Interventions addressing genetic disease burdens in the MENA region: A scoping review

Madison Grant, MPH HPCH Advisor: Dr. Tamar Kabakian Second reader: Dr. Soha Yazbek

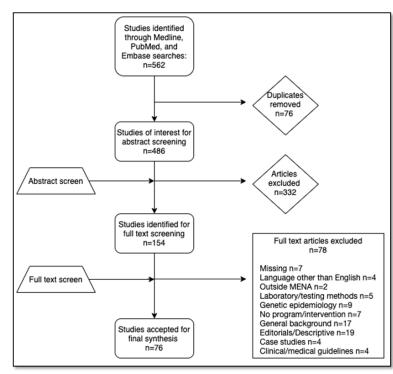
Rationale

- The MENA region is disproportionately affected by the burden of genetic disease due to the cultural practice of consanguinity
- No previous research provides a comprehensive view of all types of genetic health interventions present within this region
- The aim of this study was to identify how genetic disease burdens are being managed within MENA nations

Objectives

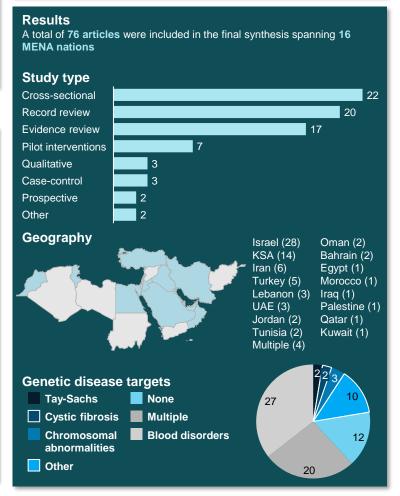
- Identify interventions in the MENA region that address genetic disease burdens linked to consanguinity
- 2. Synthesize literature to map interventions by their geography, characteristics, and any associated outcomes and impact
- 3. Identify gaps to inform potential areas for research in the region

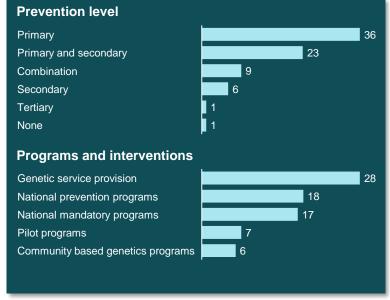
Process



Methodology

- Scoping review process followed the Arksey & O'Malley framework and the PRISMA guidelines for reporting
- Three databases utilized: Medline Ovid, Embase, PubMed
- Abstract screening performed independently, full-text screen performed with two collaborators
- Numerical analysis presented for study types, geographic regions, genetic disease targets, prevention levels, and type of interventions





Conclusions

- More research is needed to pilot effective community-based genetics programs for those most impacted by genetic disease
- Research gaps highlight the need to expand prevention efforts to more prevalent genetic diseases
- Better evaluations are needed for national level programs to realize progress on disease burden impact
- Capacity building is needed to improve genetic services such as genetic counselling

Recommendations

 A systematic review is warranted to provide evidence on program development strategies suitable to the MENA context

Ethics

 The religious ban on abortions in many MENA nations have implications for program development