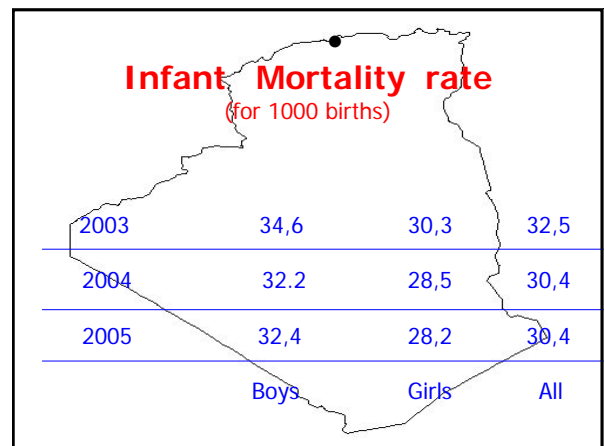
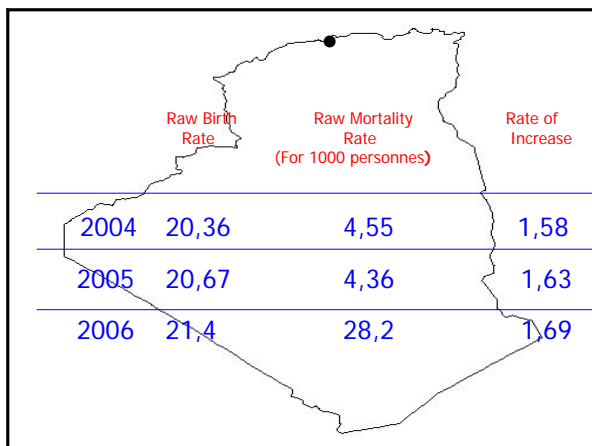
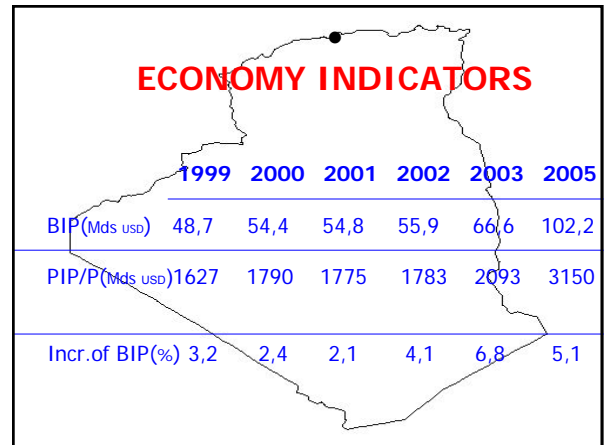
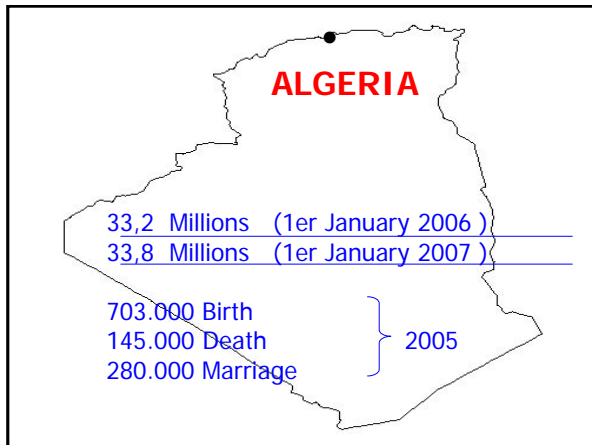
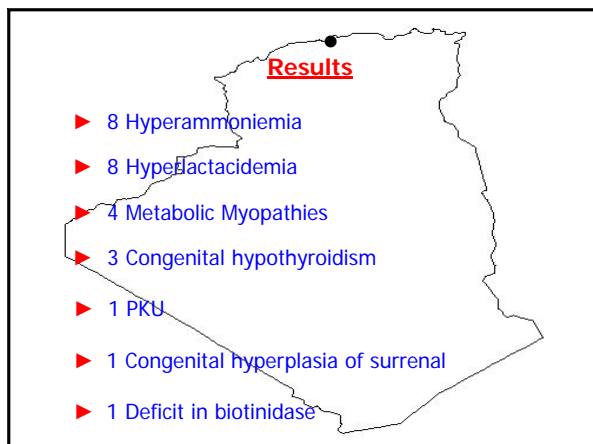
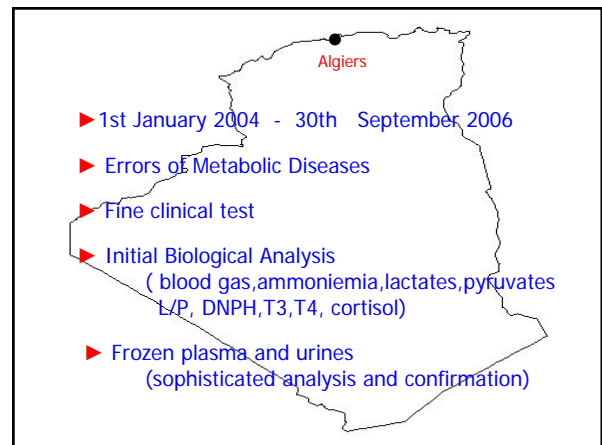
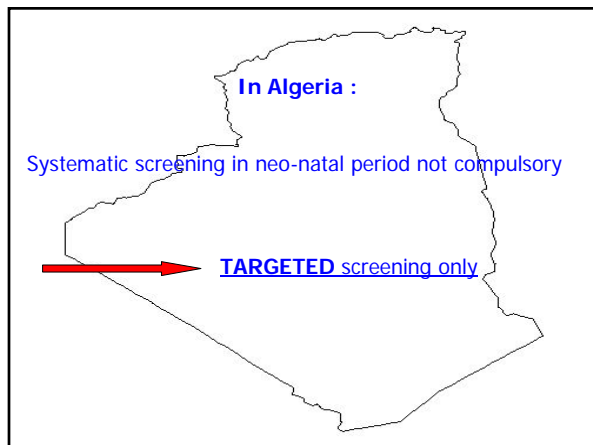
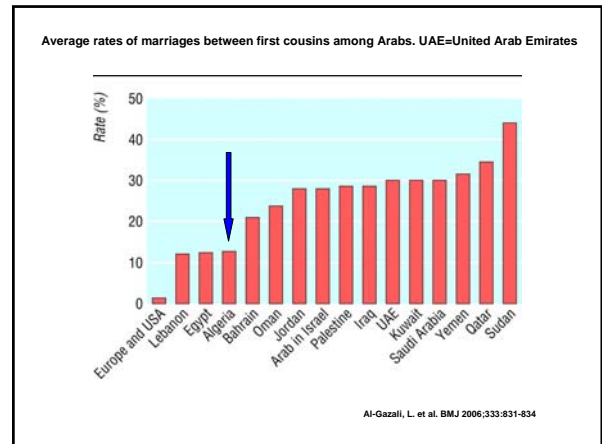
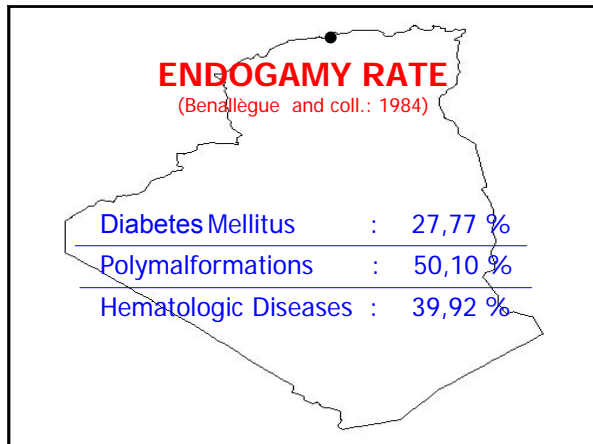


# Born without risks : utopia or reality ?

Paediatric Intensive Care  
M.T Hamlaoui Unit  
Algiers

Marrakech Nov. 2006





Mardi 13 Juin 2006 16h45  
 Hélène OGIER" <helene.ogier@rdb.ap-hop-paris.fr>  
 Re: Re: Re: BIOTINIDASE  
 Mohamed Hamlaoui"  
[hamlaouimohamed@yahoo.fr](mailto:hamlaouimohamed@yahoo.fr)

Cher Ami, je viens d'avoir le résultat de biotinidase de votre patient:

**Il est déficitaire: votre patient n'a aucune réaction.**

Il n'y a pas de chiffre car il s'agit d'un dosage 1/2 quantitatif par réaction colorée. Bravo pour ce beau diagnostic. Il faut convaincre les parents et plus tard l'enfant de la nécessité de traiter par biotine sans interruption

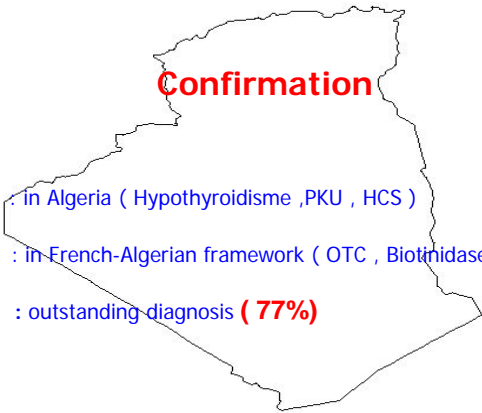
Pr H.Ogier De Beauigny  
 Chef de service Maladies Métaboliques .  
 Hop.Robert Debré

## **Confirmation**

4 : in Algeria ( Hypothyroidisme ,PKU , HCS )

2 : in French-Algerian framework ( OTC , Biotinidase )

**20 : outstanding diagnosis ( 77% )**



## **Conclusions**

- ▶ Targeted screening of metabolic diseases possible but LIMITED
- ▶ Providing active educational programme ( nurses, physicians , biochemists, genetician)
- ▶ Interprofessional educational programme
- ▶ Consensus statements by experts
- ▶ Audit and follow-up

➔ NATIONAL GENETIC MEDICAL COMITY

