A clinical guideline recommended

For use in: A&E, Medical Assessment Unit, ITU/HDU Medical and Surgical wards

By: Medical, Clinical investigation unit and Surgical staff

For: Investigation of incidental adrenal masses in adults

Division responsible for document: Medical Division (Including Emergency)

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Trust Guideline for the Investigation of Incidental Adrenal Masses in Adults
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Glossary

<table>
<thead>
<tr>
<th>Initials/term</th>
<th>Definition</th>
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<tr>
<td>Dexamethasone suppression test</td>
<td>Specific tests to determine whether adrenocortical tumours are secreting glucocorticoids</td>
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<tr>
<td>HU</td>
<td>Hounsfield Units (CT characteristic, &lt;10 HU = consistent with benign adrenal adenoma)</td>
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<tr>
<td>LDDST</td>
<td>Low dose dexamethasone suppression test</td>
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<tr>
<td>Phaeochromocytoma</td>
<td>Tumour of the adrenal medulla secreting catecholamines</td>
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<td>ODST</td>
<td>Overnight Dexamethasone Suppression Test</td>
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A. Quick Reference Guideline A

Incidental adrenal mass greater than 1cm in absence of known malignancy

- No worrying radiological features, lesion <4cm
- Lesion >4cm or any worrying or indeterminate features radiologically

Arrange CXR and non-contrast CT adrenals
Consider FDG-PET, MRI with chemical shift, washout CT

Referral to Endocrinologists for establishment of definitive diagnosis
- 1mg dexamethasone suppression test (ODST)
- Urine metanephrine x2
- Renin and aldosterone and U&E

MDT discussion at Endocrine radiology meeting

Non Functioning benign lesion <4cm or confirmed adrenal myelolipoma

- No further investigation
- Refer for surgery following confirmatory investigations and medical preparation
- Annual assessment
  - associated comorbidity (Hypertension, Diabetes, obesity, Osteoporosis, Dyslipidemia)
  - Cortisol excess (See Text)

Discharge with no routine follow up tests recommended

Referral for discussion at urology MDT to consider surgery if increasing size (20% and >5mm)

Functioning (phaeochromocytoma, overt Cushing’s, Conn’s) or malignant features

Possible autonomous cortisol secretion (ODST- cortisol >50nmol/l)
*see main text 1.7, figure 2

SEE FIGURE 2 Page 5

Annual assessment
- associated comorbidity (Hypertension, Diabetes, obesity, Osteoporosis, Dyslipidemia)
- Cortisol excess (See Text)

Consider additional investigation, surgery or follow up

Radiologically Indeterminate Mass

Repeat non-contrast CT adrenals or MRI x 2 at 6-12 month intervals and discuss at endocrine radiology meeting

4 year follow up

** for lesion >4cm : individualised approach and surgery may be considered
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Objective of Guideline

The safe and consistent management of patients detected as having unexpected or incidental adrenal lesions on imaging performed for unrelated reasons or in the staging of known malignancy.

B. Rationale for the recommendations

Adrenal incidentalomas are common and their investigation presents a large clinical problem with a high associated financial cost.

Biopsy of undiagnosed or unsuspected Phaeochromocytomas can be fatal. Safe and rational investigation of adrenal nodules is therefore of paramount importance.

C. Detailed Recommendations

1. INCIDENTAL ADRENAL MASSES IN THE ABSENCE OF KNOWN MALIGNANCY OR ENDOCRINE DISEASE

1.1 Initial CT or MRI suggests benign adrenal myelolipoma or adrenal adenoma <4 cm with no concerning features

1 Refer to endocrinology for biochemical and clinical assessment
2 Endocrinology will organise overnight dexamethasone suppression test, urine metanephrines, renin and aldosterone level, and baseline blood tests prior to first clinic appointment to determine whether the adrenal is functioning or not.
3 Review at dedicated weekly endocrine radiology meeting to confirm benign characteristics
4 Endocrine clinic review to confirm functional status, risk of malignancy and other relevant co-morbidities.
5 Patients with radiologically benign and biochemically and clinically non-functioning adrenal lesions <4cm can be safely discharged with no routine follow up investigations required.
6 Functioning adenosomas and phaeochromocytomas require specific medical management prior to consideration of surgery.
7 Autonomous cortisol secretion

Patients with benign adenosomas with possible autonomous cortisol production (ODST result >50) or in whom there is a clinical suspicion of Cushing’s syndrome require further specialist assessment with urinary free cortisol and LDDST testing. These patients then require yearly endocrine follow-up to assess for

a. cortisol excess – LDDST and urinary free cortisol

b. associated CV risk (and comorbidities): diabetes, hypertension, osteoporosis, hypertension and dyslipidemia.

Possible autonomous cortisol secretion is defined by a cortisol of 51-138 after ODST. Autonous cortisol secretion is defined by a cortisol >138nmol/L. Consider
surgery for autonomous cortisol secretion with deteriorating comorbidities, but surgery is rarely indicated in patients with milder disease. See figure 2.

2. >4cm on initial imaging OR unusual / not clearly benign features / growth of lesion on follow up scan at any stage

A. Chest X-Ray in all patients with lesions >4cm.
B. Repeat adrenal imaging (non contrast CT adrenals or MRI) at 6-12 months according to MDT discussion.
C. If at any stage lesion exceeds 4cm, is significantly increasing in size, or has atypical imaging features, review at endocrine radiology meeting, perform functional assessment as above (if not done already), start medical treatment as necessary and refer to urology MDT for consideration of surgery.

3. PATIENTS WITH A NEWLY DIAGNOSED ADRENAL MASS AND A POSSIBLE OR KNOWN EXTRA ADRENAL MALIGNANCY

A. Indeterminate lesions in this group has up to 70% of malignancy, consider PET/CT and discussion in relevant MDT.
B. Measure urine and or plasma metanephrine even if adrenal mass is likely to be metastasis.
C. Benign CT feature need no further investigation for adrenal mass.
D. Consider performing a biopsy of adrenal mass only if all following criteria are met.
   1. Pheochromocytoma is excluded*.
   2. Not benign feature on CT.
   3. Management of underlying malignancy would altered by knowledge of histology of the adrenal mass**.

E. Assess for adrenal insufficiency in large bilateral adrenal potential metastasis.

*Biopsy of an undiagnosed Phaeochromocytoma can lead to fatal complications. For this reason, a Phaeochromocytoma must be ruled out before proceeding to an adrenal procedure in all patients.

1. Arrange plasma metanephrines. Interfering medications are listed in appendix 1 Safe to proceed to biopsy or adrenalectomy if clinically indicated and plasma metanephrines are normal.

2. In all other cases, refer to endocrinology for further investigation and possible pre operative alpha blockade.

**Patients with widespread metastatic disease and adrenal nodules do not need adrenal biopsy Ensure radiology and management is reviewed at appropriate MDT.

E. Clinical Audit Standards derived from guideline

1. Baseline clinical assessment of all patients with adrenal nodules.

2. Exclusion of Phaeochromocytoma or treatment for Phaeochromocytoma in all patients undergoing adrenal biopsy or adrenalectomy.

F. Summary of development and consultation process undertaken before registration and dissemination

The authors listed above on behalf of a guideline development group within the endocrinology directorate, which has agreed the final content, drafted the guideline. During its development it has been circulated for comment to all endocrine consultants, endocrine specialist nurses, and consultants in urology, oncology, anesthetics and radiology. This guideline was approved by the clinical governance committee of the endocrine directorate.

G. Distribution list/ dissemination method

Trust Intranet

H. References/ source documents

Management of Adrenal Incidentalomas- a European Society of Endocrinology Clinical Practice: Guideline in collaboration with European N: Guideline in collaboration with European Network for the study of Adrenal Tumours

Aghili A, Chitale S, Armitage J, Swords F, Burgess N, Dhaturiya K.
Referral pattern & biochemical work-up of adrenal lesions and the role of endocrinologists in managing patients undergoing surgical adrenalectomy: A single centre audit of 10 years of laparoscopic adrenalectomies. Society for Endocrinology, Harrogate 2012

ACP best practice, Adrenal incidentaloma: evaluation and management
P K Singh, H N Buch


Postgrad Med J doi:10.1136/postgradmedj-2013-132386
Adrenal incidentalomas: management in British district general hospitals
Emily Davenport, Pitsien Lang Ping Nam, Michael Wilson, Alastair Reid, Sebastian Aspinall

Appendix 1

Metanephrine collections:

Paracetamol should be avoided for 48 hours prior to and throughout urinary collections for metanephrines. Antidepressants, snoring, untreated sleep apnoea, and major illness will also lead to transient elevations in metanephrines.

Renin and aldosterone measurements:

Angiotensin converting enzyme inhibitors prevent angiotensin production, and angiotensin 2 receptor blockers prevent its action, such that both classes of drug may lead to falsely low ratios (increased renin and decreased aldosterone). These should ideally be stopped for 4 weeks prior to sampling.

Spironolactone and other aldosterone antagonists prevent its action and so potentially increase aldosterone. These should ideally be stopped for 4 weeks prior to sampling.

Beta blockers prevent renin release and also lead to suppression of renin and a falsely increased renin:aldosterone ratio. These should ideally be stopped for 2 weeks prior to sampling.

Diuretics potentially reduce fluid delivery to renal tubules. This leads to increased renin production and so increased renin and aldosterone. This also makes interpretation of renin:aldosterone ratios difficult. These should ideally be stopped for 2 weeks prior to sampling.