Appendix 1. General Description of Included Studies

Level of						Condition					
evidence	Study	Design	No. of pts	Sex	Age, y	treated	Treatment; dose; duration	Comparator	Efficacy results	AEs	Follow-up
	Biologics										
4	Chang et al	Case series	7	F	47-71	Erosive MLP	Alefacept; 15 mg IM; 12 wk	Placebo (saline)	2 pts achieved notable improvement	Cystitis, UTI	24 wk
	(1)							IM	in PGA of disease severity, MP		
									severity, and IS		
5	Fivenson et	Case report	2	F	25, 57	Generalized LP	Alefacept; 15 mg/wk IM; 12	None	Both pts reported considerable	None	20 wk
	al (2)					(including OLP)	wk		diminution of itching and noticeable		
									improvement within 4 wk		
5	Cheng and	Case report	1	F	54	Erosive OLP	Efalizumab; initial 0.7 mg/kg,	None	Resolution of oral erosions, erosive	None	Not mentioned
	Mann (3)						then 1.0 mg/kg/wk; 10 wk		gingivitis, and dysphagia		
1c	Heffernan et	Prospective	4	F	52-71	Erosive OLP	Efalizumab; 0.7 mg/kg	None	Mean reduction in affected mucosal	Urticaria,	20 wk
	al (4)	pilot study					subcutaneously at wk 0, then		surface area 71.1%; mean	staphylococcus abscess	
							1.0 mg/kg/wk; 12 wk		improvement in VAS for pain of	of artificial hip joint (1	
									82%; mean improvement in OHIP-14	pt), drug-induced	
									questionnaire of 69.3%	subacute cutaneous	
										lupus (1 pt)	
5	Parmentier et	Case report	1	F	53	Mucocutaneous	Rituximab; 375 mg/m²/wk; 4	None	Dramatic improvement at 3 and 6-mo	None	10 mo
	al (5)					LP w/ esophageal	IV courses		follow-up; control esophagoscopy at		
						involvement			3 mo, no active LP lesions or		
									stenosis; immunohistochemistry of		
									esophageal mucosa, disappearance of		
									CD20+ cells		
5	Goni Esarte	Case report	1	F	59	ELP	Rituximab; 4 IV doses (375	None	Endocscopic improvement of lesions;	None	9 mo
	et al (6)						mg/m ² each) per wk; duration		decreased intensity of dysphagia and		
							not mentioned		odynophagia		
2b	Sartori-	Retrospective	19	15F	Mean, 57	Otic LP	Topical tacrolimus (1 pt also	None	One pt w/ severe LP of the ear, oral	Not mentioned	4.9 y
	Valinotti et al	chart review		4M			received rituximab); dose and		cavity, esophagus, and genital area		
	(7)						duration not mentioned		reported remarkable relief w/		
									rituximab prescribed primarily for		
									Sjögren syndrome		

5	Rebora et al	Case report	1	F	67	Erosive LP (oral	Basiliximab; 2 bolus IV	None	Basiliximab cleared erosions and	Appearance of	Not mentioned
	(8)					and vaginal)	infusions (20 mg) 4 days apart;		abated symptoms temporarily	antimitochondrial	
							duration not mentioned		(rebound effect after 1 mo)	antibodies and increase	
										in gamma-glutamyl	
										transferase	
5	Ho et al (9)	Case report	1	F	67	Recalcitrant	Adalimumab; 160 mg	None	Notable improvement of genital and	None	Not mentioned
						vulvovaginal	subcutaneously, 80 mg 2 wk		oral lesions (edema and erosions)		
						gingival	later, then 40 mg every other		after 4 wk and complete resolution		
						syndrome	wk; 12 wk		after 12 wk		
5	Chao et al	Case report	1	F	52	Cutaneous LP	Adalimumab; 40 mg	None	Almost clear response by 6 wk and	None	Maintained
	(10)					and MLP (oral	subcutaneous injections every		complete clearance, including oral		almost clear
						and vulvar)	other wk; 22 wk		lesions, at 8 wk		response of
											oral/vulvar LP
											beyond wk 50
5	Yarom et al	Case report	1	F	56	OLP	Etanercept; 25 mg twice/wk;	None	Notable symptom relief (up to 90%)	Mild-to-moderate	3 y
	(11)						10 wk		2 wk after start of therapy;	tenderness at injection	
									keratinization of eroded mucosa after	site	
									4 wk		
	MMF										
2b	Ashack et al	Retrospective	53	37F	Mean, 60	MLP	Topical or oral corticosteroids,	None	Average number of lesions reduced	MMF: fatigue, anemia,	Not mentioned
	(12)	review		16M		(oral/genital)	MMF, or cyclosporine; MLP		from 3.77 to 1.67 (P<.001); average	diarrhea, elevated	
							algorithm:		disease activity reduced from 2.73 to	blood pressure, UTI	
							Step 1, TCS or tacrolimus		0.90 (P<.001); average pain		
							b.i.d.		decreased from 2.03 to 1.03		
							Step 2, burst and taper of		(P<.001).		
							prednisone				
							Step 3, MMF				
							Step 4, oral cyclosporine + MMF				
							and TCS or calcineurin inhibitor;				
	1						MMF mean duration 1.7 y,				
							Cyclosporine, 3-7 mo				
5	Frieling et al	Case report	3	2F	17-54	Disseminated	Cyclosporine, 3-7 mo MMF; 2 g/d; 5-12 mo	None	Complete remission in 2 pts, and	None	19-65 mo
5	Frieling et al (13)	Case report	3	2F 1M	17-54	Disseminated and erosive LP		None	Complete remission in 2 pts, and substantial improvement in the other	None	19-65 mo

						(including oral in all pts and genital in 1)					
2b	Wee et al (14)	Retrospective review	10	9F 1M	16-54	Recalcitrant erosive OLP (vulvovaginal- gingival, penogingival, oral)	MMF; 500 mg/d increased according to tolerance and disease activity, aiming for 2 g b.i.d. by 2 mo; mean 3.7 y	None	6 pts achieved remission, 1 had well- controlled disease, and 3 had partially controlled disease.	Headache and tiredness (2 pts)	4.2 y
5	Deen and McMeniman (15) Azathioprine	Case report	1	F	66	Erosive genital LP	MMF; 500 mg b.i.d. increased to 1.5 g/d after 6 mo; 3 mo	None	Improvement in pain, dysuria, and pruritus within 4 wk of treatment initiation	None	Several months
4	Verma et al (16)	Case series	9	5F 4M	Mean (range), 32 (5-54)	Severe erosive oral or generalized LP	Azathioprine; 50 mg b.i.d. orally (about 2 mg/kg/d); 3-7 mo (mean, 5 mo)	None	7 pts, excellent response; 1 pt, good response; 1 pt, poor response; response considered excellent if 75%-100% improvement in lesions and itching/irritation, good if 50%-75%, and poor if <50%	Bleeding from the gums due to gingivitis	6-9 mo
5	Lear and English (17) MTX	Case report	2	F	60, 74	OLP and skin LP	Azathioprine; 50 mg b.i.d., after 2 mo, dose reduced to 50 mg/d; 2 mo and 5 mo	None	Decrase in size of erosions within 1 mo; complete healing after 2 mo; no recurrence at 6 mo	None	6 то
1b	Lajevardi et al (18)	Prospective open trial	18	13 F 5 M	NA	Erosive OLP	MTX; 15 mg/wk; 12 wk	None	Partial response or better in 15 (83.3%) pts. Statistically significant reduction in Thongprasom scale and VAS scores (<i>P</i> <.001)	.Skin eruption, nausea, epigastric pain, elevated liver enzymes	12 wk
1b	Chauhan et al (19)	Prospective observational study	45	Group: A, 12F 3M B, 9F 6M C, 8F	Group, mean (SD): A, 44.47 (13.30) B, 46.33	OLP	Group A, topical triamcinolone; 0.1% oral paste 3 times/d; 16 wk or until complete clinical remission	Group: B, MTX (0.3 mg/kg/wk) C, triamcinolone + MTX	Pts in combination group had better reduction in outcomes (clinical severity score, VAS and quality of life impairment questionnaire) compared to other 2 groups	MTX: nausea/vomiting, anemia; triamcinolone: telangiectasia/ atrophy at site of application	Not mentioned

				7M	(10.78)						
					C, 45.53						
					(17.79)						
					(=,,,,						
4	Jang et al	Case series	4	F	23-61	Erosive	Oral MTX, clobetasol, and	None	All pts experienced improvement in	None	4-6 mo
	(20)					vulvovaginal LP	tacrolimus; MTX 2.5-7.5		symptoms and healing of lesions		
							mg/wk + topical clobetasol		within 4-8 weeks		
							dipropionate 0.05% ointment				
							and tacrolimus 0.03%-0.10%				
							ointment; 4-6 mo				
4	Torti et al	Case series,	50	35F	41-80	Erosive OLP	Therapeutic ladder of sequential	None	Best responses in treatment naive pts;	Oral candidiasis	27 mo
	(21)	retrospective		15M			treatments (TCS, topical		most achieved substantial response	(corticosteroid)	
		review					immunomodulator, HCQ,		w/ limited AEs		
							systemic retinoids, MTX, and				
							thalidomide); dose and duration				
							not mentioned				
4	Nylander	Case series	4	3F	30, 45,	Erosive LP of	MTX; 10-15 mg/wk; 14-21 mo	None	Complete healing of oral and genital	None	17-48 mo
	Lundqvist et			1M	55, 60	mouth,			mucosa. No repeat esophageal		
	al (22)					esophagus, and			dilation for stenosis required since		
						genitalia			starting MTX		
	Cyclosporin										
5	Boyce et al	Case report	1	F	58	Erosive MLP	Cyclosporin A; 150 mg b.i.d.;	None	Reduction in genital and oral	None	Not mentioned
	(23)					(oral, genital,	6 wk		erosions/ulceration, improvement of		
						esophageal, and			dysphagia and odynophagia,		
						ocular w/			resolution of epiphora		
						epiphora)					
5	Chaklader et	Case report	1	F	54	ELP	Cyclosporin; dose and duration	None	Improvement of dysphagia	None	Not mentioned
	al (24)						not mentioned				
2b	Brewer et al	Retrospective	11	9F	43-75	Ocular LP	Dapsone, doxycycline,	None	Symptom improvement and stable	Not mentioned	3.3 y
	(25)	chart review		2M			prednisone, and cyclosporine;		physical findings		
							dose and duration not				
							mentioned				
	Tacrolimus			1							
			<u> </u>								

Control of the cont	5	Yeo et al (26)	Case report	2	F	63, 70	Erosive LP	Oral tacrolimus; 2-4 mg b.i.d.	None	Improvement in vulval and oral	None	Not mentioned
Note Cheer et al. Case series 3 F Moste, OPP C2 pro) and Decreases Decrease Dec							(vulvovaginal/	(0.05-0.15 mg/kg); at least 6		symptoms within 6 weeks		
Chee et al. Chee et al. (27) Case series 3 P Mean, Oi P (2 pity) and Taccoliums; 0.5-1.0 mg ht.d.; 4 None inprovement of oral unake, pain, gingsed cyclema, and oral bosons of necotions; 1 prefured upon cessation of necotions; 1 and followed. 1 remains stable on maintenance dose RCP ID B Secherel et al. Prospective 7 OF Mean, Ol. Chronic consiste of the part of							oral, vulvar, and	wk				
S2.3 Onlivegral LP 100, 3 no. and 1.5 y gingival crythems, and oral lesions spon cestation of teachings 1 no. 5 no. and 1.5 y gingival crythems, and oral lesions spon cestation of teachings 1 nor followed; 1 noralises 1 nor followed; 1 noralises 1 noralise							skin)					
Case series 12 10F 40.74 Erosive OLD P (8 genital LP) 20 20 20 20 20 20 20 2	5	Chen et al	Case series	3	F	Mean,	OLP (2 pts) and	Tacrolimus; 0.5-1.0 mg b.i.d; 4	None	Improvement of oral intake, pain,	Fatigue	1 pt flared
Case series 12 10F 40.74 Ecosive OLP (8 ps alto late 12 ps alto late 13 ps alto late 1		(27)				52.3	oral/vaginal LP	mo, 5 mo, and 1.5 y		gingival erythema, and oral lesions		upon cessation
Decrease of Color extracted of the Septiment of the Complete remission and (25) partial remission. Septiment (25) 25 26 27 26 27 27 28 28 28 28 28 28							(1 pt)					of tacrolimus;
Becheric et al Prospective 7 OF Mean, Chronic crossive ECP, monomodelar cells extracted, soluble 8-MOP added to cytopheresis product (280) smaly 1 M 61.4 LP added to cytopheresis product (200 ngmL), cells irradiated w/UVA and reinfused into pts; twice/wk for 3 wk, then tripend according pt needs 2M 2M 2M 2M 2M 2M 2M 2												1 not followed;
ECP Description Descripti												1 remains
ECP 1b Becherel et al (28) Study 1 M 61.4 LP Extracted, soluble 8-MOP added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pes; twice/wk for initial 3 wk, then reduced to once every 4 mo 2 M Case report 2 M Case report 1 F 50 Refractory ECP, mononuclear cells ECP, mononuclear cells extracted, soluble 8-MOP added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pes; twice/wk for initial 3 wk, then reduced to once every 4 mo Ecose of Case report 1 F 50 Refractory ECP, mononuclear cells extracted, soluble 8-MOP added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pes; twice/wk for initial 3 wk, then reduced to once every 4 mo Ecose of Case report ECP, mononuclear cells ECP, mononuclear cells extracted, soluble 8-MOP added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pes; twice/wk for initial 3 wk, then reduced to once every 4 mo Ecose of Case report ECP, mononuclear cells None All pss, decrease of crosive surface; 9 Decrease of (75%) archieved complete remission In prodominantly CD4+ of 8 ps followed for 33 y had recurrence of erosions when ECP sessions slowed or stopped; after resumption of ECP, partial or complete remission achieved None No												stable on
Becherol et al Prospective 7 6F Mean, Chronic erosive ECP; mononuclear cells cxtracted, soluble 8-MOP added to eyupheresis product (28) study 1M 61.4 LP extracted, soluble 8-MOP added to eyupheresis product (200 g/mL), cells irradiated w.UVA and reinfused aftor pts; twice/wk for 3 wk, then tapend according pri needs (29) 2M 2M Erosive OLP (8 ECP; mononuclear cells cxtracted, soluble 8-MOP (29) 2M 2M Erosive OLP (8 ECP; mononuclear cells cxtracted, soluble 8-MOP (29) added to cytapheresis product (200 g/mL), cells irradiated (200 g/mL), cells irradi												maintenance
Becherel et al. Prospective 7 6 F Mean, Chronic erosive ECP, monomuclear cells extracted, soluble 8-MOP added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pts: twice/wk for 3 wk, then tapered according pt needs 4 Gayot et al. Case series 12 10F 40-74 Erosive OLP (8 pt salso had genital LP) added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pts: twice/wk for 3 wk, then tapered according pt needs 4 Case series 12 10F 40-74 Erosive OLP (8 pt salso had genital LP) added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pts: twice/wk for initial 3 wk, then reduced to once every 4 mo 5 Zingoni et al. Case report 1 F 5 SD Refractory erosive oral, genital, and skin LP 6 Promonomuclear cells w/UVA and reinfused into pts: twice/wk for initial 3 wk, then reduced to once every 4 mo 6 Progressive decrease in 24 mo w/UVA and reinfused into pts: twice/wk for initial 3 wk, then reduced to once every 4 mo 7 S pts followed complete remission achieved 8 Decrease of 12 yraphocytes subsets, predominantly CD2+ T I I/mphocytes. 9 Decrease of 12 yraphocytes subsets, predominantly CD2+ T I/mphocytes when ECP existing a slowed or stopped; after resumption of ECP, partial or complete remission achieved 9 S Zingoni et al. Case report 1 F 5 SD Refractory erosive oral, genital, and skin LP												dose
Case series 12 10F 20D		ECP										
Case series 12 10F 20D	1b		Prospective	7	6F	Mean.	Chronic erosive	ECP: mononuclear cells	None	All pts had complete remission.	Progressive decrease in	24 mo
added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pts; twice/wk for 3 wk, then tapered according pt needs 4 Guyot et al (29) Case series 12 10F 40-74 Erosive OLP (8) genital LP) added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pts; twice/wk for 3 wk, then tapered according pt needs ECP; mononuclear cells of 8 pts followed complete remission and 3 (25%) partial remission. Seven of 8 pts followed for >3 y had recurrence of erosions when ECP sessions slowed or stopped; after reduced to once every 4 mo w/UVA and reinfused into pts; twice/wk for initial 3 wk, then reduced to once every 4 mo Solved or stopped; after resumption of ECP, partial or complete remission achieved Solved or stopped; after resumption of ECP, partial or complete remission of skin lesions, w/re-epithelization of vulvar erosions and those on tongue, cheeks, and lips												
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w/UVA and reinfused into pts; twice/wk for 3 wk, then tapered according pt needs A										,		
twice/wk for 3 wk, then tapered according pt needs 4 Guyot et al (29) 2M Erosive OLP (8 ECP; mononuclear cells pts also had genital LP) 2M Erosive OLP (8 ECP; mononuclear cells of extracted, soluble 8-MOP added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pts; twice/wk for initial 3 wk, then reduced to once every 4 mo 5 Zingoni et al (30) 2 Zingoni et al (30) 2 Erosive OLP (8 ECP; mononuclear cells of ECP; mononuclear cells of RDP (75%) achieved complete remission and 3 (25%) partial remission. Seven of 8 pts followed for >3 y had recurrence of erosions when ECP sessions slowed or stopped; after resumption of ECP, partial or complete remission achieved complete remission achieved or complete remission achieved or complete remission achieved or complete remission of skin lesions, w/re-epithelization of vulvar erosive oral, genital, and skin LP										required	subsets	
tapered according pt needs Guyot et al Case series 12 10F 40.74 Erosive OLP (8 ECP; mononuclear cells pts also had pts also had pts also had genital LP) added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pts; twice/wk for initial 3 wk, then reduced to once every 4 mo ECP; a consecutive days every None Complete remission of skin lesions, None Not mentioned Not menti												
Guyot et al (29) Case series 12 10F 40-74 Erosive OLP (8 pts also had genital LP) Added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pts; twice/wk for initial 3 wk, then reduced to once every 4 mo Zingoni et al (30) Case report 1 F 50 Refractory erosive oral, genital, and skin LP Refractory erosive oral, genital, and skin LP												
2M	4	Count at al	G	12	100	40.74	Ei OI D (0		None	All standards of succine success 0	Decreesed	2
genital LP) added to cytapheresis product (200 ng/mL), cells irradiated w/UVA and reinfused into pts; twice/wk for initial 3 wk, then reduced to once every 4 mo 5 Zingoni et al Case report 1 F 50 Refractory erosive oral, genital, and skin LP Refractory genital, and skin LP and 3 (25%) partial remission. Seven of 8 pts followed for >3 y had recurrence of erosions when ECP sessions slowed or stopped; after resumption of ECP, partial or complete remission achieved Complete remission achieved None Complete remission of skin lesions, w/ re-epithelization of vulvar erosions and those on tongue, cheeks, and lips	4		Case series	12		40-74	· ·		None			3 yrs
Case report 1 F 50 Refractory ECP; 2 consecutive days every (30) 3 weeks; 16 courses (8 mo) Case rejoint and those on tongue, cheeks, LP Case rejoint and lips Case rejoint an		(29)			2M					-		
w/UVA and reinfused into pts; twice/wk for initial 3 wk, then reduced to once every 4 mo resumption of ECP, partial or complete remission achieved 5 Zingoni et al Case report 1 F 50 Refractory erosive oral, genital, and skin LP Sources (8 mo) w/ re-epithelization of vulvar erosions and those on tongue, cheeks, and lips							genital LP)			· · · · ·		
twice/wk for initial 3 wk, then reduced to once every 4 mo sessions slowed or stopped; after resumption of ECP, partial or complete remission achieved 5 Zingoni et al Case report 1 F 50 Refractory erosive oral, genital, and skin LP Sources (8 mo) sessions slowed or stopped; after resumption of ECP, partial or complete remission achieved None Complete remission of skin lesions, w/ re-epithelization of vulvar erosions and those on tongue, cheeks, and lips											Tlymphocytes	
reduced to once every 4 mo resumption of ECP, partial or complete remission achieved 5 Zingoni et al Case report 1 F 50 Refractory erosive oral, genital, and skin LP Refractory erosive oral, genital, and skin LP reduced to once every 4 mo resumption of ECP, partial or complete remission achieved Complete remission of skin lesions, None Not mentioned w/ re-epithelization of vulvar erosions and those on tongue, cheeks, and lips								_				
Case report 1 F 50 Refractory ECP; 2 consecutive days every None Complete remission achieved								·				
5 Zingoni et al Case report 1 F 50 Refractory ECP; 2 consecutive days every None Complete remission of skin lesions, None Not mentioned (30) w/re-epithelization of vulvar erosions and those on tongue, cheeks, and lips								reduced to once every 4 mo				
erosive oral, genital, and skin LP genital, and skin LP w/ re-epithelization of vulvar erosions and those on tongue, cheeks, and lips												
genital, and skin LP genital, and skin and lips	5		Case report	1	F	50	Refractory		None		None	Not mentioned
LP and lips		(30)					erosive oral,	3 weeks; 16 courses (8 mo)				
							genital, and skin			erosions and those on tongue, cheeks,		
HCQ							LP			and lips		
		HCQ										

				44M	(CD)						
					(SD),	erosive, atrophic		mg/d, reduced to	expression of Tregs, IL-8, TGF-β1,		
					44.5	atypical		5 mg/d when	and IL-10 higher in OLP pts than		
					(13.3)	reticular, and		50% reduction	healthy controls before tt. Frequency		
						atypically		in lesion size	of Tregs was downregulated after 2		
						erosive)		achieved	wk of HCQ, whereas prednisone had		
									no effect on Tregs levels		
1c Ye	Yeshurun et	Clinical trial	21	15F	Mean	OLP	HCQ sulphate; 400 mg/d; 1-36	None	Five (24%) pts, complete remission;	Blurred vision, visual	Not mentioned
al	1 (32)			6M	(range),		mo		12 (57%), moderate to marked	field defects, rash,	
					55 (30-				improvement; 3 (14%), no	hyperpigmentation,	
					82)				improvement; for 1 pt therapy	elevated kidney function	
									terminated after 1 mo due to AEs;	tests	
									response to therapy observed after 2-		
									4 mo; 3 of 6 pts who responded to		
									therapy flared on stopping		
1c Eis	Eisen et al	Clinical trial	10	9F	Mean	OLP	HCQ; 200-400 mg/d; 6 mo	None	Nine of 10 pts had excellent response	None	6 mo
(33	33)			1M	(range),				to therapy; 3 of 6 pts w/ erosions at		
					59 (40-				baseline had complete healing; pain		
					66)				relief and reduced erythema observed		
									after 1-2 mo of therapy, but erosions		
									required 3-6 mo to resolve		
5 De	De Argila et	Case report	1	F	51	OLP on lower	Chloroquine phosphate; 250	None	Excellent response within 3 mo;	None	Not mentioned
al	1 (34)					lip (solitary	mg b.i.d. for 3 mo, then		symptoms disappeared completely		
						lesion)	reduced to 250 mg b.i.d. for 3		and only mild erythema remained		
							mo; 3 mo				
2b Ve	Vermeer et al	Retrospective	15	15F	55 years	ELP of the vulva	HCQ 200 - 800 mg for 23.8	None	60% of patients responded to	GI disturbance,	38 mo
(35	35)	chart review			(range	and vagina	months		HCQ, with almost half experiencing	dizziness, infection,	
					23–82)				long-term effect – clinical response	headache	
									was defined as a decrease in PGA		
									score		
Th	Thalidomide										
5 Ca	Camisa et al	Case report	1	M	70	Recalcitrant	Thalidomide; 100 mg/d; 1 y 5	None	Complete resolution of desquamative	Dizziness, low	Not mentioned
(30	36)					erosive OLP	mo		gingivitis after 11 mo	extremity edema,	

										erythematous-	
										•	
										squamous rash of face	
										and trunk	
5	Petropoulou	Case report	1	М	NA	Erosive oral and	Thalidomide; 50 mg/d for 2	None	Complete healing of erythematous	Muscle cramps and	Not mentioned
	et al (37)					genital LP	wk, 25 mg/d for 1 yr, 25 mg		and erosive areas on penis and	weakness, numbness,	
							every other d for 6 mo; 18 mo		whitish mouth lesions	and lower extremity	
										burning	
	IVIG										
5	Nakashima et	Case report	1	M	57	Refractory OLP	IVIg; 400 mg/kg/d for 5 d; 2	None	VAS decreased by 30% after 1 wk of	None	Not mentioned
	al (38)						cycles		IVIG and lip erosions and ulcers		
									improved after 2 mo		
1c	Bender et al	Clinical trial	3	F	53-78	Refractory OLP	Adjuvant IVIG + acitretin;	None	Pts showed mixed responses to	IVIG-induced	Not mentioned
	(39)						IVIG 2 g/kg/m cycle, acitretin		adjuvant IVIG, ranging from	leukopenia	
							0.3-0.5 mg/kg/d (ie, 30 mg/d);		therapeutic efficacy to no response		
							at least 7 mo				
	BCG-PSN										
1c	Nasr et al	Clinical trial	11	6F	7-69	OLP	BCG-PSN; intradermal	None	Significant differences in decrease of	Swelling at injection	3 mo
	(40)			5M			injections of 0.5 mL (0.453		lesion areas, NRS, REU, and VAS	sites	
	(10)			3111			105-15 3 105 cfu); twice/wk		scores; most pts achieved CR after 3	Sico	
							for 3 wk		wk of treatment		
	D " (<u> </u>				101 3 WK		wk of treatment		
		inocycline/eryth	romycin								
5	Kandula et al	Case report	1	М	56	Ulcerative LP	Oral and TCS, topical	None	Marked clinical improvement of the	Not mentioned	Not mentioned
	(41)					on plantar foot	tacrolimus, and oral		cutaneous and oral mucosal lesions		
						and lateral	doxycycline; dose not				
						tongue	mentioned; 4 wk				
2b	Cooper et al	Prospective	114 (31	F	Mean,	Erosive vulvar	Minocycline, erythromicine,	None	Minocycline: 33% good, 33% partial,	Not mentioned	72 mo
	(42)	cohort	systemic		56.9	LP	prednisone, acitretin,		and 33% poor response		
			treatment)				cyclosporine, azathioprine,		Erythromicine: 50% good, 33%		
							HCQ, thalidomide, or colchicine;		partial, and 17% poor response		
							dose and duration not mentioned		Prednisone: 33% good, 33% partial,		
									and 33% poor response		
									Acitretin, cyclosporine, azathioprine,		
									HCQ, thalidomide, and colchicine:		
			<u> </u>		<u>l</u>						

									100% poor response		
	Dapsone										
	Dapsone										
5	Beck et al	Case report	1	F	74	Erosive LP, oral	Dapsone; 50 mg/d, gradually	None	Buccal mucosa and toe lesions	None	7 mo
	(43)					and toes	increased to 150 mg/d; 7 mo		completely healed after 7 mo, tongue		
	(- /										
									erosions reduced to ¼ of pretreatment		
									size, itching mouth pain disappeared		
5	Basak et al	Case report	1	M	9	Generalized LP,	Dapsone; 50 mg daily (1.5	None	Itching subsided within 1 mo and	None	Not mentioned
	(44)					oral and nail	mg/kg/d) for 8 mo, increased		disappeared in 2 mo; trunk lesions		
	()										
							to 75 mg/d (2.5 mg/kg/d) for 5		started flattening after 3 mo; at 8 mo,		
							mo; 13 mo		still active lesions on limbs, so		
									dosage was increased and lesions		
									cleared at 5 mo; oral mucosal lesions		
									·		
									were cleared		
1c	Chopra et al	Clinical trial	75	NA	NA	LP	Dapsone and chlorpheniramine	Local	Total efficacy of dapsone regimen	Not mentioned	Not mentioned
	(45)						maleate; dose and duration not	corticosteroids and	was 18% higher than corticosteroid		
							mentioned	chlorpheniramine	regimen		
								maleate			
	Metronidazole	;									
1c	Buyuk et al	Clinical trial	20	14F	Mean	Generalized LP	Metronidazole; 500 mg b.i.d.;	None	15 (78.9%) pts responded to	Mild headache, nausea	5-36 mo
10		Cinnear una	20					None		wind neadache, nausea	3-30 mo
	(46)			6M	(range),	(7 pts w/ OLP)	20-60 d		metronidazole (13 complete and 2		
					40.7 (14-				partial); worsening of lesions in 1 of		
					62)				4 nonresponders; 3 of 7 oral lesions		
									(all in CR pts) cleared after		
									metronidazole		
1c	Rasi et al	Clinical trial	49 (2	25F	Mean	Cutaneous LP	Metronidazole; 250 mg/8 h; 3	None	Overall response for MLP, 66.6%;	Metallic taste (1 pt)	3 mo
	(47)		mucosal)	24M	(SD), 48.3	and MLP	mo		overall response for itching, 75%; pts		
					(12.3)				w/ CR were symtpom-free in <3 mo		
					(12.3)				w/ CK were symponi-nee in <3 ino		
		1									

	Griseofulvin										
4	Aufdemorte	Case series	3	F	68, 63, 52	Erosive OLP	Griseofulvin; 500 mg b.i.d.,	None	Pt 1: complete remission of oral	None	15 mo and 9
	et al (48)						reduced to 250 mg/d after 3		lesions at 8 wk; Pt 2: course		mo (NA for Pt
	et all (10)						mo; 12 mo and 6 mo (duration		paralleled first pt, but responses more		3)
							not mentioned for Pt 3)		rapid, complete remission at 10 wk;		3)
							not mentioned for Pt 3)				
									Pt 3: marked clinical improvement at		
									10 wk w/ few persistent erosions,		
									complete remission not achieved;		
									response of oral lesions was		
									dramatic; response intervals varied		
2b	Massa et al	Chart review	29	20F	23-76	OLP only (11	Griseofulvin; 500 mg/d; 3 wk-	None	11 pts w/ OLP only: 3 complete	Rash, nausea,	Not mentioned
	(49)			9M		pts) and LP of	12 mo		remission, 3 marked improvement	constipation, diarrhea	
						skin, genitalia,			between 3 wk and 3 mo; overall		
						nails, and scalp			response rate 54.5% in cutaneous LP;		
						(18 pts)			benefit from griseofulvin much less		
									likely		
1c	Matthews et	Clinical trial	23 (11	NA	NA	OLP	Griseofulvin; 500 mg b.i.d.; 3	None	Symptomatic benefit in 21% of 23	Headaches, nausea,	Not mentioned
	al (50)		completed)				mo		pts, but no clinical improvement; half	vomiting, diarrhea	
									of pts withdrew due to AEs or lack of		
									symptom improvement; possible		
									placebo effect, since no clinical		
									improvement noted by observation or		
									photos		
1c	Bagan et al	Clinical trial	7	5F	34-68	OLP (4 erosive	Griseofulvin; 500 mg b.i.d.;	None	No improvement; in 4 pts (2 w/	Headaches	Not mentioned
	(51)			2M		and 3 reticular)	mean, 2.5 mo		erosive and 2 w/ reticular) condition		
	(51)			2171		and 3 reticular)	moun, 2.5 mo		worsened; in 2 pts, lesions remained		
									unchanged from baseline		

4	Naylor (52)	Case series	4	2F	43-67	Erosive OLP	Griseofulvin; 125 mg 4	None	Pt 1: erosion increased in size during	None	12 mo
				2M			times/d; 8 wk		8-wk regimen, disappeared 3 mo		
									after completion; Pts 2 and 3: erosion		
									disappeared during 8-wk regimen,		
									recurred 2 wk after completion; Pt 4:		
									erosion decreased in size during 8-wk		
									regimen, recurred 3 mo after		
									completion; griseofulvin had little/no		
									effect on pain, pigmentation, or		
									resolution of lesions, did not protect		
									from recurrences		
	Etretinate										
1b	Hersle et al	Randomized,	28	18F	Group,	Severe OLP	Etretinate; 75 mg/d (25 mg 3	Placebo	93% of buccal lesions treated w/	Keratoconjunctivitis,	3 mo
	(53)	double-blind		10M	mean	(atrophic and	times/d), mean (range) dose		retinoid improved (erosion clearance	exanthematic rash,	
		comparative			(range):	erosive or	0.98 (0.75-1.25) mg/kg/d; 2		or ≥50% reduction, erythema	headache, dry skin and	
		trial			treatment,	plaque and	mo		reduction by at least 2 steps on 4-step	mucosa, itchiness, hair	
					54 (35-	reticular)			scale) compared to 5% of placebo-	loss	
					77)				treated lesions; blind evaluation of		
					placebo,				photos agreed w/ clinical evaluation;		
					58 (52-				etretinate provided effective		
					75)				symptomatic relief for severe OLP,		
									but 6 pts stopped due to AEs		
4	Gorsky and	Case series	6	4F	Mean	OLP	Etretinate; 75 mg/d (25 mg 3	None	50% of pts became asymptomatic	Dry skin and mucosae,	12 mo
	Raviv (54)			2M	(range),		times/d); 2 mo		(atrophic or erosive lesions	skin sloughing, skin	
					61 (49-				disappeared or converted into	rash and itching, hair	
					77)				keratotic asymptomatic stage); 4 pts	loss, alanine	
									(66%) improvement of at least 70%;	aminotransferase and	
									same frequency of recurrence and	triglyceride elevation	
									discomfort after stopping etretinate,		
									requiring topical/ systemic		
									corticosteroids for disease control		
	Alitretinoin										
		<u> </u>									

5	Brehmer et al	Case report	3	F	78, 77, 58	Mucocutaneous	Oral alitretinoin; 10 mg/d; 4	None	Oral lesions and pain disappeared	Hypertriglyceridemia,	Not mentioned
	(55)					and cutaneous	wk		completely, pts remained symptom-	headache and dizziness	
						LP			free after discontinuation		
5	Kolios et al	Case report	1	F	54	Cutaneous, oral,	Oral alitretinoin; 30 mg/d; 2	None	Oral/skin changes and dysphagia	Mild sporadic	12 mo
	(56)	and literature				and ELP	cycles of 6 mo each, 12 mo		completely resolved within 4 wk; nail	headache	
		review					total		changes within 6 mo		
	Isotretinoin										
4	Camisa et al	Case series	6	3F	35-72	Erosive OLP	Systemic isotretinoin; 10-50	None	4 of 5 evaluable pts were improved	Cheilitis, dry skin,	4 wk
	(57)			3M			mg/d (0.5 mg/kg/day, raised to		after 8 wk, but none were completely	headache, rash, joint	
							a maximum of 1 mg/kg/d); 8		cleared	pain, pruritus (all pts)	
							wk				
	Vitamin A										
5	Chopra and	Case report	1	M	44	Hyperkeratotic	Vitamin A; 3 tablets/d (50,000	None	Positive response when vitamin A	None	Not mentioned
	Kaur (58)					OLP	IU each); 4 wk		was added to corticosteroid and		
									dapsone regimen; regression in size		
									and thickness of lesion after 1 mo		
	Corticosteroid	s									
2b	Harewood et	Retrospective	4	F	40-65	ELP	Prednisone; 60 mg/d (1 pt), 40	None	3 pts responded dramatically within 1	None	Not mentioned
	al (59)	chart review			(symptom		mg/d (3 pts), then rapidly		mo; 4th pt flared every time		
					onset)		tapered; 2-3 wk		perdnisone tapered below 10 mg,		
									remained on 10 mg w/o symptoms		
5	Kumar et al	Case report	1	M	58	Atrophic OLP	Oral mini pulse therapy with	None	Gradual and consistent reduction in	Decrease in white	4 mo
	(60)	and literature					betamethasone; 5 mg/d for 2		burning sensation, complete	blood cell count and	
		review					d/wk for 3 wk, tapered by 0.5		remission of oral lesions	hemoglobin, increase	
							mg every wk; at 15th wk, pt			in erythrocyte	
							was taking 0.5 mg/d,			sedimentation rate	
							maintained for 3 wk; 18 wk				
4	Wedgeworth	Case series	5	F	49-58	ELP	Combination of balloon	None	Most cases, full resolution of	Pneumomediastinum (1	18-54 mo
	et al (61)						dilatation and intralesional		dysphagia generally sustained for	pt)	
							triamcinolone; each stricture		several months		
							injected w/ 40-60 mg of				
			Ì		1						
							triamcinolone (10 mg/mL				

							quadrants of the stricture;				
							graduated balloon				
							dilatation performed through				
							scope, graduated balloons for				
							30 s; Average interval between				
							treatments was 8.3 mo; pts did				
							not become tolerant to				
							procedure and time between				
							procedures tended to lengthen				
2b	Fahy et al	Retrospective	100	F	Mean,	Genital (vulval)	Corticosteroids (22), MTX (17),	None	Most pts who received systemic	Dermatomyositis and	24-42 mo
	(62)	chart review			60.3	LP	MMF (10), acitretin (1),		treatments did not achieve remission;	myositis induced by	
							isotretinoin (3), HCQ (12),		of pts w/ disease remission,	oral corticosteroid	
							dapsone (3), colchicine (1),		approximately ½ were prescribed	therapy (1 pt)	
							azathioprine (5), cyclosporine		systemic medications; 82% w/		
							(4), or IVIG (2); dose and		remission were receiving care		
							duration not mentioned		through the dermatology service		
4	Franco et al	Case series	6	5F	NA	ELP	Topical fluticasone (3),	None	Symptoms resolved w/ treatment, but	Not mentioned	1-7 y
	(63)			1M			prednisone, intralesional		recurred after 10 mo (2 pts), 1 yr (1		
							triamcinolone (2), or		pt), and 2 y (1 pt)		
							budesonide (1); fluticasone				
							b.i.d, prednisone 20 mg/d,				
							intralesional triamcinolone 20				
							mL (10 mg/mL), budesonide 3				
							mg b.i.d, then 3 mg/d;				
							fluticasone 8 wk, prednisone 2				
							wk, budesonide 10 mo				
5	Sheehan-	Case report	1	F	50	ELP	Prednisolone; 20 mg/d;	None	Rapid symptomatic improvement	Not mentioned	Not mentioned
	Dare et al						duration not mentioned		within days; when prednisolone		
	(64)								reduced to 5 mg/d, dysphagia		
									returned, resolved when increased to		
									20 mg/d; repeated attempts to		
						1	i e		1		
									withdraw corticosteroids resulted in		

									symptom-free on prednisolone 10		
									mg/d		
5	Ynson et al	Case report	1	F	63	ELP	Fluticasone propionate; 220 µg	None	Resolution of symptoms at 4-wk	Not mentioned	Not mentioned
	(65)						b.i.d.; 6 wk		follow-up; 15-wk follow-up		
									endoscopy: small light pink plaques		
									in mid-esophagus, normal looking		
									mucosa in rest of esophagus		
5	Sato et al	Case report	1	F	85	ELP	Prednisone; 20 mg daily;	None	Symptoms improved within 1 wk,	Not mentioned	2 yrs
	(66)						duration not mentioned		oral corticosteroid tapered by 5 mg		
									every 2 wk until reached 5 mg/d;		
									endoscopic and histologic		
									improvement 3 mo after prednisone		
									initiation; clinical remission		
									remained for 2 y		
2b	Bradford and	Retrospective	131	F	Mean, 57	Vulvovaginal	Oral prednisolone (22), oral	None	98% saw improvement of symptoms	Prednisolone: mood	1 mo-15 y
	Fischer (67)	chart review				LP	prednisolone + TCS (31), oral		and examination findings in a mean	disturbance	(mean, 6.4 y)
							prednisolone + MTX (1), or		of 7.5 wk; on follow-up, 77% had		
							MTX + TCS (1); prednisolone		stable disease, while 23% had		
							5-50 mg/d, MTX 5-7.5 mg/wk;		unstable disease and progressive		
							duration not mentioned		tissue destruction		
5	Teixeira et al	Case report	1	M	50	ELP	Prednisolone; 40 mg; duration	None	Rapid improvement of odynophagia	Not mentioned	Not mentioned
	(68)						not mentioned		and dysphagia		
2b	Kern et al	Prospective	20 (of 32	12F	Mean	ELP	Prednisone, azathioprine,	None	Treatment with topical budesonide	Not mentioned	1.5-6 y (mean,
	(69)	cohort	with LP)	8M	(range),		topical budesonide, or		formulation or systemic		3.6 y)
					55 (27-		acitretin; prednisone 50 mg/d,		corticosteroids was successful in		
					74)		azathriopine up to 2 mg/kg/d,		most pts, w/ proven ELP and		
							and topical budesonide 0.5 mg		reversed functional esophageal		
							b.i.d.; acitretin not mentioned;		stenosis		
							duration not mentioned				
					1			1			

5	Menges et al	Case report	1	F	56	ELP	Methylprednisolone and	None	Immunosuppressive therapy with	Not mentioned	5 y
	(70)						budesonide solutions; 20 mg/d;		systemic and local corticosteroid		
							3 wk		application did not prevent recurrent		
									stenosis, endoscopic dilatation had to		
									be performed 5 times in 5 y		
1b	Singh et al	RCT	40	20F	Mean	OLP	Topical triamcinolone, oral	Not mentioned	In all groups, significant	Mild tingling in oral	Not mentioned
	(71)			20M	(SD),		dapsone, topical tacrolimus, or		improvement in symptoms and sign	cavity in pts treated	
					32 (10.5)		topical retinoid; triamcinolone		scores, steroidal and non-steroidal	with topical agents	
							buccal paste 0.1% b.i.d.,		agents had equal efficacy; dapsone		
							dapsone 100 mg b.i.d. + iron		had greater efficacy than topical		
							and folic acid tablets, topical		retinoid, but no significant		
							tacrolimus 0.1% b.i.d., topical		differences for dapsone vs topical		
							retinoid b.i.d.; 3 mo		tacrolimus or topical retinoid vs		
									topical tacrolimus		
1b	F Agha-	RCT	27	17F	Mean	OLP	Combination of	Triamcinolone	Rate of symptom recurrence was	None	6 mo
	Hosseini et al			10M	49.81 (± 9		HA/triamcinolone or	alone	74.1% on control side and 11.1% on		
	(72)				.63)		triamcinolone alone was		test side. Group treated with		
							injected intralesionally, with		combination of HA and		
							1 ml of the medication injected		triamcinolone experienced		
							for each 2 cm ² surface area of		significantly better resolution of		
							the lesion		lesions and symptoms		
4	Kurt et al	Case series	3	2F	22/49/52	OLP	Intra-lesional corticosteroid	None	Resolution of symptoms of pain,	None	1 mo
	(73)			1M			injection: 0.4 mL of a 10		burning and sensitivity and healing of		
							mg/mL solution of		lesions in 1 month		
							triamcinolone acetonide				
							(TA) applied directly into the				
							lesion twice at a 2-week				
							interval				
	Diode laser										
1b	Agha-	Randomized	28	21F	Mean,	Erosive-atrophic	Diode laser; WL 633/890 nm,	CO ₂ laser	Laser-treated group showed better	None	3 mo
	Hosseini et al	controlled		7M	50.7	OLP	energy density 0.3-0.5 J/cm ² ,	surgery	improvement than CO2 group		
	(74)	trial					irradiation 5 s; 5 sessions every		(P<.001)		
							other day				
	1		1		1			1			1

1b	Dillenburg et	Randomized	42	35F	Mean,	Erosive-atrophic	Diode laser (red); WL 660 nm,	Clobetasol	Laser was more effective (P<.001)	None	2 mo
	al (75)	controlled		7M	58.2	reticular OLP	energy density 6 J/cm ² , power				
		trial					density 1,000 mW/cm ² ,				
							irradiation 6 s; 3 sessions/wk				
							(12 sessions)				
1c	El Shenawy	Controlled	24	18F	Mean,	Erosive-atrophic	Diode laser; WL 970 nm,	Triamcinolone	Corticosteroid group showed	None	Not mentioned
	et al (76)	trial		6M	53.6	OLP	power output 3 W, irradiation		significantly lower mean VAS scores		
							8 min; twice/wk (maximum 10		than laser group (P=.02)		
							sessions)				
1c	Othman et al	Controlled	24	18F	35-70	Erosive-atrophic	Diode laser; WL 970 nm,	Triamcinolone	Improvement in signs of disease w/	None	Not mentioned
	(77)	trial		6M		reticular OLP	power output 2 W, irradiation		no difference between 2 groups		
							8 min; twice/wk (10 sessions)				
1b	Kazancioglu	Randomized	120	64F	Mean,	Erosive-atrophic	Diode laser; WL 808 nm,	Group:	Improvement in all groups, but	None	6 mo
	et al (78)	controlled		56M	42.6	OLP	energy density 1.5 J/cm ² ,	1, ozone	significantly better in ozone and		
		trial					power density 10 mW/cm ² ,	2, dexamethasone	corticosteroid groups		
							irradiation 2.5 min; twice/wk	3, placebo	8		
							(10 sessions)	e, p			
	PDT						(,				
4	Rakesh et al	Case series	10	8F	20-70	Erosive OLP	5-ALA-mediated PDT; 4% (4	No treatment on	Clinical response to PDT similar in	None	Every 6 mo for
	(79)	Case series		2M	20 70	LIGHTO GEN	mg) 5- ALA gel applied twice	contralateral	both men and women, for treatment	Tione	4 y
	(17)			2.11			(2 mg each application) at 1 h	side	and controls; greater reduction was		.,
							interval; red light emitted by	side	noted in buccal mucosal and tongue		
							diode laser WL 600–670 nm,		lesions than in gingiva		
							energy density of 80 J/cm ² ;		icsions than in gingiva		
							duration not mentioned				
1b	Lundquist et	RCT	18	13F	Mean, 59	OLP	Photochemotherapy with 8-	No treatment on	Marked improvement in 9 pts, slight	Nausea, dizziness, eye	12 mo
10	al (80)	KC1	10	5M	ivicali, 39	JL1	MOP and long-wave UV-A;	contralateral	improvement in 4 pts, no	symptoms, numbness,	12 1110
	ai (60)			Sivi			WL 320-400 nm w/ UV-A	side (buccal		headache	
							irradiance 17.5 mW/cm², UV-		improvement in 3 pts	neadache	
							A dose 0.75 J/cm ² , increased	mucosa)			
							by 0.25 J/cm ² every other				
							session, 8-MOP 0.6 mg/kg				
							orally 2h before irradiation; 12				

							times at intervals of 2-3 d, total				
							dosage 16.5 J/cm ²				
							dosage 10.5 J/cm				
1b	Jajarm et al	Randomized	25	Study,	Group,	Erosive-atrophic	Toluidine blue-mediated PDT;	Corticosteroid	Traditional corticosteroid therapy	None	12 mo
	(81)	controlled		8F 3M	mean:	OLP	WL 630 nm, energy density	(dexamethasone	showed better results than toluidine		
		trial		control,	study,		1.5 J/cm ² , power density 10	mouthwash)	blue-mediated PDT		
				9F 5M	48.71		mW/cm ² , irradiation 2.5 s;				
					control,		twice/wk (10 sessions)				
					43.73						
1b	Mostafa et al	RCT	20	17F	Group,	Erosive OLP	MB-PDT; WL 660 nm,	Conventional	MB-PDT much more effective in	MB-PDT: edema and	2 mo
	(82)			3M	mean		intensity 100-130 mW/cm ² ,	TCS (kenakort	pain reduction and lesion regression	mild burning sensation	
					(SD):		5% MB-PDT once/wk, thin	A-orabase)	than TC	during application	
					A, 47.0		layer of TCS 3 times/d; 2 mo				
					(6.94)						
					B, 48.6						
					(5.25)						
2b	Cosgarea et	Prospective	20	17F	62 ± 8.66	OLP	PDT was performed within 14	None	Reduction of clinical parameters	None	6 weekskurt
	al (83)	case-		3M			days on the most extensive oral		(lesion size, ABSIS, Thongprasom-		
		controlled					lesion in 4 sessions (day 1, 3,		score), improvement		
		pilot study					7, 14)		of all quality-of-life (QOL) items and		
									significant decrease of relative		
									number of CD4+ and CD8+ T cells		
									in mucosal OLP-lesions		
	PRGF										
4	Pinas et al	Case series	4	F	43-59	Refractory	PRGF; 1 infiltration (2 pts) or	None	Complete healing of lesions (80%-	None	6 mo
	(84)					ulcerative	2 infiltrations (2 pts), 1 pt		90% size reduction) after first PRGF		
						erosive OLP	needed 3 rd infiltration at 6 mo		infiltration in 2 pts, other 2 needed a		
							posttreatment due to flare-up;		2 nd infiltration to achieve complete		
							duration not mentioned		healing; after 1st infiltration, pain was		
									reduced by 5.75 points, further		
									reduced by 2 points after 2 nd		

2b	Pinas et al	Retrospective	10	F	Mean	Erosive OLP	PRGF; 1 infiltration (8 pts) or	None	Complete remission	None	Mean (SD), 13
	(85)	study			(SD), 48		2 infiltrations (2 pts); duration				(1) mo
					(12)		not mentioned				
	SAFG										
5	Arcuri et al	Case report	1	F	69	OLP	SAFG; purified fat graft was	None	Improvement in pain according to	None	1 wk, 4 wk, 3
	(86)						injected with blunt cannulas in		VAS, chewing, swallowing, mouth		mo, and 6 mo
							lips and cheek with 10 mL of		opening, and appearance of lesions		after treatment
							pure centrifuged and purified				
							fat; duration not mentioned				
	Curcuminoids										
1b	Chainani-Wu	Randomized,	33 (28	23F	Mean,	OLP (atrophic	Curcuminoids (+ prednisone	Placebo	First interim analysis: no significant	None	Not mentioned
	et al (87)	placebo-	completed)	10M	60.6	or erosive)	60 mg/d for 1st week); 2,000		difference between placebo and		
		controlled,					mg/d; 7 wk		curcuminoids; study ended early for		
		double-blind							futility; reaching a conclusion		
		clinical trial							regarding efficacy of curcuminoids		
									based on this study was not possible;		
									curcuminoids at this dose were well		
									tolerated and results suggest that for		
									future studies, an RCT of shorter		
									duration w/ a larger sample size,		
									using higher curcuminoid dose, w/o		
									initial course of prednisone should be		
									considered		
1b	Chainani-Wu	Randomized,	20	10F	Group,	OLP	Curcuminoids; 2,000 mg 3	Placebo	Percent reduction in NRS, eryhtema,	Diarrhea (most	2 wk
	et al, 2011	placebo-		10M	mean:		times/d (6,000 mg/d); 12 d		ulceration, and total MOMI scores	frequent), constipation,	
	(88)	controlled,			treatment,				not significant in placebo group, but	abdominal pain,	
		double-blind			60.8				significant in curcuminoid group;	heartburn, nausea	
		clinical trial			placebo,				AEs uncommon in both groups		
					56.2						
2b	Chainani-Wu	Descriptive	43 (25/33	NA	NA	OLP	Curcuminoids; mean daily	None	72% from 1st RCT and 100% from	Mild abdominal	1st RCT: mean,
	et al, 2011	retrospective	from 1st				dose 2,137.5 mg (1st RCT) and		2 nd took over-the-counter	discomfort, diarrhea;	68.2 mo; 2 nd
	(89)	cohort	RCT and				5,058 mg (2 nd RCT); mean		curcuminoids after trial completion;	occurrence was dose	RCT: mean,
			19/20 from				duration 30 mo (1st RCT) and		60% reported reduction of symptoms	related	15.8 mo

			2 nd RCT)				9.6 mo (2 nd RCT)		w/ curcuminoids, 35% were unsure,		
									and 5% reported it did not reduce		
									symptoms		
	TGPC										
1b	Zhou et al	Prospective,	73/81 (3	42F	41-49	OLP (44	Topical or systemic	Topical or	Effective rates of combined treatment	Diarrhea	6 mo
	(90)	randomized,	dropped	31M		reticular and 37	corticosteroids + TGPC; oral	systemic	were statistically higher versus		
		controlled	out, 5			erythematous/	prednisolone 15 mg/d for 2	corticosteroids	control groups (efficacy assessed		
		clinical trial	absconded)			erosive)	wk/mo, 1,200 mg TGPC; 4 mo		through VAS and clinical signs)		
	Apremilast										
5	AbuHilal et	Case report	1	F	44	Erosive OLP w/	Apremilast; 30 mg/d; 3 mo	None	Marked reduction in erythema and	Nausea	Not mentioned
	al (91)					desquamative			erosions on upper and lower		
	, ,					gengivitis			gingivae, considerable reduction in		
									pain and discomfort with		
									improvement of quality of life w/		
									easier speaking, eating, and chewing		
	YY C 1	0 1			7.4	1.0	1 . 20 /1 4 1	N.		N	NY
5	Hafner et al	Case report	1	F	74	LP mucosae-	Apremilast; 20 mg/d; 4 wk	None	Complete clinical remission (of	None	Not mentioned
	(92)					associated			dysphagia and erosive stomatitis);		
						stenotic			control esophagoscopy, marked		
						esophagitis			recovery of the esophageal mucosa		
						(refractory to			w/ no recurrence of former stenosis		
						immuno-					
						suppressive					
						treatment w/					
						pulsed IV					
						methyl-					
						prednisolone)					
4	Bettencourt	Case series	3	F	73, 71, 66	Recalcitrant	Apremilast; 30 mg b.i.d.; 6 mo	None	At 2 to 4 wk follow-up, pts pain-free	Nausea, diarrhea	6 mo
	(93)					OLP	w/ tapering to once/d after		and sores cleared completely; 1 pt		
	, ,						complete resolution of lesions		received a course of oral prednisone		
									(40 mg) while on apremilast and		
									another short course (20 mg for 3		
									days) 2 mo later for a mild flare,		
									currently on low-dose prednisone (5		

			1	1				I	mg daily) and apremilast to prevent	I	T
									flares		
4	Paul et al	Case series	20	7F	NA	LP (1 OLP)	Apremilast; 20 mg b.i.d.; 12	None	OLP pt: oral lesions improved from	Headache, nausea	4 wk
	(94)			3M			wk		40% to 12% involvement of bilateral		
									buccal mucosa; on day 15, PGA		
									mucosal disease improved to		
									"marked resolution" and stabilized to		
									"moderate improvement" at end of		
									study		
	Levamisole										
1c	Lin et al (95)	Clinical trial	89	NA	Mean,	OLP	Levamisole; 50 mg b.i.d. in pts	None	1 TGA positive, 48 TMA positive;	None	Not mentioned
					55.1		w/ 30-50 kg body weight or 50		significant reduction in serum		
							mg 3 times/d for pts w/ 50-70		TGA/TMA levels and improvement		
							kg body weight, for 3		in signs/symptoms of OLP (reduction		
							consecutive days at beginning		in lesion size and pain, healing of		
							of each 2-wk interval; 12 mo		erosive lesions)		
1c	Lin et al (96)	Clinical trial	79 (all	NA	55	OLP (71	Levamisole; 50 mg b.i.d. in pts	None	Reduced high serum ANA to	None	Not mentioned
			ANA			erosive, 8	w/ 30-50 kg body weight or 50		undetectable level, improvement in		
			positive)			nonerosive)	mg 3 times/d for pts w/ 50-70		signs/symptoms of OLP (reduction in		
							kg body weight, for 3		lesion size and pain, healing of		
							consecutive days at beginning		erosive lesions)		
							of each 2-wk interval; 2-38 mo				
	Cyclophosphar	nide									
5	Paslin (97)	Case report	3	2F	50, 63, 74	Generalized LP	Cyclophosphamide; 50-100	None	Total resolution of LP, including oral	Leukopenia,	3-15 mo
				1M		(including OLP)	mg/d; 6 wk		lesions at end of wk 10; pts were	thrombocytopenia	
									disease free 3-15 mo after completion		
									of treatment		
									disease free 3-15 mo after completion		

Abbreviations: 8-MOP, 8-methoxypsoralen; AE, adverse event; ALA, aminolevulinic acid; ANA, antinuclear antibody; BCG-PSN, polysaccharide nucleic acid of Bacillus Calmette-Guérin vaccine; b.i.d., twice a day; cfu, colony-forming unit; CR, complete response; ECP, extracorporeal photochemotherapy; ELP, esophageal lichen planus; F, female; HCQ, Hydroxychloroquine; IL, interleukin; IM, intramuscular; IS, itch severity; IV,

intravenous; IVIG, intravenous immunoglobulin; LP, lichen planus; M, male; MB-PDT, methylene blue—mediated photodynamic therapy; MLP, mucosal lichen planus; MMF, mycophenolate mofetil; MOMI, modified oral mucositis index; MP, mucosal pain; MTX, methotrexate; NA, not available; NRS, numeric rating scale; OHIP-14, Oral Health Impact Profile-14; OLP, oral lichen planus; PDT, photodynamic therapy; PGA, Physician Global Assessment scale; PRGF, plasma rich in growth factors; pt, patient; PVAS, pain visual analogue scale; RCT, randomized clinical trial; Retro, retrospective; REU, reticulation, erythema, ulceration; SAFG, Submucosal autologous fat grafting; TCS, topical corticosteroids; TGA, anti-thyroglobulin autoantibody; TGF, transforming growth factor; TGPC, total glucosides of paeony capsules; TMA, anti-thyroid microsomal autoantibody; Tregs, regulatory T cells; UVA, ultraviolet A; VAS, visual analogue scale; WL, wavelength

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