

# THERAPEUTIC MANAGEMENT OF LEAD POISONING: SATURNISM



DI-079

Muñoz-García I<sup>1</sup>, Fernández-Lobato B<sup>1</sup>, Hernández-Roca JJ<sup>2</sup>, Pedregosa-Díaz J<sup>3</sup>, García-Márquez A<sup>1</sup>, Viney AC<sup>1</sup>, Núñez-Bracamonte S<sup>1</sup>, Conesa-Nicolás E<sup>1</sup>, Pérez-Pérez IG<sup>1</sup>, García-Simón MS<sup>1</sup>.

<sup>1</sup>Servicio de Farmacia Hospitalaria. <sup>2</sup>Servicio de Medicina Interna. <sup>3</sup>Servicio de Análisis Clínicos. Hospital General Universitario Santa Lucía. Cartagena. Murcia. Spain (iris.munoz@carm.es)

**Background:** Lead poisoning has been reported infrequently in recent years in developed countries due to the implementation of legislative measures aimed at reducing environmental lead. The few cases that appear correspond mainly to occupational exposure.

# Purpose: To describe the therapeutic management in a case of chronic lead poisoning

### Material and methods:



•Male

•53 years

•Dedicated to the exploitation of leas in abandoned mine

Back ground:
dyslipemia,
30 cigarettes/day

He goes to internal medicine for

Joint pain
Dyspnea minimum-moderate efforts
Asthenia and dysthermina sensation
Scarce cough
Dar k urine
Abdominal pain
Nausea
Vomiting

Unquantified weight loss

Initial Analysis  $8.2 \, \mathrm{g/dL}$ Hemoglobin (Hb) Hematocrit (Ht) 25,1% Peripheral blood smear with anisocitosis, rounded red spherocyte-like cells, element with basophilic stippling  $0.67 \, \text{mg/dL}$ Plasmatic Creatinine (Cr.)  $109.7 \text{ ml/min/m}^2$ Glomerular filtration 8 mg/dLUrobilinogen in urine Blood Lead  $946 \, \mu g/dL$ 

 $2024 \mu g/24h$ 

Urinary lead

excretion

Diagnosis: HEMOLYTC ANEMIA FOR LEAD POISONING

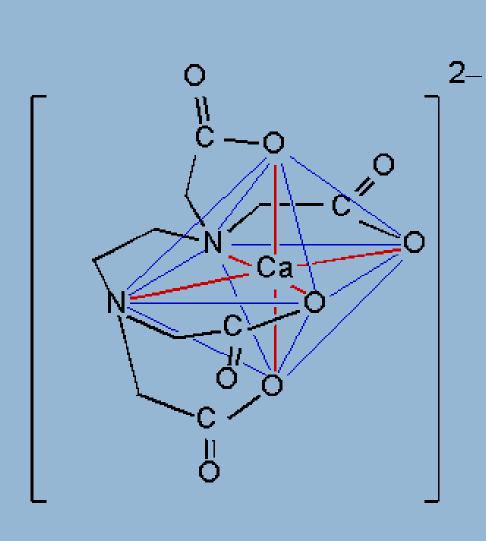
#### Results:

**Pharmacist** 

#### Due to:

•Few cases with lead blood levels above  $100\mu g/dL$  in the literature •No neurological symptoms despite higher than  $900\mu g/dL$ 

Internal Clinic



Conservative approach and hospitalization for chelation therapy:

## <u>Dimercaprol IM 200 mg</u>

(every 4h days +1,+2; every 6h days +3,+4 and every 12h day +5)

EDTA calcium 1500mg every12h

(administered for 6h) (first dose administered 4 h after Dimercaprol IM)

HSOH

Premedication

Dexclorfeniramine 5mg IV every 8h 500mL Bicarbonate 1/6M every 12h

Clinical course was
satisfactory, without
deterioration of renal
function and no complications

During treatment	+1	+2	+3	+4	+5	+6	+7
Hb (g/dL)	7.6	9.6	10.9	8	9.1	8.6	9
Ht (%)	23.1	29.1	32.9	24.5	28.5	25.3	27.7
Cr (mg/dL)	0.59	0.84	0.55	0.71	0.48	0.54	nd

One month after the chelating treatment:

Blood lead: 70 µg/dL

Urinary lead excretion: 162 µg/24h

# Conclusions:

- •Chronic lead poisoning is unusual.
- •Pharmacological management is generally little know and the literature does not reach a clear consensus, requiring a multidisciplinary team.
- •Pharmacys play an essential role in adquisicion, management, dispensation and validation of the treatment, in order to achieve the therapeutic goal in the shortest time and with an optimum result.