Metastatic Adrenocortical Carcinoma: A Case Report

Annalisa Montebello¹, Ruth Caruana^{1,2} & Sandro Vella^{1,2}

¹Department of Medicine, Mater Dei Hospital, Malta; ²Department of Medicine, University of Malta Medical School, Malta.



Introduction

• Adrenocortical carcinomas (ACC) are rare malignant tumours. An incidence of 1 to 2 per million per year is reported. We present a case of newly diagnosed metastatic ACC.

Case Report

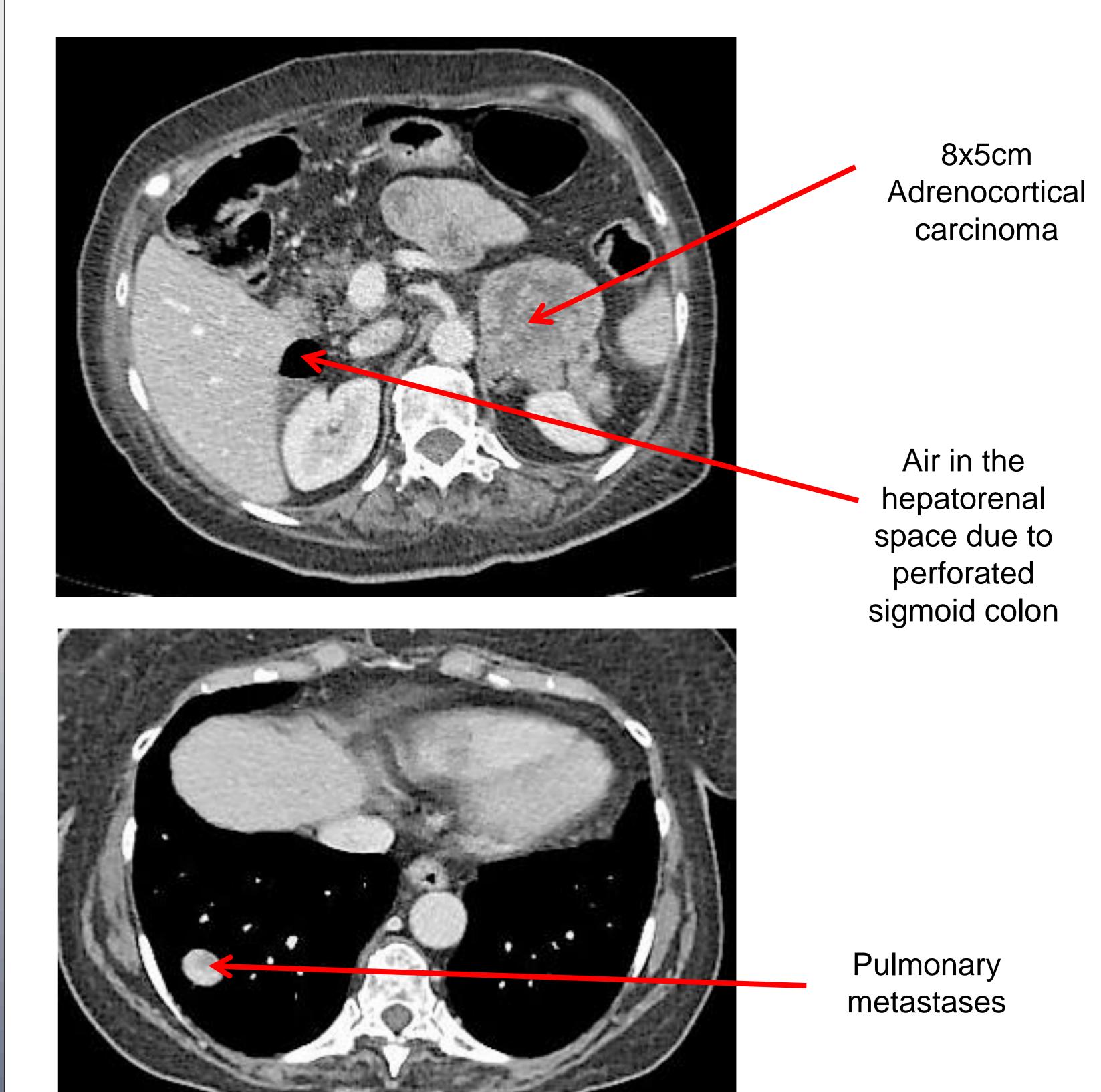
- 70 yr old lady was admitted with a **one month** history of **new onset** hypertension, hypergylcaemia, **rapidly progressive** hirsutism and generalised weakness.
- O/E: Cushingoid appearance with facial plethora, severe hirsutism, central obesity and severe proximal myopathy.
- CT trunk: a large, lobulated, inhomogeneous, solid left adrenal mass **8x5cm in size** with enlarged local and paraortic lymph nodes. Pulmonary metastases were noted.
- She was diagnosed with metastatic adrenocortical carcinoma secreting cortisol and androgens. Aldosterone rennin ratio, plasma metanephrines and catecholamines were normal.
- A few days later she complained of severe abdominal pain and was diagnosed with sigmoid bowel perforation needing emergency laparotomy.
- Post operatively she developed severe hypokalaemia of (1.94mmol/L [3.5-5 mmol/L]) which was difficult to manage with oral potassium supplementation and aldosterone antagonist treatment.
- She became dependent on continuous intravenous potassium replacement therapy.
- Her post operative course was complicated by abdominal wound dehiscence. Wound healing was unsuccessful despite treatment with multiple antibiotic therapy. She additionally developed a thrombosis of the right femoral vein.
- •She was given one shot of 50% doxorubicin-etoposide and cisplatin based chemotherapy. Unfortunately this resulted on neutropenic sepsis which needed treatment with antibiotics and granulocyte colony stimulating factor.
- •Her wounds did not heal and she passed away a few weeks later.



Figure 1: Cushingoid appearance: moon face, facial plethora and severe hirsutism

| Hormone | Result | Range |
|--------------------|-------------|------------------|
| Random Cortisol | 1209nmol/L | 145-619nmol/L |
| ACTH | <5 pg/mL | 10-48 pg/mL |
| Total testosterone | 46 nmol/L | ND-1.49nmol/L |
| Oestradiol | 507 pmol/L | ND-118nmol/L |
| Progesterone | 5.15 nmol/L | ND-3.2 nmol/L |
| 17 OH progesterone | 21.6 ng/mL | 0.13-0.6ng/mL |
| Androstenedione | 19.4 ng/mL | 0.35-2.49ng/mL |
| DHEAS | 23.3 umol/L | 0.95-11.67umol/L |

Figure 2: Biochemistry shows a cortisol and androgen co secreting tumour



Conclusion

• ACC are rapidly progressive and aggressive. Our patient developed complications of hypercortisolaemia i.e. sigmoid bowel perforation, femoral vein thrombosis, severe hypokalaemia and poor wound healing.



