

Comparison of Thymus size in normal versus malnourished children

By Jaishree Vasudevan

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Revathi Muthu, Jaishree Vasudevan, Jeffrey Skaria Joseph

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Department of Radiology, Chettinad Hospital & Research Institute (Chettinad academy of research and institute), Kelambakkam, Chengalpattu district, Chennai, Tamil Nadu, India

Corresponding author

Jaishree Vasudevan

vsvdnjshr@gmail.com

ABSTRACT

Background. Undernutrition has a detrimental effect on lymphoid organs like thymus that are responsible for production of T cells. Severe thymus gland atrophy has been observed in children with severe acute malnutrition. Hence, this study was aimed to measure thymus size in children with normal, moderate, and severe acute malnutrition who were hospitalized to the pediatric ward.

Methodology. A prospective observational study was conducted in pediatric ward with 60 children aged between 6 to 59 months. Based on the inclusion and exclusion criteria 20 children each were randomly selected as normal, moderate acute malnourished child and severe acute malnourished child. Anthropometric and thymic measurements were taken. Data was analyzed in SPSS v 29. ANOVA test and Post hoc tukey HSD were done to find the difference.

Results. Thymic index was also high among normal children when compared to malnourished children. This infers statistically significant thymus atrophy among the malnourished when compared to healthy children. Pairwise comparison of means using post-hoc tukey HSD revealed significant difference between normal children and those children with malnourishment ($p < 0.001$)

Conclusion. The present investigation concluded that malnourished patients have thymic atrophy and recurrent infections. Hence thymic measurement can be used as a tool to detect immunological modifications in children with malnourishment.

Keywords: Malnutrition, thymic index, immunity, children, immunodeficiency, thymus

INTRODUCTION

A nutritional deficit brought on by an insufficient intake of calories or protein is known as malnutrition [1]. The American Society of Parenteral and Enteral Nutrition (ASPEN) defines pediatric malnutrition as "an imbalance between nutrient requirement and intake, resulting in cumulative deficits of energy, protein, or micronutrients that may negatively affect growth, development, and other relevant outcomes [2]."

In 2022, it was anticipated that 37 million children under the age of five were overweight or obese, 149 million were stunted and 45 million were wasted on a global scale [3]. According to NHFS-5 (India 2019–21), malnourishment (wasting, stunting and underweight) in children under the age of five has decreased from 21% to 19.3%, 38.4% to 35.5% and 35.8% to 32.1%, respectively, in comparison to NHFS-4 (India 2015–16) [4].

Undernutrition is a contributing factor in over half of fatalities in children under the age of five in low- and middle-income nations. Notably, India bears one-third of the worldwide burden of undernutrition [5].

Undernourished children are more susceptible to acute infections because undernutrition weakens the immune system and causes varying degrees of immunodeficiency. Undernutrition has a detrimental effect on the primary and secondary lymphoid organs that are responsible for responding to associated infections through immunosuppression. One such organ affected is the thymus, the main lymphoid organ that produces T cells [6].

The thymus is a retrosternal organ located in the superior mediastinum. It is bilobed with two components namely the cortex and medulla [7]. Thymopoiesis is a lifelong process that slows down with age as aging causes the thymus to shrink and lose some of its functions. However, the proliferation of intrathymic T cells is a continuous activity, hence the cessation of a

particular harmful stimulation (malnourishment) leads way to restore the function back to normal steady state [8].

Severe thymus gland atrophy has been observed in children with severe acute malnutrition [9]. According to studies by Garly et al patients with more severe atrophy may also have more severe immunological impairment [10]. This finding thus helps the clinicians to identify the underlying unidentified immunological impairment. Hence, this study was aimed to measure thymus size in children with normal, moderate, and severe acute malnutrition who were hospitalized to the pediatric ward at Chettinad Hospital and Research Institute.

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MATERIALS AND METHODS

The is a cross sectional study and was conducted between June 2023 and June 2024 in a tertiary care hospital, Chengalpattu. The Institutional Ethics Review Board authorized the study procedure, which was carried out in accordance with good clinical practice. The study population comprises of normal children and those children with moderate and severe acute malnutrition aged between 6 to 59 months admitted in pediatric ward. A total of 60 such children (20 normal children, 20 moderate acute malnourished and 20 severe acute malnourished children) were enrolled after obtaining informed written consent from their parents. Moderate acute malnutrition (MAM), also known as wasting, is defined by a weight-for-height indicator between -3 and -2 z-scores (standard deviations) of the international standard or by a mid-upper arm circumference (MUAC) between 11.5 cm and 12.5 cm [11]. Severe Acute Malnutrition (SAM) was defined as mid-upper arm circumference (MUAC) < 11.5 cm or weight for height Z score (WHZ) \leq -3 SD or bipedal pitting edema [12]. We also include apparently healthy children but whose WHZ was between 2 to -2 z score were used as the reference thymus size for non-malnourished children in the same setting. Using convenient sampling method those fulfilling the eligibility criteria were included in the study.

Data of previous medical history, demographic profile was collected using a questionnaire. Anthropometric measurements like Length or height will be measured using an infant length board and MUAC using a measuring tape, both to the nearest 1 mm. Body weight was measured using a digital weighing scale to the nearest 0.1 kg. The WHZ were calculated using the Child Growth Standards of the World Health Organization (WHO) [13]. Thymus size was measured by the radiologist using ultrasonography. The procedure was done with the child lying on the back or in the mother's lap. The area of thymus was visualized as a homogenous opacity by

placing the transducer at 90 degrees to the upper sternum. The circumference of the largest lobe of the thymus was noted and the size measurement was taken. Thymic index (multiplying the transverse diameter by the sagittal area) was calculated (Figure 1,2).

9 Statistical analysis

The data collected were entered in Microsoft Excel spreadsheet and analyzed using IBM SPSS v 29. The descriptive statistics of categorical variables were expressed in terms of frequency and percentages, for continuous variables it is expressed as mean and standard deviation. To find the significant difference in the multivariate analysis the one - way ANOVA with Tukey's Post-Hoc test was used and p value < 0.05 was considered statistically significant.

Figure 1 shows the ultrasonographic measurement of thymus length and width

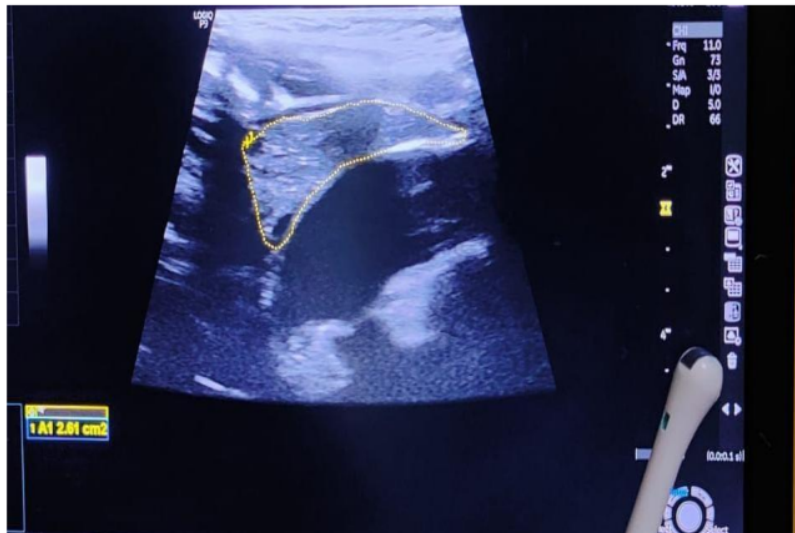
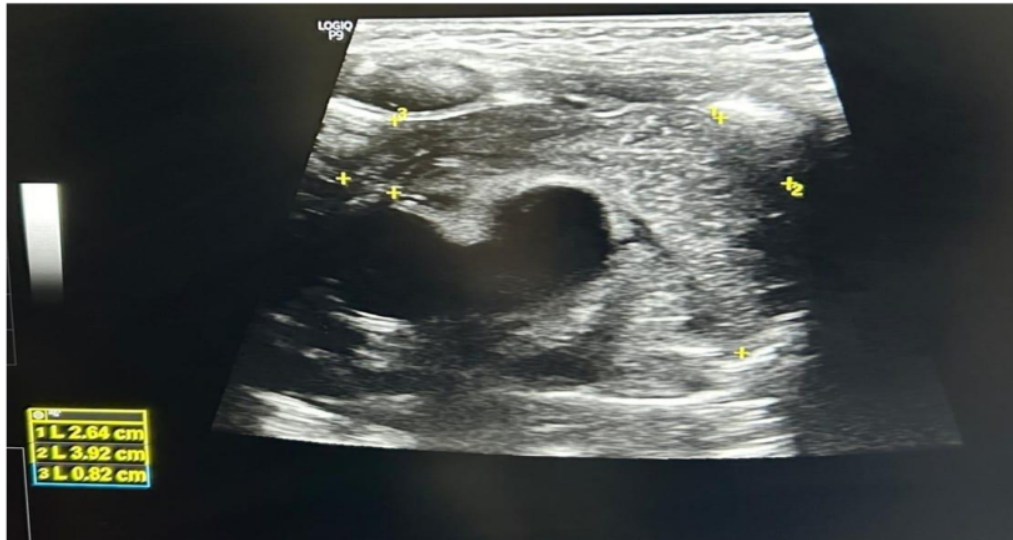


Figure 2 shows ultrasonographic measurement of thymic area



RESULTS

The mean age of the study participants is 26.2 ± 15.9 months. Majority of those participated were between the age of 1 to 3 years. It was noted that children with malnourishment had increased history of recurrent infection. The mean MUAC was found to be 13.4 ± 1.5 cm (Table 1,2).

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Table 1: Demographic characteristics of the participants

Characteristics	Number	Percentage
Age		
Upto 1 yr	18	30.0
1 - 3 yrs	24 12	40.0
3 - 5 yrs	18	30.0
Gender		
Male	31	51.7
Female	29	48.3
Recurrent infection		
No	45	75.0
Yes	15	25.0

Table 2: Anthropometric details of the participants

Characteristics	Mean ± SD
Weight	9.8 ± 2.9
Height	83.5 ± 13.4
Mid Upper Arm Circumference	13.4 ± 1.5
Head circumference	45.6 ± 3

Table 3 depicts the thymic measurements recorded among the study participants. After adjusting for age, the mean thymus gland size was found to be $3.7 \pm 1.3 \text{ cm}^2$. The mean thymic index was found to be 13.7 ± 7.1 .

Table 3: Thymus measurements of the participants

Characteristics	Mean ± SD
Thymus length	2.9 ± 0.8
Width	1.9 ± 0.9
Area	3.7 ± 1.3
Thymic index	13.7 ± 7.1

Table 4: Association between malnutrition and thymus size

Characteristics	Normal	MAM	SAM	P value
Thymus length	3.76 ± 0.36	2.55 ± 0.69	2.53 ± 0.60	0.0005
Width	2.69 ± 0.69	1.8 ± 0.78	1.33 ± 0.79	0.0005
Area	4.6 ± 0.73	3.37 ± 1.10	3.17 ± 1.41	0.0005
Thymic index	19.51 ± 4.72	11.24 ± 5.74	10.41 ± 6.9	0.0005

*ANOVA test, $p < 0.05$ is significant

1 The mean thymus gland size among the healthy children was $4.6 \pm 0.73 \text{ cm}^2$ when compared to $3.37 \pm 1.10 \text{ cm}^2$ in MAM and $3.17 \pm 1.41 \text{ cm}^2$ in SAM children. Similarly, Thymic index was also high among normal children when compared to malnourished children. This infers statistically significant thymus atrophy among the malnourished when compared to healthy

children (Table 4). Pairwise comparison of means using post-hoc tukey HSD revealed significant difference between normal children and those children with malnourishment ($p < 0.001$)

DISCUSSION

Several postmortem studies and ultrasound findings have documented thymus atrophy in malnourished children [14]. The present study showed thymic index of $10.41 \pm 6.9 \text{ cm}^3$ whereas it was $19.51 \pm 4.72 \text{ cm}^3$ among the normal children. The mean thymus gland size among the healthy children was $4.6 \pm 0.73 \text{ cm}^2$ when compared to $3.37 \pm 1.10 \text{ cm}^2$ in MAM and $3.17 \pm 1.41 \text{ cm}^2$ in SAM children. Also, history of recurrent infection was higher among the SAM children when compared to normal children. Similar results with decreased thymus size among malnourished were reported in studies by Nabukeera B et al [15], Nassar MF et al [16], Chevalier et al [17], Rytter MUH et al [18] and Parent G et al [19].

It has been suggested that thymus atrophy in malnourished children may represent the immunological inadequacy brought on by malnutrition, making them more vulnerable to infections. Children with a small thymus had a greater mortality risk in community-based research [20]. This suggests that thymus size could be a marker of competent immune system, or simply an indicator of good health thereby validating the concept of the thymus as a "barometer of malnutrition" [21].

However, the process of thymus atrophy in malnutrition is unclear. Hormonal factors appear to have a role, according to animal research. A number of hormones, controlled by insulin-like growth factor 1 include prolactin, leptin and growth hormone. These hormones are known to control the growth and functioning of the thymus according to animal research [22]. Additionally, thymic atrophy may be exacerbated by zinc insufficiency, which is common in malnutrition [23]. Thymic atrophy has also been observed to be exacerbated by infections, albeit it is unclear if this is the result of an infection or its cause [24].

The limitation of the study was i) as it is a cross-sectional study the role of causality cannot be demonstrated. Hence a prospective study is recommended to know the causality. ii) small sample size makes it difficult to generalize the study results to community settings.

CONCLUSION

The present investigation concluded that malnourished patients have thymic atrophy. Malnourished child's immune systems will probably be impacted by these alterations, which could be harmful given their early age. Though they could take longer to resolve than physical recovery, immunological modifications are fortunately reversible with nutritional therapy. Hence thymic measurement can be used as a tool to detect immunological modifications in children with malnourishment. As a conclusion, we advise a thorough evaluation of malnourished child's immune system through thymic parameters both during and after nutritional rehabilitation, until they completely recover.

REFERENCES

1. Dipasquale V, Cucinotta U, Romano C. Acute Malnutrition in Children: Pathophysiology, Clinical Effects and Treatment. *Nutrients*. 2020 Aug 12;12(8):2413. doi:10.3390/nu12082413. <https://pubmed.ncbi.nlm.nih.gov/32806622/>
2. Mehta, N.M.; Corkins, M.R.; Lyman, B.; Malone, A.; Goday, P.S.; Carney, L.N.; Monczka, J.L.; Plogsted, S.W.; Schwenk, W.F. Defining pediatric malnutrition: A paradigm shift toward etiology-related definitions. *JPEN J. Parenter. Enteral. Nutr.* 2013, 37, 460–481. doi:10.1177/0148607113479972. Epub 2013 Mar 25. <https://pubmed.ncbi.nlm.nih.gov/23528324/>
3. World Health Organization. Fact sheets - Malnutrition [Internet]. [cited 2024 Jul 30]. Available from: <https://www.who.int/news-room/fact-sheets/detail/malnutrition>
4. Malnutrition-Free India [Internet]. [cited 2024 Jul 30]. Available from: <https://pib.gov.in/pib.gov.in/Pressreleaseshare.aspx?PRID=1781673>
5. World Health Organization. Nutrition [Internet]. [cited 2024 Jul 30]. Available from: <https://www.who.int/india/health-topics/nutrition>
6. Keusch GT. Malnutrition and the Thymus Gland. In: *Nutrient Modulation of the ImmuneResponse*. CRC Press; 1992. <https://www.taylorfrancis.com/chapters/edit/10.1201/9781003066644-21/malnutrition-thymus-gland-gerald-keusch>
7. Remien K, Jan A. Anatomy, Head and Neck, Thymus. In: *StatPearls* [Internet]. Treasure Island (FL): StatPearls Publishing; 2024 [cited 2024 Jul 30]. Available from: <http://www.ncbi.nlm.nih.gov/books/NBK539748/>

8. Thapa P, Farber DL. The Role of the Thymus in the Immune Response. *Thoracic Surgery Clinics*. 2019 May 1;29(2):123–31. doi: [10.1016/j.thorsurg.2018.12.001](https://doi.org/10.1016/j.thorsurg.2018.12.001) .
<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6446584/>
9. Savino W, Durães J, Maldonado-Galdeano C, Perdigon G, Mendes-da-Cruz DA, Cuervo P. Thymus, undernutrition, and infection: Approaching cellular and molecular interactions. *Front Nutr*. 2022;9. <https://doi.org/10.3389/fnut.2022.948488> .
<https://www.frontiersin.org/journals/nutrition/articles/10.3389/fnut.2022.948488/full>
10. Garly M-L, Trautner SL, Marx C, Danebod K, Nielsen J, Ravn H, et al. Thymus size at 6 months of age and subsequent child mortality. *J Pediatr*. 2008;153: 683–8, 688–3. doi:10.1016/j.jpeds.2008.04.069. Epub2008Jun27.<https://pubmed.ncbi.nlm.nih.gov/18589444/>
11. World Health Organization. “Supplementary Foods for the Management of Moderate Acute Malnutrition in Infants and Children 6–59 Months of Age.” Technical Note, WHO,Geneva;2014.DOI: 10.1596/978-1-4648-0348-2_ch11
<https://www.ncbi.nlm.nih.gov/books/NBK361900/>
12. Guideline: Updates on the Management of Severe Acute Malnutrition in Infants and Children.Geneva:WorldHealthOrganization;2013. <https://www.who.int/publications/i/item/9789241506328>
13. WHO Multicentre Growth Reference Study Group. WHO Child Growth Standard based on length/height, weight and age. *Acta Paediatr Suppl*. 2006;450:76–85.
14. Mugerwa JW. The lymphoreticular system in Kwashiorkor. *J Pathol*. 1971;105:105–9. DOI: [10.1002/path.1711050204](https://doi.org/10.1002/path.1711050204) . <https://pubmed.ncbi.nlm.nih.gov/4109254/>
15. Nabukeera-Barungi N, Lanyero B, Grenov B, Friis H, Namusoke H, Mupere E, et al. Thymus size and its correlates among children admitted with severe acute malnutrition: a cross-sectional study in Uganda. *BMC Pediatr*. 2021;21(1):1. doi: [10.1186/s12887-020-02457-3](https://doi.org/10.1186/s12887-020-02457-3). <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7780382/>
16. Nassar MF, Younis NT, Tohamy AG, Dalam DM, El Badawy MA. T-lymphocyte subsets and thymic size in malnourished infants in Egypt: a hospital-based study. *East Mediterr Health J Rev Santé Méditerranée Orient Al-Majallah Alihḥīyah Li-Sharq Al-Mutawassit*.2007;13(5):1031–42.DOI: [10.26719/2007.13.5.1031](https://doi.org/10.26719/2007.13.5.1031) .
<https://pubmed.ncbi.nlm.nih.gov/18290395/>
17. Chevalier P, Sevilla R, Sejas E, Zalles L, Belmonte G, Parent G. Immune recovery of malnourished children takes longer than nutritional recovery: implications for treatment

- anddischarge.1998;44(5):304–7.DOI: [10.1093/tropej/44.5.304](https://doi.org/10.1093/tropej/44.5.304) .
<https://pubmed.ncbi.nlm.nih.gov/9819496/>
18. Rytter MJH, Namusoke H, Ritz C, Michaelsen KF, Briend A, Friis H, et al. Correlates of thymus size and changes during treatment of children with severe acute malnutrition: a cohort study. *BMC Pediatr.* 2017;14(1): 70. 17. doi: [10.1186/s12887-017-0821-0](https://doi.org/10.1186/s12887-017-0821-0) .
<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5348758/>
19. Parent G, Chevalier P, Zalles L, et al. In vitro lymphocyte-differentiating effects of thymulin (Zn-FTS) on lymphocyte subpopulations of severely malnourished children. *AmJClinNutr.*1994;60(2):274–8.DOI: [10.1093/ajcn/60.2.274](https://doi.org/10.1093/ajcn/60.2.274) .
<https://pubmed.ncbi.nlm.nih.gov/8030607/>
20. Garly M-L, Trautner SL, Marx C, Danebod K, Nielsen J, Ravn H, et al. Thymus size at 6 months of age and subsequent child mortality. *J Pediatr.* 2008;153: 683–8, 688–3. DOI: [10.1016/j.jpeds.2008.04.069](https://doi.org/10.1016/j.jpeds.2008.04.069) . <https://pubmed.ncbi.nlm.nih.gov/18589444/>
21. Prentice AM. The thymus: a barometer of malnutrition. *Br J Nutr.* 1999;81:345–7. <https://pubmed.ncbi.nlm.nih.gov/10615206/>
22. Bartz S, Mody A, Hornik C, Bain J, Muehlbauer M, Kiyimba T, et al. Severe Acute Malnutrition in Childhood: Hormonal and Metabolic Status at Presentation, Response to Treatment, and Predictors of Mortality. *J Clin Endocrinol Metab.* 2014;99(6):2128–37. DOI: [10.1210/jc.2013-4018](https://doi.org/10.1210/jc.2013-4018) . <https://pubmed.ncbi.nlm.nih.gov/24606092/>
23. Rytter MJH, Kolte L, Briend A, Friis H, Christensen VB. The immune system in children with malnutrition—a systematic review. *PloS One.* 2014;9(8):e105017. DOI: [10.1371/journal.pone.0105017](https://doi.org/10.1371/journal.pone.0105017) . <https://pubmed.ncbi.nlm.nih.gov/25153531/>
24. Savino W, Dardenne M. Nutritional imbalances and infections affect the thymus: consequences on T-cell-mediated immune responses. *Proc Nutr Soc.* 2010;69(4):636–43. DOI: [10.1017/S0029665110002545](https://doi.org/10.1017/S0029665110002545) . <https://pubmed.ncbi.nlm.nih.gov/20860857/>