



HEMOPHILIA NATIONAL PROGRAM

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This plan is part of the Ministry of Health (MoH) Action Plan 2008-2012 / which consider non-communicable disease as a national priority. It is the result of a collective work of different stakeholders in the area of hemophilia.

OBJECTIVE

To reduce morbidity, mortality and disability associated with coagulation disorders including hemophilia.

Axe I- Comprehensive care

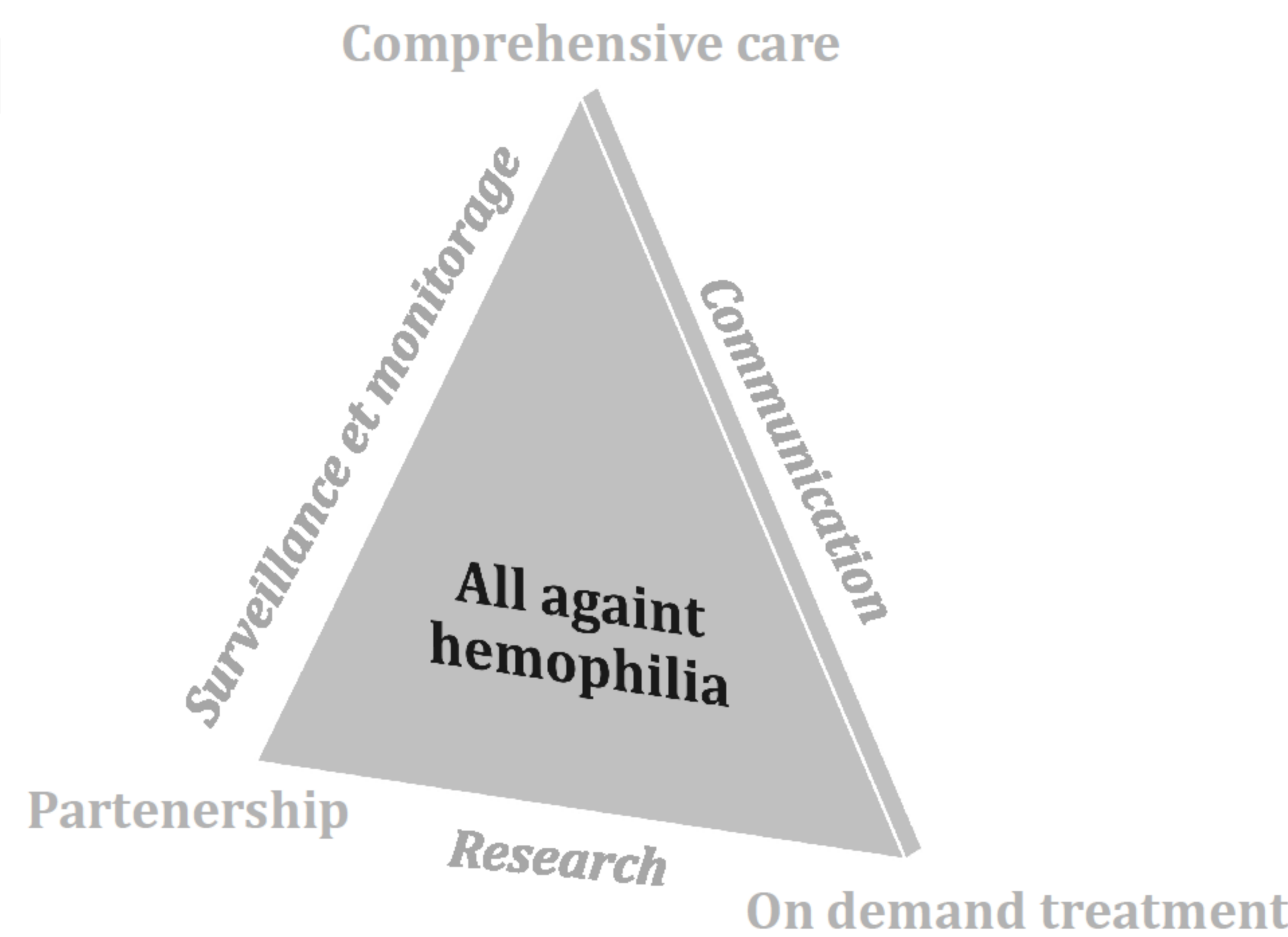
- . Access to health care is improved
- . Hemophilia care is Standardized according to national guidelines
- . Health professional skills are improved

Axe II- On demand treatment

- Appropriate treatment is available and well-managed

Axe III- Monitoring

- Information System developed



Axe IV- PARTENERSHIP

- Partnership with national and international organizations is strengthened

Axe VI- Research

- Improve knowledge of hemophilia in Morocco

Achievements:

Existing Hemophilia Treatment Centers in Morocco:

- 4 Centers within university hospitals
- 2 régional hospitals

- We have identified 9 Regional hospitals where to develop hemophilia care unit

- Elaboration of National guidelines (**moroccan society of hematology**)

- 3 days Training session for all medical staff working in hemophilia area

- 81 health professionals + 10 from MHA

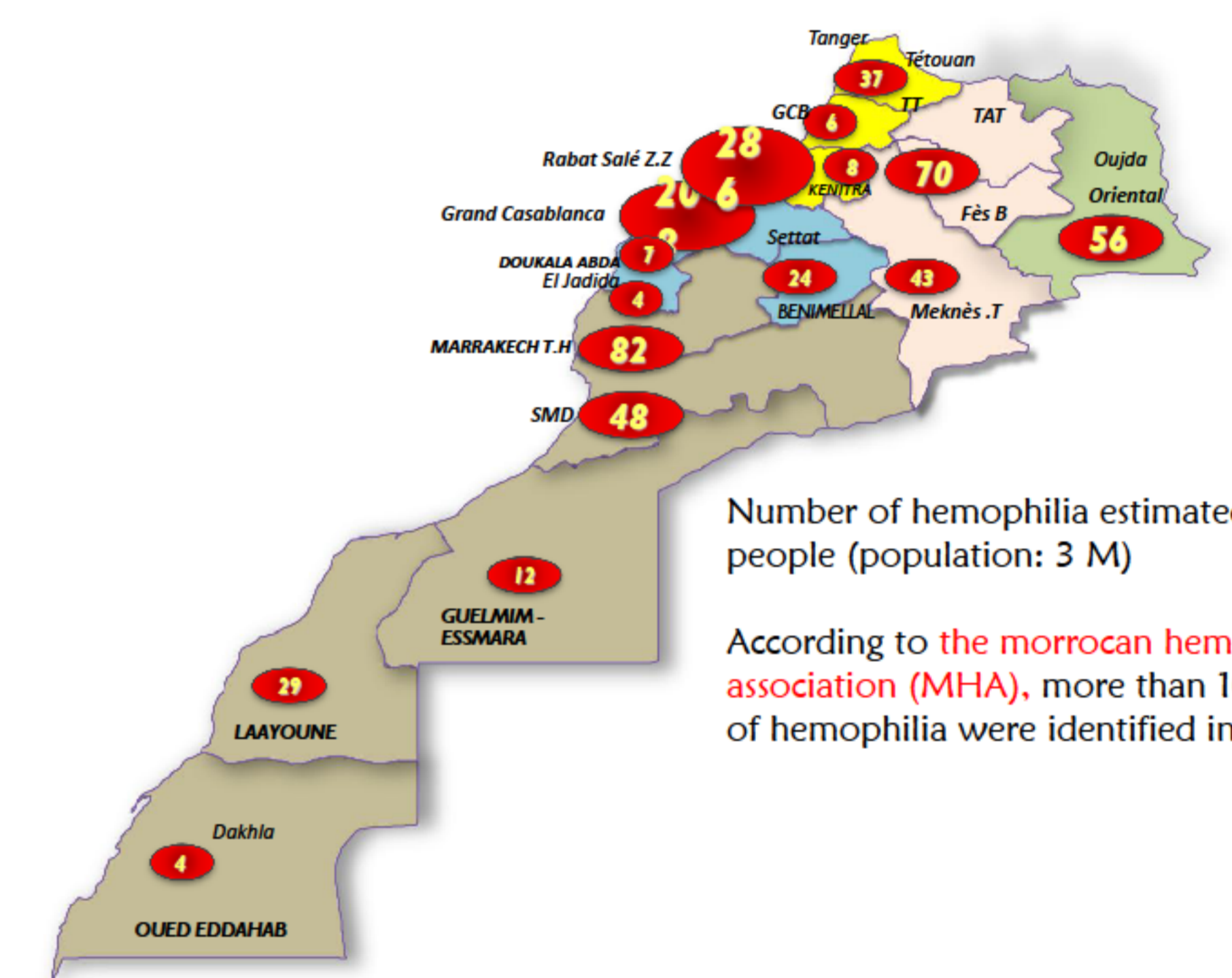
- Workshop to prepare surveillance system (national register)

- Medical insurance coverage

- AMO (Working/active population)
- RAMED december 2011 (economically disadvantaged)

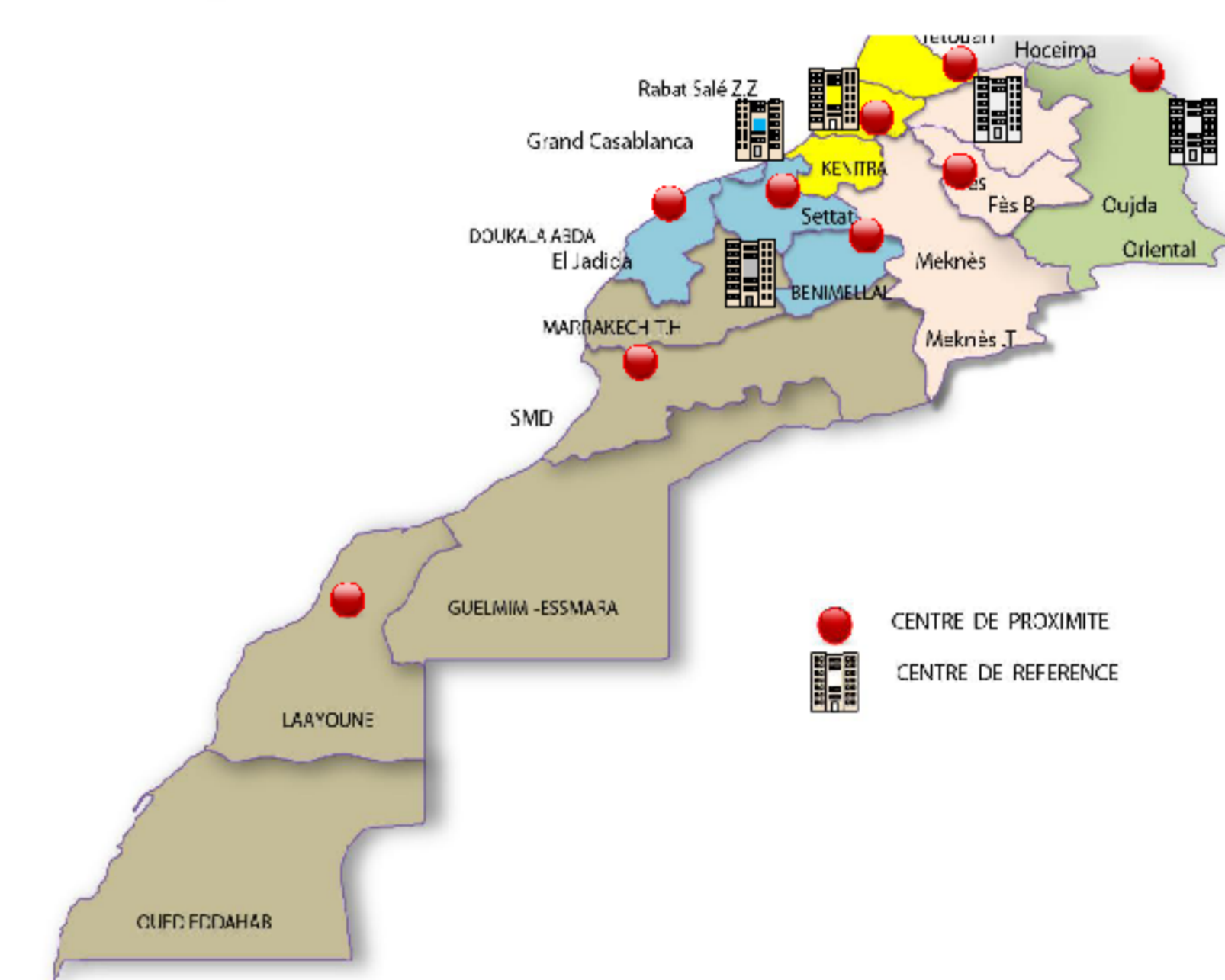
Partnership :

- MoH & WFH : Capacity building:
 - training of health professional
 - Development of laboratories : external quality control
 - technical support
- MoH & Private sector



Number of hemophilia estimated at 3000 people (population: 3 M)

According to the morrocan hemophilia association (MHA), more than 1000 cases of hemophilia were identified in 2011.



Budget:

- 25.000.000 DH (3M.US\$) FOR recombinant Factor VIII
- Plasma factor VIII and IX provided by national transfusion center
- Before 2009: 0,02 UI per Capita
- In 2010/ 0.1 UI per capita
- Objective 0,5 UI per capita 2012

Conclusion:

This ambitious plan can not be achieved without the collaboration of all stakeholders in the field of Hemophilia

