

# Endocrinology

## ICD-10 documentation

### Seven key impacts to ICD-10 documentation

1. Disease or disorder site
2. Acuity and/or encounter status of treatment
3. Etiology, causative agent, or disease type and injury/  
poisoning cause, intent, activity at the time of the event  
and place event occurred
4. Underlying and associated conditions
5. Manifestations
6. Complications or adverse events
7. Supporting info such as lab values or socioeconomic key  
impacts to ICD-10 documentation

### Most commonly used diagnoses

ICD-9 Name	ICD-9 Code	ICD-10 Code	ICD-10 Name	Documentation tips
Abnormal weight gain	<b>783.1</b>	<b>R63.5</b>	Abnormal weight gain	
Adrenogenital disorders	<b>255.2</b>	<b>E25.0</b>	Congenital adrenogenital disorders associated with enzyme deficiency	Endocrine and Metabolic Disorders <ul style="list-style-type: none"> <li>• List the specific disorder, disease, defect, deficiency or syndrome</li> <li>• Identify any underlying condition (e.g. feeding difficulties)</li> <li>• Specify the significance of abnormal lab finding</li> <li>• Detail disorders as being congenital, hereditary, primary, idiopathic, secondary, pseudo, and familial when applicable (e.g. hypopituitarism)</li> </ul>
		<b>E25.8</b>	Other adrenogenital disorders	
		<b>E25.9</b>	Adrenogenital disorder, unspecified	
Delay in sexual development and puberty, not elsewhere classified	<b>259.0</b>	<b>E30.0</b>	Delayed puberty	
Dwarfism, not elsewhere classified	<b>259.4</b>	<b>E34.3</b>	Short stature due to endocrine disorder	
Panhypopituitarism	<b>253.2</b>	<b>E23.0</b>	Hypopituitarism	
Pituitary dwarfism	<b>253.3</b>	<b>E23.0</b>	Hypopituitarism	
Precocious sexual development and puberty, not elsewhere classified	<b>259.1</b>	<b>E30.1</b>	Precocious puberty	
		<b>E30.8</b>	Other disorders of puberty	



ICD-9 Name	ICD-9 Code	ICD-10 Code	ICD-10 Name	Documentation tips
Chondroectodermal dysplasia	<b>756.55</b>	<b>Q77.6</b>	Chondroectodermal dysplasia	Refer to 7 key impacts to ICD-10 documentation
Congenital hypothyroidism	<b>243</b>	<b>E00.0</b>	Congenital iodine-deficiency syndrome, neurological type	
		<b>E00.1</b>	Congenital iodine-deficiency syndrome, myxedematous type	
		<b>E00.2</b>	Congenital iodine-deficiency syndrome, mixed type	
		<b>E00.9</b>	Congenital iodine-deficiency syndrome, unspecified	
		<b>E03.0</b>	Congenital hypothyroidism with diffuse goiter	
		<b>E03.1</b>	Congenital hypothyroidism without goiter	
Delayed milestones	<b>783.42</b>	<b>R62.0</b>	Delayed milestone in childhood	
Diabetes insipidus	<b>253.5</b>	<b>E23.2</b>	Diabetes insipidus	Diabetes <ul style="list-style-type: none"> <li>• Identify the type (i.e. Type I or Type II)</li> <li>• Clarify any cause and effect relationship between diabetes and other conditions (e.g. diabetic peripheral vascular disease)</li> <li>• Specify long-term use of insulin for type II diabetes</li> <li>• List the underlying condition, drug, or chemical responsible for secondary diabetes (e.g. steroid-induced)</li> <li>• Detail insulin underdosing or overdosing related to an insulin pump malfunction</li> <li>• Differentiate when diabetes is accompanied by 'hypo' or 'hyper' glycemia <ul style="list-style-type: none"> <li>◦ State if hypoglycemia is or is not causing a coma</li> </ul> </li> </ul>
Diabetes mellitus without mention of complication, type II or unspecified type, not stated as uncontrolled	<b>250.00</b>	<b>E11.9</b>	Type 2 diabetes mellitus without complications	
		<b>E13.9</b>	Other specified diabetes mellitus without complications	
Diabetes mellitus without mention of complication, type I [juvenile type], not stated as uncontrolled	<b>250.01</b>	<b>E10.9</b>	Type 1 diabetes mellitus without complications	
Failure to thrive	<b>783.41</b>	<b>R62.51</b>	Failure to thrive (child)	Refer to 7 key impacts to ICD-10 documentation
Gonadal dysgenesis	<b>758.6</b>	<b>Q96.0</b>	Karyotype 45, X	
		<b>Q96.1</b>	Karyotype 46, X iso (Xq)	
		<b>Q96.2</b>	Karyotype 46, X with abnormal sex chromosome, except iso (Xq)	
		<b>Q96.3</b>	Mosaicism, 45, X/46, XX or XY	
		<b>Q96.4</b>	Mosaicism, 45, X/other cell line(s) with abnormal sex chromosome	
		<b>Q96.8</b>	Other variants of Turner's syndrome	
		<b>Q96.9</b>	Turner's syndrome, unspecified	

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Glucocorticoid deficiency	<b>255.41</b>	<b>E27.1</b>	Primary adrenocortical insufficiency	<p>Malignant neoplasm</p> <ul style="list-style-type: none"> <li>• Identify the site (e.g. cortex, medulla, cerebellum)</li> <li>• State the morphology (e.g. malignant, primary/secondary)</li> <li>• Indicate the stage and any metastatic site</li> <li>• List any related exposure to smoke (second hand smoke)</li> <li>• Detail when presented for treatment related to neoplasm (e.g. chemotherapy)</li> </ul>
		<b>E27.2</b>	Addisonian crisis	
		<b>E27.3</b>	Drug-induced adrenocortical insufficiency	
		<b>E27.40</b>	Unspecified adrenocortical insufficiency	
		<b>E27.49</b>	Other adrenocortical insufficiency	
Lack of normal physiological development, unspecified	<b>783.40</b>	<b>R62.50</b>	Unspecified lack of expected normal physiological development in childhood	
		<b>R62.59</b>	Other lack of expected normal physiological development in childhood	
Malignant neoplasm of cerebellum NOS	<b>191.6</b>	<b>C71.6</b>	Malignant neoplasm of cerebellum	
Malignant neoplasm of adrenal gland	<b>194.0</b>	<b>C74.00</b>	Malignant neoplasm of cortex of unspecified adrenal gland	
		<b>C74.01</b>	Malignant neoplasm of cortex of right adrenal gland	
		<b>C74.02</b>	Malignant neoplasm of cortex of left adrenal gland	
		<b>C74.10</b>	Malignant neoplasm of medulla of unspecified adrenal gland	
		<b>C74.11</b>	Malignant neoplasm of medulla of right adrenal gland	
		<b>C74.12</b>	Malignant neoplasm of medulla of left adrenal gland	
		<b>C74.90</b>	Malignant neoplasm of unspecified part of unspecified adrenal gland	
		<b>C74.91</b>	Malignant neoplasm of unspecified part of right adrenal gland	
<b>C74.92</b>	Malignant neoplasm of unspecified part of left adrenal gland			
Multiple epiphyseal dysplasia	<b>756.56</b>	<b>Q78.3</b>	Progressive diaphyseal dysplasia	Refer to 7 key impacts to ICD-10 documentation
Nonspecific abnormal results of other endocrine function study	<b>794.6</b>	<b>R94.7</b>	Abnormal results of other endocrine function studies	
Obstructive sleep apnea (adult) (pediatric)	<b>327.23</b>	<b>G47.33</b>	Obstructive sleep apnea (adult) (pediatric)	
Osteogenesis imperfecta	<b>756.51</b>	<b>Q78.0</b>	Osteogenesis imperfecta	
Osteopetrosis	<b>756.52</b>	<b>Q78.2</b>	Osteopetrosis	
Osteopoikilosis	<b>756.53</b>	<b>Q78.8</b>	Other specified osteochondrodysplasias	
Other congenital osteodystrophies	<b>756.59</b>	<b>Q77.3</b>	Chondrodysplasia punctata	
		<b>Q78.5</b>	Metaphyseal dysplasia	
		<b>Q78.6</b>	Multiple congenital exostoses	
		<b>Q78.8</b>	Other specified osteochondrodysplasias	

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Other congenital osteodystrophies	<b>756.59</b>	<b>Q77.3</b>	Chondrodysplasia punctata	
		<b>Q78.5</b>	Metaphyseal dysplasia	
		<b>Q78.6</b>	Multiple congenital exostoses	
		<b>Q78.8</b>	Other specified osteochondrodysplasias	
Other dyspnea and respiratory abnormality	<b>786.09</b>	<b>R06.00</b>	Dyspnea, unspecified	
		<b>R06.09</b>	Other forms of dyspnea	
		<b>R06.3</b>	Periodic breathing	
		<b>R06.83</b>	Snoring	
		<b>R06.89</b>	Other abnormalities of breathing	
Other specified hypoglycemia	<b>251.1</b>	<b>E08.649</b>	Diabetes mellitus due to underlying condition with hypoglycemia without coma	
		<b>E16.0</b>	Drug-induced hypoglycemia without coma	
		<b>E16.1</b>	Other hypoglycemia	
Polyostotic fibrous dysplasia of bone	<b>756.54</b>	<b>Q78.1</b>	Polyostotic fibrous dysplasia	
Short stature	<b>783.43</b>	<b>R62.52</b>	Short stature (child)	
Toxic diffuse goiter without mention of thyrotoxic crisis or storm	<b>242.00</b>	<b>E05.00</b>	Thyrotoxicosis with diffuse goiter without thyrotoxic crisis or storm	
Unspecified hypothyroidism	<b>244.9</b>	<b>E03.9</b>	Hypothyroidism, unspecified	

The ICD-10 Provider Specialty Tool Kit is meant to assist providers with documentation best practices necessary for the transition from ICD-9-CM to ICD-10-CM. This tool kit includes the most commonly used diagnoses for a specialty using 2012 claims data for top encounters and top number of dollars per diagnosis. The tool kit can be used as a quick reference guide for providers and can also be shared with Practice Managers. This tool kit is not meant to be a coding cross walk and therefore should not be used as such. Questions regarding this tool kit and/or ICD-10 should be submitted via email to ICD10@choa.org.