Baring the facts to uncover the naked truth: the stubborn puzzle

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“In endeavoring to trace the clinical features of a given disease to their source, use is made of the physical signs and modern laboratory methods of investigating disease, as well as of all data relating to the anamnesis. The marks of disease are often decidedly obscure, and for their detection the diagnostician must call into requisition the various instruments of precision contributed by science, e.g., the microscope, stethoscope, hemomanometer, and many others, as well as the helps furnished from the laboratory expert.”

“Gut feeling”

• [..] an “inexplicable gut feeling is an important diagnostic sign and a very good reason for seeking the opinion of someone with more paediatric expertise or performing additional testing”

• Gut feeling should make three things mandatory:
  – conducting a full and careful examination;
  – seeking advice from a more experienced clinician;
  – providing the parent with safety netting advice.

• Clinicians should not ignore gut feeling and use it in decision making

http://www.bmj.com/press-releases/2012/09/25/doctors%E2%80%99%E2%80%9Cgut-feeling%E2%80%9D-should-not-be-ignored
Ms. G.S., 50 y.o. woman

• 1st visit (Mar-2011) to general practitioner (GP) for mild respiratory viral infection
• GP asks the patient to undress – noticed hyperpigmentation of the back
• Referred for hip skin biopsy – no iron deposits

DOI: http://dx.doi.org/10.15605/jafes.030.02.12
Ms. G.S., 50 y.o. woman (W 44 kg, H 151 cm, BMI 19.3 kg/m²)

• Mild diarrhea (May-2011) – severe fatigue
• GP remembers adrenals → endocrinologist
Ms. G.S.,
50 y.o. woman
(W 44 kg, H 151 cm, BMI 19.3 kg/m²)
First visit (Jun 2011)

- Darker face and palms, nationality – European background
- No significant family history
- Single, no children, menopause since 2008
- Postal operator (indoors), no smoker, no EtOH
- **Medications** – risedronate, OTC Ca
- First examination – no other pathology
- HR 74x’, **TA 85/60 mmHg**, no orthostatic changes
First lab tests (Jun-2011)

• ACTH – 218.1 pg/ml (7.2-63.3)
• Cortisol (8AM) – 6.1 µg/dl (3.7-19.4)
• 24h urinary cortisol – 22.5 µg/24h (4.3-176) (volume – 1.05 l)

• **Diagnosis:** Primary adrenal insufficiency
Confirmation of diagnosis

ACTH stimulation tests:
• No need for insulin hypoglycaemia test
• No need and possibility for tetracosactide/cosyntropin (*Synacthen* test)
Years of investigations and consultations behind...
...and then came an internist...
Additional history

• Frequent recurrent respiratory infections and paranasal sinusitis (her mother’s words: “ты гнилая” = “you are foul”)

• Failure to thrive, small stature, underweight

• Abdominal bloating, diarrhea exacerbated by fatty food

• Tolerates milk
History (2002-2011)

  – atrophic gastritis
  – duodenal reactive lymphoid hyperplasia
  – low mucosal plasma cell count
  – lymphoma excluded (immunochemical staining)

• Common variable immunodeficiency (2003):
  – IgA↓↓, IgG, IgM, C3, CD4 ↓,
  – CD3, CD8, CD19 ↑

• Colonoscopy (2011) – similar findings to FGS
History (2008-2011)

• Osteoporosis (2008):
  T-score -2.8SD in lumbar vertebrae

• Hypocalcaemia (2008, 2011)

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<tr>
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<th>Apr-2008</th>
<th>Jun-2011</th>
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<tbody>
<tr>
<td><strong>Ca</strong>²⁺ (2.1 - 2.55 mM/l)</td>
<td>1.17</td>
<td>1.6</td>
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<td><strong>PTH</strong> (15-68 pg/ml)</td>
<td>95</td>
<td>135.8</td>
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• ECG – no pathology (QTc 437ms)

• ANA, ENA, dsDNA, SAA – neg.
Lab tests (Jun-2011)

• Full blood count: Leu 16.000, Eo 0.887 (<0.4), Hb 9.89 g/dl, slight macrocytic anemia
• Urinalysis – no pathology
• Biochemistry (no changes in other electrolytes):
  – Ca 1.6 mM/l (2.1-2.55), PTH 135.8 pg/ml (15-68)
    – No changes in lipids, liver enzymes, creatinine
• 24 h urinary calcium – 4.2 mg/24h (100-300)
Lab tests (Jun-2011)

- $\text{B}_12$ vitamin **77.03** pg/ml (191-663)
- 25(OH) $\text{D}_3$ vitamin **4** ng/ml (20-55)
- Glu **3.41** mM/l, HbA1c **4.6%**, C-peptide **0.694** ng/ml
- IgA **0.03** (0.7-4), IgM **0.1** (0.4-2.3), IgG **1.78** (7-16) g/l
Hormones and antibodies (ab)

- **TSH** – 2.64 µIU/ml (2.34-4.94)
- **FT4** – 1.05 ng/dl (0.7-1.48) - **euthyroid**
- **LH** – 61.57 µIU/ml (7.7-58.5), **FSH** – 92 µIU/ml (25.8-134.8), **menopause** – 3 years

- **Anti-thyroid peroxidase (TPO) ab** – negative
- **Anti-adrenal ab** – negative
- **Anti-ovarian ab** – positive

- **Transglutaminase IgA/IgG** - negative
Radiology

- **Abdominal, thyroid USG** – no pathology
- **Chest X-ray** – bronchiectasis, bilateral hilar calcification
- **Head CT** – no pathology
- **Abdominal CT:**
  - small adrenals (left< right)
  - no calcification, no tumors
  - mesenterial lymphadenopathy
Diagnosis

- Primary adrenal insufficiency
- Hypocalcemia with secondary hyper-PTH (vitamin D deficiency, intestinal malabsorption)
- Secondary osteoporosis
- Common variable immunodeficiency (CVID)

- Autoimmune polyglandular syndrome?
Treatment

• **Hydrocortisone** – 10 mg (8AM) + 5 mg (3PM)

• **Calcitriol** \((1,25(\text{OH})_2 \text{D}_3 \text{ vit.})\) – 0.5 \(\mu\text{g}\) x2 1 week followed by **cholecalciferol** 2000IU=3 drops

• **Calcium** (citrate) – 1000 mg/day

• **Risedronate** – 35 mg/week

• **Diet, activities**

• Main worry – “but my skin?”
Stubborn puzzle?

- CVID only?
- CVID + adrenal insufficiency?
- CVID + APS-4? (APS-2?)
- CVID + adrenal insufficiency + celiac disease?

- ACTH/cortisol/Ca/PTH/D vit level monitoring
- To do: Antiparietal cell ab? Endomysial ab?
Autoimmune Polyendocrine Syndromes

- **APS-1** — hypoAdren + hypoPTH + candidiasis
- **APS-2** — hypoAdren + chrThyr + T1DM + hypoGonad
- **APS-3** — chrThyr + 2 other except hypoAdr/T1DM
- **APS-4** — 2 or more other organ-specific AID

**APS-3/4 = APS-2**
APS and CVID

• Coexistence of APS-3 and CVID may be due to autoimmunity and the association of both conditions with human leukocyte antigen (HLA).

• **Clinical tip** - antibodies return negative due to CVID!
Literature list (for further reading)

- [www.eje-online.org/content/161/1/11.full.pdf](www.eje-online.org/content/161/1/11.full.pdf) (review on APS)
- [http://www.worldallergy.org/professional/allergic_diseases_center/suspected-immune-deficiency/](http://www.worldallergy.org/professional/allergic_diseases_center/suspected-immune-deficiency/) (review on immunodeficiencies)
- [http://www.bloodjournal.org/content/99/8/2694](http://www.bloodjournal.org/content/99/8/2694) (autoimmunity in human primary immunodeficiency syndromes)
Take home messages

• Listen to your patients
• *A capite ad calcem* investigation
• Trust your “gut feelings”
• “Take your hat, but don’t take your head off in front of all consultants”
• Use the internist broader view to problems

Thank you!