Clinico-epidemiological Study on Dengue Pocket Outbreak at the AWMCH from July 2022 to May 2023

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#### Introduction

- Dengue an arthropod-borne viral infection of humans
- Endemic to tropical and subtropical regions of the world and represents an important public health problem.
- Dengue viruses are transmitted by the bite of Aedes Aegypti mosquito primarily and to a lesser degree by Aedes Albopictus
- Infected by the one of the four dengue serotypes DEN 1,2,3,4.

### Aims of our study

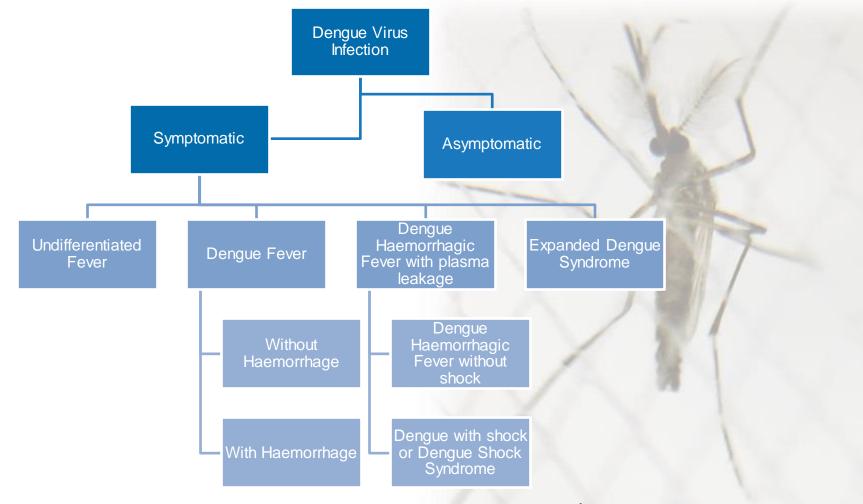
- Understand the clinical and epidemiological dimensions of the dengue outbreak
- Analyze the impact of demographic factors on the occurrence and severity of dengue
- Evaluate the frequency, pattern, and causes of disease in specified populations and risk factors associated with them, variation in clinical presentation, differences in treatment modality

### Objectives

- Identify the sero-immunological prevalence of the dengue syndrome cases
- Document the demographic and socio-economic variable
- Assess the clinical signs and symptoms among the specified group
- Understand the changing pattern of dengue syndrome this year

### Methods & Methodology

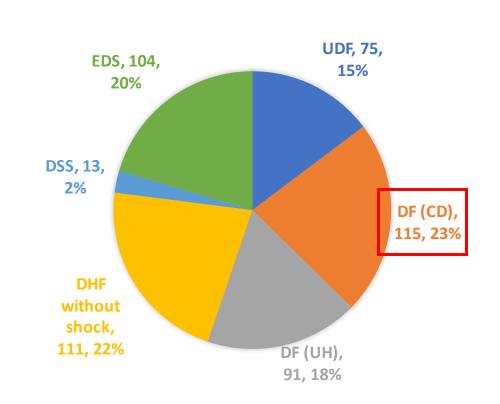
- Study Design: Cross-sectional Prospective Study
- Place of Study: Ad-din Women's Medical College & Hospital
- Study Population: 509
- Study Period: from July 2022 to May 2023



Ref: National Guideline For Clinical Management of Dengue Syndrome, 4th Edition 2018 (Revised)

Table 1: Distribution of Dengue Syndrome among the study population (n=509)

Dengue Syndrome		Number of
		Patients
	1 aticitts	
Undifferentiated F	ever (UDF)	75
Dengue Fever	DF without	115
(DF)	haemorrhage	
(DI)		
	DF(CD)	
	DF with unusual	91
	haemorrhage	
	DF(UH)	
Dengue	DHF without	111
Haemorrhagic	shock	
Fever (DHF)	SHOCK	
rever (Diff)	Dengue Shock	13
		1)
	Syndrome (DSS)	
Expanded Dengue	104	
Total	500	
TOTAL	509	



*UDF*: those who develop a simple fever indistinguishable from other viral infections, especially those who have been infected with the dengue virus for the first time.

Table 2: Association of patients' demographic characteristics with dengue syndrome

Danis amandia amidda			Type of Dengue Syndrome (n=509)						
Demographic	variables	UDF	DF(CD)	DF(UH)	DHF	DSS	EDS		
		(n=75)	(n=115)	(n=91)	withou	(n=13)	(n=104)		
					t shock (n=111)				
Sex of patient	Male- 42% (n=214)	31(14%)	51(24%)	48(22%)	47(22 %)	6(3%)	31(14%)		
	Female- 58% (n=295)	44 (15%)	64 (22%)	43 (14%)	64 (22%)	7 (2%)	73 (25%)		
	≤19years (n=163)	49	30	51	19	8	6		
Age of patient	20-39 years (n=255)	21	61	30	70	3	70		
	≥40years (n=101)	5	24	10	22	2	38		

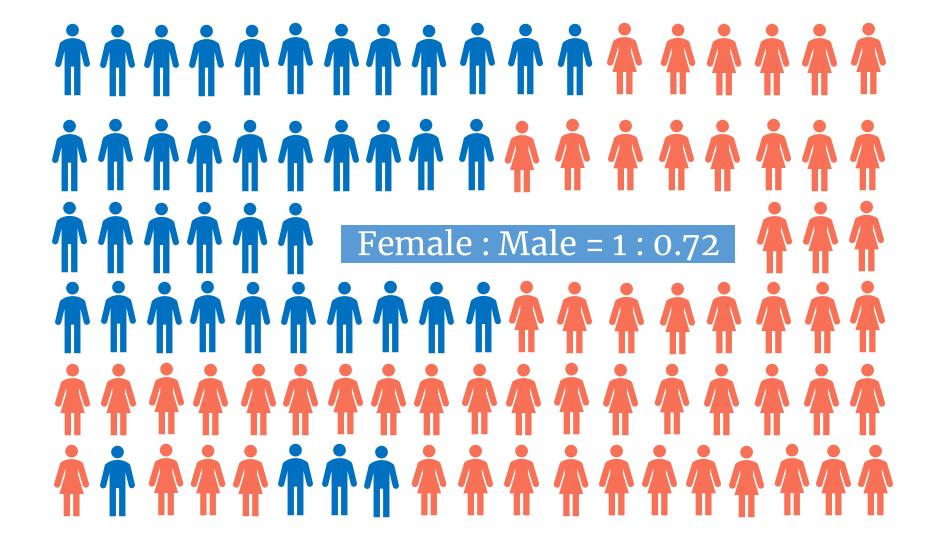


Table 2: Association of Sero-immunological findings with Dengue Syndrome

		Type of Dengue Syndrome (n=509)					
Sero-im	nmunological findings	UDF (n=75)	DF (CD) (n=115)	DF(UH) (n=91)	DHF without shock (n=111)	DSS (n=13)	EDS (n=104)
	Positive(60%) (n=261)	0(0%)	77(30%)	71(27%)	71(27%)	10(4%)	32(12%)
NS1	Negative (40%)(n=181)	75	38	20	40	3	5
	Positive(16%) (n=69)	0(0%)	22(32%)	7(10%)	36(52%)	1(1%)	3(4%)
IgM	Negative(84%) (n=373)	75	93	84	75	12	34
	Positive(n=57)	0	20	7	27	1	2
IgG**	Negative (n=385)	75	95	84	84	12	35

Table 3: Composite table showing association of dengue syndrome with socio-economic and environmental status

		Type of Dengue Syndrome(n=509)				
Socio-Economic Variables		Mild dengue infection(n=281) Undifferentiated Fever(75 cases)+ DF(UH) (91 cases)+ DF(CD) (115 cases)	Severe dengue infection(n=228) DHF (111 cases)+DSS (13 cases)+EDS(104 cases)			
Income Group	Total Monthly Income					
Low-income	≤9737BDT(n=84)	46	38			
Lower-mid income	9738- 38184BDT(n=126)	92	34			
Upper-mid income	38185 - 118500BDT(n=165)	108 (38.4%)	57(25%)			
High-income	≥118501BDT(n=134)	102	32			
Following World Bank,2023 per capita Gross Net Income: GNI; HH=Household Economy: Total monthly HH-income						

of families in BD Currency(Tk.)

Table 3: Composite table showing association of dengue syndrome with socio-economic and environmental status

Environmental	Variables	Mild dengue	Severe dengue infection(n=228)
		infection(n=281)	DHF (111 cases)+DSS (13
		Undifferentiated Fever(75	cases)+EDS(104 cases)
		cases)+ DF(UH) (91 cases)+	
		DF(CD) (115 cases)	
	Ground floor(n=131)	82	49
	1st floor(n=178)	138	40
Which Floor	2 <sup>nd</sup> + 3 <sup>rd</sup> floor(n=158)	106	52
	4 <sup>th</sup> +Others floor(n=42)	22	20
Water	Yes(n=270)	206	64
Container	No(n=239)	142	97
Maggita Nat	Yes(n=342)	249	93
Mosquito Net	No(n=167)	99	68
Plant Flower	Yes(n=298)	192	106
Vase	No(n=211)	156	55

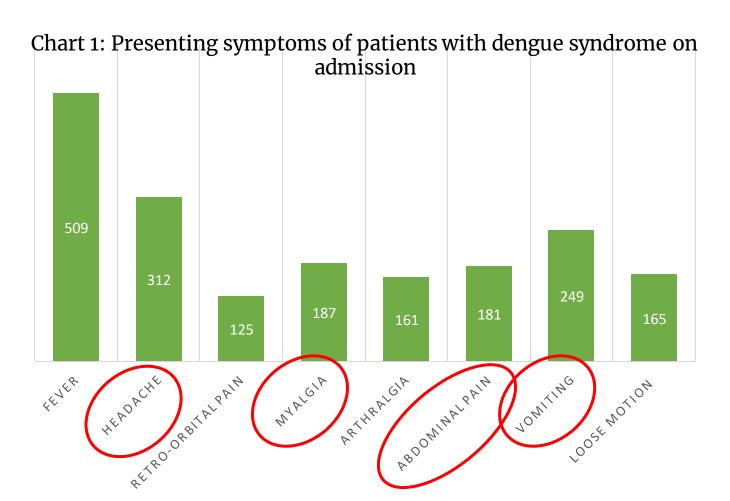


Table 4: Haemorrhagic manifestations of patients with Dengue Syndrome on admission

Haemorrhagic ma	nifestations		Type of Dengu	ie Syndrome (1	n=509)	
Types	Number of Patients	UDF (n=75)	DF with unusual hemorrhage (n=91)	DHF without shock (n=111)	DSS (n=13)	EDS (n=104)
Haematuria	100	0	35	50	3	13
Purpura	91	0	44	27	5	19
Epistaxis	80	0	65	9	1	3
Hematemesis	72	0	37	16	5	14
Hemorrhage spots inside lips/tongue	69	0	37	20	4	10
Hemoptysis	81	0	47	18	4	11
Gum Bleeding	102	0	57	39	5	4
PV Bleeding/Pre- date menstruation	152	0	77	42	5	23
Melena	60	0	33	18	2	2
Hematochezia	52	0	27	17	1	3

Chart 2: Haemorrhagic manifestations of patients with Dengue Syndrome on admission

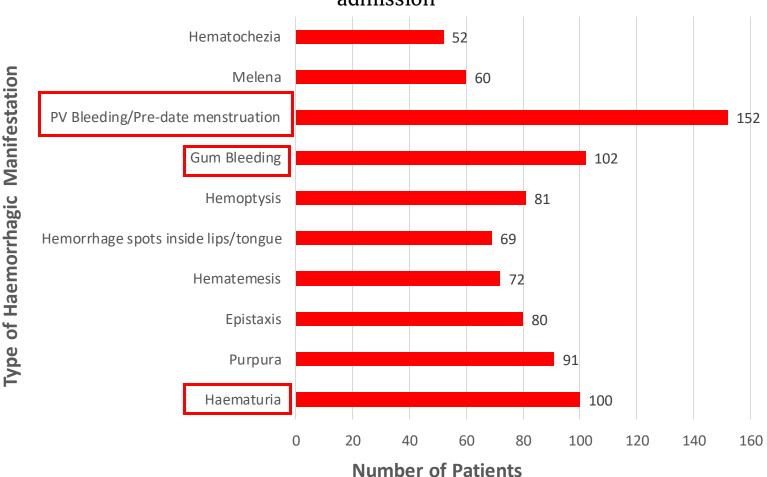


Table 5: Signs of patients with Dengue Syndrome on admission Type of Dengue Syndrome (n=509) UDF DF(CD) DF(UH) DSS Clinical signs DHF

EDS(n=104)

83

34(25%)

70

	O	(n=75)	(n=115)	(n=91)	without shock (n=111)	(n=13)	
Tourniquet	Positive (n=109)	0	0	0	81 (73%)	3(9%)	25
Test	Negative (n=400)	75	115	91	30 (27%)	10 (91%)	79
Capillary	Normal(n=444)	75	115	152	83(75%)	2(15%)	84
Refill Time	Prolonged(n=65)	0	0	6	28(25%)	11(85%)	20
Body Rash	Present(n=160) (31%)	16(10%)	51(32%)	22(14%)	36(23%)	7(4%)	32(20%)
bouy Kasii	Absent(n=349) (69%)	59	64	136	75	6	72
Ascites	Present (n=135) (27%)	0(0%)	0(0%)	0(0%)	103(75%)	11(8%)	21(15%)
Ascites	Absent (n=374)	75	115	01	8	2	82

115

0(0%)

115

91

0(0%)

91

75

0 (0%)

75

(73%)

(26%)

Absent

(74%)

Pleural

Effusion

Present (n=134)

(n=375)

8

91(68%)

20

2

9(7%)

4

Table 5: Haematological, Serological, Radiological Investigation Findings

				Туре о	f Dengue Syn	drome (n=50	09)	
		Parameters measured	UDF	DF (CD)	DF (UH)	DHF	DSS	EDS
of			(n=75)	(n=115)	(n=91)	without	(n=13)	(n=104)
sd						shock(n=		
no						111)		
e gr	Adolescents males (13—	Low (<38%) (n=12)	1	4	2	0	4	1
& age groups	18 years)	Normal (38-50%) (n=18)	3	1	12	0	1	1
⊗	(n=30)	High (>50%) (n=0)	0	0	0	0	0	0
nder	Adolescent	Low (<34%) (n=19)	2	1	9	7	0	0
to gender a	females (13- 18 years) (n=21)	Normal (34-44%) (n=2)	1	1	0	0	0	0
to	, , , ,	High (>44%) (n=0)	0	0	0	0	0	0
ling	Adult male	Low (<41%) (n=52)	3	7	8	29	0	5
HcT according	(≥19 years) (n=124)	Normal (41-50%) (n=58)	5	25	16	9	0	3
асс	, ,,	High (>50%) (n=14)	0	6	1	1	4	2
Ę	Adult female	Low (<36%) (n=88)	11	14	25	29	0	9
H	(≥19years)	Normal (36-44%) (n=82)	6	29	15	21	0	11
	(n=192)	High (>44%) (n=22)	1	6	5	4	1	5

*Limitations of finding baseline HcT:* 

HcT on admission was recorded as the baseline, as patients came to Ad-din after rehydration (oral/parenteral)

Table 5: Haematological, Serological, Radiological Investigation Findings

		UDF	DF (CD)	DF (UH)	DHF	DSS	EDS
		(n=75)	(n=115)	(n=91)	without	(n=13)	(n=104)
					shock(n=		
					111)		
Platelet Count	Low (86%) (n=439) *<100,000 cells/mm <sup>3</sup>	63 (14%)	99 (22%)	86 (20%)	104 (24%)	12 (3%)	75 (17%)
	Normal (14%) (n=70)	12	16	5	7	1	29
Total count of White Blood Cells	Normal (14%)(n=346)	46	82	92	88	10	28
White Blood delis	Leucopenia (86%) (n=162) *<5000 cells/mm <sup>3</sup>	29 (18%)	33 (20%)	66 (41%)	23 (14%)	3(2%)	8(5%)

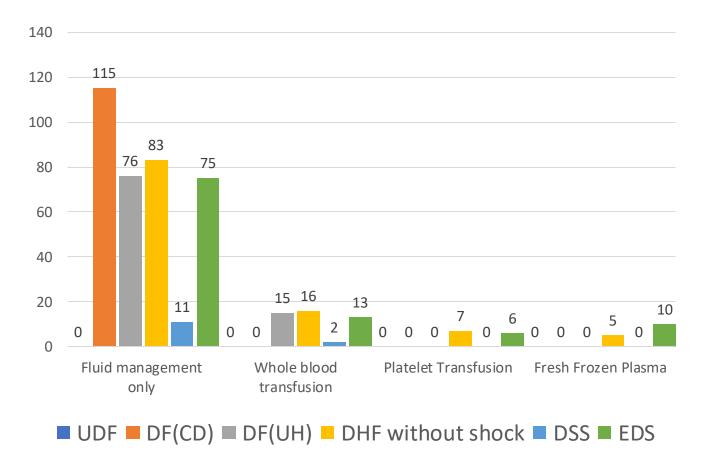
Table 5: Haematological, Serological, Radiological Investigation Findings

		UDF	DF (CD)	DF (UH)	DHF	DSS	EDS
		(n=75)	(n=115)	(n=91)	without	(n=13)	(n=104)
					shock		
					(n=111)		
USG Findings	Ascites +/- edematous or thickened gall bladder/ acalculous cholecystitis (n=172)	0(0%)	0(0%)	0(0%)	110(64%)	11(7%)	51(29%)
Chest X-ray findings	Pleural effusion (n=138)	0(0%)	0(0%)	0(0%)	93(67%)	10 (7%)	35(25%)
ALT	<40 IU/L (n=361)	75	93	85	59	7	42
ALI	40-400 IU/L(n=106)	0	22	6	49	5	24
	>400 IU/L (n=42)	0(0%)	0(0%)	0(0%)	3(7%)	1(2%)	38(91%)

#### Table 6: Transfusion History of patients with Dengue Syndrome on admission

Transfusion			Type of D	engue Syndro	me(n=509)	
History	UDF	DF(CD)	DF(UH)	DHF	DSS	EDS
	(n=75)	(n=115)	(n=91)	without shock (n=111)	(n=13)	(n=104)
Fluid management only (n=360) (71%)	0(0%)	115 (32%)	76 (21%)	83(23%)	11(3%)	75(21%)
Whole blood transfusion(n=46)(9 %)	0(0%)	0(0%)	15(33%)	16(35%)	2(4%)	13(28%)
Platelet Transfusion(n=13)(3 %)	0(0%)	0(0%)	0(0%)	7(54%)	0(0%)	6(46%)
Fresh Frozen Plasma(n=15)(3%)	0(0%)	0(0%)	0(0%)	5(33%)	0(0%)	10(67%)
No transfusion/infusion required (n=75)(14%)	75(100%)	0(0%)	0(0%)	0(0%)	0(0%)	0(0%)

#### Chart 3: Transfusion History of patients with Dengue Syndrome on admission



## Mortality & Mode of Death

- 3 out of 509 dengue patients died
- Mortality rate was approximately 0.6%

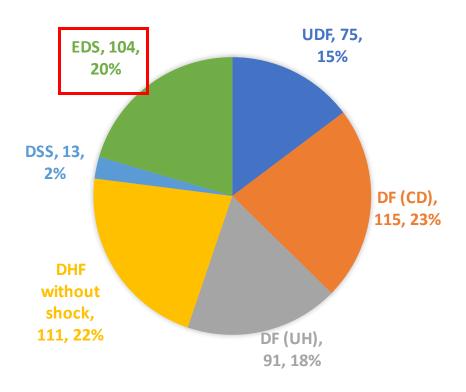
#### Mode of Death:

- Acute Liver Failure 1 out of 3 patients
- Myocarditis with cardiogenic shock 2 out of 3 patients

### Persistent Thrombocytopenia in Convalescent Phase of DENGUE: ITP in disguise

**Prof Dr Richmond Ronald Gomes Department of Internal Medicine** 

#### Distribution of Dengue Syndrome among the study population (n=509)



### Expanded dengue syndrome

The World Health Organization, in 2012, coined the term "expanded dengue syndrome" to describe patients that do not fit into either DHF or DSS but **show atypical symptoms in vital organs systems** such as the cardiovascular system, neurological system, kidneys, gut, and hematological system.

## Expanded dengue syndrome

	<b>O</b> ,
System	Unusual or atypical manifestations
Neurological	Febrile seizures in young children.
	Encephalopathy.
	Encephalitis/aseptic meningitis.

Intracranial haemorrhages/thrombosis. Subdural effusions.

Acute renal failure.

Hemolytic uremic syndrome.

Renal

Mononeuropathies/polyneuropathies/Guillane-Barre Syndrome. Transverse myelitis. Gastrointestinal/hepatic Hepatitis/fulminant hepatic failure. Acalculous cholecystitis. Acute pancreatitis. Hyperplasia of Peyer's patches.

Acute parotitis.

# Expanded dengue syndrome

Myositis with raised creatine phosphokinase (CPK).

Infection associated haemophagocytic syndrome.

IAHS or Haemophagocytic lymphohistiocytosis (HLH), idiopathic

Post-infectious fatigue syndrome, depression, hallucinations, psychosis,

System	Unusual or atypical manifestations			
Cardiac	Conduction abnormalities.			

Myocarditis.

Pericarditis.

Respiratory Acute respiratory distress syndrome.

Pulmonary haemorrhage.

thrombocytopenic purura (ITP). Spontaneous splenic rupture.

Lymph node infarction.

Macular haemorrhage. Impaired visual acuity.

Optic neuritis.

alopecia.

Rhabdomyolysis.

Musculoskeletal

Eye

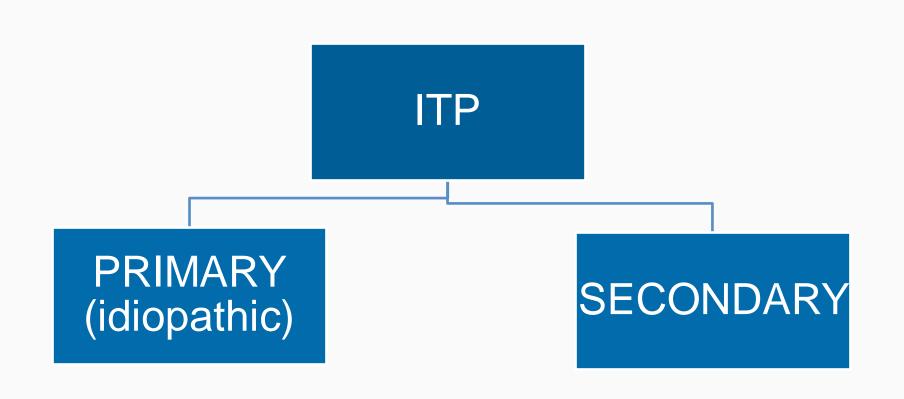
Others

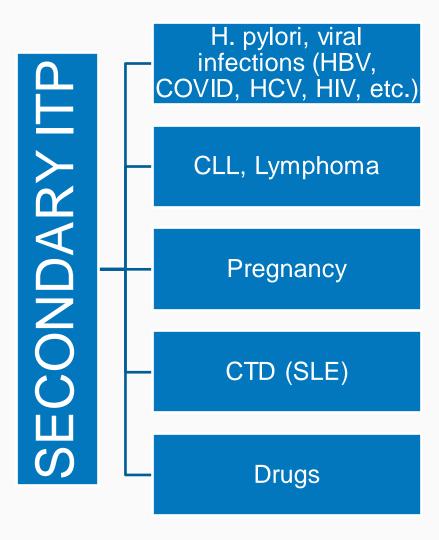
Lymphoreticular/bone marrow

#### Immune Thrombocytopenic Purpura

Idiopathic thrombocytopenic purpura (ITP) is immune mediated with involvement of autoantibodies most often directed against the platelet membrane glycoprotien IIb/IIIa, which sensitise the platelet resulting in premature removal from the circulation by cells of the reticulo-endothelial system.

\* Davidson's Principles and Practice of Medicine, 24th Edition





Distribution of dengue syndrome among the study population (n=509) Hepatitis+ Acalculous 38 36.5% cholecystitis ITP 34 32.6% **DHF** without Cholecystitis 11.5% shock **Pancreatitis** 4 3.8% DSS 22% Acute Kidney Injury 3.8% DF(UH) Myocarditis +ALF 3.8% **EDS** 18% **RPGN** 1.0% 20% SIADH 1 1.0% Non Immune Hemolytic 1 1.0% Anemia UDF 3 2.9% Meningitis DF(CD) 15% 23% Encephalitis 1.9% Total 104

# Pathophysiology of Thrombocytopenia in Severe Dengue

- Direct virus induced platelet destruction
- Virus induced bone marrow suppression
- Auto-immune destruction of platelets
- Increased platelet consumption due to trapping of platelet in sticky and damaged endothelium

# Objectives of the study

- ☐ Find the seropositive dengue cases during this ongoing outbreak with persistent thrombocytopenia (below 50,000 cells/mm³) from post critical phase (convalescent phase) (Beyond 10 days of illness)
- ☐ To rule out the secondary causes of ITP (HBV,HCV,HIV,SLE) to determine if dengue is the underlying etiology in these cases
- To determine the role of corticosteroids in treating secondary ITP which is not categorically prescribed in dengue by monitoring the preand post-treatment platelet count.

### Methods & Methodology

- Study Design: Case Control Study
- Place of Study: Ad-din Women's Medical College & Hospital
- Study Population: 50
- Study Period: from July 2022 to May 2023

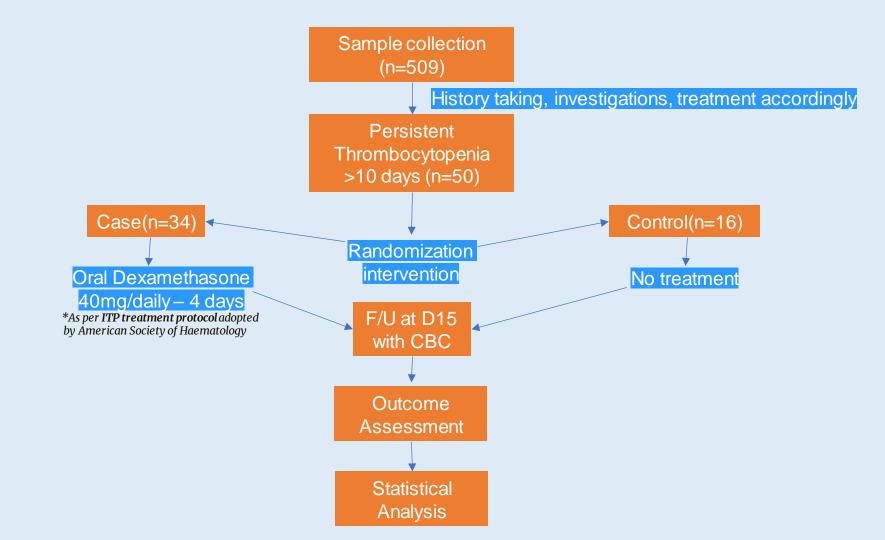


Table 1: Comparison between the case and control (N=50)

Dengue IgM positive 13 (26%)

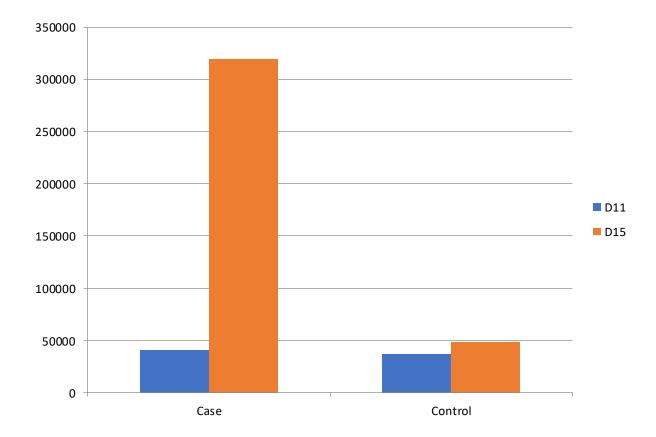
Demographic and clinical profile	Case	Control
(N=50)	(34)	(16)
Age in years	37.55 ± 14.94	35.87 ± 13.02
(Mean ± SD): 37.02 ± 14.24		
Sex		
Male - 32 (64%)	19 (55.88%)	13(81.25%)
Female – 18 (36%)	15 (44.115)	3 (18.75%)
Comorbidities		
DM 7 (14%)	4(11.765)	3 (18.75%)
HTN 5 (10%)	4 (11.76%)	1(6.25%)
DM+HTN 1 (10%)	1(2.945)	0
Bronchial asthma 6 (12%)	3 (8.82%)	3 (18.75%)
Multiple comorbidity (DM, HTN,	3 (8.82%)	3 (18.75%)
IHD,CKD) 6 (12%)		
NS 1 Positive 37 (74%)	27 97.41%)	10 (62.5%)

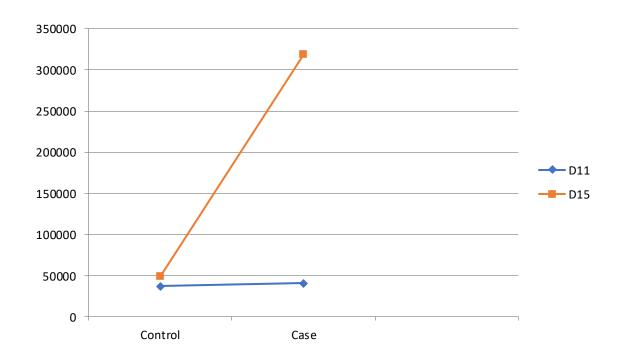
7 (20.58%)

6 (37.5%)

#### Results of our study

	Case	Control	P value
	(34)	(16)	
Platelet			
After 11 days	40470.58± 4775.15	37062.50 ±6597.66	0.043
After 15 days	319352.94 ± 93717.13	48437.50 ± 7907.53	0.000





#### Conclusion

- Persistent Thrombocytopenia beyond 10 days of dengue illness in the convalescent phase is rare.
- If present, we can suspect that the patient is having this thrombocytopenia due to dengue-induced secondary ITP as a part of expanded dengue syndrome. To prove this, as we have no provision of anti-platelet antibody, we can conduct a therapeutic trial with short term steroid therapy (dexamethasone) after excluding common causes of secondary ITP.

#### Conclusion

• In our study, all the patients responded in the therapeutic trial, whereas the control group had little or no improvement in platelet count, so we can conclude that cases with persistent thrombocytopenia in dengue fever beyond 10 days of illness maybe a suitable candidate for short-term steroid therapy.

• Our trial also proves that dengue is one of the underlying etiology for secondary ITP.

# THANK YOU