

## A CLINICOEPIDEMIOLOGICAL STUDY ON IGA VASCULITIS IN ADULTS AND CHILDREN

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### Abstract

**Background:** The IgA vasculitis is a small vessel vasculitis. It is more common in children. It is formerly known as Henoch Schonlein purpura. Palpable purpura is the first and most common sign. It is underreported in adults. The aim of the study is to compare the Clinical and epidemiological features of IgA vasculitis in adults with that of IgA vasculitis in Children. **Material and Methods:** The study was conducted in the Department of Dermatology, Government Rajaji Hospital, Madurai Medical college among the patients attending dermatology OPD with cutaneous small vessel vasculitis. The study was done for a period from June 2020 to May 2021. 65 study participants recruited based on inclusion and exclusion criteria. The enrolled study participants were randomized and allocated into two groups (Adult-34, children-31). Baseline characteristics of the study participants were obtained. Occupation history is also elicited. Clinical examination both general and systemic were done. Detailed Skin examination was done. Laboratory test, throat swab and Ultrasound abdomen was also done. Data collected were entered in MS Excel and the statistical analysis was done in SPSS 23. p value <0.05 is considered as statistically significant. **Results:** Majority of the study participants were in 19-40 years 21(62%) of age in adults and less than 10 years in children. Female preponderance were more in our study. Disease manifestation is observed more in summer for adults and winter for children. For majority of the diseases the etiology was unknown. Joint pain, pedal edema and itching over the lesion is significant in adults compared to children. In 30(96%) of the study participants present with palpable purpura whereas in adults only 25(73%) were with palpable purpura followed by ecchymotic patches 4(12%). Thrombocytosis was seen more in Children 5(16%) and 2(6%) in adults and it was found to be statistically significant. ASO titre was found to be increased in adults. Microscopic hematuria was noted in 3(9%) of adults and 2(7%) in children. Throat swab was found to be positive only in 2(6%) of adults. Recurrence was more in adults. **Conclusion:** Our study concluded stating that IgA vasculitis is adults is not uncommon and it is frequent as in children. Hence IgA vasculitis should be considered as one of the differential diagnosis for adults with vasculitis. Longer follow up has to be done in adults with elevated renal parameters to uncover the delayed systemic complications.

## INTRODUCTION

IgA vasculitis is a disease which involves deposition of immune complexes containing IgA along the joints, kidneys, gastrointestinal tract. It less commonly affects the central nervous system and testicles. It is a subset of vasculitis. Small vessel vasculitis affects both children and adult. It predominantly occurs in 3-15 years of age. More

than half of the cases occurs in less than 5 years of age. Male: Female ratio of the disease is 1.5:1. The disease most commonly peaks in winter.<sup>[1,2]</sup> The incidence ranges from 6-24 per 100,000.<sup>[3]</sup>

It is either primarily arising denova or may occur secondary to infection, drugs, autoimmune diseases and malignancy. It is also suggested that environmental factors, lifestyle, economic status and genetic factors also associated with incidence of the

disease. The presentation varies widely as the signs and symptoms depends on the system involved and affected.<sup>[4,5]</sup> It is characterised by palpable purpura, arthritis, abdominal pain and renal involvement. The pathology behind IgA vasculitis is indicated by increased concentration of IgA and deposition of IgA in the renal glomeruli and vessel walls.<sup>[6]</sup> IgA vasculitis might be more common in children but due to underdiagnosis and couldn't differentiate between the essential differences between children and adults.<sup>[7,8]</sup>

IgA adults associated with severe diseases and therefore it is essential to focus on its management.<sup>[9]</sup> IgA vasculitis is otherwise known as Henoch Schonlein pupura (HSP). It is frequently a self limiting disease with favourable diagnosis if it not involves the renal system. It takes around four weeks for recovery in two thirds of the patients. But in one half the patients the recurrence occurs within 4 weeks to two years. More often in Adults IgA is idiopathic and 12.5% to 26% is reported as drug implicated whereas in 23% to 35% it is due to infections. Relapses occurs in 30% of the study participants and mostly of cutaneous. In adults recalcitrant is more common. Mortality is more in older patients with IgA vasculitis. Though many studies were done among children ,paucity of studies seen in adults.<sup>[10,11,12,13,14]</sup>

#### Aim Of the Study

To compare the Clinical and epidemiological features of Ig A vasculitis in adults with that of IgA vasculitis in Children.

## MATERIALS AND METHODS

**Study Setting:** This study was conducted in the Department of Dermatology, Government Rajaji Hospital, Madurai Medical college among the patients attending dermatology OPD with cutaneous small vessel vasculitis. The study was done for a period from June 2020 to May 2021.

**Study Design:** Comparative cross sectional study

#### Inclusion Criteria

- All patients with palpable purpura fulfilling the diagnostic criteria of IgA vasculitis.
- Age >18 years and both sexes

#### Exclusion Criteria

- Patients not fulfilling diagnostic criteria
- Patient who were unwilling for the study

#### Sample Size

The study participants fulfilling the inclusion and the exclusion criteria were included in the study throughout the study period. The final attained sample is 65.

**Data Collection Method:** After obtaining the Institutional Ethical Committee clearance, the study was started after obtaining patients informed consent. Based on inclusion and exclusion criteria the study participants recruited during the study period is 65. A detailed history was taken which includes name, sex, presenting symptoms, duration, joint pain, systemic symptoms, history suggestive of malignancy and collagen vascular diseases, drug intake history, past history of sore throat infection. Occupation history is also elicited. Clinical examination was done .Both general examination and systemic examinations. Detailed Skin examination was done i.e the morphology, distribution, symmetry, tenderness .Diascopy was also done.

Baseline Laboratory investigations were done which includes complete hemogram, serum urea, serum creatinine, liver function tests. chest X ray, urine (routine and microscopic haematuria), stool occult blood test, CRP, ASO titre, throat swab culture, ANA, USA Abdomen and pelvis. Patients were screened for HIV, Hepatitis B and C. Punch biopsies were done from the early tender skin lesions and sent to H & E. Direct immunofluorescence were done for the early skin lesions.

#### Statistical Analysis

The obtained data was entered in MS Excel Windows 10. Statistical analysis was done with the help of SPSS 23. Continuous data was expressed in terms of mean and standard deviation. Categorical data was expressed in terms of Numbers and Percentages. Test of association for Categorical data was Chi square test and for Continuous data was t test and Anova test. p values <0.05 is considered as statistically significant.

## RESULTS

**Table 1: Demographic profile of the study participants**

Baseline Characteristics	Adult (N=34)	Children (N=31)	P value
Age			
<10 years	0(0%)	19(61%)	<0.001
11-18years	0(0%)	12(39%)	
19-40 years	21(62%)	0(0%)	
41-60 years	13(38%)	0(0%)	
Sex			
Male	14(41%)	13(42%)	0.54
Female	20(59%)	18(58%)	

Majority of the study participants were in 19-40 years 21(62%)of age in adults and less than 10 years in children. Female preponderance were more in our study.

**Table 2: Epidemiological profile of the study participants**

Variables	Adult (N=34)	Children (N=31)	P value
Duration of illness			
<5	20(59%)	29(94%)	<0.001
>5	14(41%)	2(6%)	
Seasonal variation			
Summer	19(56%)	10(32%)	<0.001
Winter	7(20.5%)	17(55%)	
Monsoon	8(23.5%)	4(13%)	
Etiological factors			
Unknown	27(79%)	19(61%)	8.172
URI	5(15%)	12(39%)	
Drugs	2(6%)	0(0%)	
Koernisation			
Positive	5(15%)	0(0%)	0.07
Negative	29(85%)	31(100%)	

Disease manifestation is observed more in summer for adults and winter for children. For majority of the diseases the etiology was unknown followed by URI. Koernisation was seen more in adults (15%).

**Table 3: Clinical profile of the study participants**

Clinical features	Adult (N=34)	Children (N=31)	P value
Sore throat	5(15%)	1(3%)	0.24
Fever	16(47%)	8(26%)	0.13
Itching	14(41%)	2(6%)	0.003*
Burning sensation	5(15%)	1(3%)	0.24
Joint pain	25(74%)	9(29%)	<0.001*
Joint swelling	4(12%)	1(3%)	0.41
Abdominal pain	6(18%)	7(23%)	0.85
Melena	1(3%)	0(0%)	0.96
Diarrhoea	3(9%)	2(6%)	0.91
Pedal edema	7(21%)	0(0%)	0.02*
Past history of similar lesions	7(21%)	3(7%)	0.38

Joint pain, pedal edema and itching over the lesion is significant in adults compared to children. Other clinical symptoms are marginally more in adults compared to children.

**Table 4: Laboratory parameters of the study participants**

Variables	Adult (N=34)	Children (N=31)	P value
Hemoglobin			
<11 grams	8(23.5%)	2(6%)	0.09
11-13 grams	18(53%)	20(65%)	
>13 grams	8(23.5%)	9(29%)	
Platelets (Lakhs)			
<2	3(9%)	2(7%)	0.003*
2.1-4.5	29(85%)	24(77%)	
>4.5	2(6%)	5(16%)	
ASO			
Positive	5(15%)	0(0%)	0.07
Negative	29(85%)	31(100%)	
RFT			
Elevated	7(21%)	0(0%)	0.01*
Normal	27(79%)	31(100%)	
CRP			
Positive	5(15%)	0(0%)	0.07
Negative	29(85%)	31(100%)	
Urine for RBC			
Positive	3(9%)	2(7%)	0.914
Negative	31(91%)	29(93%)	

Majority of the study participants have haemoglobin of 11-13 grams (Adult 18(53%) and Children 20(65%). Thrombocytosis was seen more in Children 5(16%) and 2(6%) in adults and it was found to be statistically significant. ASO titre was found to be increased in adults and none detected in children. RFT was found to be elevated in 7(21%) of the adults whereas in children it was in normal range. CRP was found to be elevated only in adults 5(15%). Microscopic hematuria was noted in 3(9%) of adults and 2(7%) in children.

**Table 5: Ultrasound and throat swab results of the study participants**

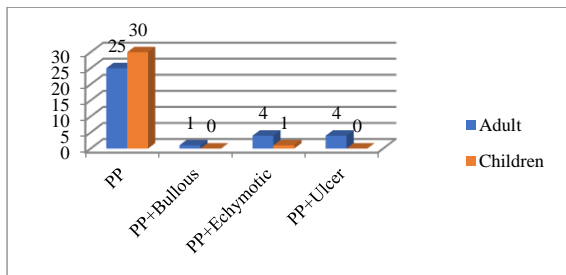
Variables	Adult (N=34)	Children (N=31)	P value
USG abdomen			
Fatty liver	2(6%)	0(0%)	0.41
Bowel wall edema	1(3%)	1(3%)	
B/L ovarian mass	1(3%)	0(0%)	
Nil	30(88%)	30(97%)	
Throat swab			
Staphylococcus/Klebsiella	2(6%)	0(0%)	0.51
Nil	32(94%)	31(100%)	

In adult group the most common USG finding was fatty liver 2(6%) followed by bowel wall edema 1(3%) and B/L ovarian mass 1(3%). Whereas in children only 1(3%) bowel wall edema was reported. Others are normal. Throat swab was found to be positive only in 2(6%) of adults whereas no throat swab was found to be positive in children.

**Table 6: Recurrence**

Variables	Adult (N=34)	Children (N=31)	P value
Recurrence			
Yes	12(35%)	3(10%)	0.03*
No	22(65%)	28(90%)	

Recurrence was found to be more in adult 12(35%) compared to children 3(10%) and the difference was found to be statistically significant.

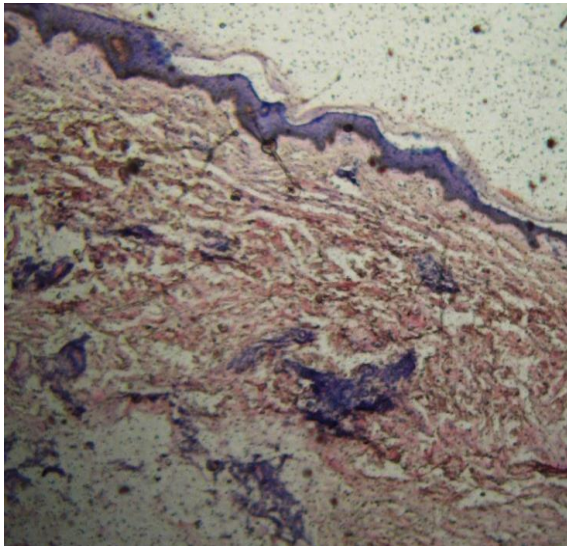
**Figure 1: Type of skin lesions**

In 30(96%) of the study participants present with palpable purpura whereas in adults only 25(73%) were with palpable purpura followed by echymotic patches 4(12%) followed by ulcers 4(12%).

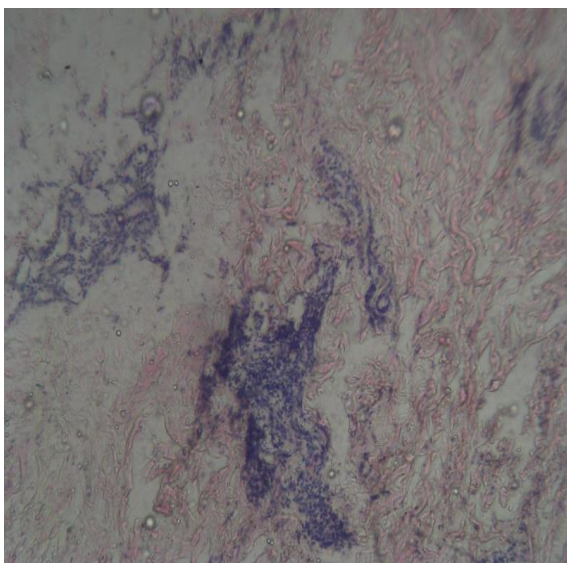
**A) Palpable purpura in adults****B) Palpable purpura in children****C) Koebernisation in adults**



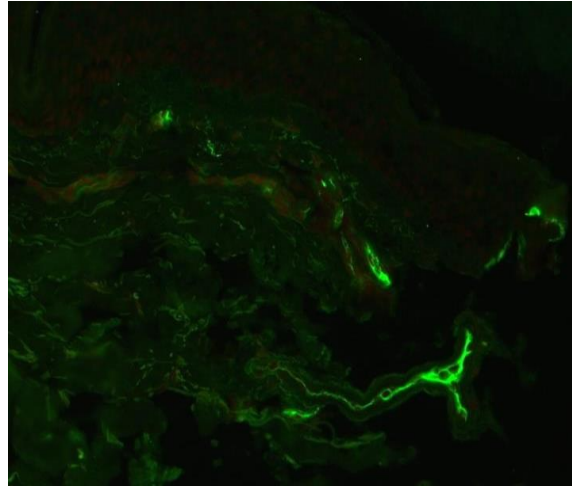
**D) Bullous lesion in adults**



**E) Low power view showing epidermis with Perivascular infiltrate**



**F) High power view shows leucocytoclastic vasculitis**



**G) DIF showing blood vessel vessel wall staining with granular IgA deposits**

## DISCUSSION

In our study majority of the study participants were in 19-40 years 21(62%) of age in adults and less than 10 years (61%) in children. Similar results were also seen in uppal et al<sup>[15]</sup> study. Female preponderance were more in our study. Disease manifestation is observed more in summer for adults and winter for children. For majority of the diseases the etiology was unknown followed by URI. These results were similar to the Carlos Garcia et al study.<sup>[16]</sup> This is due to the reason of upper respiratory infection is common in children during winter and it is twice common in children compared to adults. For majority of the diseases the etiology was unknown followed by URI. Kang et al<sup>[17]</sup> also shown similar results.

In both the groups purpura was seen in all patients in the lowest extremities. Arthralgia was present in adults than in children. Joint pain was more in adults 79%. Ankle joints swelling seen in adults 11%. Gracia et al study showed similar results. Gastrointestinal symptoms was most common in paediatric group. Similar results was also seen in Y et al study.<sup>[18]</sup> Renal involvement is noted in 20% of the adult patients with elevated renal parameters whereas in children all patients have normal renal function. Hung et al<sup>[19]</sup> showed similar results.

During the disease course three children relapsed and 12 adult patients relapse. This results was similar to previous studies.

## CONCLUSION

Our study concluded stating that IgA vasculitis in adults is not uncommon and it is frequent as in children. Hence IgA vasculitis should be considered as one of the differential diagnosis for adults with vasculitis. When compared with the children the duration and the severity was more in adults. Longer follow up has to be done in adults with elevated

renal parameters to uncover the delayed systemic complications.

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#### **Competing Interest**

There is no competing interest

#### **Authors Contribution**

All authors in our study contributed to the data collection of the patients

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## **REFERENCES**

1. Abushaiwia A.M, Rhuma N.R, Zletni M.A, Ebtisam S.Sm Gr up Clinico Epidemiological profile of research children with Henoch Schonlein Purpura at Tripoli Children's Hospital.SM J.Arthritis.2018;2:1006
2. Lardhi A.A. Henoch-Schonlein Purpura in children from the eastern province of Saudi Arabia.Saudi Med.J.2012;33:973-978
3. Chen J-Y, Mao J. Henoch schonlein purpura nephritis in children: Incidence ,pathogenesis and management.World J Pediatrics.2014;11:29-34
4. Asad. S, Smith AG, Cutaneous vasculitis:a retrospective study. J Am Acad Dermatology.2004;50(3)(Suppl):113
5. Florentino DF:Cutaneous vasculitis.J Am Acad Dermatol.2003;48(3):311-40
6. Trapani S, Micheli A, Grisolia F, Resti M, Chiappini E, Falcini F, De Martino M. Henoch Schonlein purpura in childhood: Epidemiological and clinical analysis of 150 cases over a 5 year period and review of literature. Semin. Arthritis Rheum.2005;35:143-153
7. Hocevar A, Rotar Z, Ostrovrsnik J. Incidence of IgA vasculitis in the adults Slovenian Population.Br J Dermatol .2014;171:524-527
8. Nossent J, Raymond W, Keen H, Inderjeeth C, Preen D.B.Hospitalisation rates and characteristic for adult and childhood immunoglobulin A Vasculitis in Western Australia.Intern Med J.2019;49:475-481
9. Pillebout E.A.V.Alexandra.Efficacy of cochlincine to prevent skin relapses in adult IgA vasculitis.National library of Medicine.2019
10. Calvo-Rio.V.Loricera J,Mata C.Henoch-Schonlein Purpura in northern spain:Clinical spectrum of the disease in 417 patients from single center.Medicine (Baltimore).2014;93(2):106-113
11. Kang Y, Park J.S, Ha YJ.Differences in clinical manifestations and outcomes between adults and child patients with Henoch-Schonlein Purpura.J Korean Med Sci:2014;29(2):198-203
12. Pillebout E.A.V,Alexandra.National library of Medicine.:Efficacy of cochlincine to prevent the relapses in adult IgA vasculitis.2019
13. Hitomi K,Izaki S,Teraki Y.Characterization of adult type IgA vasculitis :A retrospective study of 122 cases.Open Dermatol J.2014;8:51-59.
14. Nossent J, Raymond W, Keen H.I, Preen D, Inderjeeth C.Morbidity and mortality in adult onset IgA vasculitis:a long term population based cohort study.Rheumatology.2021;61(1):291-298
15. Uppal SS,Hussain MA,Al-RAqum HA,Nampoory MR,Al-Saeid K,AlAssousi A,Abraham M,Malaviya AN.Henoch-Schonlein's purpura in adult versus children/adolescents:a comparative study.Clin Exp Rheumatol 2006;24:S26-30
16. Carlos Garci a Porru a, Maria C, Calvin O, Javier Llorca Jose M, Couselo and Miguel A, Gonza lez-Gay.Henoch-schonlein purpura in children and adults:Clinical differences in a defined population
17. Kang Y, Park JS, Ha YJ, Kang MI, Park HJ, Lee SK, Park YB.Differences in clinical manifestations and outcomes between adults and child patients with Henoch-Schonlein purpura.J Korean Med Sci.2014;29(2):198-203
18. Ilan Y, Naparstek Y.Schonlein-Henoch Syndrome in adults and children.Semin Arthritis Rheum.1991;21(2):103-9
19. Hung SP, Yang YH, Lin YT, Wang LC, Lee JH, Chiang BL. Clinical manifestations and outcomes of Henoch schonlein purpura:comparison between adultls and children.Pediatr Neonatol.2009;50(4):162-8.