Supplemental eTable 1. Clinical presentation and final pathology in patients with incidental versus non-incidental mode of adrenal tumor discovery [a]

Variable	Incidentalomas	Non-incidentalomas	*P* value
N, %	472 (67%)	233 (33%)	
Sex Female, n (%)	249 (53%)	124 (53%)	0.90
Age at diagnosis years, median (range)	60 (19-88)	54 (18-91)	**<.001**
Adrenal mass diameter cm, median (range)	5 (4-20)	6 (4-24.4)	**<.001**
Diagnosis n (%)			
Adrenocortical Adenoma (ACA)	184 (39%)	32 (14%)	**<.001**
Other benign	104 (22%)	12 (5%)	
Pheochromocytoma	86 (18%)	72 (31%)	

Other malignant	61 (13%)	66 (28%)	
Adrenocortical carcinoma (ACC)	37 (8%)	51 (22%)	

[a]Range was defined as minimum to maximum value.

Supplemental eTable 2. Clinical presentation of patients with adrenal tumors based on tumor size cutoff of 6 cm[a]

	Tumor size ≤ 6 cm	**Tumor size > 6cm**	***P* value**
N, %	449 (64%)	256 (36%)	
Sex Female, n (%)	234 (52%)	139 (54%)	0.60
Age at diagnosis years, median (range)	60 (18-91)	56 (19-87)	**<.001**
Mode of discovery (n (%)) Incidental Hormone excess Cancer staging Other[b]	 326 (73%) 62 (14%) 43 (10%) 18 (4%)	 145 (57%) 46 (18%) 18 (7%) 47 (18%)	 **<.001**
Adrenalectomy yes, n (%)	274 (61%)	183 (72%)	**.005**

Diagnosis, n (%)			
Adenoma	194 (43%)	24 (9%)	**<.001**
Pheochromocytoma	112 (25%)	46 (18%)	
Other malignant	76 (17%)	52 (20%)	
Other benign	53 (12%)	60 (23%)	
ACC	14 (3%)	74 (29%)	
Bilateral n (%)	74 (16%)	30 (12%)	.08

[a]Range was defined as minimum to maximum value

[b]Other: mass effect and B symptoms

Abbreviations used: ACC, adrenocortical carcinoma;