

- 22q11.2 Allele in 1,053 Subjects with 22q11.2 Deletion Syndrome Reveals Modifiers of Conotruncal Heart Defects [J]. *Am J Hum Genet*,2019,106(1): 26-40.
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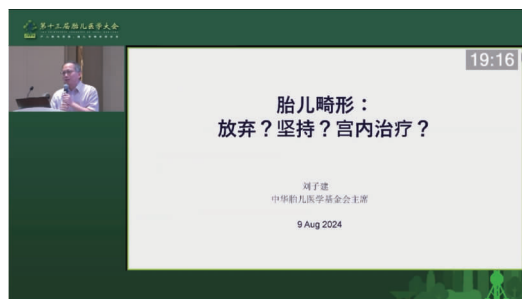
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· 视频导读 ·

胎儿畸形:放弃? 坚持? 宫内治疗?

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在第十三届中国胎儿医学大会上,针对从事产前诊断的专业人士在每天的临床咨询工作中常常经历的“灵魂拷问”,家庭在面对胎儿畸形时如何抉择,是放弃? 坚持? 还是考虑宫内治疗? 中华胎儿医学基金会(CFMF)主席刘子建教授以此为主题做了生动的讲座。随着产前诊断技术的发展,临床医生对“胎儿异常”的态度逐渐发生改变。在遗传咨询过程中,需综合考虑胎儿预后、家庭社会因素以及出生后可能面临的社会压力。

可以通过建立罕见病的非政府组织,帮助罕见病患者更好地融入社会。

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