

Master of Science Thesis
The University of Western Ontario



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Chapter 1

Introduction

Antimicrobial resistance (AMR) is the natural phenomenon by which microorganisms acquire defences against harmful chemicals within their environment. All microorganisms have the potential to acquire AMR under various environments, including terrestrial and aquatic ecosystems, agri-food, hospitals, and the human body. In the context of an infection, resistant microorganisms limit available treatment options. Acquired antimicrobial resistance (AMR) is estimated to have caused 5,400 Canadian fatalities in 2015 — a number which is expected to climb to 13,700 deaths per year by 2050, while disproportionately affecting populations that are at a greater risk of acquiring infections (Council of Canadian Academies, [2019](#)).

AMR is acquired in response to any chemical that imposes a selective pressure on the microorganism. AMR is a huge burden to global healthcare systems, economies, and societies. As Buongermino Pereira et al. ([2020](#)) has said...

Chapter 2

Methods

2.0.1 C1I gene cassette sequencing

2.0.1.1 DNA isolation from soil microplots

Total genomic DNA was isolated from 2019 Day 30 post-application soil microplots using the DNeasy PowerSoil Kit (Qiagen) with an input of 250 mg of dry soil. The DNA was eluted in 100 uL of 10 mM Tris-HCl, and Nanodrop readings were taken to assess quality. The isolated DNA was stored at -20°C until it was used as template for gene cassette PCR.

2.0.1.2 Gene cassette PCR

Class 1 integron gene cassettes were amplified using primers and thermocycling conditions described by Stokes et al. (2001), except with 33 and 34 bp oligonucleotides ligated onto the 5' ends (Figure XX). The purpose of these oligonucleotides was to provide more distance between the 5' end of the PCR product (lost during library preparation) and the desired gene cassette sequence. DNA from each soil microplot was used as template for five technical replicates of 25 uL reactions (125 uL total). PCR product was analyzed by gel electrophoresis and visualized under UV light. Technical replicates were pooled together and PCR product was purified using the GenepHlow PCR Cleanup Kit (Geneaid). Cleaned PCR product was quantified using the Qubit dsDNA HS Assay Kit (Fisher Scientific). PCR and cleanup were repeated as described above using the same template DNA for a total of two technical replicates for each soil microplot.

2.0.1.3 Library preparation

The Nextera XT DNA Library Preparation Kit (Illumina) was used to prepare the library for sequencing by following the manufacturer's protocol. The indexed PCR product was quantified using the Qubit dsDNA HS Assay Kit and sized using the High Sensitivity DNA Kit on a Bioanalyzer 2100 (Agilent). Individual libraries were diluted to 10 nM, and 15 uL of each diluted library was pooled into a single library for 2 x 125 bp sequencing at Sick Kid's Hospital (Toronto, ON) on a HiSeq 2500 (Illumina).

2.0.1.4 Bioinformatics analysis

The quality of the demultiplexed HiSeq reads, and the quality of the reads following sequencing adapter trimming with Cutadapt (v2.10) was assessed using FastQC (v0.11.9) and MultiQC (v1.8) (Martin, 2011; Andrews, 2010; Ewels et al., 2016). The adapter-trimmed reads were assembled into contigs using MEGAHIT (v1.2.9), assembly statistics were generated using QUAST (v4.0) (Li et al., 2015; Gurevich et al., 2013), and ARGs were identified using CARD-RGI (v5.1.0) (Alcock et al., 2019).

Chapter 3

Results

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Chapter 4

Discussion

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Chapter 5

Conclusion

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