Type Diseased Carriers

No. identified Incidence Prevalence (%) No. identified Incidence Prevalence (%)

Table 5 Prevalence of confirmed sickle-cell diseases, non-sickle haemoglobinopathies and haemoglobinopathy carriers among

Sickle Hb	231ª	1:238	0.04	4481	1:121	0.83	
Non-sickle Hb	80	-	-	-	-	-	
HbC	11	_	_	69	_	0.02	

a Sickle-cell diseases: included 203 cases confirmed as homozygous SS and 28 cases confirmed as sickle/beta-thalassaemia; b diagnosed as HbO Arab by family study;

592

19

11<sup>b</sup>

15°

0.11

11	-	-	
65	_	0.01	

4

newborns (January 2002 to December 2011) (n = 542 286)

<sup>c</sup>β-thalassaemia and hereditary persistence of fetal haemoglobin.

HbD

HbE

Fetal Hb

Unidentified band

Hb = haemoglobin.