

Original Research Article

EVALUATE THE ADC VALUES IN PROBABLY BENIGN AND SUSPICIOUS MALIGNANT BREAST LESIONS

ABSTRACT

Aim: To characterize probably benign and suspicious breast lesions with non invasive MRI techniques of diffusion weighted imaging using ADC values and to correlate the values of ADC with histopathological findings of breast lesions.

Study Design: Observational study.

Place and Duration of Study: The study was conducted in Department of Radiology of Himalayan Institute of Medical Sciences, SRH University, Dehradun from September 2016 to June 2018.

Methods: In this observational study, 54 patients were included with diagnosis of BIRADS III and BIRADS IV on mammography and sonomammography. The DWI MRI was done and ADC values were calculated and results were correlated with histopathological outcome.

Results: Comparison between the DWI analysis and histopathological findings reveals that the majority of the lesions 58.7% with ADC value $\leq 1.03 \times 10^{-3} \text{ mm}^2/\text{s}$ were for ($P < .005$). DWI analysis showed a sensitivity of 73.68%, a specificity of 88.88%, a PPV of 83.25%, an NPV of 82.75% and an accuracy of 82.60%.

Conclusion: DWI MRI is a non invasive technique used to discriminate between benign and malignant lesions and helps in reducing unnecessary interventions.

Keywords: ADC value, BIRADS, DWI,

INTRODUCTION

Breast is a modified sweat gland, comprising of fibrous, fatty and glandular tissue. It can be a site for various lesions ranging from mastitis to invasive carcinoma, over a wide range of age. It becomes essential to differentiate between inflammatory and benign lesions from early carcinoma, especially in women predisposed to carcinoma of breast.

42 One of the leading causes of cancer death in women is breast carcinoma (1). It has been ranked
43 number one cancer in Indian females with age adjusted rate of 25.8 per 100,000 with mortality
44 rate of 12.7 /100,000 women(2). The increasing rate of carcinoma breast is an alarming area in
45 perturbation to the clinicians and researchers (3). Breast imaging has proven to detect breast
46 cancer in its early stage. However, in females under 40 years of age with dense breast, other
47 technologies pertaining to early detection such as sonomammography and MRI breast may also
48 contribute to the early detection of breast cancer, for whom the X-ray mammography is less
49 sensitive (4). Magnetic resonance technique have shown great potential to ameliorate the
50 sensitivity and specificity in diagnosing breast malignancy. Dynamic contrast enhance (DCE) MRI
51 is a crucial imaging tool in diagnosis and management of breast masses. It gives precise
52 information about the extent of the lesion and elaborate information about the multifocal or
53 multicentric disease which influences the treatment decisions (5). MRI was established as an
54 imaging technique in medicine over 20 years but only in the last few years it is being used
55 consistently to image the breast (6). Using routine MRI sequences there is difficulty in
56 ascertaining the benign from malignant lesions, as these two categories may share certain
57 morphology and contrast enhancement characteristics. In the era of fast improving technology
58 the MRI techniques have also sequences with excellent spatial resolution and soft tissue contrast
59 which contribute in differentiating the nature of the masses. Diffusion weighted MRI (DWI)
60 imaging might be of value in assessment as it has the ability to provide tissue contrast based on
61 molecular diffusion (7). Diffusion weighted MRI is highly sensitive for breast malignancy allowing
62 its detection that is occult on physical examination, X-ray mammography and sonomammography
63 (8). DWI can easily be embraced as an adjunction for standard clinical imaging protocols and has
64 been reported to achieve higher pick-up rates than X-ray mammography.
65 Breast MRI with special sequences may be used to discriminate benign and malignant lesions
66 which may minimize the number of breast biopsy performed in probably benign lesions (9). The
67 patient is always concerned with such lesions.
68 DW-MRI generates images that are sensitive to water displacement at the diffusion scale and
69 quantifies such diffusion according to a quantitative index reflecting the apparent freedom of
70 diffusion (apparent diffusion coefficient (ADC) (10).This sequences appears to be an effective
71 tool for tumour detection and characterization as well as for monitoring and speculating treatment
72 response (11). DWI is a non-contrast sequence that has shown potential for discriminating the
73 nature of breast lesions. In our study we will be using this single MRI sequence in the probably
74 benign and suspicious breast masses on routine investigations and validate its usefulness in
75 terms of its non invasiveness in discriminating the nature of the breast lesions.

81 **MATERIALS AND METHODS**

82 The study was conducted in the Department of Radiology, Himalayan Institute of Medical
83 Sciences (HIMS), Swami Ram Nagar, Dehradun from September 2016 to June 2018. Patients
84 who were diagnosed to be having breast masses were recruited from department of Surgery
85 (cancer centre), Himalayan institute of medical sciences, Dehradun after taking a proper written
86 informed consent and clearance from ethical committee. The study included 54 patients. The
87 inclusion criteria was female patients above 30years and who were diagnosed with BIRADS III
88 and BIRADS IV on mammography and sonomammography. Exclusion criteria was patients with
89 ferromagnetic implants and pacemaker and all post operative patients who underwent surgery for
90 breast mass .

91 The study tools included :

- 92 1. Conventional mammography, both cranio caudal and oblique views of bilateral breast, was
- 93 performed on SIEMENS 3000 NOVA mammography machine.
- 94 2.Sonomammography was done on Machine Philips EPIQ 7G with high frequency (5-18 MHz)
- 95 Linear transducer.
- 96 3. Magnetic resonance imaging (DWI) of both breasts was performed on Machine 1.5 Tesla MR
- 97 Unit: AVANTO, SIEMENS (Germany) using dedicated breast coil for optimal signal acquisition.
- 98 4. FNAC / Biopsy reports were analyzed.

99
100 Study protocol included:

- 101 1. Informed consents was taken
- 102 2. Conventional mammographic examination was done (mediolateral oblique and craniocaudal
- 103 views were obtained).
- 104 3. Sonomammography was done.
- 105 4. On the basis of combined mammography and sonomammography lesions were assessed and
- 106 higher category was assigned using fifth edition of the American College of Radiology (ACR)
- 107 Breast Imaging Reporting and Data System (BI-RADS) lexicon.
- 108 5.Further diffusion weighted MR images were obtained and ADC values were calculated by
- 109 manually placing the ROI within lesion on the ADC map and record the mean value in that ROI.
- 110 6. FNAC / Biopsy reports were analyzed.

111
112 The outcome on histopathology was considered as final diagnosis and compared with

113 DWI ADC findings. All data was analyzed with SPSS software version 22.0. The data was

114 presented as mean±SD for continuous variables and as frequency or percentage for categorical

115 variables. Categorical data has been represented as frequency (number) and proportions

116 (percentages). Continuous data has been presented as mean ± standard deviation (SD). The chi-

117 square test and student's test were used for statistical comparison of qualitative and quantitative

118 variables. *P* values <.005 was considered statistically significant.

119
120
121

122 **RESULTS AND DISCUSSION**

123
124 One of the major malignant killer in women is the carcinoma breast. The basic modalities

125 for the detection of the breast lesions are mammography, sonomammography and the breast

126 MRI. Each of these modalities have their strengths and weakness .The sensitivity and specificity

127 of picking the breast lesions alone by the single modality is less. However when used in

128 combination increases the detection rate.

129 There have been improvement in the detection of breast cancer with wide spread

130 application of mammography and ultrasound. However still it remains difficult to diagnose and

131 characterize the lesion specially in dense fibroglandular breast. The limitation of the

132 mammography is the overlapping of tissue which hides the lesions mainly in dense breast.

133 However the sensitivity of picking microcalcification, the early sign of malignancy is markedly

134 reduced in ultrasound. The strength of the ultrasound lies in characterization of the solid or cystic

135 masses. With the advancement of the MRI Breast specially the DWI sequence which do not

136 require IV contrast, is an important tool in differentiating the benign and malignant lesions as

137 proved by the various studies.

138 In the present study we included 54 patients with 56 breast lesions. The lesions were
139 clinically palpable and film screen mammography and ultrasound were done. The lesions were
140 categorized on basis of BIRADS classification (ACR V edition). A combination of mammographic
141 and sonomammographic BIRADS category III, IVA, IVB, IVC were included in our study and
142 higher category was assigned. DWI was done for these lesions and ADC values were calculated.

143 In our study all the patients were female with the mean age group of (48.81± 9.53). The
144 majority 46.3% of patients evaluated were between 41-50 years, 22.2% in 51-60 years, 20.4% in
145 31-40 years and 11.1% in 61-70 years of age group. Fernanda Philadelpho and Arantes and
146 Pereira et al conducted a study which also showed that mean age group of female patients were
147 46.1 (12).

148 Analysis of data from more than 150,000 women who participated in 54 epidemiological
149 studies (National cancer institute in United states) showed that overall women who had ever used
150 oral contraceptive had a 7% increase in the relative risk of breast cancer as compared with
151 women who had never used oral contraceptives (13). In the present study 68.5% of patients had
152 history of oral contraception, there was increase in the percentage of carcinoma in females who
153 had history of oral contraception (57.1%).

154 It is a well known fact that the carcinomas present with breast pain in the later stages
155 (14). The same was not found true in our study where 81.48% patients had no pain while 18.52%
156 had pain. As we have included BIRADS category III and BIRADS category IV and there were no
157 advanced cases.

158 Most of the breast cancers are unilateral and are found in upper outer quadrant. The
159 favored site because of increase fibroglandular tissue in this quadrant. Siwa Chan and Jeon-Hor
160 Chen et al in their study also reported that upper outer quadrant is the most favored site (15). Our
161 study also favored this fact as 96.3% of lesions were unilateral and 3.7% were bilateral, 48.21%
162 were present in upper outer quadrant, followed by upper inner quadrant (26.79%), lower inner
163 (10.71%) ,lower outer quadrant (7.14%), retroareolar region (3.57%) and (3.57%) in upper inner
164 and outer quadrant.

165
166
167 Mostly the benign lesions are well defined on film screen mammography with a peripheral halo
168 while the malignant lesions have irregular margins as stated by Haixia Li and XianjingMeng et al
169 in their study (16). In our study most of the lesions showed indistinct margins (73.21%) followed
170 by circumscribed margins (26.79.%). Majority are indistinct as (57.1%) cases histologically
171 malignant.

172 The clinically palpable masses may be seen as mass or asymmetry. In our study 48
173 mammograms showed masses while 7 mammograms showed asymmetry. This asymmetry was
174 further seen as mass lesions on sonomammography, thus favoring the fact that combined
175 imaging increases the detection rate.

176 In the malignancy the cells are compactly packed than in the benign lesions thus casting
177 high density. In our study the mammogram showed increased density in 98.21% lesions. It is
178 because our study comprise of lesions mainly in the BIRADS category IV.

179 The malignant calcifications are the hallmark of malignancy on the lesions as stated by
180 Yojana V Nalawade in his study (17). In our study 8.93% had suspicious calcification while 3.57%
181 had benign calcification. The pick up rate of calcification was less because the study was
182 conducted using film screen mammography which is less sensitive than digital mammography.

183 Architectural distortion may be seen in the malignant and the inflammatory lesions, we

184 encountered 1.8% cases showing architectural distortion. This could be because of the film
185 screen mammography used for imaging.

186 Ultrasound plays an important role in further characterization of the mammographic
187 masses. It acts as an adjuvant and increases the confidence rate of reporting. The malignant
188 lesions are usually taller than wider and the benign are wider than taller. Sudheer Ghokhale also
189 stated the same fact in his study (18). In our study it was observed that 32 (57.15%) had oval
190 shape, followed by irregular 18(32.14%) and round in 6(10.71%). Since we had not included
191 BIRADS category V, so most of the lesions maintained their shape.

192 Ultrasound has a strength to discriminate cystic, solid and mixed echotexture masses.
193 Most of the lesions in our study were hypoechoic (83.9%), (12.5%) mixed and (3.6%) isoechoic
194 pattern. The purely cystic lesions were not included in our study.

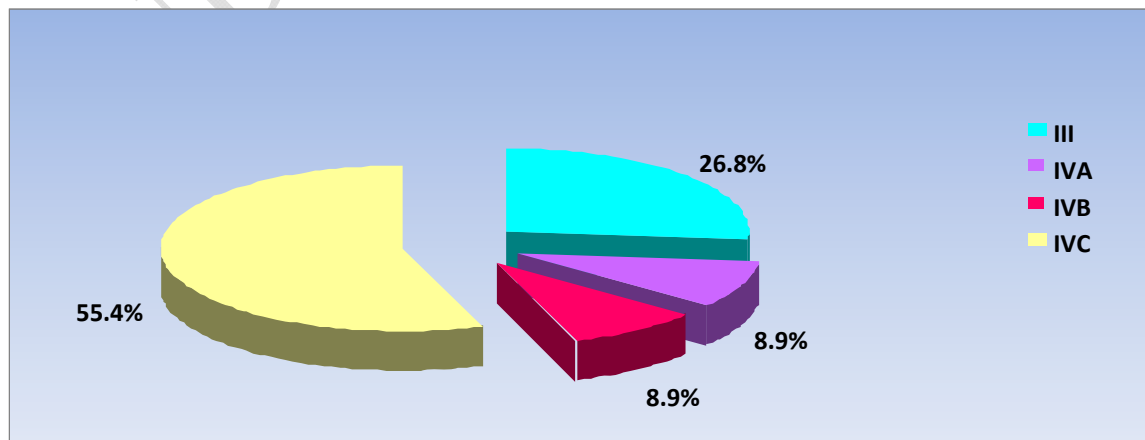
195 The margins are better appreciated on Ultrasound than the mammography, which further
196 helps in characterization of the masses. In our study it was observed that 44.64% of the lesions
197 had indistinct margins, 39.3% circumcised margins, 5.4% indistinct with spiculated margins, 5.4%
198 microlobulated margins ,3.6% angular and 1.8% had indistinct and angulated margins. Most of
199 these margins suggested malignancy. It is in concordance with the findings as majority of the
200 study cases (57.1%) are malignant.

201 Ultrasound is a good modality to evaluate the infiltration in the surrounding tissue around
202 the mass. This is helpful to label the mass as malignant, however one has to be cautious in
203 differentiating from inflammation. We observed that adjacent Parenchyma was hyperechoic in
204 58.93% and normal in 41.07% as our lesions spectrum mainly included BIRADS category IV
205 masses.

206 Evaluation of the skin over the breast mass is important in characterizing the masses. The
207 pure benign masses do not produce any change in the skin, however usually the advanced
208 malignant and inflammatory masses do so. We found in our study on the basis of combined
209 mammography and sonomammography the overlying skin was seen normal in (89.3%) and
210 affected in (10.7%). This was because the masses included in the study are BIRADS III and IV. It
211 was found that nipple was also retracted in (10.7%) because of the same reason.

212
213 All the lesions were categorized on the combined mammography and sonomammography
214 findings and the higher category was awarded. Of the BIRADS IV category lesions, 55.4% of the
215 patients had Category IVC, 8.9% category IVA and 8.9% category IVB. While 26.8% had BIRADS
216 Category III lesions (Fig 1).

217
218

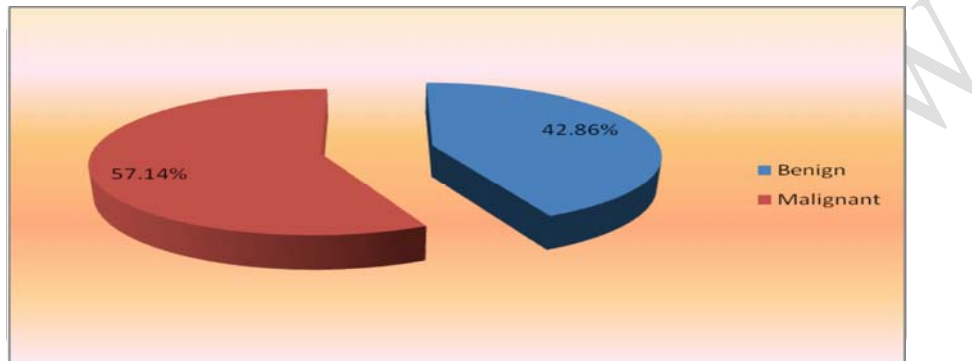


219
220

221
222
223
224
225
226
227
228
229

Fig.1. Showing BIRADS category of the lesions on combined mammography and sonomammography

As the histopathology was the gold standard investigation in our study. It was found that on the basis of histopathology 57.14% of the lesions were malignant and 42.86% were benign (Fig 2).



230
231
232
233
234
235
236
237
238
239
240
241
242
243
244

Fig.2. Showing histopathological diagnosis of lesions.

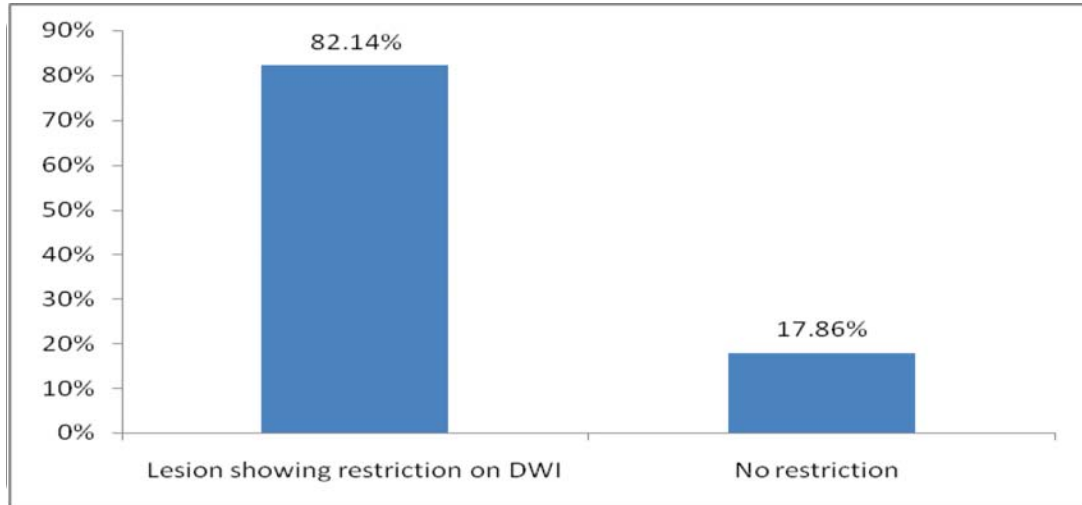
A study conducted by I Trop and Lalonde et al, in 2009 , concluded that the sensitivity and specificity of CBE alone was 17% and 95.9%, that of mammography was 58% and 95.4%, and that of ultrasonography was 42% and 93.8%. Combined sensitivity and specificity of CBE, mammography and US was 67% and 90.3% (19).

In our study we included the clinical breast examination, mammography and sonomammography to increase the sensitivity and specificity of the lesions.

245
246
247
248
249
250
251
252
253
254
255
256
257
258

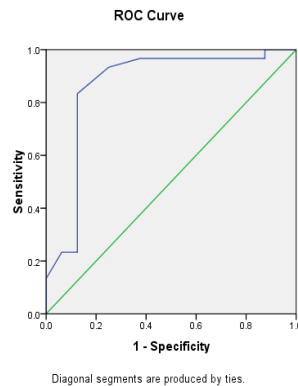
Diffusion weighted MRI was the main objective of study to evaluate the ADC values of breast masses. DWI is a technique where no IV contrast is used and in the various studies conducted by Fernanada philadelpho and Arantes Pereira et al in 2007 (12), Richa Bansal and Viral Shah et al in 2013 (20) , Wasan Ismail AL Saadi et al in 2014 (21), Hongmin Cai and Lizhi Liu et al (22) and Uma Sharma and Rani G. Sah et al (23) showed the efficacy of DWI in characterizing the benign or malignant lesion. In our study, DWI showed restricted diffusion in (81.6%) of the 56 lesions and 10(17.86%) showed no restriction (Fig 3). Majority of the masses showing restriction were the solid masses. The ADC value was calculated and by using the ROC curve, the cut off value came out to be $1.03 \times 10^{-3} \text{ mm}^2/\text{s}$ (Fig 4), so all the lesions in our study having the ADC more than this were histologically proven to be benign that helps in discriminating benign from malignant lesions. In our study comparison between the DWI analysis and histopathological findings reveals that the majority of the lesions (58.7%) with $\text{ADC value} \leq 1.03 \times 10^{-3} \text{ mm}^2/\text{s}$ were found to be malignant and 41.3% with $\text{ADC value} > 1.03 \times 10^{-3} \text{ mm}^2/\text{s}$ were found to be benign ($P < .005$) (Table 1). The ADC values of malignant lesions were lower that

259 ranges from 0.6 to 1.0 x 10⁻³ mm²/s and the ADC value of benign lesions were higher that
260 ranges from 1.1 to 2 x10⁻³ mm²/s. In our study for the distinction between benign and malignant
261 lesions, DWI analysis showed a sensitivity of 73.68%, a specificity of 88.88%, a PPV of 83.25%,
262 an NPV of 82.75% and an accuracy of 82.60% (Fig 5).
263



264
265
266
267
268
269
270
271

Fig.3. Showing distribution of the patients that showed restricted diffusion on DW MRI



272
273
274
275
276
277
278
279
280
281

Fig.4. Receiver Operating Curve showing the cut off value of ADC

Table 1 Shows correlation between the ADC values and histopathology results.

282

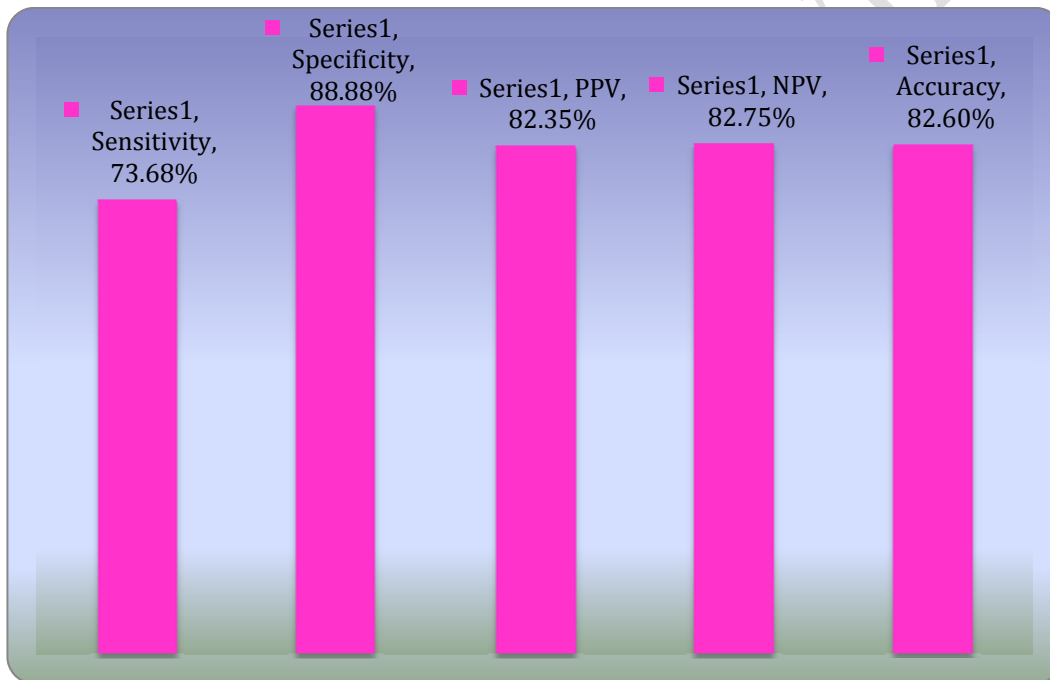
		ADC		Total	P value
		Benign	Malignant		
HISTO	Benign	14	3	17	0.005
		82.4%	17.6%	100.0%	
	Malignant	5	24	29	
		17.2%	82.8%	100.0%	
Total		19	27	46	
		41.3%	58.7%	100.0%	

283

284

285

286



287

288

289

290

291

292

Fig.5. Correlation between the ADC values and histopathology results.

293

294

295

296

297

The cut off ADC value was taken as $1.03 \times 10^{-3} \text{ mm}^2/\text{s}$. 27(48.21%) showed ADC values $\leq 1.03 \times 10^{-3} \text{ mm}^2/\text{s}$ and 19(33.39%) showed ADC value $> 1.03 \times 10^{-3} \text{ mm}^2/\text{s}$ and 10(17.86%) showed no restricted diffusion (Fig 6).

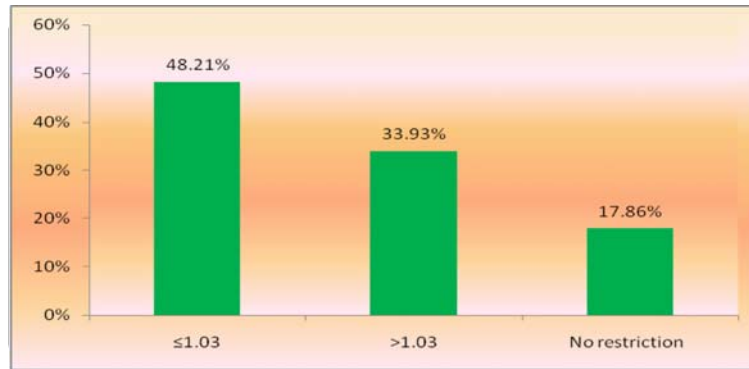


Fig .6. Shows cut off value of ADC in various lesions

298
 299
 300
 301
 302
 303
 304
 305
 306
 307
 308
 309
 310
 311
 312
 313
 314
 315
 316
 317
 318
 319
 320
 321
 322
 323
 324
 325
 326
 327
 328
 329
 330

The considerable variation was explained by the different protocols used in the studies. The cut off ADC values obtained in the differentiation between benign and malignant lesions were dependent upon the respective b value chosen. In our study we use b value of 800s/mm², in terms of the ADC values , cut off value , sensitivity and specificity, were in agreement with those found in literature.

Despite the promising capacity of ADC values to differentiate between benign and malignant lesions, the ADC values for benign and malignant lesions can overlap leading to false positive and false negative results. In our study false negative cases i.e 5 out of 32 lesions showed the ADC values >1.03x10⁻³mm²/s came out to be malignant on histopathology and all were ductal carcinomas and there was only 1 of 32 lesion that shows no restriction but diagnosed as ductal carcinoma on histopathology. 2 out of 24 benign lesions show ADC <1.03x10⁻³mm²/s, however came out to be chronic abscess on histopathology .

The results of the present study should be considered in the context of certain limitations. Firstly our patient population comprised of individuals referred mainly from our surgery department (cancer centre) in the institute, featured a predominance of malignant pathological findings. Secondly, patients with suspected benign lesion usually undergo US and mammography was not done, thereby limiting the cases.

The single sequence of DWI is a non invasive technique and has high sensitivity and specificity and is a great tool that helps us in discriminating benign from malignant breast lesions and can reduce the intervention.

331 CONCLUSION

332 In present day scenario breast cancer is the most common cause of cancer related death in
 333 females. Early detection of malignancy is essential to decrease the morbidity and mortality.
 334 Various imaging modalities are used to detect breast lesions, which includes Mammography,

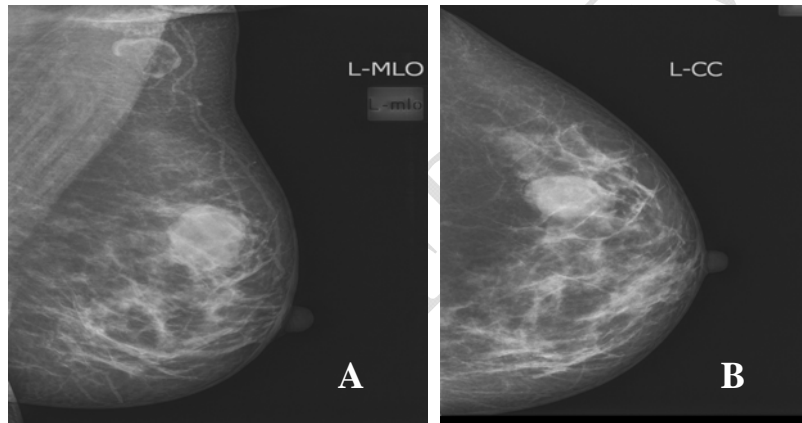
335 sonomammography and breast MRI. However mammography is the basic modality for screening
336 and ultrasound (US) is an adjuvant to it. These modalities are known to have high false positive
337 rates because of their own limitations. DWI MRI is a technique based on diffusivity of water
338 molecules and is quantified by ADC value. High cell proliferation in malignant tumors increases
339 cellular density, creating more barriers to the extracellular water diffusion, reducing the ADC, and
340 resulting in signal loss and vice versa occurs in benign lesions and showed high value. This
341 parameter is used in our study to discriminate between benign and malignant lesions and helps in
342 reducing unnecessary interventions.

343
344
345
346

347 Appendix A. Cases

348 A.1. Case (1)

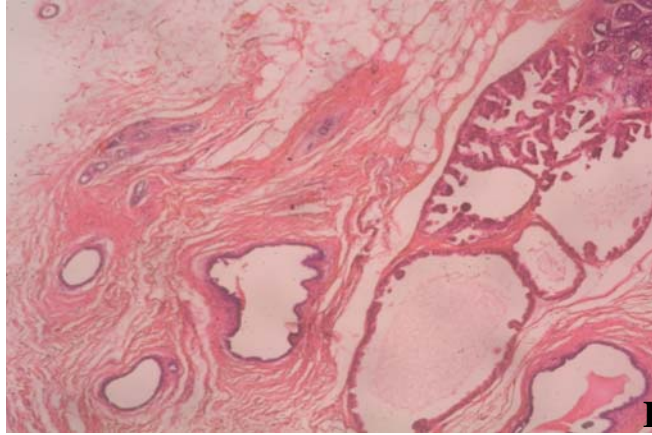
349



350



351



352

353

354

355

356

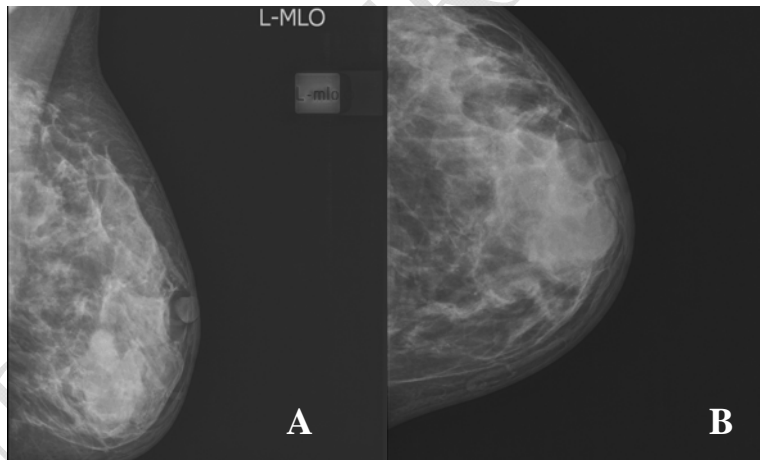
357

358

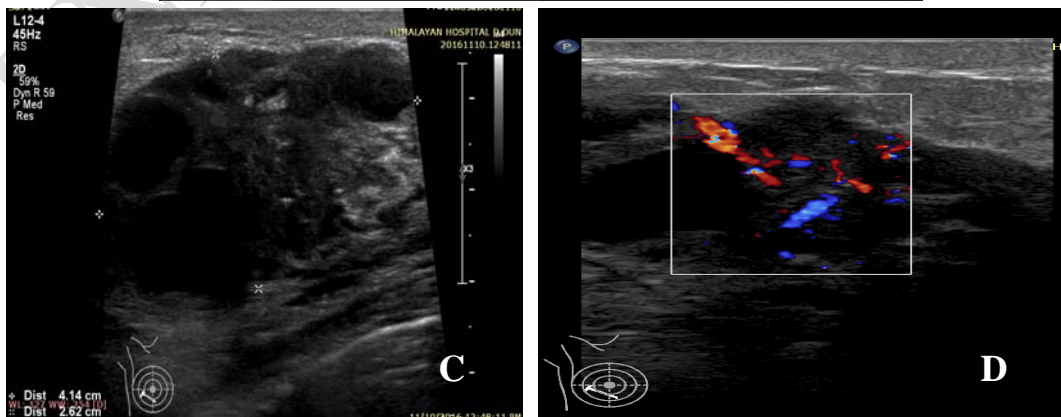
46 years old female with lump left breast. Mammography, (A)MLO and (B)CC show soft tissue density mass in upper outer quadrant with smooth margins. (C)Ultrasound shows hypoechoic mass with posterior wall enhancement in upper outer quadrant, categorised as BIRADS category III on combined mammography and sonomammography. (D) DWI MRI with ADC mapping at $b=800$ and ADC value of 1.6×10^{-3} . (E) Histopathology H and E section reveals fibroadenoma(10X).

359

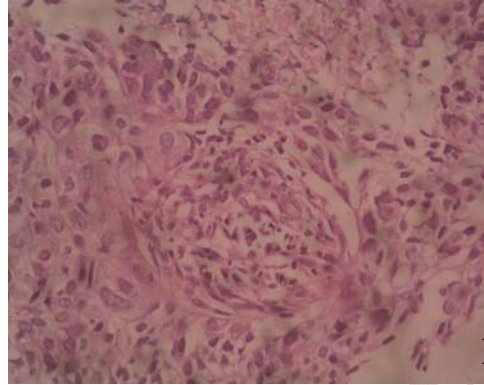
CASE 2



360



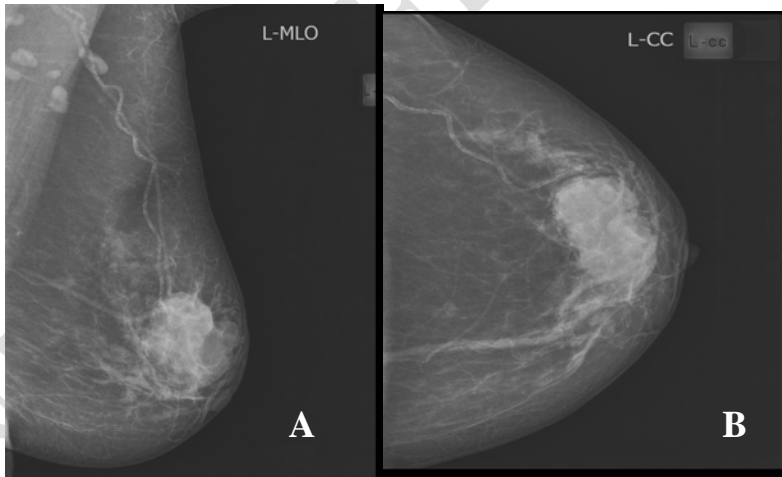
361



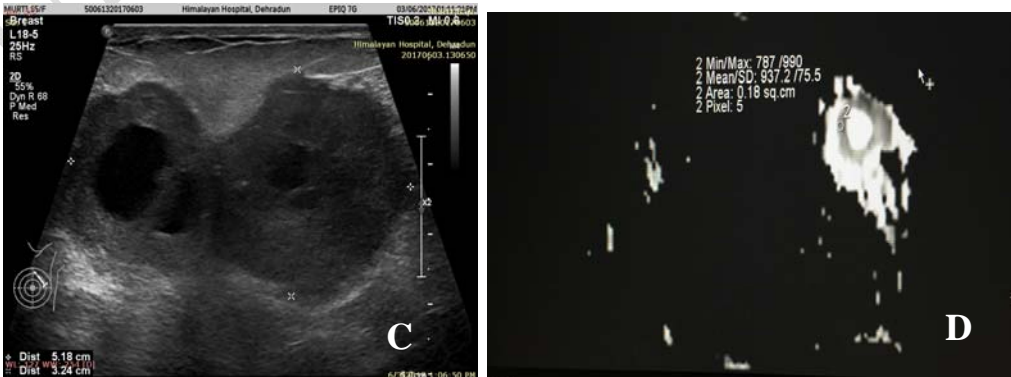
362
363
364
365
366
367
368
369
370
371
372
373
374

35 years old female with lump left breast breast. Mammography, (A)MLO and (B)CC show soft tissue density mass in lower inner quadrant with ill defined margins. (C)and (D)Ultrasound shows complex mass with solid and cystic areas, solid component shows vascularity on color doppler and calcification, categorised as BIRADS category IVC on combined mammography and sonomammography. (E) DWI MRI with ADC mapping at b= 800 and ADC value of 0.8×10^{-3} . (F) Histopathology H and E section reveals infiltrating ductal carcinoma (40 X).

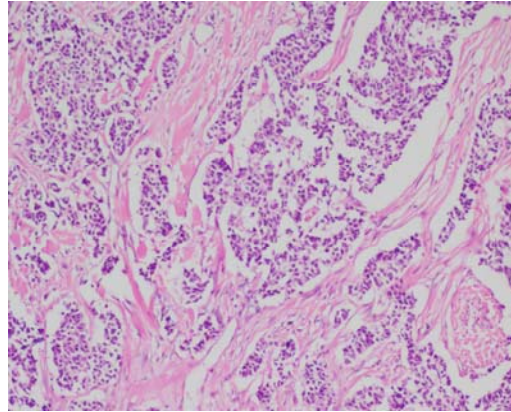
CASE 3



375



376



E

377
378
379
380
381
382
383
384

65 years old female with painless lump left breast. Mammography, (A)MLO and (B)CC show soft tissue density mass in upper outer quadrant with irregular margins. (C)Ultrasound shows hypoechoic mass with anechoic areas within and smooth lobulated margins in upper outer quadrant, categorised as BIRADS category IVC on combined mammography and sonomammography. (D) DWI MRI with ADC mapping at $b=800$ and ADC value of 0.9×10^{-3} . (E) Histopathology H and E section reveals infiltrating ductal carcinoma (40X).

385

386

COMPETING INTERESTS

387
388
389
390
391
392

We have no conflict of interest with anybody working in the area.

393

CONSENT

394
395
396
397
398

"All the authors declare that written informed consent was obtained from the patient for publication of this paper and accompanying images. A copy of the written consent is available for review by the Editorial office/Chief Editor/Editorial Board members of this journal."

399

ETHICAL APPROVAL

400
401
402
403
404

"All authors hereby declare that all experiments have been examined and approved by the appropriate ethics committee and have therefore been performed in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki."

405

406

407

408

409

410
411

REFERENCES

- 412 1. What are the key statistics about breast cancer? American Cancer Society.
413 [http://www.cancer.org/cancer/breastcancer/detailedguide/breast-cancer-key-](http://www.cancer.org/cancer/breastcancer/detailedguide/breast-cancer-key-statistics)
414 [statistics](http://www.cancer.org/cancer/breastcancer/detailedguide/breast-cancer-key-statistics). Last revised December 31, 2013. Accessed March 20, 2013.
- 415 2. Babu GR, Lakshmi SB, Thiyagarajan JA. Epidemiological correlates of breast cancer
416 in South India. *Asian Pac J Cancer Prev* 2013; 14: 5077–83.
- 417 3. Guo A. Role of diffusion-weighted (DWI) in magnetic resonance (MR) of the breast. *J*
418 *MagnReson Imaging* 2002;16:172–3.
- 419 4. Saslow D, Boetes C, Burke W. American cancer society guidelines for breast
420 screening with MRI as an adjunct to mammography. *Am Cancer Soc*
421 2007;57(3):185.
- 422 5. Barker P, Salkowski L. Diffusion-weighted imaging may improve accuracy of breast
423 MRI, American Roentgen Ray Society (ARRS) annual meeting. Abstract
424 2009;129:35–43.
- 425 6. Guo A. Role of diffusion-weighted (DWI) in magnetic resonance (MR) of the breast. *J*
426 *MagnReson Imaging* 2002;16:172–3.
- 427 7. Le Bihan D, Breton E, Lallemand D, Aubin ML, Vignaud J, Laval- Jeantet M.
428 Separation of diffusion and perfusion in intravoxel incoherent motion MR imaging.
429 *Radiology*. 1988;168(2):497–505.
- 430 8. Wang LC, DeMartini WB, Partridge SC. MRI-detected suspi- cious breast lesions:
431 predictive values of kinetic features measured by computer-aided evaluation. *Am J*
432 *Roentgenol* 2009;193(9):826–31.
- 433 9. Wax A. Breast cancer and MRI, web MD, breast cancer guide. *Am SocClinOncol*2009.
- 434 10. Marini C, Iaconi C, Giannelli M, Cilotti A, Moretti M, Bartolozzi C: Quantitative
435 diffusion-weighted MR imaging in the differential diagnosis of breast lesion.
436 *EurRadiol* 2007, 17(10):2646–55.
- 437 11. Koh DM, Collins DJ. Diffusion-weighted MRI in the body: applications and challenges
438 in oncology. *AJR* 2007; 188:1622–35.
- 439 12. Fernanda Philadelpho, Arantes Pereira. Assessment of Breast Lesions With
440 Diffusion-Weighted MRI: Comparing the Use of Different Values. *AJR* 2009;
441 193:1030–35.
- 442 13. Marchbanks PA, McDonald JA, Wilson HG, Folger SG, Mandel MG, Daling JR,
443 Bernstein L, Malone KE, Ursin G, Strom BL, Norman SA. Oral contraceptives and
444 the risk of breast cancer. *New England Journal of Medicine*. 2002;346(26):2025-32.
- 445 14. Minjoung Monica Koo and Christian von Wagner et al. Typical and atypical
446 presenting symptoms of breast cancer and their associations with diagnostic
447 intervals: Evidence from a national audit of cancer diagnosis. *Cancer Epidemiology*.

- 448 2017;48:140–6.
- 449 15. Siwa Chan, Jeon-Hor Chen. Evaluation of the association between quantitative
450 mammographic density and breast cancer occurred in different quadrants. BMC
451 Cancer. 2017;17:274.
- 452 16. Haixia L, Xianjing M. Breast masses in mammography classification with local
453 contour features. Bio Med EngOnLine. 2017;16:44-52.
- 454 17. Nalawade YV. Evaluation of breast calcifications. The Indian Journal of Radiology &
455 Imaging. 2009;19(4):282-6.
- 456 18. Gokhale S. Ultrasound characterization of breast masses. The Indian Journal of
457 Radiology & Imaging. 2009;19(3):242-7.
- 458 19. Trop I, Lalonde L. Multimodality breast cancer screening in women with a familial or
459 genetic predisposition. Current Oncology. 2010; 17, (3): 28-36.
- 460 20. Bansal R, Shah V, Aggarwal B. Qualitative and quantitative diffusion-weighted
461 imaging of the breast at 3T - A useful adjunct to contrast-enhanced MRI in
462 characterization of breast lesions. Indian J Radiol Imaging. 2015;25(4):397-403.
- 463 21. Wasan Ismail Al-Saadi, EhabNaeemShallab. Diffusion weighted MRI in the
464 characterization of solitary breast mass. The Egyptian Journal of Radiology and
465 Nuclear Medicine. 2015;46:1337–41.
- 466 22. Hongmin C, Lizhi L. Diagnostic assessment by dynamic contrast-enhanced and
467 diffusion-weighted magnetic resonance in differentiation of breast lesions under
468 different imaging protocols. BMC Cancer. 2014, 14:366-72.
- 469 23. Sharma U, Sah RG. Potential of Diffusion-Weighted imaging in the characterization
470 of Malignant, Benign, and healthy Breast Tissues and Molecular subtypes of Breast
471 cancer. Front. Oncology. 2016;6:126-33.

472

473

474

475

476

477

478

479

480

481