

## Congenital anomalies

Deskripsi Lengkap: <https://lib.ui.ac.id/detail?id=91142&lokasi=lokal>

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### Abstrak

Tuberous sclerosis complex (TSC) is an autosomal dominant disorder caused by mutations of either of the two tumor suppressor genes, TSC1 and TSC2, encoding hamartin and tuberin, respectively.