

Early Onset Insulin-Dependent Diabetes Mellitus (before 5 years old) Surveillance in Portugal (2001-2003)

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Context: The incidence of Type 1 Diabetes Mellitus (DM1) in children < 5 years old seems to be increasing in several countries. The reason is still obscure (*Svensson et al, Diabetes Care 2002; Charkaluk et al, Pediatric Research 2002*). This age group includes very rare situations: a) Transient neonatal DM1 (1/10⁵ – 5/10⁵, SGA, 1st-6th week, 25% isodisomy of chromosome b) DM1 in children < 1 year (no β-cell autoantibodies, no measurable C-peptide in response to glucose, 35% *de novo* mutations in the gene encoding Kir 6.2) (*Gloyn et al, NEJM 2004*).

Objectives: To collect data on the epidemiology of DM1 in children <5-years-old, in order to get clues to understand its aetiology.

Methods: Since April 2001, the PPSU national, active, voluntary surveillance system is used for monthly reporting of this condition. Notification cards are monthly sent to circa 1600 paediatricians, attending circa 1,5 million children and adolescents. The mailing list was increased with Endocrinologists who attend children. A specific inquiry on epidemiology and clinical data confirms reported cases. Results to December 2003 are presented (33 months of surveillance).

Results: Through December 2003, 139 cases were reported (circa 50/year), 59 cases are already confirmed (11 are duplicate or misreported cases, due to overage or non-resident cases).

DIAGNOSIS

Where?
Hospital (52%)
Office (36%)
Health Centre (8%)

Who?
Paediatrician (85%)
General Practitioner (10%)
Endocrinologist (2%)
Internist (2%)
Family (2%)

When?
(after presumed onset)
Two weeks (87%)
Before 1 week (25%)

First diagnosis
DM 1 (83%)
Urinary infection (8%)
Respiratory infection (5%)
Otitis (5%)



National and Regional Incidences of Reported Cases					
AHR	cases	pop<5years	cases/year	incidence*	RHA/Port
Portugal	56	566500	20,4	3,6	1
Norte	9	214250	3,3	1,5	0,4
Centro	15	86200	5,5	6,3	4,1
LVT	24	191500	8,7	4,6	0,7
Alentejo	4	23250	1,5	6,3	1,4
Algarve	1	20500	0,4	1,8	0,3
Açores	2	16000	0,7	4,5	2,6
Madeira	1	15000	0,4	2,4	0,5
AHR - Administrative Health Region				* :100.000	

CLINICAL FEATURES (presentation)

Thirst (polydipsia) **98%**
Polyuria **93%**
Weight loss **76%**
Decreased activity **42%**
Altered consciousness **10%**
Fever (opportunistic infection) **7%**
Ketoacidosis **32%**

Logistic regression analysis isolated hyperglycemia as the main risk factor for ketoacidosis, excluding age, disease duration or socio-familial data

COEXISTING CONDITIONS

One 21 Trisomy; no other neonatal pathology associated

Prevalence of Asthma **19%**
" Rhinitis **11%**
" Eczema **11%**
" Recurrent respiratory infections **46%**

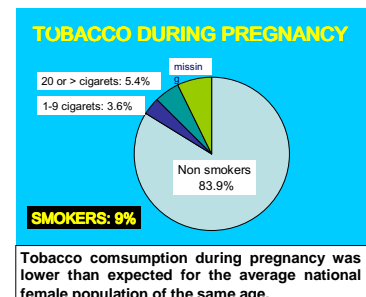
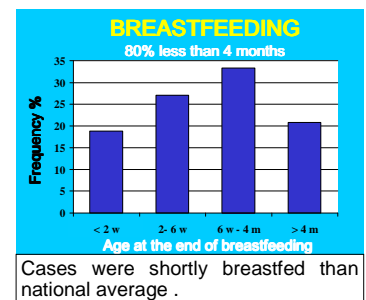
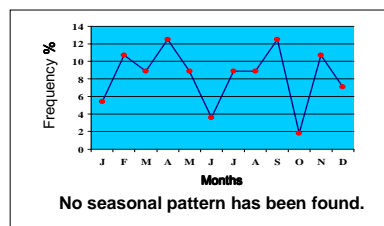
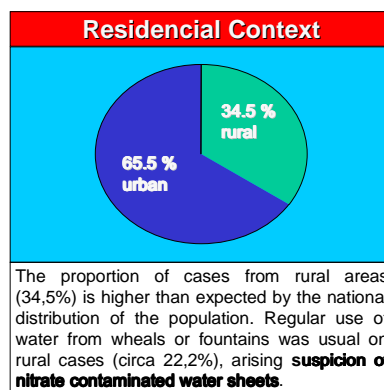
Previous stress **20%** psychological (80%)
(mainly parents separation)

DIABETES MELLITUS AMONG RELATIVES

DM Type 2
Mother family: 26%
Father family: 15%
Both families: 17%
DM Type 1
2 Fathers: 4%

62%

One Gestacional DM



Discussion: The high proportion of missing inquiries may affect the results. Regional asymmetries may be due to underreporting, but we cannot exclude they are due to real environmental or population differences. We found clues of potential interaction among environmental, toxic and immunity factors associated to DM1.

Conclusions: Data from national surveillance may provide clues for the aetiology of DM1 to be followed on later studies. Joining efforts among different surveillance programs will be useful to get more powerful associations that would eventually lead to future preventive or curative interventions.