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Use of Pegylated Interferon Alpha-2a in Cutaneous T-cell Lymphoma: A Retrospective Case Collection

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Mycosis fungoides and Sézary syndrome are rare and largely incurable types of cutaneous T-cell lymphoma with limited therapeutic options. In 1984 Bunn et al. reported that interferon alpha is an efficient monotherapy in cutaneous T-cell lymphoma and 14 years later it was shown in a prospective, randomized trial that a combination of interferon alpha and psoralen plus ultraviolet A therapy (PUVA) is most efficient in the treatment of cutaneous T-cell lymphoma. Since then interferon alpha as single agent or, most often, in combination with phototherapy and/or retinoids has been integrated as standard of care in cutaneous T-cell lymphoma guidelines worldwide. However, production of interferon alpha was discontinued recently worldwide and pegylated interferon alpha-2a (PEG-IFN α) has been used as an alternative therapy. In contrast to numerous interferon alpha studies, only a few studies focusing on PEG-IFN α are available. Therefore, the aim of this study was to conduct a retrospective data collection to report on the efficacy, adverse events and therapy regimens of PEG-IFN α in cutaneous T-cell lymphoma. In 28 patients with cutaneous T-cell lymphoma treated in Germany and in the Netherlands, 36% of patients achieved complete remission, 36% partial remission and 29% stable disease. Eighteen percent of patients developed adverse events during therapy, which led to the discontinuation of PEG-IFN α therapy in 2 patients. The most common concomitant therapies were oral PUVA phototherapy and local radiotherapy. In conclusion, PEG-IFN α , especially in combination with skin-directed therapies, is an effective treatment option for cutaneous T-cell lymphoma in clinical practice.

Key words: mycosis fungoides; pegylated interferon; Sézary syndrome.

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Mycosis fungoides (MF) and Sézary syndrome (SS) are rare and largely incurable types of cutaneous T-cell lymphoma (CTCL) with limited therapeutic options (1). Interferon alpha (IFN α) is a well-known immuno-

SIGNIFICANCE

Mycosis fungoides and Sézary syndrome are rare and largely incurable types of skin cancer, that can also affect lymph nodes and blood. Therapeutic options are limited. Recombinant interferon alpha has been used for decades as an effective medication, but was discontinued recently worldwide. Pegylated interferon alpha-2a (PEG-IFN α ; Pegasys®, zrpharma Vienna, Austria) has been used as an alternative therapy. This retrospective study analysed the response and adverse events with PEG-IFN α in 28 patients. With 36% of patients achieving complete remission, 36% partial remission, and 29% stable disease, PEG-IFN α presents an innovative treatment option for patients with early as well as advanced CTCL.

modulatory modality. Bunn et al. published for the first time, in 1984, that IFN α is an efficient monotherapy in CTCL (2). Current guidelines for the therapy of cutaneous lymphomas, including the European Organisation for Research and Treatment of Cancer (EORTC), the International Society of Cutaneous Lymphomas (ISCL), the European Society for Medical Oncology (ESMO), the British Association of Dermatologists and UK and German guidelines, recommend, among others, the use of IFN α in MF and SS (3–6). The EORTC Cutaneous Lymphoma Task Force (EORTC CLTF) recommends IFN α as first-line therapy alone in MF stage IIB and III and as second-line therapy in MF stage I and IIA (3). The ESMO recommends treatment with IFN α for patients with extensive infiltrated plaques and tumors or patients refractory to skin-directed therapies, in combination with psoralen plus ultraviolet A therapy (PUVA) or other skin-directed therapies (4). IFN α is recommended as first-line treatment in combination with PUVA in MF stage IIB to IVB, or in combination with extracorporeal photopheresis or PUVA in Sézary syndrome in the German guideline (6). In prospective and retrospective studies IFN α alone or in combination can be effective in all stages of MF and SS, with overall response rates ranging up to 80% (3, 7). However, published data has the limitation of heterogeneity in treatment schedules, patient selection and methodology and randomized controlled trials are rare (3, 7).

In early 2020 IFN α was withdrawn from the European market and pegylated interferon alpha-2a (PEG-IFN α ,

Pegasys®) is now increasingly used as an alternative. PEG-IFN α has antiviral, antitumour, immunomodulatory and antiproliferative properties. Compared with non-pegylated IFN α , it has a significantly longer plasma half-life due to its binding to polyethylene glycol, which reduces renal clearance (8, 9). PEG-IFN α is therefore administered subcutaneously only once a week (10). It has been approved for the therapy of chronic hepatitis B and C for more than 20 years (10). Adverse events include anaemia, thrombocytopaenia, hyper- and hypothyroidism, anorexia, depression, headache, poor concentration, cough, gastrointestinal complaints, hair loss, dermatitis, arthralgias and myalgias, viral and bacterial infections, fever and fatigue (10).

The aim of this study was therefore to conduct a retrospective data collection on the use of PEG-IFN α in patients with CTCL naive to interferon treatment visiting the dermatology clinics of Leiden University Medical Center (The Netherlands) and Minden University Medical Center (Germany). The study evaluates the therapeutic effect, adverse events, co-medication and treatment regimens of PEG-IFN α .

MATERIALS AND METHODS

A retrospective analysis was performed in the Department of Dermatology at the University Hospitals of Minden and Leiden. All patients treated with PEG-IFN α from January 2020 to October 2022 for CTCL in stage IB to IVAa (MF, SS) were included. The EORTC/ISCL staging system for cutaneous lymphomas was used to evaluate tumour stage (3). For evaluation of therapy response (complete response, partial response, stable disease and progressive disease) the recommendations for clinical trial design and staging update from the ISCL, USCLC, and EORTC, from Olsen et al. (11), were used. The study was conducted in accordance with the Declaration of Helsinki, and was approved by the ethics committee of Ruhr University of Bochum (approval number 2021-811) and the ethics committee of the University of Leiden (approval number N20.052).

RESULTS

Between January 2020 and October 2022, a total of 28 patients with CTCL, 11 women and 17 men, were treated with PEG-IFN α at the Departments for Dermatology in Minden and Leiden. Details of the study cohort are shown in **Table I**. Patient age at therapy initiation ranged from 43 to 82 years (range 39 years) and the median age was

Table I. Case series overview

Case	Age at start of therapy, years, Sex	Diagnosis	Stage ISCL/EORTC	Status of therapy	Duration of therapy, weeks	Reason for cessation of therapy	Therapy combination	Best response	Adverse events of PEG-IFN	PEG-IFN dose, µg/week
1	59, F	MF	IB	Ongoing	32		TCS	SD	No	135
2	56, M	MF	IB	Ongoing	11		UVB, syst. PUVA	PR	No	135
3	65, F	MF	IB	Ongoing	35		Syst. PUVA	CR	No	135
4	70, F	MF	IB	Stopped	15	CR	Syst. PUVA	CR	Fatigue	135
5	56, M	MF	IB	Ongoing	44		Syst. PUVA, chlormethine gel, TCS	SD	Leucopaenia (grade 1) Thrombocytopaenia (grade 1)	135
6	76, M	MF	IB	Stopped	68	Adverse events	TCS	SD	Thrombocytopaenia (grade 1) Lymphopaenia (grade 1) Fatigue, Weight loss	135–90
7	66, F	MF	IIB	Stopped	20	Adverse events	Syst. PUVA	PR	Gastrointestinal (diarrhoea)	135
8	59, M	MF	IIB	Ongoing	37		Syst. PUVA, TCS	PR	No	135
9	80, M	MF	IIB	Stopped	16	CR	Syst. PUVA, local RTX	CR	No	135
10	46, M	MF	IIB	Ongoing	73			CR	No	135
11	75, M	MF	IIB	Ongoing	37		Syst. PUVA	CR	No	135
12	73, F	MF	IIB	Stopped	22	CR	Syst. PUVA	CR	No	135
13	57, F	MF	IIB	Ongoing	59		Local RTX	SD	No	135
14	56, M	MF	IIB	Stopped	54	CR	Local RTX	CR	No	135
15	73, M	MF	IIB	Ongoing	21		Syst. PUVA	PR	No	135
16	66, M	MF	IIB	Stopped	33	CR	Syst. PUVA, local RTX	CR	No	135
17	64, M	MF	IIB	Ongoing	74		Syst. PUVA	PR	No	135
18	72, M	MF	IIB	Stopped	122	PR/SD	Chlormethine gel, local RTX	PR	No	135
19	76, F	MF	IIB	Stopped	95	PD	Chlormethine gel, syst. PUVA	PR	No	135
20	61, M	MF	IIB	Ongoing	28		Local RTX, chlormethine gel, TCS	SD	No	135
21	58, F	MF	IIB	Ongoing	40		TCS	PR	Elevated liver enzymes (grade 1)	135–90
22	66, M	MF	IIIA	Ongoing	64		UVB	CR	No	135
23	52, M	MF	IIIA	Ongoing	35			CR	No	135
24	45, M	MF	IIIB	Ongoing	28		Syst. PUVA, chlormethine gel	PR	No	135
25	58, M	FMF	IIB	Ongoing	19		Chlormethine gel, TCS	SD	No	135
26	82, F	FMF	IIB	Ongoing	145		Local RTX, TCS, chlormethine gel	SD	No	90
27	43, F	FMF	IVa	Ongoing	136		Local RTX, TCS	SD	No	65
28	74, F	SS	IVa	Ongoing	28		PUVA	PR	No	180

MF: mycosis fungoides; SS: Sézary syndrome; FMF: follicular mycosis fungoides; CR: complete remission; PR: partial response; SD: stable disease; PD: progressive disease; syst. PUVA: oral psoralen UVA; UVB: narrowband ultraviolet B light therapy; TCS: topical corticosteroids; RTX: radiation; PEG-IFN: pegylated interferon alpha-2a; EORTC/ISCL: European Organisation for Research and Treatment of Cancer/ International Society of Cutaneous Lymphomas.

65 years (25th percentile = 57, 75th percentile = 73). Most patients (86%; 24/28) had classic MF. Three patients were diagnosed with follicular MF. One patient had Sézary syndrome. At the start of therapy, 6/27 of the patients with MF were in stage IB, 17/27 in stage IIB, 2/27 in stage IIIA, 1/27 in stage IIIB and 1/27 in stage IVA.

The duration of therapy varied from a minimum of 11 weeks to a maximum of 145 weeks (range 134 weeks) and the median duration was 36 (25th percentile = 27, 75th percentile = 65) weeks of therapy. The majority of patients (19/28) were still on therapy at the time of data collection. Reasons for discontinuation were complete remission (56%, 5/9), adverse events (22%, 2/9), inadequate response with stable disease (11%, 1/9) or disease progression (11%, 1/9). In 89% (25/28) of patients, a starting dose of 135 µg once weekly was selected. In 1 obese patient (body mass index (BMI) > 25) a dose of 180 µg was selected, while in 2 very slim patients (BMI < 18.5) a starting dose of 65 µg and 90 µg weekly was selected. In 2 patients, the therapy was reduced from 135 µg to 90 µg once weekly during the course of therapy, due to adverse events.

Monotherapy with PEG-IFNα was used in only 7% (2/28) of patients, whereas, in all other cases, combination therapies were used (26/28). The most frequent combination therapies were PUVA (54%, 15/28) and local radiotherapy (29%, 8/28). A combination with UVB311 nm was given in 7% (2/28). In 25% (7/28) of cases, supplementary local therapy with chlormethine gel (Ledaga®; Recordati, Rare Diseases, Ulm, Germany) was used in addition to interferon therapy. 32% (9/28) patients also received topical steroids (Fig. 1).

Eighteen percent (5/28) of patients developed adverse events during therapy, including gastrointestinal discomfort with diarrhoea, leukopaenia, thrombocytopenia, lymphopaenia, increase in transaminases, fatigue and weight loss. Adverse events that led to discontinuation of therapy were thrombocytopenia, lymphopaenia, fatigue and weight loss in 1 patient and gastrointestinal symptoms in another patient. In 2 patients the therapy was reduced from 135 µg to 90 µg 1 × weekly during the course of therapy due to adverse events. Both patients had fewer adverse events after dose reduction, but in 1 of the patients MF plaques increased after dose reduction.

Complete remission was achieved in 36% (10/28), partial remission in 36% (10/28) and stable disease in 29% (8/28) of patients. Patients who received a combination of PUVA and PEG-IFNα achieved complete remission (CR) in 40% (6/15), partial remission (PR) in 53% (8/15) and stable disease (SD) in 7%

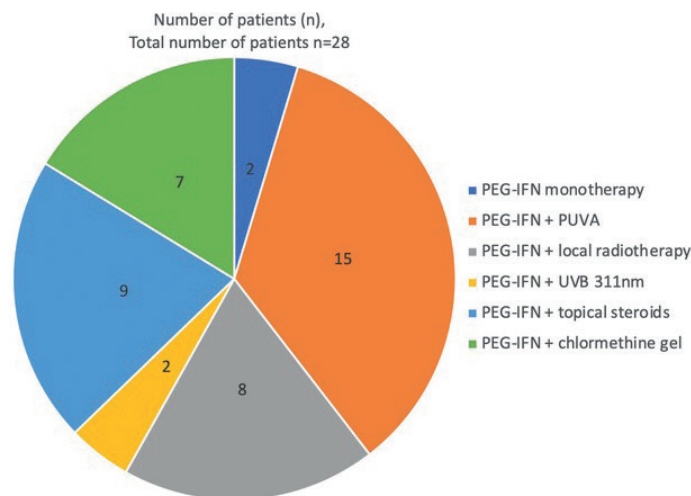


Fig. 1. Therapy combinations in the study cohort (n = 28 patients). PEG-IFN: pegylated interferon alpha-2a; PUVA: psoralen UVA therapy; UVB: ultraviolet B.

(1/15). Patients treated with local radiotherapy and PEG-IFNα achieved CR in 38% (3/8), PR in 13% (1/8) and SD in 50% (4/8) (Fig. 2).

Patients with MF stage IB achieved partial or complete response in 50% (3/6) and 50% (3/6) showed stable disease. Patients with classic MF in stage IIB showed partial or complete response in 87% (13/15) and only 13% (2/15) showed stable disease. All patients in stage III (3/3) achieved partial or complete response. All patients with follicular MF (3/3, stage IIB and IVA) achieved stable disease. The only patient with SS achieved a partial response. In the case series, especially patients in stage IIB and III, seem to profit most from the treatment with PEG-IFNα.

Combination therapy of PEG-IFNα with topical steroids, chlormethine-gel, PUVA, UVB 311-nm phototherapy, or local radiotherapy was well tolerated.

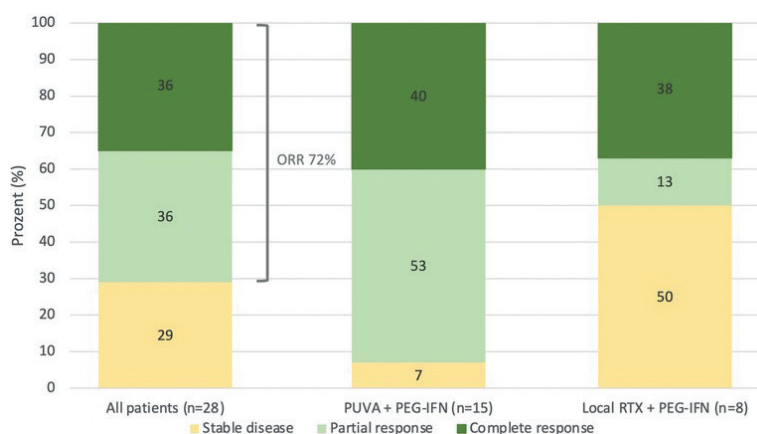


Fig. 2. Therapy response in all patients treated with pegylated interferon alpha-2a (PEG-IFN), in patients treated with a combination of PEG-IFN and psoralen UVA (PUVA), and in patients treated with a combination of PEG-IFN and local radiotherapy. ORR: overall response rate.

DISCUSSION

In this retrospective study 20/28 (71%) of patients showed a partial or complete response to PEG-IFN α combination therapy. In the large majority of cases this therapy could be continued for months or even years without major adverse events. Adverse events were relatively rare, could be managed effectively using dose reduction and led to discontinuation only in a small minority of patients. At the recent presentation at an EORTC/Cutaneous Lymphoma Tumour Group (CLTF) meeting in Madrid in 2022 Mitsunaga et al. reported on the use of PEG-IFN α in 110 patients with CTCL stage IA to IVA2. They found a lower rate of complete remission (12% vs 36%), but a higher rate of partial response (40% vs 36%) than in the current study, possibly due to the lower case numbers in the current study (12). In comparison with Patsatsi et al.'s study PEG-IFN α was only combined with skin-directed therapy in the current study, explaining the high response rates (13). They reported a 75% ORR for stage IB and 55% for all stages, underscoring that PEG-IFN α will belong to standard of care in the treatment of cutaneous T-cell lymphoma.

Prospective trials from Stadler et al. (14) and Chiarion-Seleni et al. (15) evaluated the combination of non-pegylated IFN α and PUVA therapy in patients with MF and found complete response rates from 70% to 74% that were even higher than in the current study, and partial response rates from 6% to 10% that were lower than in the current study. Overall response rate with PEG-IFN α in the current study (72%) was similar to trials with non-pegylated IFN (80%). An international multicentre retrospective follow-up study from the cutaneous Lymphoma International Consortium from 2017 found that IFN α was the second most common first-line treatment in stage IVA1 and also most widely employed as third-line treatment in patients with MF and those with SS (16).

Other haematological malignancies in which PEG-IFN α is used include myeloproliferative diseases, such as essential thrombocythemia and polycythemia vera, with overall response rates of 69% and 60% (17). Loss of response to IFN therapy has been reported in patients with hepatitis C, possibly based on the presence of IFN antibodies (18, 19). In the current study no loss of response to IFN in combination with skin-directed therapies was observed.

In a previous study adverse events to IFN α were mild or moderate and affected 55% of patients treated with a combination of IFN α and PUVA (14). Compared with the results of the current study, the number of patients experiencing adverse events (18%) was lower. However adverse events were similar, also mild to moderate, and generally not a reason to stop therapy. As the medication is administered only once a week and can be self-administered by the patient, the effort required on the part of the patient is relatively low.

When interpreting the results of the current study, the shortcomings of a retrospective data collection must be taken into account, which include, above all, a non-standardized and a smaller patient collective, as well as no defined intervals and procedures of assessment of tolerability and efficacy. The influence of co-treatment on therapy response, especially of PUVA used in more than 50% of patients, and of local radiation, must be noted.

This study shown that PEG-IFN α , particularly in combination, with a dose between 135 and 180 μ g, is an effective therapy option for treatment of MF and SS, particularly in classic MF stage IIB and III, in the real-world setting and seems not inferior to recombinant IFN α combinations.

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