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Huntington disease and other polyglutamine diseases : using CAG repeat variations to explain missing heritability

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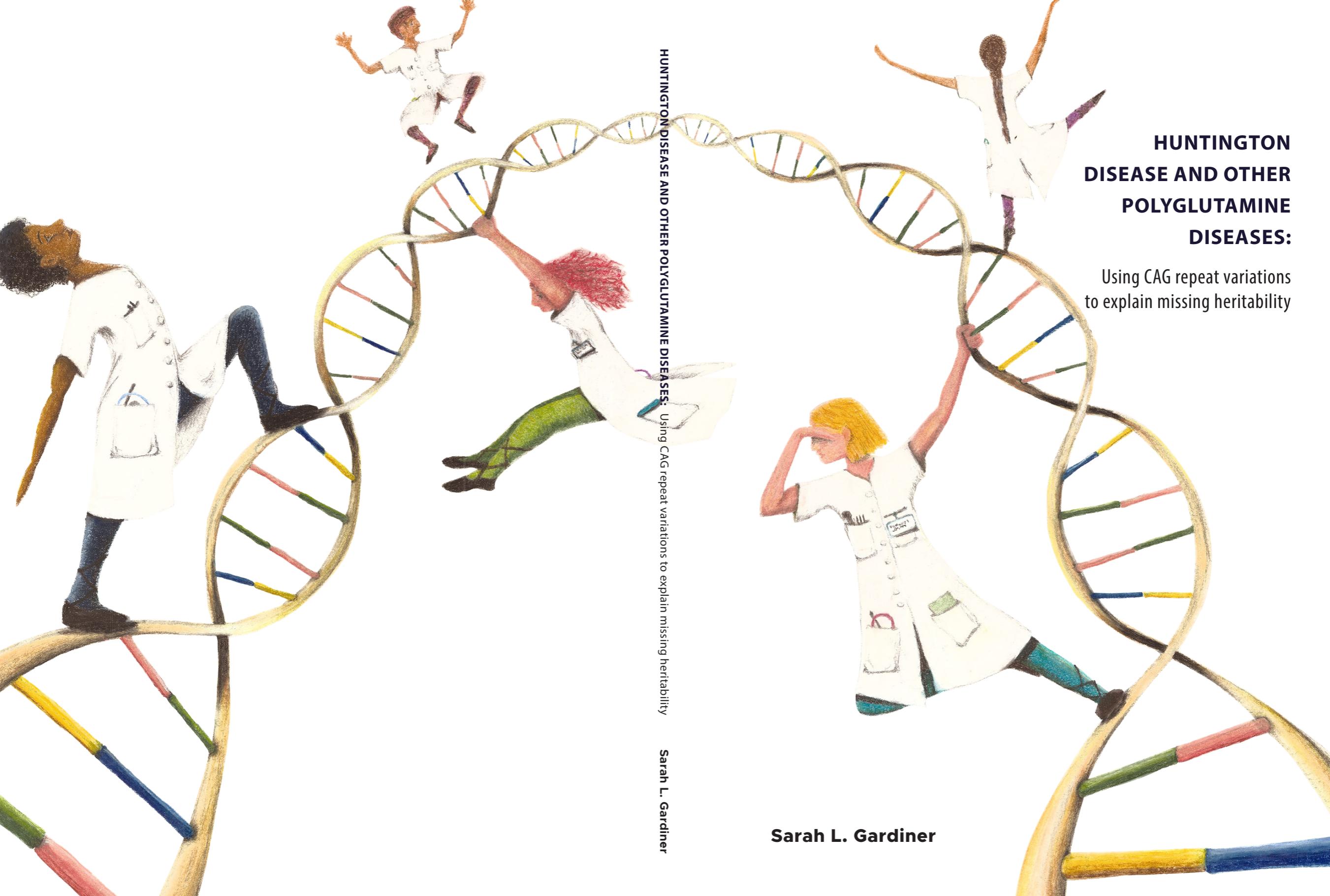


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