

**American College of Radiology  
ACR Appropriateness Criteria®**

**Clinical Condition:** Neuroendocrine Imaging

**Variant 1:** Hypopituitarism.

<b>Radiologic Procedure</b>	<b>Rating</b>	<b>Comments</b>	<b><a href="#">RRL*</a></b>
MRI head without and with contrast	8	Multiplanar thin sellar imaging. See statement regarding contrast in text under "Anticipated Exceptions."	O
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
CT head without contrast	4	Indicated if MRI not available or contraindicated.	⊕ ⊕ ⊕
CT head without and with contrast	4	Indicated if MRI not available or contraindicated.	⊕ ⊕ ⊕
MRA head with or without contrast	3	Indicated if better visualization of carotid arteries needed.	O
CTA head	2	For surgical planning or vascular detail if MRI and MRA contraindicated.	⊕ ⊕ ⊕
X-ray tomography head	1		⊕
X-ray sella	1		⊕
Arteriography cerebral	1		⊕ ⊕ ⊕
Venous sampling petrosal sinus	1		NS
<b><u>Rating Scale:</u> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate</b>			<b>*Relative Radiation Level</b>

**Clinical Condition:****Neuroendocrine Imaging****Variant 2:****Obesity/eating disorder.**

<b>Radiologic Procedure</b>	<b>Rating</b>	<b>Comments</b>	<b><u>RRL*</u></b>
MRI head without contrast	4	In carefully selected patients with high clinical likelihood of structural abnormality. Multiplanar thin sellar imaging.	O
MRI head without and with contrast	4	In carefully selected patients with high clinical likelihood of structural abnormality. Multiplanar thin sellar imaging. See statement regarding contrast in text under "Anticipated Exceptions."	O
CT head without contrast	3	Indicated if MRI not available or contraindicated. In selected patients with high clinical likelihood of structural abnormality.	☼☼☼
CT head without and with contrast	3	Indicated if MRI not available or contraindicated. In selected patients with high clinical likelihood of structural abnormality.	☼☼☼
MRA head with or without contrast	2		O
X-ray sella	1		☼
Arteriography cerebral	1		☼☼☼
Venous sampling petrosal sinus	1		NS
X-ray tomography head	1		☼
CTA head	1		☼☼☼
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

**Variant 3:****Hyperthyroidism (high TSH).**

<b>Radiologic Procedure</b>	<b>Rating</b>	<b>Comments</b>	<b><u>RRL*</u></b>
MRI head without and with contrast	8	Multiplanar thin sellar imaging. See statement regarding contrast in text under "Anticipated Exceptions."	O
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
CT head without contrast	3	Indicated if MRI not available or contraindicated.	☼☼☼
CT head without and with contrast	3	Indicated if MRI not available or contraindicated.	☼☼☼
MRA head with or without contrast	3		O
CTA head	2	For surgical planning or vascular detail if MRI and MRA contraindicated.	☼☼☼
Arteriography cerebral	1		☼☼☼
X-ray tomography head	1		☼
X-ray sella	1		☼
Venous sampling petrosal sinus	1		NS
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

**Clinical Condition:****Neuroendocrine Imaging****Variant 4:****Cushing's syndrome (high ACTH).**

<b>Radiologic Procedure</b>	<b>Rating</b>	<b>Comments</b>	<b><u>RRL*</u></b>
MRI head without and with contrast	8	Multiplanar thin sellar imaging. See statement regarding contrast in text under "Anticipated Exceptions."	O
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
CT head without and with contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
Venous sampling petrosal sinus	4	Indicated if MRI is negative or equivocal.	NS
CT head without contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
MRA head with or without contrast	3	Indicated if better visualization of carotid arteries needed.	O
CTA head	2		☼ ☼ ☼
X-ray sella	1		☼
Arteriography cerebral	1		☼ ☼ ☼
X-ray tomography head	1		☼
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

**Variant 5:****Hyperprolactinemia.**

<b>Radiologic Procedure</b>	<b>Rating</b>	<b>Comments</b>	<b><u>RRL*</u></b>
MRI head without and with contrast	8	Multiplanar thin sellar imaging. See statement regarding contrast in text under "Anticipated Exceptions."	O
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
CT head without and with contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
CT head without contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
MRA head with or without contrast	3	Indicated if better visualization of carotid arteries needed.	O
CTA head	2	For surgical planning or vascular detail if MRI and MRA contraindicated.	☼ ☼ ☼
X-ray sella	1		☼
Venous sampling petrosal sinus	1	Indicated in unusual cases in which lateralization is indeterminate.	NS
Arteriography cerebral	1		☼ ☼ ☼
X-ray tomography head	1		☼
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

**Clinical Condition:****Neuroendocrine Imaging****Variant 6:****Acromegaly/gigantism.**

<b>Radiologic Procedure</b>	<b>Rating</b>	<b>Comments</b>	<b><u>RRL*</u></b>
MRI head without and with contrast	8	Multiplanar thin sellar imaging. See statement regarding contrast in text under "Anticipated Exceptions."	O
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
CT head without and with contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
CT head without contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
Venous sampling petrosal sinus	3	Indicated in unusual cases in which lateralization is indeterminate.	NS
MRA head with or without contrast	3	Indicated if better visualization of carotid arteries needed.	O
CTA head	2	For surgical planning or vascular detail if MRI and MRA contraindicated.	☼ ☼ ☼
X-ray sella	1		☼
X-ray tomography head	1		☼
Arteriography cerebral	1		☼ ☼ ☼
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

**Variant 7:****Growth hormone deficiency, growth deceleration, panhypopituitarism.**

<b>Radiologic Procedure</b>	<b>Rating</b>	<b>Comments</b>	<b><u>RRL*</u></b>
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
MRI head without and with contrast	5	Multiplanar thin sellar imaging. See statement regarding contrast in text under "Anticipated Exceptions."	O
CT head without contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
CT head without and with contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
MRA head with or without contrast	2		O
CTA head	2	For surgical planning or vascular detail if MRI and MRA contraindicated.	☼ ☼ ☼
Arteriography cerebral	1		☼ ☼ ☼
X-ray tomography head	1		☼
X-ray sella	1		☼
Venous sampling petrosal sinus	1		NS
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

**Clinical Condition:****Neuroendocrine Imaging****Variant 8:****Diabetes insipidus.**

<b>Radiologic Procedure</b>	<b>Rating</b>	<b>Comments</b>	<b><u>RRL*</u></b>
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
MRI head without and with contrast	6	Multiplanar thin sellar imaging. See statement regarding contrast in text under “Anticipated Exceptions.”	O
MRA head with or without contrast	4	See statement regarding contrast in text under “Anticipated Exceptions.”	O
CT head without contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
CT head without and with contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
CTA head	2	For surgical planning or vascular detail if MRI and MRA contraindicated.	☼ ☼ ☼
Venous sampling petrosal sinus	1		NS
X-ray tomography head	1		☼
Arteriography cerebral	1		☼ ☼ ☼
X-ray sella	1		☼
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

**Variant 9:****Pituitary apoplexy.**

<b>Radiologic Procedure</b>	<b>Rating</b>	<b>Comments</b>	<b><u>RRL*</u></b>
MRI head without and with contrast	8	Multiplanar thin sellar imaging. See statement regarding contrast in text under “Anticipated Exceptions.”	O
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
CT head without contrast	6		☼ ☼ ☼
MRA head with or without contrast	4	Indicated if better visualization of carotid arteries needed. See statement regarding contrast in text under “Anticipated Exceptions.”	O
CT head without and with contrast	4	Indicated if MRI not available or contraindicated.	☼ ☼ ☼
CTA head	4		☼ ☼ ☼
X-ray tomography head	1		☼
Venous sampling petrosal sinus	1		NS
Arteriography cerebral	1		☼ ☼ ☼
X-ray sella	1		☼
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

**Clinical Condition:** Neuroendocrine Imaging

**Variant 10:** Postoperative sella.

Radiologic Procedure	Rating	Comments	RRL*
MRI head without and with contrast	8	Multiplanar thin sellar imaging. See statement regarding contrast in text under “Anticipated Exceptions.”	O
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
CT head without and with contrast	4	CT may be indicated to assess bony anatomy and if MRI is not available or contraindicated.	☼ ☼ ☼
CT head without contrast	4	CT may be indicated to assess bony anatomy and if MRI is not available or contraindicated.	☼ ☼ ☼
CTA head	4		☼ ☼ ☼
MRA head with or without contrast	2		O
X-ray sella	1		☼
X-ray tomography head	1		☼
Arteriography cerebral	1		☼ ☼ ☼
Venous sampling petrosal sinus	1		NS
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

**Variant 11:** Precocious puberty.

Radiologic Procedure	Rating	Comments	RRL*
MRI head without and with contrast	8	Multiplanar thin sellar imaging. See statement regarding contrast in text under “Anticipated Exceptions.”	O
MRI head without contrast	7	Multiplanar thin sellar imaging.	O
CT head without contrast	2		☼ ☼ ☼
CT head with contrast	2	If MRI not available or contraindicated.	☼ ☼ ☼
CTA head	2		☼ ☼ ☼
MRA head with or without contrast	2		O
X-ray sella	1		☼
X-ray tomography head	1		☼
Arteriography cerebral	1		☼ ☼ ☼
Venous sampling petrosal sinus	1		NS
<b>Rating Scale:</b> 1,2,3 Usually not appropriate; 4,5,6 May be appropriate; 7,8,9 Usually appropriate			<b>*Relative Radiation Level</b>

# NEUROENDOCRINE IMAGING

Expert Panel on Neurologic Imaging:  
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## Summary of Literature Review

The imaging approach to the hypothalamic pituitary axis is based on specific endocrine testing suggested by clinical signs and symptoms. Endocrine disorders are generally characterized by an excess or deficiency of specific hormones. Hormone excess is diagnosed under conditions that would ordinarily suppress hormone secretion. Endocrine deficiencies are diagnosed on the basis of hormone measurements under conditions of stimulation. Specific clinical syndromes of hormonal disorders are determined by the physiologic role of that particular hormone.

The hypothalamic pituitary axis consists of two separate neuroendocrine organs: the anterior pituitary system and the posterior pituitary system. The hormones of the anterior pituitary are thyroid-stimulating hormone (TSH), adrenal corticotrophic hormone (ACTH), prolactin (PRL), growth hormone (GH), and the gonadotropins (FSH and LH). These are secreted under the influence of hypothalamic trophic factors, corticotrophin releasing factor (CRF), thyrotropin releasing factor (TRF), and somatostatin- and gonadotropin-releasing hormone (GnRH). Prolactin release is under the control of a dopaminergic circuit. The hypothalamic-releasing hormones are transported to the pituitary gland by the hypothyseal portal system.

The posterior pituitary gland consists of axonal terminations of neurons whose cell bodies are located in the hypothalamus. The principal hormones secreted by these cells are oxytocin and vasopressin or antidiuretic hormone (ADH). The hypothalamus also participates in complex mediation of food intake, temperature

regulation, sleep and arousal, memory, thirst, and other autonomic functions.

Structural causes of obesity, anorexia, central hypothermia and hyperthermia, insomnia, and hypersomnia are only very rarely demonstrated in the hypothalamus and pituitary gland. Imaging in patients who present with these symptoms absent other specific neurological or endocrine abnormality is almost always unrewarding [1]. An exception is in children in whom the “diencephalic syndrome” of hypothalamic lesions is relatively common. Also, precocious puberty in children can result from hypothalamic lesions.

Pituitary adenomas are the most common lesions of the pituitary gland. These may secrete prolactin, TSH, GH, ACTH, or gonadotropins. Prolactinomas are the most common and are generally present as microadenomas in premenopausal females with amenorrhea and galactorrhea. PRL elevation by itself is nonspecific and may be due to a variety of medical, neurological, or pharmacological causes as well as pituitary adenoma, depending on serum hormone level. In males, prolactinomas may be entirely asymptomatic until visual symptoms occur, due to compression of the chiasm, or they may result in hypogonadotropic hypogonadism with loss of libido and impotence. Growth-hormone-secreting tumors generally present as larger lesions manifesting clinical acromegaly. Because of the gradual onset of deformity, these tumors may be present for many years and grow to substantial size prior to their detection. In a prepubertal individual the growth-hormone-secreting tumor may result in gigantism. TSH- and ACTH-secreting tumors may present at very small size because the impact of their hormone product is usually apparent more rapidly. Gonadotropin-secreting tumors are rare [2].

Precocious puberty and other neurological symptoms can be produced by hypothalamic lesions such as hamartoma. MRI is generally indicated in all patients with endocrinologically confirmed precocious puberty, especially when rapid progression of development and neurological symptoms are present [3-7].

Posterior pituitary dysfunction with loss of antidiuretic hormone results in the clinical syndrome of diabetes insipidus. This may occur as a transient phenomenon after trauma or neurosurgical procedures. The etiology is usually evident, and the phenomenon is frequently transient. Imaging is performed to search for the cause of stalk transection, which can be a manifestation of numerous sellar or parasellar pathologies, trauma, or congenital. Rarely, the hormone is absent developmentally. The syndrome of inappropriate ADH is usually due to an extracranial source. Frequently this is a paraneoplastic phenomenon related to small-cell lung carcinoma, though a variety of pulmonary diseases and pharmacological disturbances can result in SIADH.

Other common mass lesions that may affect the neuroendocrine system are germ-line tumors, meningioma, craniopharyngioma, and Rathke’s cleft cyst

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among others. Metastatic lesions may affect the sella. Sarcoid and other inflammatory processes occur in the sellar and suprasellar regions as well. Pituitary apoplexy is a syndrome of headache ophthalmoplegia and visual loss that results from pituitary hemorrhage. In the postpartum period, pituitary infarcts may occur, and hypophysitis is an uncommon disorder resulting in endocrine disturbance and other symptoms [8].

Classically, radiography and pluridirectional x-ray tomography were the mainstays of sellar imaging. Computed tomography (CT) largely replaced these modalities through the seventies and eighties. More recently, magnetic resonance imaging (MRI) has largely supplanted CT [9-18]. MRI for sellar pathology includes thin-section multiplanar imaging with slice thickness of 3 mm or less, often before and after contrast administration. Other techniques that are used for evaluating this anatomical region are computed tomography angiography (CTA), magnetic resonance angiography (MRA), direct catheter angiography, and petrosal sinus sampling.

Radiography and pluridirectional tomography are insensitive and nonspecific imaging modalities for evaluating sellar pathology. Pituitary microadenoma and even small pituitary macroadenomas are frequently associated with a normal sella size. The sella turcica can be enlarged when no neoplasm or mass is present. This is due to pulsations of cerebral spinal fluid (CSF) transmitted through a developmental or acquired dehiscence of the diaphragm sella in the empty sella syndrome. Therefore, these imaging modalities are rarely, if ever, used productively in the evaluation of endocrine disease [19].

CT revolutionized evaluation of the sella and suprasellar region. Due to the ability of CT, especially with intravenous contrast, to depict pathology within the unenlarged sella, and its ability to visualize suprasellar pathology noninvasively, this technique facilitates accurate diagnosis of neuroendocrine abnormality [11,20-23]. Pituitary microadenomas and macroadenomas are reliably detected. It is, however, difficult at times to distinguish the tumor from the optic chiasm and to diagnose cavernous invasion. Also, cystic lesions of the suprasellar region may be confused with normal CSF.

Additionally, artifact due to dental amalgam, difficulty in obtaining reliable contrast enhancement, and awkward positioning for direct coronal scanning limit the utility of this imaging modality. In the hands of experienced radiologists this technique can result in excellent diagnostic accuracy, though the examinations are sometimes hard to interpret despite excellent technique.

MRI provides excellent noninvasive evaluation of the hypothalamus and pituitary gland. It is the only imaging modality that reliably depicts the hypothalamus in a useful fashion. It depicts the anatomy of the pituitary gland, infundibulum, optic chiasm, cavernous sinuses, and neighboring vascular structures accurately and noninvasively [24-36].

The addition of gadolinium facilitates diagnosis of microadenoma and increases the confidence with which cavernous sinus invasion can be diagnosed or excluded [37,38]. The specific bony landmarks may be difficult to demonstrate, but the signal pattern of sphenoid sinus mucosa permits assessment of septa for operative planning [39,40]. Visualization of vascular structures in the parasellar region or even intrasellar carotid artery loop or aneurysm is crucial in some cases.

Angiography is reserved for those patients in whom vascular pathology is known or suspected on the basis of clinical or radiological findings. Aneurysm is the most important vascular lesion in the parasellar region, but these lesions rarely present as endocrine disorders. Knowledge of vascular anatomy guides surgery. Occasionally, a sellar lesion may grow to displace or encase the carotid arteries or other major intracranial vessels. Interventional neuroradiology procedures can be planned on the basis of CTA, MRA, or catheter angiography [41].

Petrosal sinus venous sampling is reserved for those cases in which a definite excess of pituitary hormone is present, medical management has failed, sectional imaging is negative or equivocal and surgery is planned [42]. When a significant discrepancy in hormone level, usually ACTH, exists between the vessels studied, tumor localization is very accurate. Complications occur uncommonly in experienced hands.

A significant problem encountered in CT and MRI imaging of the pituitary, particularly when endocrine findings suggest microadenoma, is the false positive examination. Since the endocrine studies confirm the presence of a lesion, and first-line therapy is usually medical, false negative examinations are less problematic once chiasmatic compression has been excluded. Approximately 20% of the population may harbor small incidental nonfunctioning adenomas or cysts. It is important, therefore, that the probability of disease be high in the target population if a positive MRI is to be relied upon for surgical planning. Additional problems are created by variations in size of the pituitary gland, which occur normally in response to physiological hormonal changes. The gland may enlarge in puberty and pregnancy. Pituitary hyperplasia in hypothyroidism may simulate a pituitary adenoma in some patients. Similar problems arise in imaging the posterior pituitary, since up to 29% of normal subjects do not demonstrate a bright posterior pituitary [43,44].

### **Anticipated Exceptions**

Nephrogenic systemic fibrosis (NSF) is a disorder with a scleroderma-like presentation and a spectrum of manifestations that can range from limited clinical sequelae to fatality. It appears to be related to both underlying severe renal dysfunction and the administration of gadolinium-based contrast agents. It has occurred primarily in patients on dialysis, rarely in patients with very limited glomerular filtration rate (GFR) (ie, <30 mL/min/1.73m<sup>2</sup>), and almost never in other patients. There is growing literature regarding NSF.

Although some controversy and lack of clarity remain, there is a consensus that it is advisable to avoid all gadolinium-based contrast agents in dialysis-dependent patients unless the possible benefits clearly outweigh the risk, and to limit the type and amount in patients with estimated GFR rates <30 mL/min/1.73m<sup>2</sup>. For more information, please see the [ACR Manual on Contrast Media](#) [45].

### Relative Radiation Level Information

Potential adverse health effects associated with radiation exposure are an important factor to consider when selecting the appropriate imaging procedure. Because there is a wide range of radiation exposures associated with different diagnostic procedures, a relative radiation level (RRL) indication has been included for each imaging examination. The RRLs are based on effective dose, which is a radiation dose quantity that is used to estimate population total radiation risk associated with an imaging procedure. Patients in the pediatric age group are at inherently higher risk from exposure, both because of organ sensitivity and longer life expectancy (relevant to the long latency that appears to accompany radiation exposure). For these reasons, the RRL dose estimate ranges for pediatric examinations are lower as compared to those specified for adults (see Table below). Additional information regarding radiation dose assessment for imaging examinations can be found in the ACR Appropriateness Criteria<sup>®</sup> [Radiation Dose Assessment Introduction](#) document.

Relative Radiation Level Designations		
Relative Radiation Level*	Adult Effective Dose Estimate Range	Pediatric Effective Dose Estimate Range
O	0 mSv	0 mSv
☼	<0.1 mSv	<0.03 mSv
☼☼	0.1-1 mSv	0.03-0.3 mSv
☼☼☼	1-10 mSv	0.3-3 mSv
☼☼☼☼	10-30 mSv	3-10 mSv
☼☼☼☼☼	30-100 mSv	10-30 mSv
*RRL assignments for some of the examinations cannot be made, because the actual patient doses in these procedures vary as a function of a number of factors (eg, region of the body exposed to ionizing radiation, the imaging guidance that is used). The RRLs for these examinations are designated as NS (not specified).		

### Supporting Document(s)

- [ACR Appropriateness Criteria<sup>®</sup> Overview](#)
- [Procedure Contrast Information](#)
- Evidence table under review

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The ACR Committee on Appropriateness Criteria and its expert panels have developed criteria for determining appropriate imaging examinations for diagnosis and treatment of specified medical condition(s). These criteria are intended to guide radiologists, radiation oncologists and referring physicians in making decisions regarding radiologic imaging and treatment. Generally, the complexity and severity of a patient's clinical condition should dictate the selection of appropriate imaging procedures or treatments. Only those examinations generally used for evaluation of the patient's condition are ranked. Other imaging studies necessary to evaluate other co-existent diseases or other medical consequences of this condition are not considered in this document. The availability of equipment or personnel may influence the selection of appropriate imaging procedures or treatments. Imaging techniques classified as investigational by the FDA have not been considered in developing these criteria; however, study of new equipment and applications should be encouraged. The ultimate decision regarding the appropriateness of any specific radiologic examination or treatment must be made by the referring physician and radiologist in light of all the circumstances presented in an individual examination.