# Assessing Historical DNA from Pocillopora museum specimens

Smithsonian Institution



Pocillopora

damicornis,

**USNM** 

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#### Background

Pocillopora is a diverse genus of stony corals used as model organism in biological studies and restoration efforts. However, they face a classification issue due to extensive morphological variation. It is problematic because accurate taxonomy is the foundation of further studies. Molecular genetic tools help solve this problem. Meanwhile, large museum collections are a valuable yet widely untapped resource for genetic data. However, no literature on coral historical DNA extraction methods from old museum specimens (aged <200) has been published so far.

Compare the performance of different methods for the extraction of *Pocillopora* historical DNA.

Objectives

Assess the pattern of DNA degradation across specimen age.

#### Methods

Sampling (0.1-1g coral fragments) Autogen GP Autogen Gene Prep Qiagen BT Qiagen DNeasy Blood & Tissue Kit **DNA Extraction** Qiagen DNeasy PowerSoil Pro Kit Qiagen PP

DNA quantification with Qubit Fluorometer Agarose Gel Electrophoresis

#### Results

concentration(ng/µL/g)

600 While the t-test showed no statistically significant difference between methods, Qiagen PP generally has lower concentration with 200 out-of-range values even in the Qubit High Sensitivity Kit.

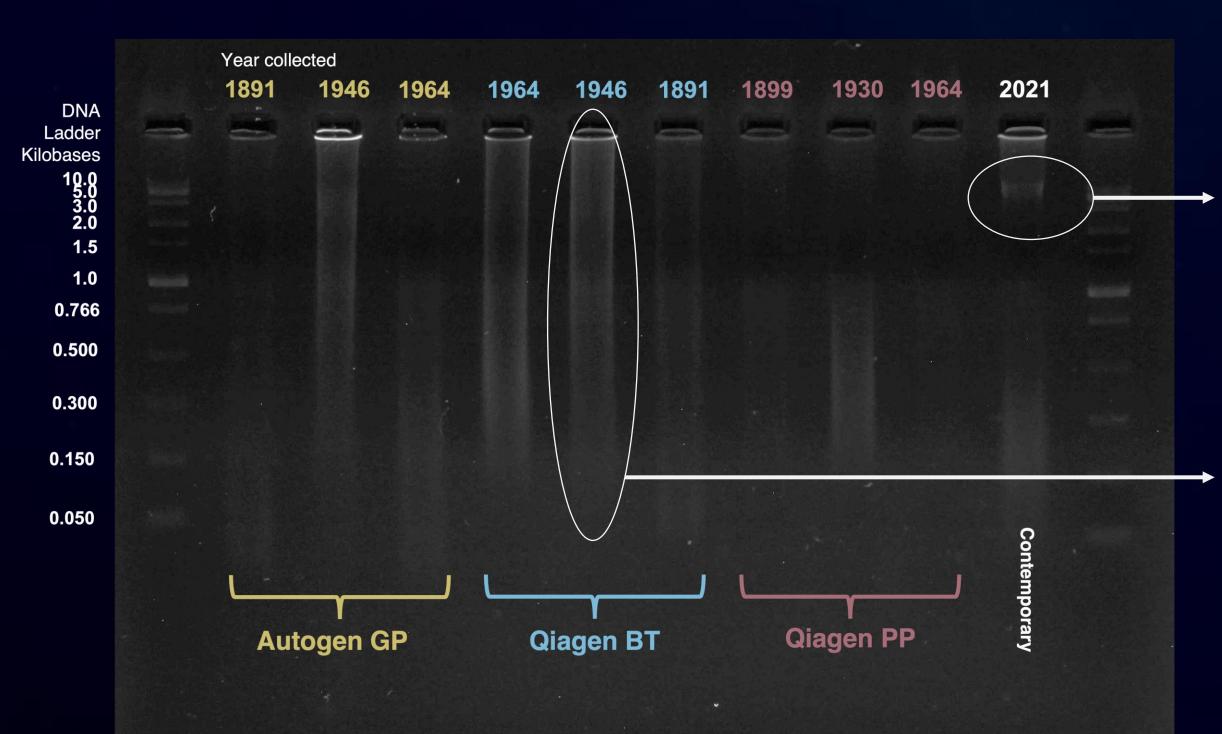
Qiagen PP

#### **Agarose Gel Electrophoresis image**

Qiagen BT

Autogen GP

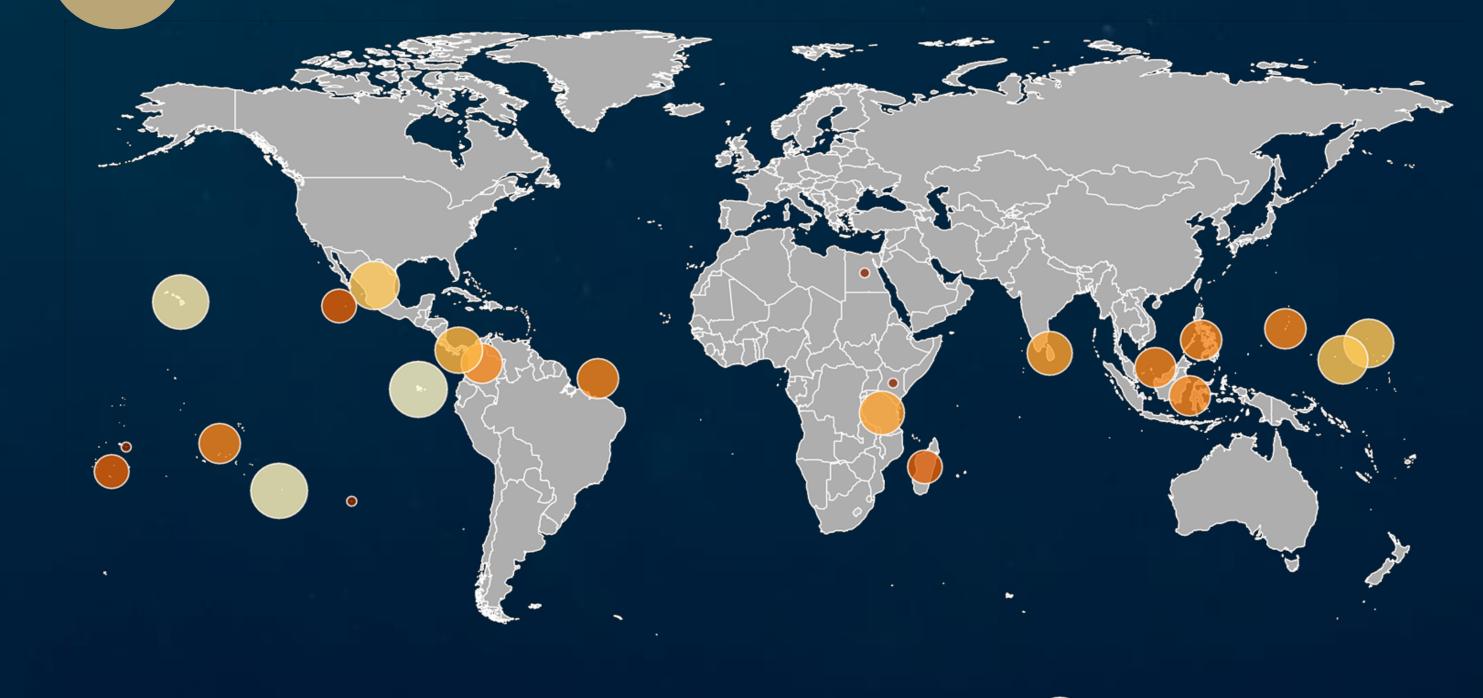
DNA concentration by extraction method



High molecular weight DNA in contemporary sample

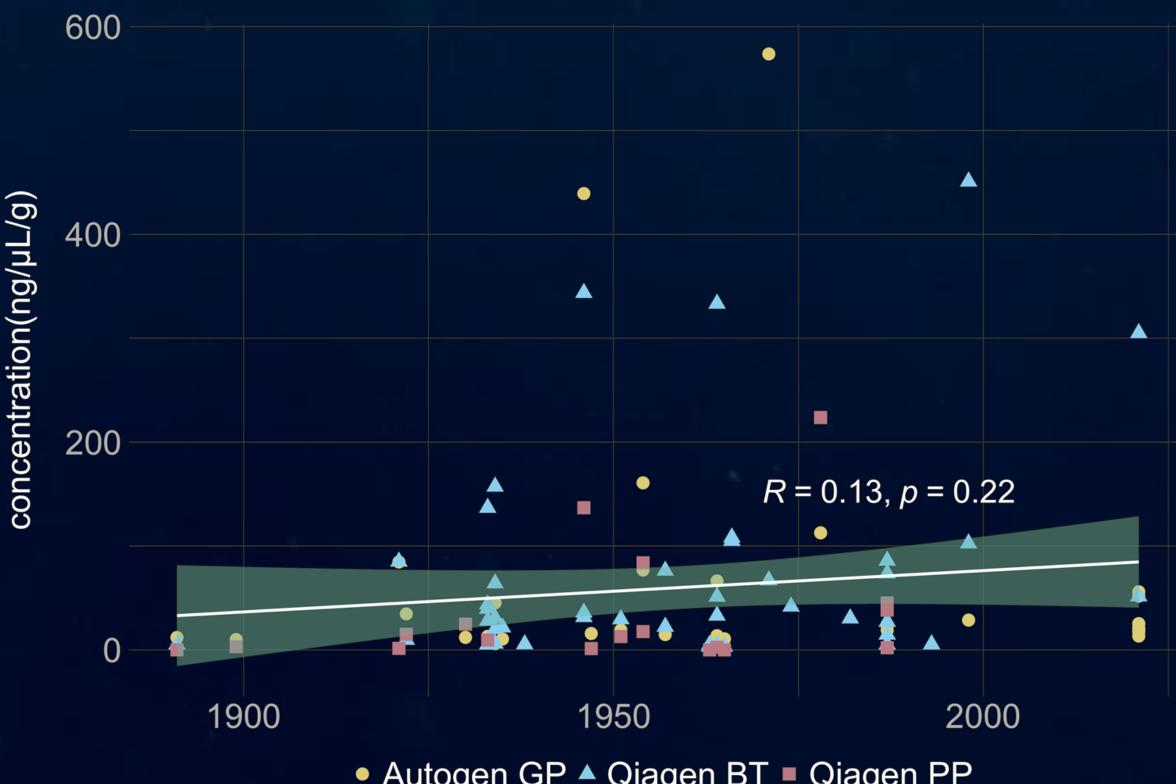
Visible DNA degradation in historical samples with the long smears of DNA bands. This is expected given the age of the samples.

## **Country locations of specimens**



Number of Specimens • 1 — 3

DNA concentration across specimen age



correlation with specimen age and DNA concentration. **DNA** quality depends not on archival storage time but likely on storage factors after collection.

No strong

Autogen GP ▲ Qiagen BT ■ Qiagen PP

#### Conclusion

- Qiagen PP is not recommended for historical samples given the generally low DNA yield. Autogen GP and Qiagen BT may be more reliable for historical DNA extraction.
- Next steps for more insightful results:
- Compare more DNA extraction methods specialized for dry coral/museum specimens.
- Sequence DNA or barcoding genes through target capture.

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Want to dive deeper?

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Visit this page for the list of references and more information about the research project.