Genomics Education Programme

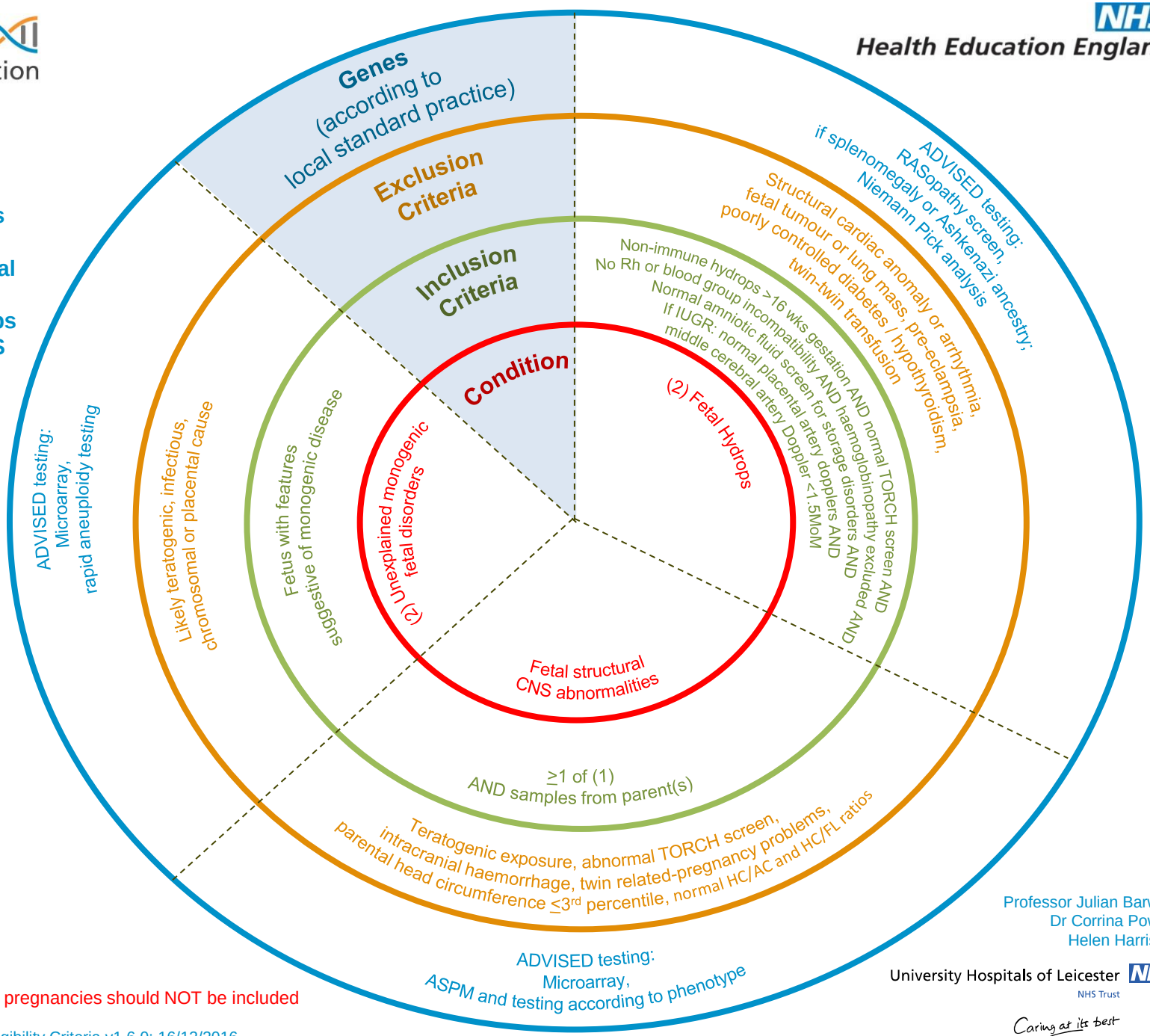
NHS
Health Education England

100,000 Genomes Project
Eligibility wheels
A tool for clinicians

Rare Diseases: Fetal Disorders – monogenic, hydrops and structural CNS abnormalities

(1)

- Bilateral neuronal migration abnormality, confirmed on MRI / pathology
- Cerebellar vermis agenesis / hypoplasia <5th percentile and / or bilateral cerebellar hemisphere hypoplasia / atrophy
- Absent cavum septum pellucidum or holoprosencephaly or complete / partial agenesis of the corpus callosum
- Severe ventriculomegaly (posterior ventricle >15mm) , non-obstructive and normal spine anatomy
- Head circumference <3rd percentile



(2) Samples from ongoing pregnancies should NOT be included

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Caring at its best