Imageological Co-relation of Pituitary Tumors with Per-op Findings

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ABSTRACT

Our study is Prospective Study about correlation of MRI Brain Findings with Per-op Findings in case of Pituitary Tumors. This study was performed in 40 patients including both males & females; new & recurrent cases, in duration of 2 ½ years. All patients were undergone MRI Brain with special sequences for Pituitary Gland evaluation with other Hormonal analysis in some cases as per requirement. All patients were undergone Endoscopic Trans-sphenoidal Endonasal Excision of Pituitary Tumor. All patients were called for follow-up after 3 months for Surgical Excision evaluation with MRI Brain study. In follow-up cases we have studied our surgical achievement in relation of tumor clearance & consider any need of adjuvant therapy like Radiation therapy.

Key Words: Pituitary, Endoscopic transphenoidal, MRI Brain, Hormonal, Adjuvant therapy.

Introduction

Pituitary gland tumors consist of 10-15% of all intracranial tumors. They are responsible for Endocrinopathies and Pressure symptoms over optic apparatus, to cause significant morbidity & mortality in general population. Recent era, MRI Brain (P+C) with different sequences and views are available to study in detail about nature of Pituitary tumors. These tumors having different Variable Consistency, so mostly are Unpredictable regarding Surgical Excision. So this study deals with use of MRI Brain (P+C) to define criteria for involvement of vital structures by tumor & tumor consistency in benefits of surgical outcome.

Materials & Methods

- This is Prospective Study.
- It involves 40 patients over 2 ½ years duration & is performed at Care Hospital Group, Hyderabad.

Use of Specific Imaging:
1.5 T 3mm slice MRI Brain (P+C)
- T 1/T 2/ ADC/DWI,
- Coronal, Sagital, Axial views

- Surgery Planned: All patients were undergone Endoscopic Endonasal
Transphenoidal Excision of Pituitary Tumor (Binostril approach)

**Inclusion Criteria**
- 5 years to 70 years age group
- Fresh and recurrent cases
- Males and females both
- Acute and chronic presentation both

**Exclusion Criteria**
- Very high risk patients with Anesthesia ASA grade 4/5
- Age < 5yr and >70 year
- Recent major nasal surgery undergone cases
- Patients with bleeding disorders (Hemophilia), coagulopathies

**Objective**
To co-relate mri brain (p+c) findings with per-op findings in terms of:
- Tumor Consistency,
- Invasive Nature of Tumor,
- Tumor Extension,
- Tumor Bleed
- Classify Tumor Accordingly.

To Plan Management of Pituitary Tumors in form:
- To decide The Extent of Surgical Excision of Tumor
- To get Histopathological Exact diagnosis & classify tumor accordingly
- To achieve as much as possible maximum Tumor Clearance So that Hormonal Imbalance Symptoms or Pressure Symptoms caused by Tumor should be subsided at most.
- Post-op need of any Adjuvant Therapy in form of Radiation

**Others investigations**
- CT Paranasal sinuses
- Hormonal study- TFTs, Prolactin, Cortisol, GH
- Visual field charting & visual acuity

- Inferior Petrous venous sampling (microadenoma)
- Through general examination of patient

**Our Hypothesis**
If a Pituitary Tumor is

<table>
<thead>
<tr>
<th>Sr.No</th>
<th>MRI Brain Findings</th>
<th>Per-op Possibility</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>T 1: Isointense-Hypointense T 2: Hyperintense</td>
<td>Tumor should be Soft</td>
</tr>
<tr>
<td>2</td>
<td>T 1: Isointense-Hypointense T 2: Hypointense</td>
<td>Tumor should be Firm to Hard</td>
</tr>
<tr>
<td>3</td>
<td>T 1: Isointense-Hypointense T 2: Hyperintense FLAIR: Hyperintense</td>
<td>Tumor should be Cystic</td>
</tr>
<tr>
<td>4</td>
<td>T 1: Isointense-Hypointense T 2: Mixed intensity (Hypointense &amp;Hyperintense)</td>
<td>Tumor should be Mixed. More Part of T 2 is Hypointense → More is tumor firmness.</td>
</tr>
</tbody>
</table>

**Clinical Presentation for Surgery**
Most of pituitary tumors patients may be found to be asymptomatic for long time till it causes notifying abnormality.
Clinical presentation is due to –
1. Hormonal Imbalance
2. Mass Effect of tumor

Mass effect produced of tumors may cause-
- Headache
- Visual disturbances
- Intracranial Hemorrhage
- Raised Intracranial Pressure Changes
- The most common symptoms caused by mass effect by tumor is Visual Impairment:
  - Bitemporal Hemianopia(62%)
  - Unilateral Temporal Hemianopia(26%)
  - Optic Atrophy & Loss of Vision (39%).
- Involvement of Cavernous Sinus by Pituitary Tumor may manifest as extra-ocular nerve palsies (3rd, 4th, 6th cranial nerves involvement)

**Hormonal Imbalance & Manifestation:**
- Hyperprolactinomas present as-
  - Females-Infertility,
  - Amenorrhoea, Galactrhoea
Males-Loss of Libido, Impotence.
- Somatotropinomas- Gigantism and Acromegaly presentation
- Corticotropinomas-usually are microadenomas, presents as Cushing’s disease manifestation.
- Thyrotropinomas- presents as secondary hyperthyroidism changes
- Gonadotropinomas- most of time they are asymptomatic, usually are microadenomas. They rarely cause testicular enlargement in males and ovarian hyperstimulation changes in females.

Analysis of Study
1. Types of Pituitary Tumors in our study:

<table>
<thead>
<tr>
<th>Sr.No</th>
<th>Type of Pituitary Tumor</th>
<th>Number of Cases</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Non-Functional Pituitary Macroadenoma</td>
<td>18</td>
</tr>
<tr>
<td>2</td>
<td>Cushing’s Disease (Microadenomas)</td>
<td>3</td>
</tr>
<tr>
<td>3</td>
<td>Rathke’s Cleft Cyst</td>
<td>3</td>
</tr>
<tr>
<td>4</td>
<td>Somatotropinomas (Acromegaly)</td>
<td>5</td>
</tr>
<tr>
<td>5</td>
<td>Giant Pituitary Tumor</td>
<td>2</td>
</tr>
<tr>
<td>6</td>
<td>Invasive Pituitary Carcinoma</td>
<td>1</td>
</tr>
<tr>
<td>7</td>
<td>Recurrent Pituitary Macroadenoma</td>
<td>5</td>
</tr>
<tr>
<td>8</td>
<td>Pure Pituitary Apoplexy</td>
<td>1</td>
</tr>
<tr>
<td>9</td>
<td>Functional Pituitary Prolactinoma</td>
<td>1</td>
</tr>
<tr>
<td>10</td>
<td>Lymphocytic Hypophysitis</td>
<td>1</td>
</tr>
</tbody>
</table>

Surgical Excision of Pituitary Tumors achieved in our study:

<table>
<thead>
<tr>
<th>Sr.No.</th>
<th>Surgical Excision Extent</th>
<th>Number of Cases</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Total</td>
<td>9</td>
</tr>
<tr>
<td>2</td>
<td>Near Total</td>
<td>12</td>
</tr>
<tr>
<td>3</td>
<td>Subtotal</td>
<td>9</td>
</tr>
<tr>
<td>4</td>
<td>Partial</td>
<td>6</td>
</tr>
<tr>
<td>5</td>
<td>Biopsy</td>
<td>3</td>
</tr>
<tr>
<td>6</td>
<td>Marsupialization</td>
<td>1</td>
</tr>
</tbody>
</table>

Pituitary Tumor Consistency

<table>
<thead>
<tr>
<th>Sr.No</th>
<th>Consistency of Tumor</th>
<th>MRI Brain Findings (No.of Cases)</th>
<th>Per-op Findings (No.of Cases)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Soft</td>
<td>15</td>
<td>13</td>
</tr>
<tr>
<td>2</td>
<td>Mixed (Firm+ Soft)</td>
<td>16</td>
<td>17</td>
</tr>
<tr>
<td>3</td>
<td>Firm to Hard</td>
<td>9</td>
<td>10</td>
</tr>
</tbody>
</table>

Statistical Analysis

<table>
<thead>
<tr>
<th>Sr. No.</th>
<th>Parameter</th>
<th>Correlation</th>
<th>Cohen's Kappa</th>
<th>Interpretation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>MRI: Consistency</td>
<td>0.871</td>
<td>0.836</td>
<td>Almost nearly perfect Agreement</td>
</tr>
</tbody>
</table>

Conclusion
1) We have found as per our hypothesis about co-relation of mri brain intensity signals with per-op findings is satisfactorily proved.
2) So around in 33 cases (83-84%) in our study hypothesis has been proved.
3) Only in 7 cases our hypothesis was not proven because of variation in per-op findings in case of tumor consistency.
4) Thus, by knowing pre-operatively we can predict about tumor consistency, its invasive nature & plan for extent of surgical excision.
5) Also on basis of surgical extent of excision we can plan about need of post-op adjuvant therapy.
6) So pre-op mri brain (p+c) with special sequences study is essential for the pituitary tumor management in relation with surgical clearance of disease.

Index Cases
1. Hypothesis proved(Soft Tumor)
I would like to place my thanks on record for everybody, including the staff of CARE Hospitals, who were important for the successful realization of this thesis. I owe my thanks to my family for providing constant support and encouragement. Last but not the least I thank all my patients who in their periods of sorrow and sufferings have made my study possible with their kind cooperation. Collective and individual acknowledgements are also owed to my colleagues in the Department of Neuro Surgery and other Departments, whose presence is perpetually refreshing, helpful and memorable.

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References


