Flat Epithelial Atypia Identified on Core Needle Biopsy Does Not Require Excision



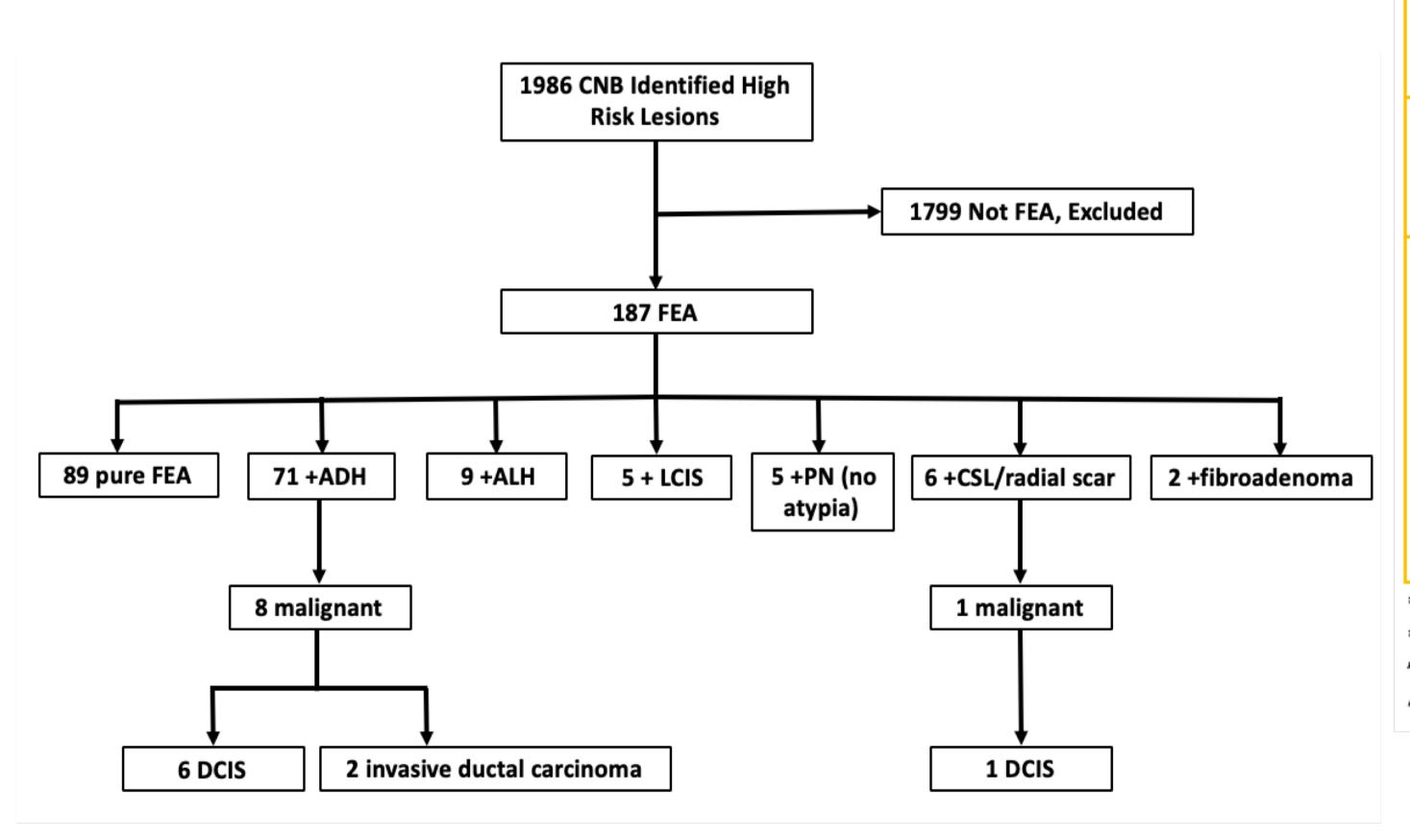
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INTRODUCTION

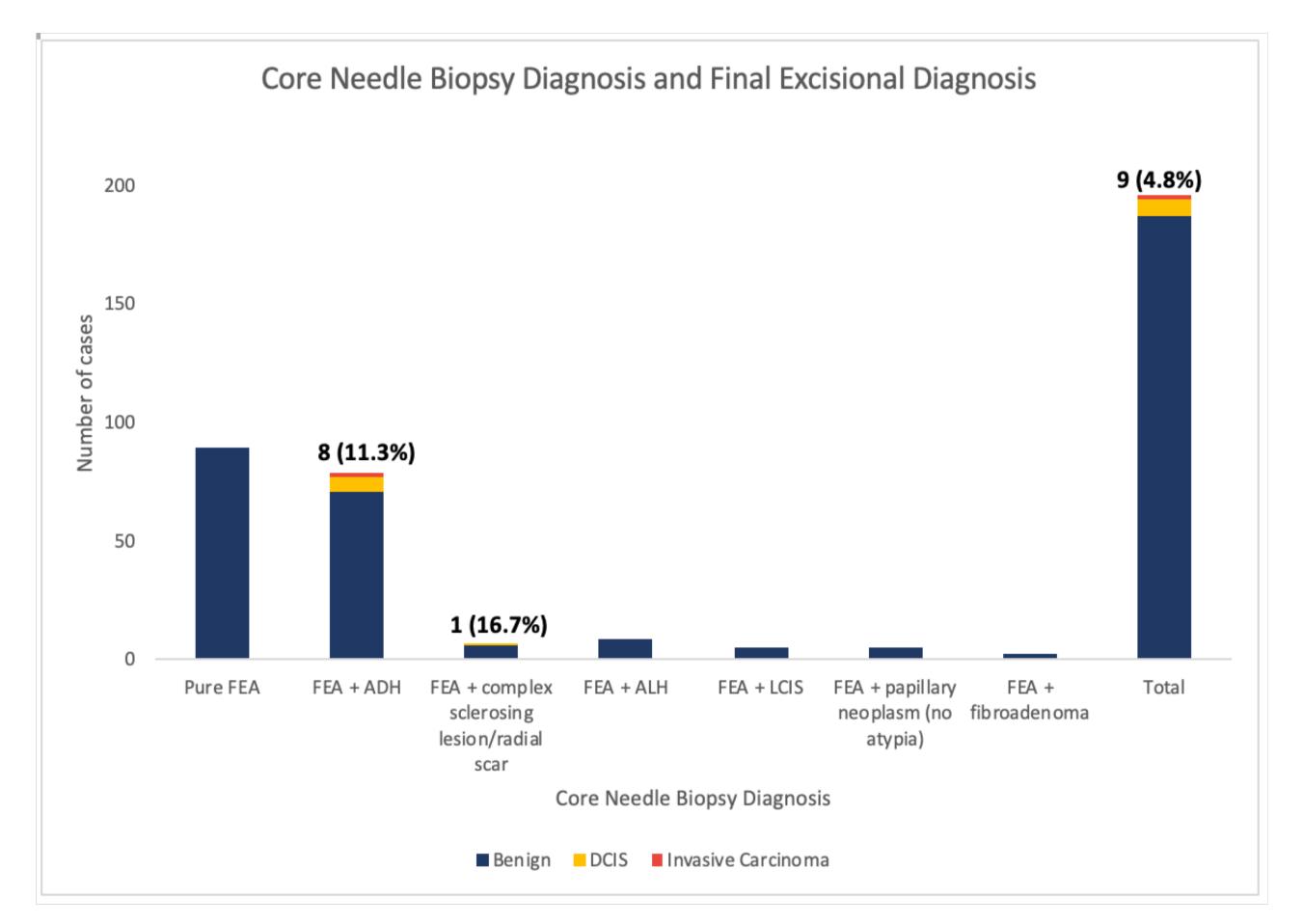
- Flat epithelial atypia (FEA, also known as "DIN1a") diagnosed on core needle biopsy (CNB) have traditionally been excised due to risk of missing a cancer
- In recent years, routine excision of FEA has been called into question
- Current literature reports variable upstage to malignancy rates ranging from 4% to 30%, and most studies report data from single diagnostic centers which limits generalizability of results
- The aim of this study was to evaluate the upstage rates of CNB diagnosed FEA from multiple diagnostic centers across Metro Vancouver, and identify factors predictive of malignancy

METHODS

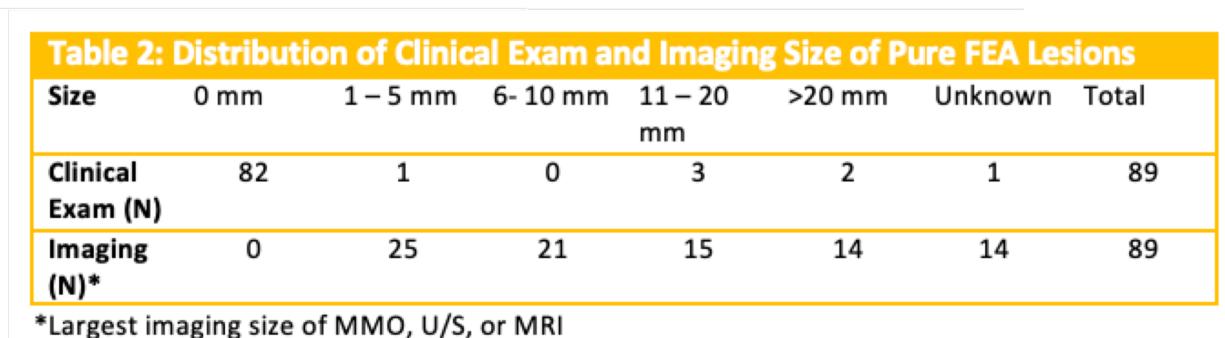
- Patients having excision of FEA at Mount St. Joseph Hospital between
 2013 and 2017 were identified from OR lists
- The primary endpoint was rate of upstage to malignancy
- The association of age, palpability, discharge, clinical exam size, imaging size, family history of breast cancer, type of CNB, and associated histology, with upstage to cancer was evaluated



RESULTS



				Dura FFA Draconting Drablem (0/)
Characteristics			Pure FEA Palpability (%)	Pure FEA Presenting Problem (%)
Age (years)	Mean	52	¹ (1.1) 6 (6.7)	4 (4.5) 1 (1.1)
	Median	50	0 (5.7)	8 (9.0)
	Range (min,	31, 85		
	max)			
Family history	Yes	28 (31.5)		
N (%)	No	52 (58.4)		
	Unknown	9 (10.1)		
Clinical exam	Mean	27		
size (mm)*	Median	20	82 (92.1)	76 (85.4)
	Range (min,	0, 40	02 (32.1)	70 (83.4)
	max)			
Imaging size	Mean	12.6		
(mm)**	Median	8.0	■ Yes ■ No ■ Unknown	■ Mass ■ Imaging abnormality ■ Breast pain ■ Unspecified
	Range (min,	2, 45		
	max)		Pure FEA Diagnostic Imaging Modality	Pure FEA Biopsy Type (%)
Location of	A'	0	1 (1 1)	1 (1.1)
CNB N (%)	B'	4 (4.5)	1 (1.1)	13 (14.6)
	C'	5 (5.6)		
	D'	49 (55.1)		
	E'	16 (18.0)		
	F^	6 (6.7)		
	G^	0		
	H^	0		
	^ O+ (1.1	1 (1.1)		75 (84 3 0)
	Other/Unknown	8 (9.0)		75 (84.3.0)
*of palpable lesions				
**largest imaging size of MMO, U/S, or MRI 74(83.1)				
'within Vancouver			- 11/C - NANAO - NADI	
^adjacent communities		■ U/S ■ MMO ■ MRI	 U/S guided CNB Stereotactic CNB Unknown 	



DISCUSSION

- These results are consistent with literature suggesting low upstaging of pure FEA lesions¹⁻⁶
 - Becker et al (2013) reported 4.2% upstage rate and follow up of nonexcised lesions showed no suspicious findings
 - Chan et al (2018) reported 0% upstage rate and noted presence of ADH was the only predictor of upstaging
- Solorzano et al (2011) reported a higher upstage rate of 14% and concluded that mammographic and sonographic presentation of FEA is not specific and recommend surgical excision
- The presence of ADH or CSL in the biopsy were the only predictors of histological upstage to malignancy (p=0.001, p=0.0001)
 - The two invasive cancers were found in lesions associated with ADH, which is a lesion previously studied at our center and found to be highrisk for upstaging. Current literature also recommends excision of most cases of ADH for this reason
 - CSL is still a controversial lesion, with variable reported upstage rates. An on-going study at our center is evaluating upstage rates in our region to better inform management
- ASBS endorses observation with clinical and imaging follow up for pure FEA lesions and excision if concurrent ADH
- ASBS recommends surgical excision of most CSL
- Majority of published upstage rates for FEA are single-institutional studies limiting generalizability of results vs. this study represents a population-based sample from across our region

CONCLUSION

- The upstage rate to malignancy after excision of CNB diagnosed pure FEA at our regional center is 0%
- Therefore, we recommend that pure FEA with radiology and pathology concordance does not require surgical excision, and can instead be followed with serial imaging
- Patients with FEA in association with other high-risk lesions should be managed as per indicated for the other high-risk lesion due to the variable associated upstage rates
- We specifically recommend the excision of FEA lesions found in association with ADH due to the higher rates of upstaging

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