

Pharmacogenomics: Considerations for aminoglycoside prescribing

AMT Network Event

16 November 2021

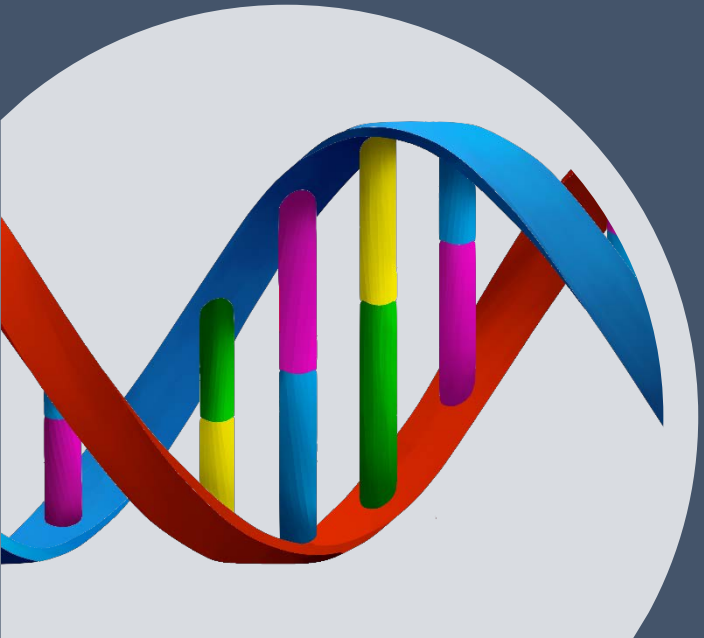
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Pharmacogenomics

the influence of genes on an individual's response to medicines



Clinical applications

Stroke medicine-
clopidogrel

Cancer chemotherapy-
DPYD

Companion diagnostics-
siponimod



Key considerations for implementation

Testing strategy – who, when, how

Equitable access to testing

Workforce development

Governance/policy

Integration of genomics data with clinical systems

Patient engagement/information needs

The aminoglycoside story.....

MHRA Drug Safety Update – Jan 2021; “Aminoglycosides: increased risk of deafness in patients with mitochondrial mutations”

m.1556G>A and ↑ risk of sensorineural deafness

Risk vs benefit – implications for clinical practice.



The m.1555G>A mutation

- Mitochondrial DNA mutation (mtDNA)
- Mutated ribosome resembles bacterial ribosome, offering binding site for AG & ↑ risk
- Most common – affects ~0.2% population
- Associated with SNHL, risk ↑ with aminoglycoside exposure
- One of at least 5 identified – association with SNHL of ototoxicity





Where an aminoglycoside is required in the management of acute or life threatening infection
TREATMENT MUST NOT BE DELAYED

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New rapid genetic test could prevent antibiotic-related hearing loss in newborns

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