SUPPLEMENTARY TABLE S2

A case report of syndromic multinodular goitre in adolescence: Exploring the phenotypic overlap between Cowden and DICER1 syndromes

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Supplementary Table S2: DICER1 syndrome-related diseases

Tumour	**Abbreviation**	**Typical age of onset**
Pleuropulmonary blastoma	PPB	~ 0-6 years
Cystic nephroma	CN	0-4 years
Ovarian sex cord-stromal tumour (esp. Sertoli-Leydig cell tumour)	SCST (SLCT)	~ 2-45 years
Multinodular goitre	MNG	~ 5-40 years
Ciliary body medulloepithelioma	CBME	~ 8-15 years
Nasal chondromesenchymal hamartoma	NCMH	~ 6-21 years
Cervical embryonal rhabdomyosarcoma	cERMS	~ 10-20 years
Pituitary blastoma	PitB	< 24 months
Pineoblastoma	PinB	~ 2-24 years
DICER1 anaplastic sarcoma of the kidney	D1 ASK	~ 1-20 years
Differentiated thyroid carcinoma	DTC	~ 5-40 years
Ovarian embryonal rhabdomyosarcoma	oERMS	?
Well-differentiated fetal lung adenocarcinoma	WDFA	?

References: Foulkes WD, Priest JR, Duchaine TF 2014 DICER1: mutations, microRNAs and mechanisms. Nat Rev Cancer **14**:662-672; de Kock et al. 2016 Germline and somatic *DICER1* mutations in a well-differentiated fetal adenocarcinoma of the lung. J Thorac Oncol **11**: e31-3.