

Genetic Disorders in Arabs

Ghazi Omar Tadmouri

Centre for Arab Genomic Studies
Dubai, United Arab Emirates

Centre for Arab Genomic Studies (www.cags.org.ae)

Impact of GDs in the Arab World

- **Chronic in nature with lifelong management and no definitive cure**
- **Common GDs in Arabs reached epidemic proportions (SCD, AT, HT, DM)**
- **Spontaneous abortion, neonatal death, morbidity and mortality in children and adults**

Centre for Arab Genomic Studies (www.cags.org.ae)

Economics of GDs in the Arab World

- **Longer and more frequent hospital admissions**
- **Higher number of surgeries**
- **Total costs paid are more expensive**
- **Come frequently from outside major cities and travel farther to obtain better treatment**
- **Families end up by paying the bill**
- **SCD: 6300\$ required annually excluding indirect and tangible costs**

Centre for Arab Genomic Studies (www.cags.org.ae)

Preventive Aspects of Genetic Disorders

- **Expertise and resources to initiate prevention programs are available in the region**
- **Newborn and carrier screening programs**
- **Abnormal findings should lead to clinical management of genetic disorders**
- **Cost effectiveness**
- **Extensive educational activities**
- **Sensitivity to cultural backgrounds**

Centre for Arab Genomic Studies (www.cags.org.ae)

Centre for Arab Genomic Studies

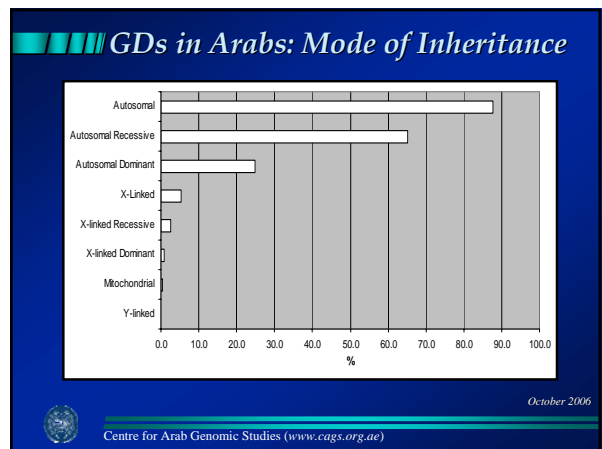
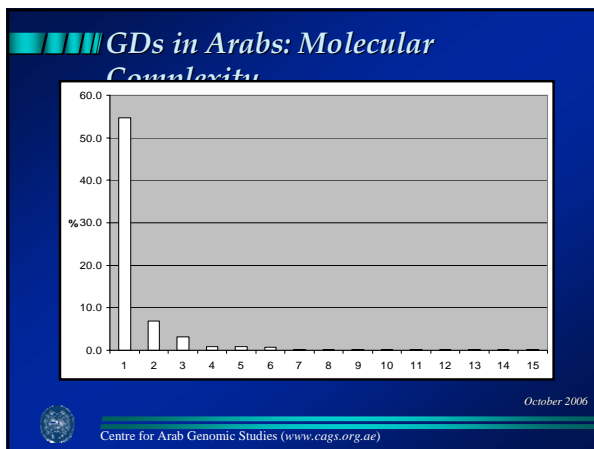
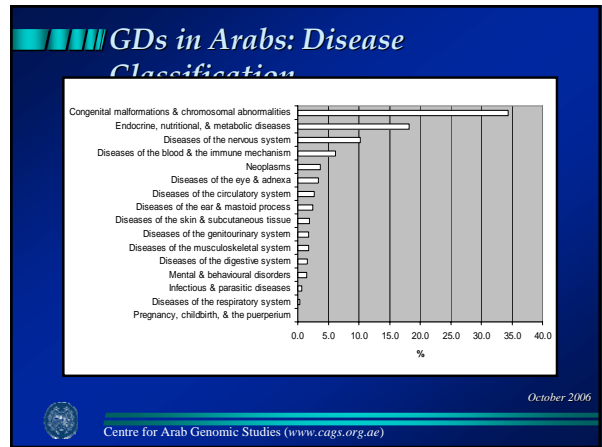
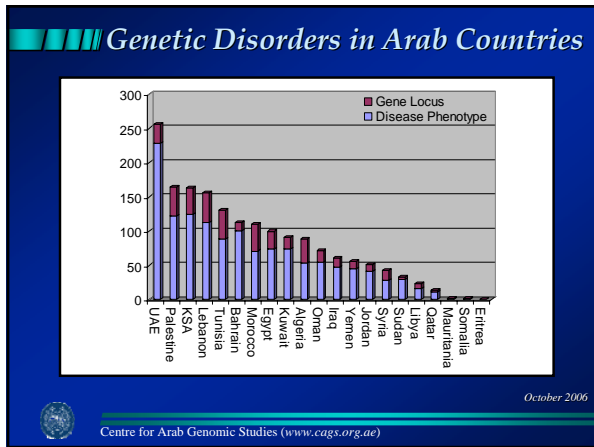
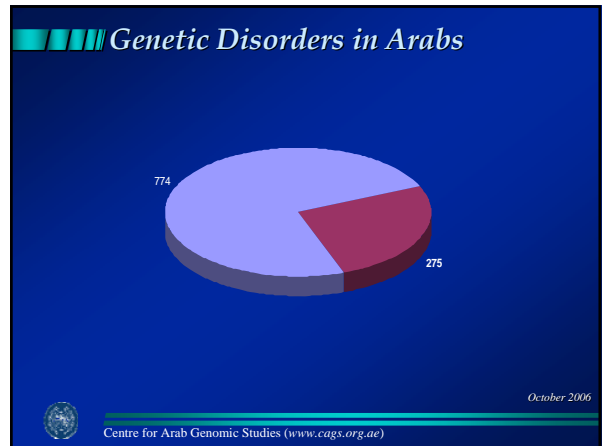
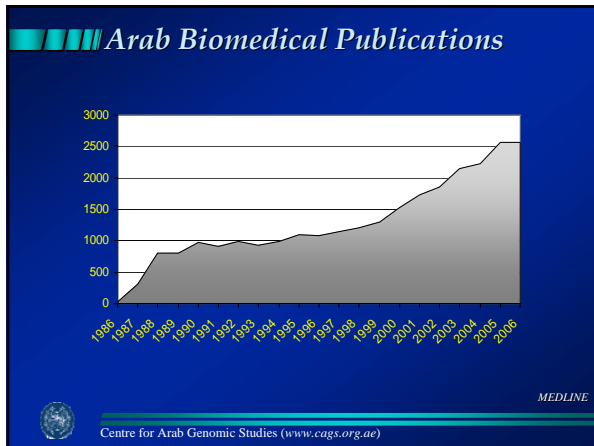
DUBAI
CAGS
UAE

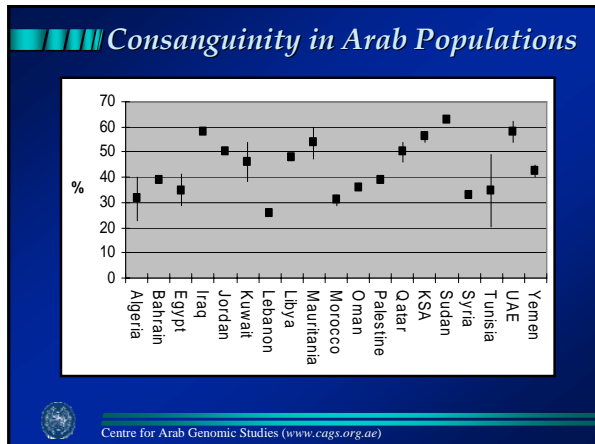
Centre for Arab Genomic Studies (www.cags.org.ae)

The CTGA Database

www.cags.org.ae

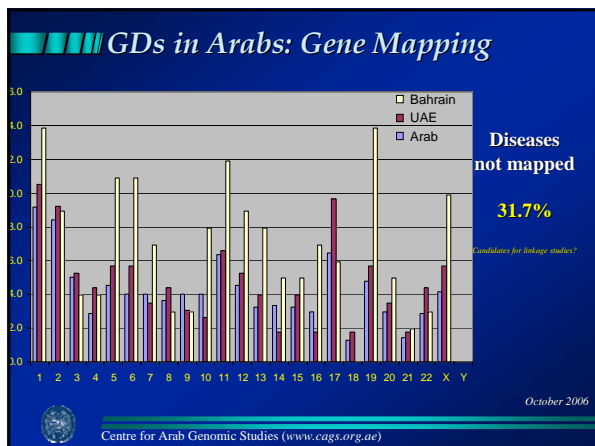
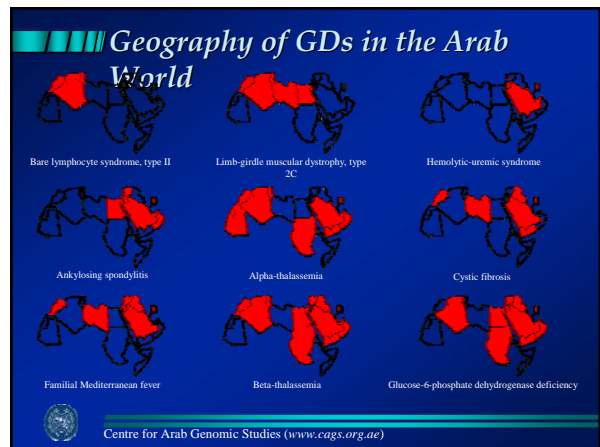
Centre for Arab Genomic Studies (www.cags.org.ae)





- ### Spectrum of Genetic Disorders in Arabs
- Teebi type of Hypertelorism (1987)
 - Teebi-Shaltout syndrome (1989)
 - Al-Gazali syndrome (1994)
 - Megarbane syndrome (2001)
- Centre for Arab Genomic Studies (www.cags.org.ae)

- ### Spectrum of Genetic Disorders in Arabs
- Lebanese type of mannose 6-phosphate receptor recognition defect (1984)
 - Algerian type of spondylometaphyseal dysplasia (1988)
 - Kuwaiti type of cardioskeletal syndrome (1990)
 - Yemenite deaf-blind hypopigmentation syndrome (1990)
 - Nablus mask-like facial syndrome (2000)
 - Jerash type of the distal hereditary motor neuropathy (2000)
 - Karak syndrome (2003)
 - Omani type of spondyloepiphyseal dysplasia (2004).
- Centre for Arab Genomic Studies (www.cags.org.ae)



- ### Challenging Issues: Scientific Reporting
- Inconsistencies in reporting (patient origin, family history)
 - Occur more in international articles
 - Unified system for clinical and molecular reporting
- Centre for Arab Genomic Studies (www.cags.org.ae)

//// Challenging Issues: Research

Directions

- **Clinical reports vs. molecular studies**
- **Lack of large-scale data of DNA or protein sequences, mutations, and SNPs data**
- **Data necessary to address urgent health needs**
- **Constructive engagement of global efforts to build sustainable research activity based upon education and improvement of human**



health for Arab Genomic Studies (www.cags.org.ae)