

Morphea in children: a retrospective study of its clinical characteristics and extracutaneous manifestations

Morfea en niños: un estudio retrospectivo de características clínicas y manifestaciones extracutáneas

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Abstract

BACKGROUND: Morphea is an inflammatory disease with diffuse thickening and hardening of the skin. It is common in children and it has been reported that as many as 40% have extracutaneous articular and neurological manifestations, vascular and ocular disorders, and gastrointestinal or respiratory symptoms.

RATIONALE: There are no studies of associated extracutaneous manifestations in Mexican children with morphea.

OBJECTIVE: Describe clinical characteristics, frequency of extracutaneous manifestations, and treatment of pediatric patients with morphea who received care at the Instituto Nacional de Pediatría.

MATERIAL AND METHODS: A retrospective study of patients with morphea, 1 to 18 years of age, who received care between January 1, 2000, and December 31, 2015.

RESULTS: Thirty-two patients with morphea were included, half (n = 16) with the linear subtype, followed by plaque-type morphea in 7 (22%). Extracutaneous manifestations were reported in 23 (72%) patients; the most common were: musculoskeletal in 23 (72%), neurological in 9 (28%), and ophthalmological in 3 (9%). Twenty-three patients (72%) required systemic treatment. Nineteen patients (60%) required orthopedic and surgical treatment for complications. The clinical evolution was toward remission in 72% of the patients (n = 23).

CONCLUSIONS: In this series of 32 cases of children with morphea, 72% had extracutaneous manifestations, principally musculoskeletal and neurological. It is important to conduct targeted questioning, and if necessary perform paraclinical studies, to identify extracutaneous manifestations which may be associated with morphea, especially when it is linear or extensive.

KEY WORDS: Morphea; children; skin; ocular; Mexican; pediatric; linear morphea; ophthalmological.

Resumen

ANTECEDENTES: La morfea es una enfermedad inflamatoria con engrosamiento difuso e induración de la piel. Es frecuente en niños y se ha reportado que incluso 40% tiene manifestaciones extracutáneas articulares, neurológicas, anomalías vasculares, afectación ocular, síntomas gastrointestinales o respiratorios.

JUSTIFICACIÓN: No existen estudios de las manifestaciones extracutáneas asociadas en niños mexicanos con morfea.

OBJETIVO: Describir las características clínicas, la frecuencia de las manifestaciones extracutáneas y el tratamiento de pacientes pediátricos con morfea atendidos en el Instituto Nacional de Pediatría.

MATERIAL Y MÉTODOS: Estudio retrospectivo de pacientes con morfea, de 1 a 18 años, atendidos entre el 1 de enero de 2000 y el 31 de diciembre del 2015.

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RESULTADOS: Se incluyeron 32 pacientes con morfea, la mitad (n = 16) con el subtipo lineal, seguido de placas en 7 (22%). En 23 (72%) pacientes se reportaron manifestaciones extracutáneas, las más comunes fueron: musculoesqueléticas en 23 (72%), neurológicas en 9 (28%) y oftalmológicas en 3 (9%). Se requirió tratamiento sistémico en 23 pacientes (72%). Diecinueve pacientes (60%) requirieron tratamiento ortopédico y quirúrgico, por complicaciones. La evolución clínica fue hacia la remisión en 72% de los pacientes (n = 23).

CONCLUSIONES: En esta serie de 32 casos de niños con morfea, 72% tuvo manifestaciones extracutáneas, sobre todo musculoesqueléticas y neurológicas. Es importante efectuar un interrogatorio dirigido, y si es necesario, estudios paraclínicos para identificar las manifestaciones extracutáneas que pueden asociarse con morfea, especialmente cuando ésta es lineal o extensa.

PALABRAS CLAVE: Morphea; niños; piel; ocular; mexicanos; pediátricos; morfea lineal; oftalmológica.

BACKGROUND

Morphea, also called localized scleroderma, is a chronic autoimmune disease, of unknown cause, which is clinically defined as an inflammatory disease, primarily of the skin and subcutaneous tissue, in which patients develop sclerosis with diffuse thickening and hardening of the skin.¹⁻⁵

Figure 1

Around 20% of patients with morphea are children or adolescents, with a mean age of onset at seven years. Linear morphea affects two thirds of patients under 18 years of age; this variant may cause retarded growth of the affected extremity, making proper treatment decisive to prevent asymmetry or permanent functional disability of the extremity.^{2,3,6}

Approximately 40% of children with morphea present extracutaneous manifestations: arthritis, neurological symptoms (headache, seizures), vascular disorders (Raynaud's phenomenon), ocular disorders (uveitis or episcleritis, mainly of the linear variety, in the head or *en coup de sabre*) and gastrointestinal and respiratory symptoms. The association between location or type of morphea lesions in the skin and the risk of extracutaneous manifestations is unknown, as are risk factors which predispose an individual to its appearance.⁷



Figure 1. Patient with linear morphea in the lower right extremity, with visible sclerosis, changes of pigmentation, and mild atrophy.

The activity of morphea lesions lasts three to five years, with a tendency for improvement, although with residual alterations. In general, more serious, mutilating, and disabling forms are observed in children than in adults.^{1,2,8,9}

The objective of this study was to describe the clinical characteristics, frequency of extracutaneous manifestations, and treatment of pediatric patients with morphea who received care at the National Institute of Pediatrics.

MATERIALS AND METHODS

This was a retrospective study of patients with morphea, 1 to 18 years of age, who received care at the National Institute of Pediatrics between January 1, 2000, and December 31, 2015. The search was conducted in the institution's archive of physical and electronic records. Criteria for inclusion were: patients with diagnosis of morphea between 1 and 18 years of age, who received care between January 1, 2000, and December 31, 2015, for whom clinical photographs of lesions were available. Criteria for exclusion were: incomplete records (less than 80% of the variables to be studied), patients with diagnosis of linear pansclerotic morphea, eosinophilic fasciitis, or systemic sclerosis.

The descriptive statistical analysis was performed in Microsoft Office Excel, version 2017, with measurements of central tendency and dispersion for the quantitative variables and ratios for the qualitative variables.

RESULTS

Fifty-nine patient files were located and 27 were discarded because they did not meet the criteria for inclusion. Thus, 32 patients with the diagnosis of morphea were included; females predominated, at 69% (n = 22). The linear morphea subtype was the most common, at 50% (n = 16) (Figure 1), followed by plaque morphea at 22% (n = 7) (Figure 2). The median age at initial manifestations was 9 years, with a range of 1 to 16 years. Morphea lesions had characteristics of lichen sclerosus and atrophicus in 41% (n = 16) of the cases; deep in 25% (n = 8), and with atrophoderma in 22% (n = 7). Table 1

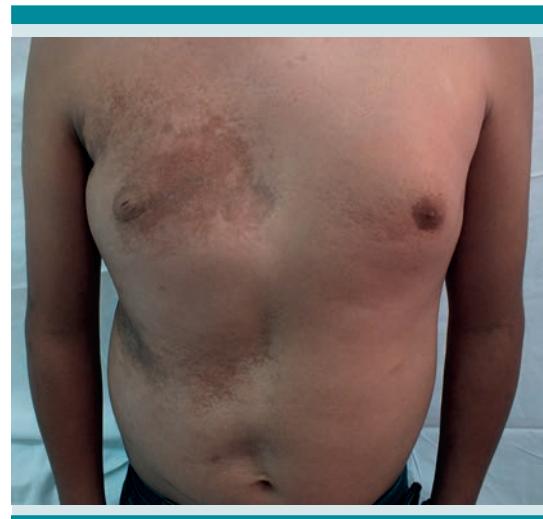


Figure 2. Patient with plaque morphea in the anterior torso, with irregular shaped plaques with dermal atrophy and visible blood vessels.

Table 1. Clinical characteristics of all patients

Characteristic	n = 32
Age of onset of morphea (median, range)	9 (1-16 years)
Gender n (%)	
Female	22 (68.75%)
Male	10 (31.25%)
Morphea subtype n (%)	
Linear	16 (50%)
Plaque	7 (21.9%)
Generalized	5 (15.6%)
Mixed	4 (12.5%)
Additional characteristics of morphea n (%)	
Lichen sclerosus	13 (40.6%)
Deep	8 (25%)
Atrophoderma	7 (21.9%)
None of the above	10 (31.2%)
Clinical evolution n (%)	
Remission	23 (71.87%)
Activity	8 (12.5%)

The diagnosis was clinical in 69% of patients (n = 22) and skin biopsies were taken from 32% (n = 10). The histological findings reported for morphological changes compatible with morphea were: hyalinization of collagen, collagen entrapment of skin appendages, and superficial and deep inflammatory perivascular infiltrate which may affect the subcutaneous cellular tissue. **Figure 3**

Extracutaneous manifestations were reported in 84% (n = 27) of patients. The most common were musculoskeletal, with arthralgias predominating

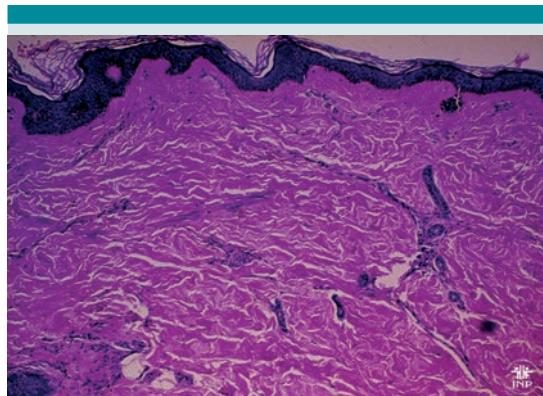


Figure 3. Histopathology of morphea showing thickening and hyalinization of collagen in the dermis, as well as decreased cutaneous appendages.



Figure 4. Sclerotic morphea plaque on the right foot, with reduction in length and joint contractures and loss of mobility in the first and fourth toes.

(34%), and joint contractures (38%) (**Figure 4**). Only nine patients (28.1%) did not report musculoskeletal manifestations. Twenty-eight percent (n = 9) of patients had neurological manifestations; the most common symptom was headache in 22% (n = 7). There was no report of targeted questioning on symptoms for 6% of the patients and related paraclinical studies were not conducted for 81%. Six patients underwent neurological studies. One patient with linear morphea in the upper left extremity, with seizures underwent: nuclear magnetic resonance (NMR), computed tomography (CT) of the skull, and electroencephalogram (EEG), documenting mild cortical atrophy and epileptic activity. Another patient with linear morphea in the hemiface suffered headaches and underwent a simple CT scan which found osseous atrophy. In most patients there were no reports of ophthalmological manifestations in notes to the clinical file (78%), only three patients had clinical alterations: hemihypertrophy, strabismus, exotropia, and keratitis. **Table 2**

Regarding treatment, the most frequently indicated topical medications were vitamin D analogs, alone or combined with steroids in 22 (69%); and calcineurin inhibitors in 13 patients (41%) with circumscribed morphea. In 23 patients (64%) with the linear subtype or extensive morphea the most commonly prescribed systemic treatment were systemic steroids as bridge or adjuvant therapy while starting on another drug. The most commonly prescribed immunosuppressants were: methotrexate in 20 (62.5%) of patients, followed by hydroxychloroquine and mycophenolate mofetil. **Table 3**

As disease activity diminished, the treatment was modified. In the last follow up visit, 12 patients (38%) continued to use 20% urea in cream to reduce residual xerosis, 6 patients (19%) continued receiving methotrexate, and 5 (16%) hydroxychloroquine as maintenance therapy.

Table 2. Extracutaneous manifestations in patients with morphea

Manifestations and studies	n = 32
Musculoskeletal n (%)	
Joint contractures	12 (37.50%)
Arthralgias	11 (34.4%)
Discrepancy in length of extremities	11 (34.4%)
Arthritis	1 (3.12%)
Others	7 (21.87%)
None	9 (28.12%)
Unknown	1 (3.12%)
Neurological symptoms n (%)	
Headache	7 (21.87%)
Migraine	1 (3.12%)
Seizures	2 (6.5%)
None	21 (65.62%)
Unknown	2 (6.25%)
Paraclinical neurological tests n (%)	
CT	4 (12.5%)
EEG	3 (9.37%)
NMR	1 (3.12%)
None	26 (81.25%)
Results of paraclinical neurological tests n (%)	
Normal	1 (16.6%)
Abnormalities incidental or not related to morphea*	4 (66.7%)
Abnormalities associated with morphea**	1 (16.6%)
Ocular manifestations n (%)	
Uveitis	0
Others (hemihypertrophy, strabismus, keratitis)	3 (9.37%)
None	25 (78.12%)
Unknown	4 (12.5%)

* A patient with morphea in the upper left extremity also had seizures, and studies (NMR, CT, and EEG) found mild cortical atrophy and epileptogenic rhythm.

** A patient with linear morphea in the hemiface had headaches, and a simple CT scan revealed atrophy of the facial bones on the affected side.

Regarding non-pharmacological treatments, 11 patients (34%) received rehabilitation therapy and 8 patients (25%) surgical treatment by orthopedics (tibia osteotomy with bone lengthening, medial hemiepiphiodesis of the right knee, plantar capsulotomy, and percutaneous tenotomy of the flexor) and plastic surgery (resection of sclerotic lesion in hairy skin, resection of fatty tissue in the lower lip, implant in the piriform fossa, and multiple lipotransfers). **Table 3**

Table 3. Treatments received by patients

Treatment	n = 32
Topical treatment n (%)	
Vitamin D analogs	22 (68.75%)
High-potency topical steroids (class I/II)	15 (46.87%)
Calcineurin inhibitors	13 (40.62%)
Low-potency topical steroids (class III/VII)	3 (9.37%)
Others	
Pirfenidone	3 (13.64%)
None	1 (3.12%)
Systemic treatment n (%)	
Systemic steroids (as bridge-adjuvant therapy while starting on another systemic treatment)	23 (71.9%)
Methotrexate	20 (62.5%)
Hydroxychloroquine	11 (34.4%)
Mycophenolate mofetil	7 (21.87%)
Oral vitamin D	2 (6.25%)
Methotrexate and mycophenolate mofetil combined	1 (3.12%)
Systemic steroids (as sole therapy)	0
NB UVB or UVA1 phototherapy	0
Others (griseofulvin)	18 (56.25%)
None	7 (21.87%)
Orthopedic and reconstructive surgical treatment n (%)	8 (25%)
Physical rehabilitation n (%)	11 (34.4%)

NB UVB= narrowband ultraviolet B radiation (311 nm)

UVA1 = ultraviolet-A1 radiation (340-400 nm)

The patients' clinical evolution, as reported in the last follow-up visit, was remission with no signs of activity in 23 patients (72%) and active disease in 4 (12%) with new lesions and 2 (6%) with reactivation of old lesions.

DISCUSSION

Morphea is an uncommon disease which often appears in childhood. In this cohort of 32 patients at the National Institute of Pediatrics, the most relevant findings were: predominance of female gender (69%), which is related to previous reports in the literature, where the female to male ratio is 2.6:1.^{1,2} The most common subtype in the patients studied was linear (50%), which also coincides with global experience.^{4,7,9} In this study, the age range was 1 to 16 years with mean age of nine years, similar to that reported in the bibliography with mean age at onset of seven years.^{2,6,8}

In literature reports, almost 50% of patients with the linear subtype also have plaque morphea (mixed morphea, common in children).^{1,2,3,4,8,10,11} In our study, only 13% of patients had the mixed subtype of morphea.

In the most relevant published series of patients with morphea, around 40% of children with the disease had extracutaneous manifestations.^{7,12-16} Of our 32 patients, 27 (84%) had some extracutaneous manifestation related to morphea. Linear morphea retards growth of the affected extremity, and causes joint contractures.^{2,6,8} In our series of cases, the most common manifestations were musculoskeletal, with predominance of arthralgias (34%), discrepancy in length of extremities (34%), and joint contractures (38%). Neurological manifestations are more common when morphea affects the head, and there are even structural alterations; in our study there were nine patients (28%) with headaches, migraine, or seizures.¹⁷ Only three patients

had ophthalmological disorders, and although uveitis is the most commonly associated with morphea, none of them presented it. In many of our patients, information for targeted questioning to check for extracutaneous manifestations, mainly neurological and ophthalmological, was not available; as a result of which their real frequency may be underestimated.

The choice of treatment should depend on the subtype and extension of morphea. Based on clinical guidelines and consensus,¹⁸⁻²² morphea which can potentially cause functional disability or deformity (like the linear subtype in the head or extremities in children) requires systemic treatment; and topical treatment is suitable for the other subtypes. In our study, the most widely prescribed topical medicines were: vitamin D analogs (69%), calcineurin inhibitors (41%), and high-potency topical steroids (47%). Although there are few clinical trials with a high degree of evidence, today calcineurin inhibitors and vitamin D analogs are the safest, most effective drugs.^{2,3,10,13} Topical steroids should be prescribed with caution in patients with morphea because they may help reduce swelling in active superficial lesions but may have little or no effect in treating sclerosis and could potentially cause greater atrophy.^{2,4}

Regarding systemic treatment, immunosuppression with systemic corticosteroids (methylprednisolone 1-2 mg/kg/day or prednisone 0.5-1 mg/kg/day) is useful in inflammatory stages of morphea, above all in patients with rapidly progressive or disabling linear morphea. Methotrexate in doses of 15-20 mg weekly (dose 10-15 mg/m² of body surface) is currently the recommended first-line treatment for the active phase of morphea, with or without adjuvant use of systemic corticosteroids.^{2,5,10,13} Mycophenolate mofetil is another well tolerated immunosuppressant which has been effective in children

with morphea who have not had a successful response to treatment with a combination of glucocorticoids and methotrexate.¹⁰ The systemic treatment most commonly prescribed in our cohort was systemic steroids as bridge or adjuvant therapy while starting on another systemic treatment (72%): methotrexate in 63%, followed by hydroxychloroquine in 34% of the patients, and mycophenolate mofetil in 23%.

Other treatments available include UVA1 phototherapy, currently the treatment of choice for deep lesions because it alters the course of the disease inducing expression of metalloproteinase;¹ a collagenase which reduces procollagen and collagen in the skin, with complete relief or noticeable improvement in at least 60-80% of patients.^{2,4} Less effective is UVB phototherapy. The National Institute of Pediatrics does not have phototherapy facilities, for which reason none of the patients received such treatment.

A limitation of this study was finding incomplete information in medical notes, which potentially prevented us from estimating the real frequency of extracutaneous manifestations (there is no way to know if the patients actually did not have certain extracutaneous manifestations, did not undergo targeted questioning, or the information was not entered in the notes).

Nevertheless, this study is the first to describe clinical characteristics and extracutaneous manifestations in Mexican children with morphea. We found a higher incidence of the linear subtype in our patients compared with the findings reported in the bibliography, and a lower incidence of the mixed subtype. A highly relevant finding in this cohort of patients is that a very high percentage had musculoskeletal and neurological alterations related to morphea. Targeted questioning and paraclinical studies to identify extracutaneous manifestations (neurological and ophthalmological alterations) was not done routinely, and thus, underreporting is likely.

CONCLUSIONS

Morphea is an uncommon disease, which often appears in childhood and predominates in females. The linear subtype is common and is associated with more musculoskeletal extracutaneous manifestations. These were reported in 84% of our patients: musculoskeletal alterations occurred in 69%, neurological disorders in 28%, and ophthalmological disorders in 9%, although the frequency of the latter two may be underestimated in our study.

As areas of opportunity for the future, we consider it important to apply targeted questioning to patients with morphea regarding the most common extracutaneous manifestations, and when they are suspected confirm them by means of paraclinical studies. Also, it is crucial to improve the amount and quality of information in medical records, in order to obtain the most complete information from retrospective studies such as this one, and strengthen our understanding of the disease and patients' medical care.

REFERENCIAS

1. Yu BD, Eisen AZ. Scleroderma. In: Freedberg IM, Eisen AZ, Wolff K, et al. ed. Fitzpatrick's Dermatology in general medicine, vol II. 6ta edición. Nueva York, EEUU: McGraw-Hill, 2003: 1709-1717.
2. Röcken M, Ghoreshi K. Morphea and Lichen Sclerosus. In: Bolognia J, Jorizzo J, Rapini R. ed. Dermatology. 2a ed. España: Mosby Elsevier; 2008: 1469-1476.
3. Weibel L, Harper J. Morphea (Localized Scleroderma). In: Irvine A, Hoeger PH, Yan AC. Ed. Harper's textbook of pediatric dermatology. 3ra ed. NJ: Wiley-Blackwell. 2011: 1651-58.
4. Careta M, Romiti R. Localized scleroderma: clinical spectrum and therapeutic update. Anais Brasileiros de Dermatologia. 2015;90(1): 62-73.
5. García de la Peña, P. Esclerodermia localizada. Protoc diagn ter pediatr. 2014;1:10: 1-6.
6. Behman R, Kliegman R, Jenson H, Agud Aparicio J. ed. Nelson, tratado de pediatría. 2a ed. Madrid: McGraw-Hill, 2002: 765-766.
7. Garcia-Romero MT, Laxer R, Pope E. Localized Scleroderma: Symptoms, Diagnosis and Treatment. In: De Winter, R. ed.

Scleroderma. Symptoms, Diagnosis and Treatment. NY: Nova Science Publishers, Inc, 2013: 87-103.

8. Arenas R. Dermatología. Atlas, diagnóstico y tratamiento. 4^a ed. México: McGraw-Hill; 2015, pag 169-174.
9. Alfaro Cordero, K. Morfea: El desequilibrio entre la producción y la destrucción del colágeno. Revista médica de Costa Rica y Centroamérica LXXIII. 2016; 619: 351-354.
10. Bielsa Marsol, I. Update on the Classification and Treatment of Localized Scleroderma. *Actas Dermo-Sifiliográficas* (English Edition). 2013;104(8): 654-666.
11. Warner Dharamsi J, Victor S, Aguwa N, et al. Morphea in Adults and Children Cohort III. *JAMA Dermatology*. 2013; 149(10): 1159.
12. Christen-Zaech S, Hakin MD, Afsar FS, Paller AS. Pediatric morphea (localized scleroderma): Review of 136 patients. *J Am Acad. Dermatol.* 2008; 59:385-96.
13. Noguera M, Saurio de Carvalho M, Lombardi V. Morfea. *Arch. Argent. Dermatol.* 2012;62: 77-84.
14. Atzeni F, Bardoni A, Cutolo M et al. Localized and systemic forms of scleroderma in adults and children. *Clin Exp Rheumatol.* 2006; 24 (suppl.40):S36-45
15. Zulian F, Vallongo C, Woo P, et al. Localized scleroderma in childhood is not just a skin disease. *Arthritis Rheum.* 2005; 52: 2873-81
16. Pequet MS, Holland KE, Zhao S, et al. Risk factors for morphea disease severity: A retrospective review of 114 pediatric patients. *Br J Dermatol.* 2014; 170(4): 895-900.
17. Chiu YV, Vora S, Kwon EK, Maheshwari M. A significant proportion of children with morphea en coup de sabre and Parry-Romberg syndrome have neuroimaging findings. *Pediatr Dermatol.* 2012; 29(6):738-48.
18. Uziel Y, Feldman BM, Krafchik BR, Yeung RS, Laxer RM. Methotrexate and corticosteroids therapy for pediatric localized scleroderma. *J Pediatr.* 2000; 136: 91-95.
19. Weibel L, Sampaio MC, Visentin MT, et al. Evaluation of methotrexate and corticosteroids for the treatment of localized scleroderma (morphea) in children. *Br J Dermatol.* 2006; 155(5):1013-20.
20. Kreuter A, Altmeyer P, Gambichler T. Treatment of localized scleroderma depends on the clinical subtype. *British Journal of Dermatology.* 2007; 156(6): 1363-1364.
21. Bielsa Marsol, I. Update on the Classification and Treatment of Localized Scleroderma. *Actas Dermo-Sifiliográficas* (English Edition). 2013; 104(8), 654-666.
22. Martini G, Campus S, Bernd R, et al. Tocilizumab in two children with pansclerotic morphea: a hopeful therapy for refractory cases? *Clin Exp Rheumatol.* 2017; 35 Suppl 106(4):211-213. Epub 2017 Sep 29. PMID: 28980909