at that time. First child died at age of 5 months due to pneumonia. Other child was alive and healthy. No significant past or medical history.

On Examination

Coneral condition was average. She was conscious cooperative, well oriented to time, place, and person Weight 70kg. Pulse 88 beats per minute. BP 170/110mm Fig. Mild pallor present. There was gross and generalised oedema all over the body. She could not open her eyes due to gross oedema. Chest and cardiovascular system were normal on clinical examination.

Abdominal examination revealed gross abdominal wall oedema along with ascites. Uterus was 32 weeks, betus was lying in longitudinal lye with cephalic presentation. Fetal heart was heard with difficulty.

Investigations Hemoglobin Platelets 2,20 000 си пъс 2) albummura Urine examination 24ms Blood urea 5.266 5. Uric acid S. Creatinine Line SGOT **SGPT** 5. Alkaline phosphate 4 K Yunii S. bilirubin 0.2002 Bilateral pleural ethision X-Ray chest Ultrasound abdomen: Single live fetus, cephalic presentation. Estimated total weight 2.1kgs. Both maternal kidneys were normal.

Diagnosis of severe pre-eclampsia with generalised ansarca was made as blood tests and ultrasound examination ruled out basic renal patholog. Patient was given 600 mg phenytom (Dilantic intravenously with sublingual nitedipine 10 mg (Deputand orally nitedipine 10 mg 8 hourly. Decision to terminate the pregnancy was made and induction of labour done with cerviprime gel followed by aminiotonic and oxytocin drip. She delivered vaginally 8 hours later a male baby weighing 2.2 kgm with Apgar 8-10 at 1 minute and 9\10 at 5 minutes.

She continued to have high blood pressure 150×110 to 160×120 postnatally along with origina and albuminuria and was continued on oral phenyton: frusemide, nitedipine and methyldopa. She improved with this treatment and oedema gradually subsided in 2 week time.

She and her baby were discharged in good condition.

Although generalised oedema can be pair of severe pre-eclampsia, such generalised anasarca mimicking a renal disease is rare. The patient should be managed energetically for severe pre-eclampsia along with termination of pregnancy and patient improvegradually.