Fine Needle Aspiration Cytology of Pediatric and Adolescent Breast Lesions

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ABSTRACT

Background: The spectrum of breast lesions in pediatric and adolescent age groups varies distinctly from that of adult breast lesions, in that it is rare and overwhelmingly benign. Histopathological examination being the gold standard in managing these cases, FNAC can be used for the accurate diagnosis in most cases. Aim of the study was to assess spectrum of breast lesions in pediatric and adolescent age groups on fine needle aspiration cytology (FNAC) and to correlate with histopathologic findings.

Methods: This is a 12 year retrospective study of cases during 2004 to 2016. 48 patients of age 19 years and below, presenting to our institute with palpable breast lumps were included. FNAC along with histopathologic examination of the excised breast lumps was performed on all the cases.

Result: Majority were females, 46 cases (95.83%). Most common age group was 16 to 18 years and the most common lesion was fibroadenoma. Other lesions were benign phyllodes, tubular adenoma, hamartoma and gynecomastia. On comparing FNAC with histopathology, concordance was seen in 45 cases (93.75%).

Conclusion: In our study all the breast lesions were benign. FNAC and histopathologic examination of these lesions aids in accurate diagnosis and their management.

Keywords: Benign Breast Lesions; Fine Needle Aspiration Cytology; Histopathology; pediatric Breast Lesion.
patients (10.4%) in 12 to 15 years. 13 patients (27%) had multiple breast lumps and 35 patients (73%) had solitary breast lumps. Most common location of breast lump was in upper outer quadrant in 24 patients (35.8%) followed by upper inner in 16 patients (23.8%), lower outer quadrant in 15 (22.3%), lower inner in 7 (10.4%) and retroareolar in 5 cases (7.4%). Most of the breast lumps ranged from 2 to 3 cm in size (47.4%) followed by 4 to 5 cm (18.6%). The largest lump in one case measured >10 cm in size. Two patients (4.16%) had rapidly progressive disease, 38 (79%) had gradually progressive disease, and eight (16.6%) had non-progressive disease. Majority of the patients had no associated symptoms (66.7%). Among symptomatic patients, most common symptom was pain in 10 patients (20.8%) followed by cyclical mastalgia in five (10.5%) and bilateral retraction of nipple in one patient (2%). 93.75% of breast lumps were firm in consistency, 4.25% were soft to cystic and 2% were hard. Two patients had recurrent breast lumps. All the breast lumps were freely mobile except one which showed restricted mobility.

On FNAC, 45 (93.75%) out of 48 cases were diagnosed as fibroadenomas, two (4.16%) as benign breast disease and one (2.08%) as gynecomastia. Among 45 diagnosed as fibroadenoma on FNAC 42 were confirmed on histopathology. Two cases were benign phyllodes tumour and one was tubular adenoma on histopathology. Among 2 cases diagnosed as benign breast disease on FNAC one case was fibroadenoma and one was hamartoma on histopathology. One case reported as gynecomastia on FNAC was confirmed on histopathology (Table 1). On correlating FNAC with biopsy 45 out of 48 cases were concordant (93.75%). Three cases were discordant (6.25%).

FNAC smears from fibroadenoma showed cellular smears with a bimodal pattern containing epithelial and stromal fragments. Large branching sheets of bland epithelial cells with background showing numerous single, bare bipolar nuclei and fragments of fibromyxoid stroma (Fig 1a). Smears from myxoid fibroadenoma showed background containing abundant myxoid material. Excision biopsy from fibroadenomas showed pericanalicular pattern with proliferation of stromal cells around ducts in a circumferential fashion and intracanalicular pattern with compression of the ducts into clefts by the proliferating stromal cells (Fig 1b). Majority of the fibroadenomas diagnosed were of classic type (36 cases). Large myxoid areas were seen in the stroma of myxoid fibroadenoma (one case). One case of cellular fibroadenoma showed highly cellular stroma. One of the fibroadenomas showed large infarcted area on biopsy. One case showed fibroadenoma with marked epitheliosis.

Excision biopsy from tubular adenoma showed well-circumscribed lesion composed entirely of small round tubules, lined by uniform epithelial cells with surrounding myoepithelial cells and separated by sparse intervening stroma (Fig 2a). This lesion was misdiagnosed as fibroadenoma on FNAC in our study. However, on reviewing the slides few three-dimensional epithelial balls in a background of naked nuclei with no stromal fragments were seen (Fig 2b).

FNAC smears from benign phylloids which were misdiagnosed as fibroadenoma were reviewed and showed few large cellular stromal fragments. The single stromal cells in the background had retained cytoplasm and appeared plumper than the typical oval bare nuclei seen in fibroadenoma (Fig 3a). The biopsy from the same showed biphasic tumor with increased stromal cellularity and epithelial hyperplasia. The spindle-cell stromal nuclei were monomorphic showing periductal condensation and mitoses < 5 per 10 high-power fields (Fig 3b).

FNAC from gynecomastia showed sparsely cellular smears with few epithelial cell clusters and small stromal fragments. Histopathology showed ductal epithelial cell hyperplasia with mild stromal edema and fibrosis.

Biopsy from a case of hamartoma showed circumscribed mass with disordered ducts. Few were dilated and showed hyperplasia. Areas of adipose tissue were seen. In this case FNAC was reported as benign breast lesion in view of few benign ductal epithelial cell clusters and adipose tissue fragments.

| Table 1: Spectrum of breast lesions on FNAC and biopsy with correlation. |
|------------------|--------|--------|--------|--------|--------|
| FNAC             | No     | HISTOPATHOLOGY |
|                  |        | Fibroadenoma | Benign Phyllodes | Tubular adenoma | Gynecomastia | Hamartoma |
| Fibroadenoma     | 45     | 42     | 02     | 01     | ---     | ---       |
| Benign breast disease | 02 | 01 | --- | --- | --- | 01 |
| Gynecomastia     | 01     | ---     | ---     | 01     | ---     | ---       |
| TOTAL            | 48     | 43     | 02     | 01     | 01      | 01        |
Fig. 1a: FNAC [Leishmans, 40x] Fibroadenoma - Cellular smear with ductal epithelial cell clusters and background of bare bipolar nuclei and stromal fragments. Fig 1b: Histopathology [H&E, 40x] Fibroadenoma showing compressed ducts surrounded by stroma (Intracanalicular pattern).

Fig. 2a: Histopathology [H&E, 400x] Tubular adenoma - Epithelial cells in small tubules and scant stroma. Fig: 2b FNAC [PAP, 400x] Tubular adenoma showing overlapping epithelial cell clusters with few bare nuclei. No stromal fragments seen.

Fig: 3a: FNAC [H&E, 400x] Benign phyllodes tumour showing cellular stromal fragment Fig: 3b Histopathology [H&E, 100x] Benign phyllodes - Increased stromal cellularity with periductal condensation and epithelial hyperplasia.
Discussion
After onset of puberty, most cases of breast enlargement arise from benign fibroadenoma in girls and Gynecomastia in boys. Other causes of breast masses include infection, trauma, and cyst formation. Hematomas most commonly result from sports or iatrogenic trauma which may present as cystic masses. Fibrocystic changes in the breast occur due to physiologic alterations that are very common in the 3rd decade of life.

In the present study most of the breast lumps (89.5%) were found in age group of 16 to 19 years age which were comparable with other studies like Chung et al (15-17 years), Devi et al (11-20 years) and Kapila et al (16-21 years). The most common lesion in our study was fibroadenoma (87.5%) similar to other studies like Knell et al (93.88%), Singh et al (89.1%) and Kaneda et al (91%).

Majority of the patients in our study had no symptoms. Most common size of fibroadenomas identified ranged between 2 and 3 cm similar to fibroadenomas identified in other studies like Fallat et al and Zang et al. The largest fibroadenoma measured was 10.4cm and was diagnosed as a cellular fibroadenoma. Usually fibroadenomas are found as solitary lesions. But 10 to 20% of conventional fibroadenomas and 25% of juvenile fibroadenomas may present as multiple or bilateral lumps. 27% of fibroadenomas were multiple in our study. Fibroadenomas in adolescents unlike in adults can recur at a site adjacent to surgical excision or in contralateral breast. Recurrence of breast lump was seen in 2 cases of our study.

The second most common lesion in our study was benign phyllodes which presented as rapidly growing breast lump with dilated superficial veins. Similarly kapila et al also showed that benign phyllodes tumour was the 2nd common lesion in this age group.

The diagnostic accuracy of FNAC in our study was 93.75%. Diagnostic accuracy in other studies were Devi et al., 94.7%, Tiwari M et al., 90%, Singh et al., 92.3% and Chandewala et al., 86.55%. On FNAC, both cases of benign phyllodes showed more epithelial components and few cellular stromal fragments and thus were misdiagnosed as fibroadenoma in our study. Such fallacies were also seen in Devi et al and also mentioned in literature.

FNAC is rapid, inexpensive, minimally invasive procedure which helps in accurate preoperative diagnosis which relieves the patient’s anxiety. The definitive treatment can be planned in advance with the informed consent of the patient. Many benign conditions can be confidently diagnosed by FNAC and surgery avoided. The need for frozen section diagnosis is reduced. However, FNAC also has some limitations. A representative sample is required and exact typing of various hyperplastic and low-grade neoplastic lesions may not be possible. When cytology does not correlate with clinical manifestations such as rapid progression or worsening of symptoms a prompt attention by histologic investigation is necessary.

Conclusion
In our study all breast lesions in children and adolescents were benign. Most common lesion was fibroadenoma. FNAC can accurately diagnose these lesions in majority of the cases with breast lumps and is the initial tool for diagnosis. When cytology does not correlate with clinical symptoms histopathologic studies are necessary.

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