Anesthetic Management of Pheochromocytoma
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ABSTRACT
Pheochromocytomas (PCCs) are rare neuroendocrine tumors originating from the chromaffin tissue.¹ These tumors produce, store, and secrete catecholamines. Most PCCs produce both epinephrine and norepinephrine, with the percentage of secreted norepinephrine being greater than that secreted by the normal gland.²

The incidence of PCCs is 2 to 8 per 1 million populations per year. They are highly vascular and catecholamine-producing tumors. They are derived from the sympathetic or parasympathetic nervous system. It is estimated that 0.1% of all hypertensives may have neuroendocrine tumors like PCC. In these patients, usual presenting signs and symptoms are tachycardia, hypertension, etc., which are the result of sympathetic stimulation. The literature also shows a high prevalence of familial and extra adrenal tumors in patients carrying germline mutations in genes.³ It may be as high as 50%.¹ Surgery and anesthesia in these patients pose a high risk of cardiovascular complications from hypertensive crisis. Here, we report a case of PCC scheduled for surgical resection.

Keywords: Catecholamines, Hypertension, Hypertensive crises, Pheochromocytoma.

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INTRODUCTION
Catecholamine-producing tumors which are derived from the sympathetic or parasympathetic nervous system are called pheochromocytomas (PCCs).³ Once diagnosed as PCCs, their removal can prevent lethal complications like hypertensive crises. Main aims of anesthetic management include proper preanesthetic checkup, intraoperative fluid management as well as management of complications like hypertension and posttumor removal hypotension. The chief presenting complaints are fluctuating blood pressure (BP), sweating, and palpitations.¹ Catecholamine release, if not controlled, can result in severe hypertension, cerebrovascular, and myocardial complications.³ This patient group poses a challenge to anesthesiologists in managing them intraoperatively as well as in ICU.

Here, we report a case of PCC for surgical resection. We discuss the preoperative evaluation, intraoperative management, and postoperative care of patients who will undergo resection of PCC.

Perioperative Management
Thorough preoperative assessments of these patients result in successful perioperative management. Roizen et al. have proposed a set of criteria to objectively gauge the efficacy of adequate preoperative alpha blockade, which are as follows:

- No-in-hospital BP of <160/90 mm Hg for 24 hours prior to surgery
- No orthostatic hypotension with BP <80/45 mm Hg
- No ST or T wave changes for 1-week prior to surgery
- Not more than five premature ventricular contractions per minute.

CASE REPORT
A 32-year-old female patient came with a history of left flank pain, dizziness, headache, and occasional palpitations since 1 month. The patient is a known case of hypothyroidism (8 years) on treatment with tab. telmisartan 40 mg + hydrochlorothiazide 12.5 mg. General random blood sugar (GRBS) was monitored every 4 hours. The patient was admitted at the Department of Surgery and anesthesia in these patients pose a high risk of cardiovascular complications from hypertensive crisis. Here, we report a case of PCC scheduled for surgical resection.

The patient was a 70 kg female with a height of 105 cm and a body mass index (BMI) of 29. She was obese, with short neck and airway of mallampati grade II. Her BP was 170/110 mm Hg, and heart rate (HR) was 95 bpm. The patient was advised to stop all antihypertensives and started on tab. phenoxybenzamine 10 mg tid. The patient was started on inj. human actrapid according to the sliding scale and started on inj. human actrapid according to the sliding scale and general random blood sugar (GRBS) was monitored every 4 hours. Preop orders included nil per orally (NPO)-6 hours, tab. alprazolam 25 mg for anxiolysis, inj. ranitidine and ondansetron, tab. thyronorm 100 µg on the morning of surgery, and skipping morning insulin dose.

The patient was continuously monitored in ICU—her BP and HR readings are shown in Table 1.
The patient’s vitals before shifting to OR were as follows: BP 170/100 mm Hg, HR 96 bpm, FBS—151 mg/dL, and serum electrolytes were normal.

**Anesthetic Management**

Preoperative counseling of the patient and attenders was done. Operation room was prepared and all drugs required including inotropes, vasodilators, emergency drugs, vasopressors, antiarrythmics, blood, and blood products were arranged. Preoperative consent was obtained.

Electrocardiogram (ECG), oxygen saturation, noninvasive blood pressure (NIBP), temperature, and urine output were monitored and two-wide bore IV cannulas (18 G and 20 G) were secured and IV infusion of crystalloids was started. End tidal CO$_2$ monitoring was commenced after intubation under all aseptic precaution—the right-hand radial artery was cannulated with a 20 G cannula; arterial BP monitoring was instituted. Patient’s vitals were: arterial BP 180/100 mm Hg (Fig. 1), HR 100 bpm, and SPO$_2$ 100%.

Injection Esmolol 30 mg in 100 mL normal saline (NS) was started IV, 10 minutes before intubation. Injection sodium nitroprusside infusion 3 mL/hour was started in view of arterial BP readings of 200/120 mm Hg. Inj. midazolam 2 mg IV and inj. fentanyl 100 µg were given as premedication. After 15 minutes, the vitals were: BP 180/100 mm Hg and HR 101 bpm.

The patient was preoxygenated with 100% oxygen, induced with inj. propofol 100 mg, sevoflurane + oxygen, inj. rocuronium 30 mg was given, the patient intubated with a cuffed endotracheal tube of size 7.0, the bilateral air entry was checked, and the tube was fixed at 19 cm: BP 150/100 mm Hg and HR 118 bpm. The patient was maintained with O$_2$ + N$_2$O + isoflurane 1% and intermittent doses of inj. vecuronium 0.5 mg. Injection paracetamol 1 g was administered IV as a part of multimodal analgesia. Under all aseptic precautions, the right subclavian central line was secured. The patient was then turned to the right lateral position and a thoracic epidural catheter was placed between T7 and T8 vertebrae and secured. The patient was then positioned for surgery in the right lateral position with adequate padding. Baseline BP was 124/67 mm Hg with a HR of 69 bpm before surgical incision. After 5 minutes of incision, BP increased to 170/100 mm Hg and HR was 85 bpm. Injection sodium nitroprusside infusion increased to 5 mL/hour and isoflurane was increased to 1.2%, inj. magnesium sulfate 1 g in 100 mL NS slow IV started as a membrane stabilizer to prevent arrythmias. When surgeons started handling the tumor, BP shot up to 233/117 mm Hg (Fig. 2), 200/118 mm Hg, and HR 103 bpm. Injection nitroglycerin infusion started at 2.5 mL/hour. Five minutes later, BP was 180/88 mm Hg and HR was 104 bpm. When surgeons excised the tumor out (Fig. 3), the BP dropped to 57/39 mm Hg with a HR of 144 bpm. Injections sodium nitroprusside and nitroglycerin were stopped. Injection noradrenaline was started at 5 mL/hour (0.08 mg/mL), inj. phenylephrine 30 µg—2 bolus doses and inj. mephentermine 6 mg—2 bolus doses were administered. Isoflurane reduced to 0.6% MAC. Injection fentanyl 50 µg was administered after tumor excision. Injection dexamethasone 8 mg and inj. hydrocortisone 100 mg were also administered.

<table>
<thead>
<tr>
<th>Time</th>
<th>BP (mm Hg)</th>
<th>HR (bpm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>4 PM (day before surgery)</td>
<td>170/110</td>
<td>95</td>
</tr>
<tr>
<td>6 PM (day before surgery)</td>
<td>166/100</td>
<td>96</td>
</tr>
<tr>
<td>8 PM (day before surgery)</td>
<td>160/100</td>
<td>95</td>
</tr>
<tr>
<td>12 AM (day before surgery)</td>
<td>156/100</td>
<td>94</td>
</tr>
<tr>
<td>4 AM (day of surgery)</td>
<td>140/100</td>
<td>90</td>
</tr>
<tr>
<td>6 AM (day of surgery)</td>
<td>156/100</td>
<td>90</td>
</tr>
<tr>
<td>8 AM (day of surgery)</td>
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<td>90</td>
</tr>
<tr>
<td>9 AM (day of surgery)</td>
<td>160/100</td>
<td>93</td>
</tr>
</tbody>
</table>

**Fig. 1:** Baseline vitals

**Fig. 2:** Hypertensive response to stimulation

**Fig. 3:** Excision of tumor
Injection noradrenaline infusion was continued at 3 mL/hour. Random blood sugar (RBS) estimation was done which was 20 mg/dL. It was corrected with 25% dextrose and maintained with dextrose in normal saline. After 20 minutes, repeated RBS showed 214 mg/dL. During closure, BP increased to 200/100 mm Hg and the HR was 118 bpm, following which epidural was activated with 6 mL of 0.2% ropivacaine with 25 µg of fentanyl. BP got stabilized at 150/74 mm Hg and the HR was 110 bpm. Injection noradrenaline infusion was continued. Arterial blood gas (ABG) analysis was done which was satisfactory. Before extubation, inj. Esmolol 10 mg slow IV was administered.

The patient was extubated after complete neuromuscular blockade reversal with inj. neostigmine 2.5 mg and glycopyrrolate 0.5 mg. After the extubation, BP was 118/70 mm Hg, with an HR of 98 bpm. The patient had received 4 pints of Ringer’s lactate, 3 pints of 0.9% saline, and 1 pint of dextrose in normal saline (DNS). Blood loss was minimal. The patient was shifted to surgical intensive care unit (SICU). One hour post-surgery, BP was 90/60 mm Hg, inj. noradrenaline infusion continued at 3 mL/hour. Twenty-four hours post-surgery, BP was 110/70 mm Hg following which noradrenaline infusion was stopped. The patient was doing well. Seventy-two hours post-surgery, the patient had a high BP of 170/110 mm Hg following which tab. Prazosin 2.5 mg and tab. Metoprolol 50 mg were started orally. The patient was discharged after the BP was stabilized.

DISCUSSION

A majority of PCCs secrete mainly norepinephrine, sometimes paroxysmal but usually sustained and often in huge quantities. The majority of patients with PCC present with sustained severe hypertension. Released norepinephrine causes vasoconstriction in arteriolar and venous beds, thereby decreasing the circulating blood volume. PCCs are usually diagnosed and confirmed by raised urinary catecholamine and vanillylmandelic acid (VMA) in 24 hours urine samples. After the diagnosis, the tumor is localized accurately by computerized tomography (CT) scan, magnetic resonance imaging (MRI), and iodine-131-meta-iodobenzylguanidine imaging (MIBG) scans.

Preoperative optimization helps in preventing the wide variation in arterial BP during the intraoperative period. This is achieved by antiadrenergic drugs, i.e., alpha (α) and beta (β) blockers. The sympathetic blockade is achieved first by an α adrenergic blocker followed by an α–β blocker. Alpha blockade results in vasodilatation and tachycardia which is controlled by beta blockers. If beta blockade is achieved first, then there may be unopposed vasoconstriction in skeletal muscles causing hypertension. Twenty-four hours ambulatory BP monitoring should show a BP of 140/90 mm of Hg and a HR of less than 100 bpm. And no more than five premature ventricular contractions per minute. Four lying and standing BP should show a postural drop up to but not less than 80/45 mm Hg in the standing position. More drops indicate inadequate hydration. ECG should be free of ST changes for at least 2 weeks. Mortality is due to ignoring subtle changes of cardiomyopathy.

CONCLUSION

Patients with PCC are at a significant risk of major adverse cardiac complications in the perioperative period. Successful management requires careful preoperative optimization, meticulous intraoperative planning, and hemodynamic management. All patients need to be monitored vigilantly in the postoperative period given the high-risk of complications.

REFERENCES