Massive cerebral air embolism in a preterm with fetal alcohol syndrome

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A mother with history of alcoholism during pregnancy delivered a baby boy weighing 1660 gms at 33rd weeks of gestation. The baby had features of fetal alcoholic syndrome (FAS) in the form of facial dysmorphic features, bilateral optic atrophy, and sensorineural hearing loss. The baby was put on continuous positive airway pressure (CPAP) as he developed respiratory distress syndrome (RDS) with which he had improvement in his respiratory status. On the seventh day he developed focal clonic seizures with secondary generalization. Computerized tomography (CT) brain revealed disseminated air collection in both the cerebral hemispheres [Figure 1] suggestive of cerebral air embolism (CAE). Review of other radiological investigations did not reveal any evidence of systemic air embolism. Repeat brain CT done two days later showed extensive bihemispherical infarcts [Figure 2]. He was given supportive care and discharged on stabilization. At 3 months follow-up the child had profound developmental delay and intractable epilepsy.

Cerebral air emboli are often micro-emboli, however massive CAE with survival have been documented. CAE has been reported rarely in the neonates receiving positive pressure ventilation for RDS. Damaged pulmonary vascular integrity due to relatively higher pulmonary inflation pressure in RDS is supposed to facilitate the entry of air into systemic circulation, including the cerebral circulation. Recent studies suggest that prenatal exposure of the fetus to alcohol may have deleterious effect on the development of cerebral vasculature. This baby had been exposed to alcohol during prenatal period. Whether such exposure had any possible role in the development of massive CAE in this baby is speculative. This theory is tempting and only neuropathological studies in FAS may through a light.

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