Usefulness of Diffusion Weighted Imaging and Apparent Diffusion Coefficient in Characterizing Pancreatitis and Pancreatic Adenocarcinoma

Prabhu Radhan\textsuperscript{1}, Anupama Chandrasekharan\textsuperscript{2}, Rajoo Ramachandran\textsuperscript{3}, Remya Baburaj\textsuperscript{4}, Poovini Soundararajan\textsuperscript{5}

\textsuperscript{1}Department of Radio Diagnosis, Sri Ramachandra Institute of Higher Education and Research, Chennai, Tamilnadu, India. \textsuperscript{2}Department of Radio Diagnosis, Sri Ramachandra Institute of Higher Education and Research, Chennai, Tamilnadu, India. \textsuperscript{3}Department of Radio Diagnosis, Sri Ramachandra Institute of Higher Education and Research, Chennai, Tamilnadu, India. \textsuperscript{4}Department of Radio Diagnosis, Sri Ramachandra Institute of Higher Education and Research, Chennai, Tamilnadu, India. \textsuperscript{5}Department of Radio Diagnosis, Sri Ramachandra Institute of Higher Education and Research, Chennai, Tamilnadu, India.

ABSTRACT

BACKGROUND
The main purpose of this article is to establish the usefulness of Diffusion Weighted Imaging (DWI) and Apparent Diffusion Coefficient (ADC) as complementary tools to conventional MRI sequences in characterizing benign (pancreatitis) and malignant (pancreatic adenocarcinoma) pancreatic pathologies in comparison to a control group.

METHODS
A prospective observational study was done over a period of two years with images obtained using Siemens Magnatom Avanto 1.5T MRI. 130 patients with pancreatic abnormalities were included in this study. An additional control group included 32 patients with normal pancreatic parenchyma. MRI findings including DWI and ADC values were calculated for all patients which was then correlated with clinical, lab and histopathological data.

RESULTS
Out of the 130 positive cases, 116 (89 \%) were benign and 14 (11 \%) were malignant lesions based on MRI features. Benign lesions include acute pancreatitis (47\%), acute on chronic pancreatitis (11\%) and chronic pancreatitis (31\%). Rest of the cases were pancreatic adenocarcinomas (11\%). There was significant reduction of ADC values in patients with acute pancreatitis (mean ADC - 1.016 \times 10^{-3}) , acute on chronic pancreatitis (mean ADC - 1.122 \times 10^{-3}) and chronic pancreatitis (mean ADC - 1.277 \times 10^{-3}) as well as pancreatic carcinoma (mean ADC - 0.934 \times 10^{-3}) in comparison with the control group (mean ADC - 1.363 \times 10^{-3}).

CONCLUSIONS
DWI with ADC mapping can play a significant role in the evaluation of pancreatic diseases. It can help in differentiating benign and malignant disease (ADC values of malignancy are significantly lower than benign disease).

KEY WORDS
Pancreatidis, DWI, MRI, ADC, Adenocarcinoma

Corresponding Author:
Dr. Poovini Soundararajan,
7/4, Logaya Colony, 4\textsuperscript{th} Cross Street,
Saligramam, Chennai-600093,
Tamilnadu, India.
E-mail: spoovini@gmail.com

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**BACKGROUND**

Diffusion Weighted Imaging (DWI) and Apparent Diffusion Coefficient (ADC) are functional imaging tools in Magnetic Resonance Imaging (MRI) that help define tissue characteristics on the basis of random Brownian movement of water molecules. DWI provides qualitative information about the cellularity of the tissues and helps in differentiating abnormal tissue from normal tissue while ADC is a quantitative entity derived from DWI.

Although DWI was initially widely used only in neuroimaging, subsequent advances in MR imaging software and hardware such as “ultrafast” MR imaging techniques and multicoil receiver systems have made diffusion weighted (DW) MR imaging a practicable tool in imaging of abdominal organs such as pancreas overcoming initial limitations related to image artefacts.[1]

In general, restricted diffusion and lower ADC values are seen in highly cellular tumours, fibrotic tissue, abscess and cytotoxic oedema. Conversely, tissues with low cellularity & disrupted membranes show fast unrestricted movement of water molecules with facilitated diffusion and high ADC values.[2]

Though ultrasound is the first-line imaging modality used in imaging patients with abdominal complaints, sonographic imaging of pancreas can sometimes be challenging or even suboptimal due to factors like bowel gas shadows and obesity because of the inherent retroperitoneal location of pancreas. Hence, cross sectional imaging plays a major role in detecting pancreatic pathologies and the use of MRI for diagnosis and further characterization of pancreatic diseases is increasing day by day.[3] Our aim was to study the DWI and ADC values of benign and malignant pancreatic pathologies in comparison to a control group and therefore assess the utility of DWI and ADC as complementary tools to conventional MR sequences in diagnosing and classifying pancreatic pathologies.

**METHODS**

This prospective observational study was done over a period of two years after obtaining clearance from the institutional ethics committee. The images were obtained using Siemens Magnatom Avanto 1.5 T MRI. The study group included 130 patients with pancreatic abnormalities and an additional control group of 32 patients with normal pancreatic parenchyma. MRI findings including DWI and ADC values were studied to differentiate benign and malignant lesions which were further correlated with clinical, lab and histopathological data.

**MRI Protocol**

Diffusion weighted images were obtained in the axial plane using a single-shot echo planar imaging sequence (TR = 1000 ms/TE - 74 ms; slice thickness 6 mm; FOV 36-42 cm; matrix, 384 x 25, b-value of 1,000 s/mm²). Conventional MRI images and DWI imaging data including ADC values were then analysed.

**RESULTS**

During the study period, a total of 130 cases with pancreatic pathologies were identified on MRI abdomen / MRCP with a wide range of age of presentation ranging from 12 to 93 years. Among these, males constituted 64% of the study population. Additionally, normalized ADC values were calculated for a control group of 32 patients who were free of pancreatic disease after averaging ADC values of head, body and tail. The mean ADC value of this control group was found to be 1.36326 × 10⁻³ mm²/s.

Out of the 130 positive cases, based on MRI features 116 (89%) cases were classified as benign and 14 (11%) were classified as malignant lesions. The benign lesions included were acute pancreatitis (47%), acute on chronic pancreatitis (11%) and chronic pancreatitis (31%). Rest of the cases were pancreatic adenocarcinomas (11%). As anticipated, all the above mentioned pathologies demonstrated hyperintense signal on DWI images, following which ADC values were then calculated from areas of true diffusion. The mean ADC value in the control group and the various pancreatic pathologies were charted as shown in Table 1.

**Image Analysis**

Following the generation of ADC map in grey scale, the Region of interest (ROI) was placed in areas of true restriction without necrosis or haemorrhage. Two ROI of size 50 +/- 10 mm² were placed following which ADC was computed automatically by the pre-set software (Argus in Siemens). Mean ADC was then calculated by averaging of the ADC values. The findings on Conventional MRI were correlated with findings on DWI (with corresponding ADC values). The mean ADC values were compared between the groups using N par Mann-Whitney test.
Hypothesized that the decrease in ADC values in acute pancreatitis was multifactorial occurring secondary to acinar cell death and subsequent pathological changes such as intercellular fibrin deposition.

Additionally, in our study, ADC values were also found to be lower than the control group in 15 patients with acute on chronic pancreatitis with a mean of 1.122 x 10^-3 mm^2/s ± 0.14.

In 40 patients diagnosed with chronic pancreatitis ADC values ranged from 0.745 x 10^-3 mm^2/s to 1.566 x 10^-3 mm^2/s with a mean value of 1.277 x 10^-3 mm^2/s. The overlap with control group ADC values seen in few patients may be attributed to the complete resolution of inflammation. Although chronic pancreatitis usually results in diffuse atrophy of pancreatic parenchyma, it can occasionally cause focal enlargement of the gland, simulating a neoplasm. In such cases, similar to acute focal pancreatitis, ADC values can serve as problem solving tools in addition to conventional MR sequences to distinguish mass-forming pancreatitis from a malignant mass lesion.

The importance of ADC values in diagnosis of malignancies of liver, breast, uterus, cervix have been widely studied. Few similar studies have explored the limitations and diagnostic challenges faced while using DWI/ADC to detect pancreatic malignancies. In our study, the ADC value in pancreatic malignancies (n=14) ranged from 0.850 x 10^-3 - 1.14 x 10^-3 mm^2/s with a mean ADC value of 0.934 x 10^-3 mm^2/s ± 0.07 which was found to be pointedly lower than the normal control group of 1.363 x 10^-3 mm^2 ± 0.16 (p<0.0005). (Figure 1c) thus clearly defining the utility of DWI/ADC in such cases. All cases of suspected malignancies were followed up and the imaging diagnosis was confirmed with correlation of histopathological findings.

Compared to the ADC values of benign pathologies like acute (focal/diffuse) pancreatitis and chronic pancreatitis, the ADC values in pancreatic malignancies were found to be significantly lower than the normal parenchyma (Figure 2). Statistical analysis obtained from 130 patients included in our study revealed significant p value (<0.0005) between ADC values in different subgroups of pancreatitis versus pancreatic adenocarcinoma.

As described above, DWI images also aided diagnosis of segmental pancreatitis by demonstrating focal hyperintense signal even in cases where the pancreatic parenchyma appeared apparently normal on conventional MR sequences.

Few limitations of DWI/ADC imaging include poor spatial resolution, lack of standardized imaging parameters/protocols and questionable reproducibility of ADC values. Additionally inevitable errors in ADC measurements can sporadically occur in body imaging due to MR artefacts despite optimization of imaging protocols. In patients with extensive parenchymal necrosis and in mass lesions with significant haemorrhage, T2 shine through confounds DWI images and accurate ADC values cannot be attained. Another inherent issue encountered in the analysis of DWI images was the occurrence of signal voids due to intercellular fibrin deposition.

In our study, among the 130 patients imaged, diagnosis of acute interstitial oedematous pancreatitis was found in 61 patients which was then further classified into focal/segmental pancreatitis (n=39) and diffuse pancreatitis (n=22) on the basis of disease distribution.

The mean ADC values for focal and diffuse acute pancreatitis were found to be 1.0165 x 10^-3 mm^2/s ± 0.13 and 1.323 x 10^-3 mm^2/s ± 0.34 respectively (Figure 1a and 1b). These values were found to be significantly lower (p < 0.0005) than the mean ADC value of the control group which was 1.363 x 10^-3 mm^2/s ± 0.16. Moreover, in 7 patients with focal pancreatitis, although conventional MR sequence images were found to be normal, segmental increased signal on DWI images facilitated the diagnosis of focal/segmental pancreatitis, as confirmed by elevated levels of serum amylase/ lipase. Isolated head involvement was the predominant pattern in segmental pancreatitis accounting for 51% followed by the tail region (30%).

Our findings correlated with a study by Yencilek E et al conducted with a smaller study population which also found that "the mean pancreatic ADC values in acute pancreatitis (1.19 x 10^-3 mm^2/s ± 0.32 ) was significantly lower than in the normal group (1.78 x 10^-3 mm^2/s ± 0.29) (p<0.001)". It is hypothesized that the decrease in ADC values in acute pancreatitis is multifactorial occurring secondary to acinar cell death and subsequent pathological changes such as intercellular fibrin deposition.

In the development of imaging protocols, DWI/ADC values were found to be very useful in differentiating focal pancreatitis from chronic pancreatitis. A study by Dent et al 

Table 1. ADC Value of Normal Pancreas and Various Pancreatic Pathologies vs Normal Pancreas

| Pathology                  | N    | Minimum (× 10^-3 mm^2/s) | Maximum (× 10^-3 mm^2/s) | Mean (× 10^-3 mm^2/s) | S.D.  |
|----------------------------|------|-------------------------|--------------------------|-----------------------|-------|
| Normal Pancreas            | 32   | 1.17                    | 1.763                    | 1.36326               | 0.164294 |
| Focal Pancreatitis         | 39   | 0.918                   | 1.45                     | 1.01652               | 0.138499 |
| Diffuse acute Pancreatitis | 22   | 0.852                   | 1.28                     | 1.12239               | 0.324532 |
| Acute on chronic Pancreatitis | 15  | 0.863                   | 1.55                     | 1.12333               | 0.163605 |
| Chronic Pancreatitis       | 40   | 0.745                   | 1.566                    | 1.277963              | 0.353202 |
| Pancreatic adenocarcinoma  | 14   | 0.85                    | 1.14                     | 0.934071              | 0.076194 |

DISCUSSION

In our study, among the 130 patients imaged, diagnosis of acute interstitial oedematous pancreatitis was found in 61 patients which was then further classified into focal/segmental pancreatitis (n=39) and diffuse pancreatitis (n=22) on the basis of disease distribution.
limitation of our study is that our results were obtained with 1.5 T scanner similar to few other studies on DWI/ADC in abdominal imaging. Therefore the probability that results obtained with a MR machine of higher magnetic field (3 T) may differ cannot be ruled out.

**CONCLUSIONS**

In our study, although the various pancreatic pathologies show reduced ADC values compared to normal parenchyma, pancreatic malignancies were found to have significantly lower ADC values. DWI also served as a problem-solving tool in diagnosis of select cases of pancreatitis when conventional MR images were apparently normal. In conclusion, DWI/ADC play an important role in distinguishing benign from malignant pancreatic entities and serve as useful complementary tools to conventional MR sequences in diagnosing and classifying pancreatic pathologies.

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