



## STUDY TO SEE THE USE OF ULTRASOUND IN THE DIAGNOSIS AND MANAGEMENT OF DENGUE FEVER (DF)

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### ABSTRACT

This study was conducted to evaluate ultrasonic (USG) finding in Dengue fever cases and effect of climate changes. 109 patients (age 5 to 70 year) who were serologically diagnosed as having DF between July, 2014 and October, 2014 were referred for USG, the imaging findings and the effect of climate analyzed. Out of 109 patients, all patients showed G.B. wall thickening, 72% honeycomb pattern, 69% hepatomegaly, 77% have splenomegaly, 88% Rt. Pleural effusion was seen and B/L pleural effusion was in 23%, 58% having ascites and pericardial effusion observed in 9%. Sonographic features of thickened GB wall, pleural effusion, ascites, hepatomegaly and splenomegaly strongly favour the diagnosis of DF in patients presenting with fever and associated symptoms, especially correlation with climatic or seasonal changes either due to decrease in virus virulence or herd immunity.

**KEYWORDS:** Dengue fever, Ultra sound, Management, Diagnosis.

### INTRODUCTION

One of the common causes of fever in the tropics is dengue virus. In 2008, South-east Asia and Western Pacific accounted for 70% of the global burden of dengue fever. The countries with a high incidence are Indonesia, Thailand, Myanmar, Sri Lanka, Bangladesh and India. Dengue is transmitted by mosquito *Aedes aegypti*, widely distributed throughout tropical and sub-tropical areas of the world.<sup>[1-3]</sup> There are four known serotypes of dengue, but severe form of dengue fever is caused by infection by more than one serotype. Clinically dengue

manifests with sudden onset of high fever with chills, intense headache, muscle and joint pain, retro-orbital pain and severe backache. Fever usually lasts for about 5 days, rarely for more than 7 days. Hemorrhagic diathesis and thrombocytopenia with concurrent hemoconcentrations are a constant finding.<sup>[4-7]</sup>

In mid 2014 there was an outbreak of DF in Lucknow, India. USG become an important adjunct to clinical & Laboratory profile in diagnostic DF. The hall mark of the pathogenesis of DF is the loss of endothelial integrity, which is assumed to be the result of an abnormal immune response against the virus. Clinical studies have shown that the level of cytokines and immune mediators are significantly increase in patients suffering from DF.<sup>[8, 9]</sup>

Mosquito –borne disease transmission is climate sensitive for several reasons; mosquitoes require standing water to breed, and a warm ambient temperature is critical to adult feeding, behavior and mortality, the rate of larval development, and speed of virus replication.<sup>[10]</sup> If the climate is too cold, viral development is slow and mosquitoes are unlikely to survive long enough to become infections. Although a suitable climate is necessary for disease transmission, other factors are needed for an epidemic to take place, including a source of infection, vector population and a susceptible human population.<sup>[11-12]</sup>

## MATERIAL AND METHODS

All ultrasound examinations were performed with a machine using 3.5 MHz and 5 MHz probes. Abdominal scanning was done after 6 h of fasting to allow better distension of gall bladder. GB wall thickening, which was the consistent finding in all the serologically positives cases, was measured by placing the calipers between the two layers of anterior wall. Thoracic scanning was done in either sitting or supine posture.<sup>[13]</sup> Both the pleural spaces were evaluated through an intercostals approach. Pericardial space was also evaluated for effusion subcostally. In all the patients ultrasound was performed prior to serology.<sup>[14]</sup> Serology tests using NS1 was performed to confirm the diagnosis in all the 122 patients which revealed 109 patients to be serologically positive for DF. The remaining 13 patients were serologically negative and not included in the study. GB wall thickening was considered when it in more than 3 mm, hepatomegaly was considered when cranio-candal measurement exceeds 13 cm in mid clavicular line and splenomegaly was taken when volume exceeds 180ml. The 13 cases which were serologically negative, 8 cases shows, mild thickening of GB wall (4-5 mm) and mild hepatosplenomegaly, But none have polyserositis The remaining 5 cases of fever shows no significant finding in USG.<sup>[15]</sup>

**RESULT**

Out of 109 patients, 32 patients admitted in July-14, 45 Patients in August, 26 Patients in September and only 6 patients in October -14. GB wall thickening was observed in all the patients and ranges from 4 mm to 11mm. patients seen in July and August GB wall thickening was 8-11 mm in compression with patients seen in September & October having GB thickening 4-8 mm. Perichole cystic fluid seen in 100% cases of DF in July and August, while in September 80% & in October 33%. Honeycomb pattern of GB walls seen in 93% cases in July and 95% cases of August while only 23% cases of September and in October no cases shows honeycomb pattern. Average incidence of hepatomegaly seen in 69% and splenomegaly 77%, polyserositis shows a decreasing trend in cases of DF admitted in months of September and October in comparison to patients admitted in month of July & August (Table-1).

**Table-1: Ultra-sonographic investigations**

Date	GB Wall Thickening	Honey comb	pericholecystic fluid	Hepato-megaly	Spleen	Rt Pleural effusion	B/L Pleural effusion	Ascites	Pericardial effusion
Jul-14	n=32 100%	n=30 93%	n=32 100%	n=22 70%	n=25 78%	n=27 85%	n=7 21%	n=22 70%	n= 5 15%
Aug-14	n= 45 100%	n =43 95%	n = 45 100%	n=34 75%	n=38 85%	n=37 82 %	n=12 27%	n=32 72%	n=5 12%
Sep-14	n= 26 100%	n=6 23%	n=21 80%	n=16 60%	n=18 69%	n=11 42%	n=5 19%	n=8 31%	n =nil
Oct-14	n=6 100%	n= nil	n=2 33%	n=3 50%	n=3 50%	n=2 34%	n=1 16%	n=1 16%	n=nil
<b>Total</b>	<b>109</b>	<b>79</b>	<b>100</b>	<b>75</b>	<b>84</b>	<b>97</b>	<b>25</b>	<b>63</b>	<b>10</b>
<b>Total %</b>	<b>100%</b>	<b>72%</b>	<b>92%</b>	<b>69%</b>	<b>77%</b>	<b>88%</b>	<b>23%</b>	<b>58%</b>	<b>9%</b>

**DISCUSSION**

Dengue is an acute febrile viral disease caused by flavivirus. It occurs in two forms: DF, a milder form of the disease and DHF. It is now endemic in more than 100 countries and threatens the health of 40% of the world's population. It is estimated that 50 million dengue infections occur each year with 5000000 cases of DHF and at least 12000 deaths annually mainly among children.<sup>[16]</sup> The increase of DF is due to uncontrolled population growth and urbanization in the absence of appropriate water management, global spread of dengue strains via travel and trade and due to erosion of vector control programmes.<sup>[17]</sup> In India the problem is even more acute because since 1963, more than 50 outbreaks have been reported by the National Institute of Communicable Disease, New Delhi.

Dengue viruses are transmitted to humans through the bite of infective female *Aedes mosquito*. The incubation period of the disease is 3-14 days. The onset of the disease is recognized by the sudden onset of high fever, retro-orbital pain, thrombocytopenia and hemorrhagic manifestations. Common laboratory findings include pancytopenia and prolonged bleeding time. These findings are consistent with increased vascular permeability, plasma leakage, abnormalities of haemostasis and protein losing shock syndrome, which commonly occur in DF pathogenesis.<sup>[18]</sup>

Serology is the mainstay in the diagnosis of DF. Haemagglutination inhibition antibodies usually appear at detectable level by day 5 to 6 of febrile illness. The diagnosis of DF is often delayed owing to time taken for availability of results. The aim of our study was to evaluate the ultrasound findings in DF, to find whether ultrasound of the abdomen is an important adjunct to clinical and laboratory profile in diagnosis.<sup>[19]</sup>

We have observed a seasonal climatic change in the USG findings of patients of DF in the month of July & August, showing severe forms of findings in terms of greater GB wall thickness (7-11mm), honeycomb pattern, pericardial effusion & pleural effusion, while the patients came in the month of September, & October show a milder form of DF, consisting of a mild increase in G.B. wall thickness, decrease incidence of poly serositis & absence of honeycomb pattern in wall of GB.<sup>[20-23]</sup>

Many studies have shown that humidity defined vapour pressure of specific humidity is high only when rainfall and temperature are high and these conditions that are conducive to breeding and survival of vector population and rapid replication of the virus.<sup>[24-27]</sup> Month of July and August on having high humidity level in comparison to month of September & October. However due to small sample size the effect of humidity, could not be certainly asserted.

## CONCLUSION

Sonographic features of thickened GB wall, pleural effusion, ascites, hepatomegaly and splenomegaly strongly favour the diagnosis of DF in patients presenting with fever and associated symptoms, especially correlation with climatic or seasonal changes either due to decrease in virus virulence or herd immunity.

**REFERENCES**

1. New Delhi: Regional Office for SEAR; 2008. WHO: Health situation in South East Asian region 2001-2007.
2. New Delhi: Ministry of Health and family welfare; 2006. Internet, Government of India. National Vector Borne Disease Control programme.
3. Joshi P, Rathnam VG, Sharma S. USG Finding in dengue Hemorrhagic fever –Our experience in the recent epidemic. *Indian Imaging* 1997; 7: 189-92.
4. Weekly epidemiological record No 6,8<sup>th</sup> February 2002: 41-3.
5. Singh B. Dengue outbreak in 2006: Failure of public health system? *Indian J Community Med* 2007; 32: 99-1010.
6. New- Delhi: DGHS, Ministry of health and family Welfare; 2010 Governments of India. Annual Report 2009-2010.
7. National vector Borne Disease Control Programme. Annual Report 2011-12. 2012.
8. Ukey P, Bondade S, Paunipager P, Power R, Akulwar S. Study of seroprevalence of dengue fever in central india. *Indian J Community Med* 2010; 35: 517-9.
9. Jawetz Melnick 24<sup>th</sup> ed. McGraw Hill, Lange Publications, Adelbergs Medical Microbiology, 2007; 350-5.
10. Lal M, Aggarwal A, Oberoi A, Dengue fever- An emerging viral fever in Ludhiana, North India. *Indian J Public Health* 2007; 51: 198-9.
11. Venkata Sai PM, Dev B, Krishnan R. Role of ultrasound in dengue fever. *Br J Radiol* 2005; 78: 416-8.
12. Shlaer WJ, Leopold JR, Scheible FW. Sonography of thickened gallbladder wall: a non-specific finding. *Am J Roentgenol* 1981; 136: 337-9.
13. Health Situation in the south East Asia Region 1994-1997, Regional office for SEAR, New Delhi; WHO (1999).
14. Pramuljo HS, Harun SR. Ultrasound findings in Dengue haemorrhagic fever, *Pediatric Radiol J* 1991; 21: 100-2.
15. Joshi P, Rathnam VG, Sharma S. USG findings in dengue haemorrhagic fever –our experience in the recent epidemic. *Ind J Radiol imag* 1997; 7: 189-92.
16. Intergovernmental panel on climate, Change (IPCC). Climate change 2001: Impacts, adaptation and vulnerability. Contribution of Working Group I to third assessment report of the Inter governmental panel on climate change. Cambridge: Cambridge University press, 2001.

17. Committee on climate ecosystems infectious diseases and human health. Under the weather. Climate, ecosystems, and infectious disease. Washington: National Academy press, 2001.
18. World Health Organization. WHO report on global surveillance of epidemic – prone infectious diseases Geneva: WHO, 2000.
19. Focks D, Haile D, Daniels E, Mount G. Dynamic life table model for *Aedes aegypti* (L) (Diptera culicidae). Analysis of the literature and model development. *J Med Entomol* 1993; 30: 1003-17.
20. Patz J, Martens W, Focks D, Jetten T. Dengue fever epidemic potential as projected by general circulation models of global climate change. *Environ Health Perspect* 1998; 106: 147-53.
21. Martens P, Jetten T, Focks D. Sensitivity of malaria, schistosomiasis and dengue to global warnings. *Climatic Change* 1997; 35: 145-56.
22. Martens P, Kovats R, Nijhof S, et al. Climate change and future populations at risk of malaria. *Global Environ Change* 1999; 9: S59-S107.
23. Hales S, Weinstein P, Soares Y, Woodward A. El Nino and the dynamics of vector born disease transmission. *Environ Health Perspect* 1999; 107: 99-102.
24. Rogers D, Randolph S. The global Spread of malaria in a future, warmer world. *Science* 2000; 289: 1763-66.
25. Reiter P. Climate change and mosquito borne disease. *Environ Health Perspect* 2001; 109(Suppl1): 141-61.
26. Bos E, Vu M, Massiah E, Bulatao R. World population projection 1994-95: estimates and projections with related demographic statistics. New York: Johns Hopkins University press, 1994.
27. Carter T, Hulme M, Lal M. Guidelines on the use of scenario data for climate impact and adaptation assessment. Geneva: Inter-governmental panel on climate change, 1999.