

# NOT EVERY OBESE CHILD HAS TYPE 2 DIABETES MELLITUS

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## CASE PRESENTATION

- 11.84 years female
- Gradually gaining weight after 8 years
- Presented at 11.03 years with obesity
- BMI: 31.5kg/m<sup>2</sup>; BMI z-score: 3.55
- No significant past health issues

### Clinical examination Signs of metabolic syndrome

- Mild acanthosis nigricans
- Arterial pressure: 129(>99<sup>th</sup>%ile) /80 mmHg (>95<sup>th</sup>%ile)
- Arterial pressure: 111 (<90<sup>th</sup>%ile) /80 mmHg (>95<sup>th</sup>%ile)

## FOLLOW-UP

- Following 3 months dietary/lifestyle intervention
- OGTT repeated
  - 120' glucose: 225mg/dl
  - T1DM antibodies negative, C-peptide normal
- Started on Metformin (850 mg BD)

OGTT 29/7/2020	0'	60'	120'
Glucose (mg/dl)	98	229	225
Insulin (μUI/ml)	31.2		
HbA1c (%)	5.5		
Glucose in urine	Negative		

- At 11.84 years BMI progressively increases (BMI z-score: 3.98 from )
- On metformin HbA1c unchanged but normal (5.5%)
- A new OGTT was performed to better assess response on metformin

OGTT 19/3/2021	0'	30'	60'	90'	120'	150'	180'
Glucose (mg/dl)	123	125	170	227	234	130	216
Insulin (μUI/ml)	32.87	29.91	63.19	106.0	102.1	70.14	98.63

Unchanged  
➢ Question of adherence  
➢ Increased metformin to 1000mg BD

## WAS IT TYPE 2 DIABETES OR SOMETHING ELSE?

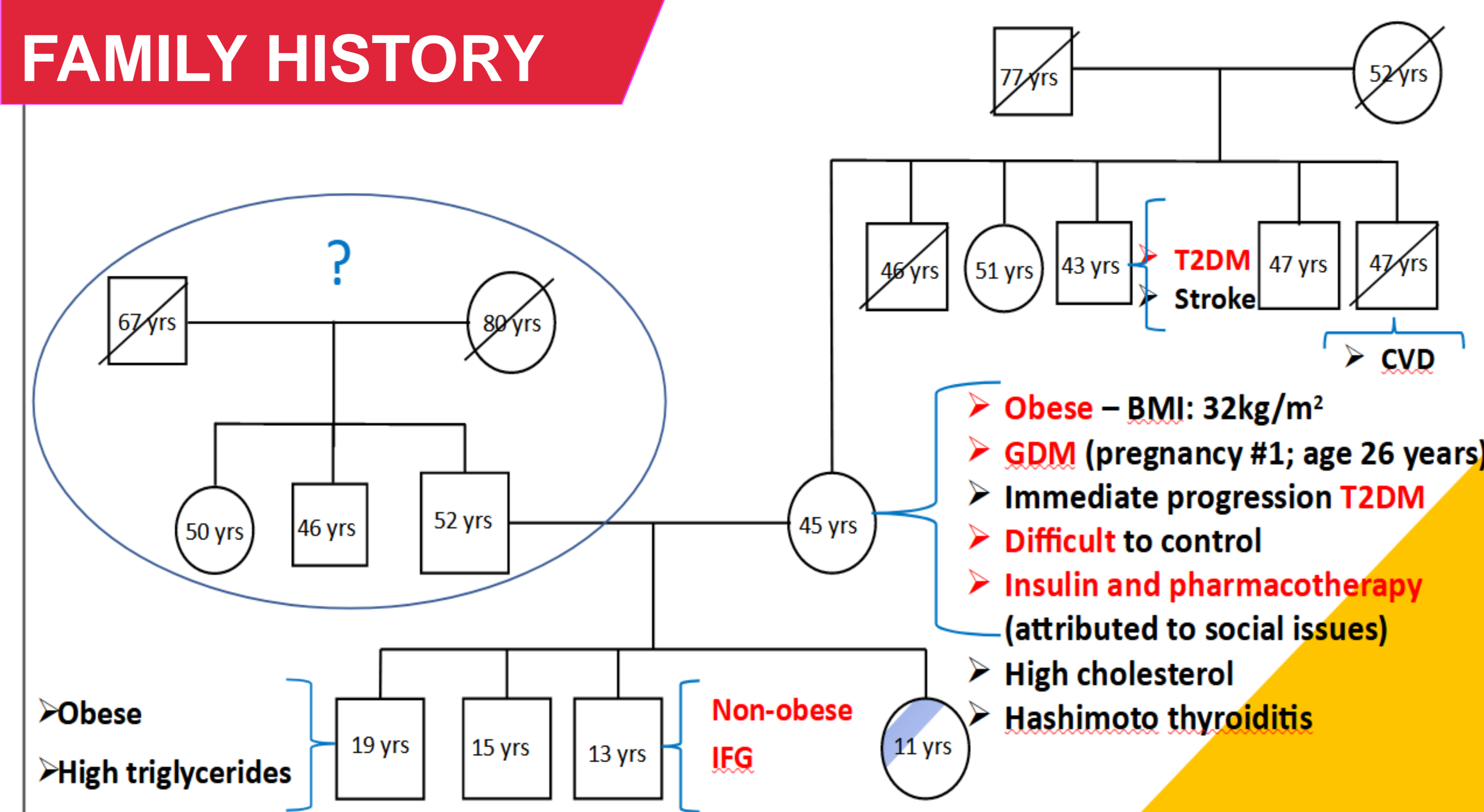
**Young people with T2DM**

- Present around puberty
- Majority are obese

**Patients with monogenic diabetes**

- May also be obese
- Can be difficult to distinguish from T2DM

## FAMILY HISTORY



## REVIEW OF MANAGEMENT

### Features that suggest T2DM:

- Signs of metabolic syndrome
- Severe obesity
- Acanthosis nigricans (mild)

NGS panel for MODY

Mutation p.R200Q/N of HNF1A mother and patient

HNF1A-MODY (MIM# 600496) or MODY3

### Features that suggest monogenic diabetes:

- Family history
  - Early onset diabetes in the mother
  - Obese, GDM and T2D at 26 years
  - Patient's brother non-obese - IFG
  - Mother's brother T2D and stroke
  - Strong family history of CVD

### Molecular diagnosis

- Choice of most appropriate treatment
- Optimize blood glucose control
- Reduce long-term complications
- Proper genetic counseling

## SUMMARY

- Cases of MODY may be difficult to distinguish from T2DM
  - Importance of Family History – Low threshold perform genetic testing
  - Importance of Genetic testing
    - Can drive the choice of treatment
    - Assess the risk for complications
    - Provide genetic counseling
  - Treatment adherence requires regular follow-up – Clinical interview skills
- Patient AND affected relatives!**

## CONCLUSIONS

- Cases of MODY may be difficult to distinguish from T2DM.
- Studies suggest that 5% of subjects diagnosed with diabetes before the age of 45 years have MODY, with 80% of them having been incorrectly diagnosed as having T1DM or T2DM.
- Importance of Family History – Low threshold perform genetic testing.
- Genetic testing allows confirmation of the correct diagnosis and leads to optimal treatment.

## REFERENCES

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## CONTACT INFORMATION

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