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Case report

Alopecia as the presenting symptom of syphilis

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Abstract

Alopecia can be one of the many symptoms of secondary syphilis and the clinical presentations include essential syphilitic alopecia or symptomatic syphilitic alopecia. In this report, we present a case of a patient with essential syphilitic alopecia whose sole presenting symptom of syphilis was alopecia. Despite an initial negative rapid plasma reagin (RPR) test, he was ultimately found to have syphilis on scalp biopsy. His alopecia improved following treatment with benzathine penicillin. This presentation serves as a reminder to clinicians to be cognizant of alopecia as a presenting sign of syphilis. A review of the specificity and sensitivity of the typical tests used for the diagnosis is presented.

Case synopsis

A 50-year-old man presented to the UC Davis Dermatology Department with a diagnosis of telogen effluvium associated with a three-month history of diffuse hair loss as well as diarrhea and a twelve-pound weight loss. His alopecia was otherwise asymptomatic. He reported no hospitalizations, surgeries, or changes in medications over the past 6 months. He reported being in a monogamous relationship for 12 years and denied lesions on his genitalia, rashes, or any urethral discharge. He denied any history of illicit drug use or blood transfusions.



Physical exam revealed patchy, non-scarring, non-erythematous alopecia that diffusely involved his entire scalp (Figure 1). His eyebrow, facial, and body hair were preserved. There was no pitting of his fingernails or toenails. The results of his hair pull test were negative.

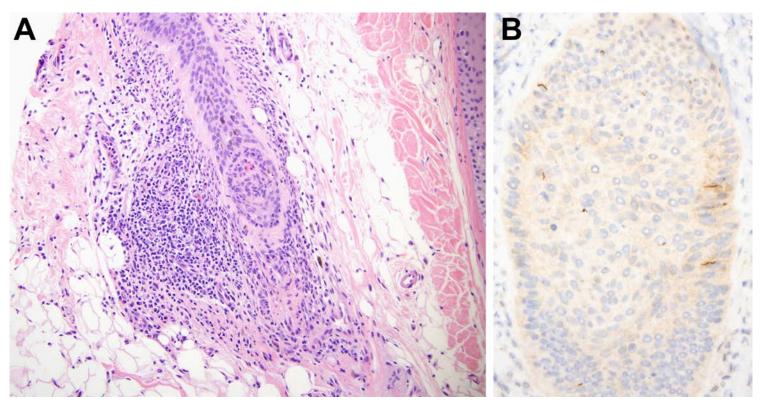


Figure 2. A) P Peribulbar inflammatory infiltrates with lymphocytes and plasma cells are present. B) Immunohistochemical stain for *Treponema pallidum* highlights corkscrew-shaped organisms within the hair follicle epithelium.

The following laboratory tests were within normal limits: ferritin, thyroid stimulating hormone, total and free testosterone, vitamin B12 level, vitamin D level, tissue transglutaminase level, and inflammatory bowel disease antibody screen. His rapid plasma reagin (RPR) test was initially negative. He had a mildly elevated aspartate amino transferase at 44 U/L and mild anemia with a hematocrit of 39%. The hepatitis B surface antigen and hepatitis C antibody were negative. The HIV antigen and antibody screeens were also negative. A scalp biopsy revealed increased catagen/telogen hairs and peribulbar inflammatory infiltrates with lymphocytes and plasma cells. Immunohistochemical staining for *Treponema pallidum* demonstrated the presence of corkscrew-shaped organisms in the hair follicle epithelium (Figure 2).

The patient was diagnosed with syphilitic alopecia and was treated with benzathine penicillin 2.4 million units. Upon further questioning, he revealed an unprotected sexual encounter 5 months prior to the onset of his alopecia. After the patient was diagnosed, his partner also tested positive for syphilis and received treatment with benzathine penicillin. RPR was repeated after the patient completed treatment and was elevated at 1:32. Also shortly after treatment, a *treponema pallidum* particle agglutination assay (TPPA) was drawn and found to be reactive. The patient was advised to return for repeat RPR 3 months, 6 months, and 12 months following the completion of treatment.

Discussion

Syphilis is a sexually transmitted disease that is caused by the sphirochete *Treponema pallidum* and mimics various dermatologic conditions. Syphilis infection can spread across years over multiple stages (Table 1).

According to a classification made by McCartney in 1940, syphilitic alopecia can be classified into symptomatic syphilitic alopecia and essential syphilitic alopecia [1, 2]. Symptomatic syphilitic alopecia is associated with other manifestations of secondary syphilis, whereas essential syphilitic alopecia has no other manifestations of secondary syphilis [1]. This case is consistent with essential syphilitic alopecia. Symptomatic syphilitic alopecia shows histopathological findings similar to

papulosquamous lesions of secondary syphilis. Essential syphilitic alopecia is characterized by either a telogen effluvium or alopecia areata –like pattern, similar to the patient's appearance in this case. The main difference between these two entities, syphilitic alopecia and alopecia areata, is the demonstration of *Treponema pallidum* in syphilis. Reports of detection of organisms in the hair follicle epithelium, as in the present case, are rare [1].

The patient described here presented with no symptoms other than alopecia, diarrhea, and weight loss; he had a negative RPR. According to CDC guidelines, the diagnosis of syphilis can be made with one positive non-treponemal serological test or an anticardiolipin test and one positive treponemal serological test as seen in table 2 [3]. If a patient has a positive treponemal serological test, then an anticardiolipin test should follow [3]. If this test is negative, then a second treponemal test should be performed. If the result of this second trenponemal test is positive then treatment should be administered. However, if the result is negative then no further treatment is necessary [3]. In low prevalence populations, the CDC recommends initially performing an anti-cardiolipin serological test and confirming positive results with a treponemal sergological test [4].

In accordance with the CDC guidelines, an RPR test was peformed [4], but initially was found to be negative. Although the sensitivity of the RPR test is fairly high at 86.4% [5], it is still less sensitive than other treponemal serological tests (Table 2). It is possible that this falsely negative result was the result of the prozone effect, which occurs when high antibody titers interfere with the antigen antibody lattice network formation necessary for obtaining a positive RPR result [6]. Although this effect can be avoided with dilutions of serum [6], dilutions are not routinely performed at this institution, further supporting the possibility of a prozone effect in this patient.

Despite advances in antibiotic development, penicillin remains the mainstay of treatment for syphilis. The recommended regimen for primary, secondary, and early latent infection is 2.4 million U of benzathine penicillin G given in a single intramuscular injection. Late latent infections can be treated with 3 weekly injections of 2.4 million U of benzathine penicillin G. Neurosyphilis can be treated with 12-24 million U/day via IM or IV for a total of 10-14 days [3, 7, 8]. Doxycycline, ceftriaxone, and tetracycline can be used as alternative treatments for syphilis when penicillin cannot be used, such as when a patient has a penicillin allergy [3, 7, 8].

Table 1. Stages of Syphilis

Stage	Timeline After Infection	Symptoms	
Primary	2-6 weeks	Local: Primary Chancre	
Secondary	4-10 weeks after infection	Hematogenous and lymphatic spread: fever, headache, anorexia,	
		lymphadenopathy, alopecia	
Tertiary	Years to decades	Dilated aorta, aortic regurgitation,	
		carotid ostial stenosis, gummas	

Table 2. Sensitivity and specificity of diagnostic tests for syphilis

	Sensitivity	Specificity	
PCR			
Primary Syphilis	89.1% [9]	99.1% [9]	
Secondary Syphilis	50% [9]	100% [9]	
Treponemal tests			
T. pallidum particle agglutination	99.8% [10]	99.6% [10]	
assay (TPPA)			
Enzyme Immunoassay (EIA)	100% [11]	99.5% [11]	
Anticardiolipin tests			
Rapid plasma reagin (RPR)	86.4% [5]	94.3% [5]	
Venereal Disease Research	70.8% [12]	99% [12]	
Laboratory (VDRL			

Table 3. Summary of treatments for syphilis

	Treatment	Dosage	Duration
Primary	Benzathine Penicillin	2.4 million U, IM	Single Dose
Secondary	Benzathine Penicillin	2.4 million U, IM	Single Dose
Early Latent	Benzathine Penicillin	2.4 million U, IM	Single Dose
Late latent (unknown duration)	Benzathine Penicillin	2.4 million U, 1x/week, IM	3 consecutive weeks
Neurosyphilis	Crystalline Penicillin	12-24 million IM, U daily 3-4 million U every 4 hours IV	10-14 days

Conclusion

Alopecia is a symptom associated with syphilis, and as in the case we presented here, can be the sole presenting symptom of syphilis. Although, the current screening tests used for the detection of syphilis have relatively high sensitivities and specificities, they are not 100% effective in the detection of syphilis. It is therefore important to be cognizant of the various potential causes of alopecia and proceed to a biopsy if clinical suspicion still remains high in a case of non-scarring alopecia.

References

- 1. Nam-Cha, S.H., et al., Alopecia syphilitica with detection of Treponema pallidum in the hair follicle. J Cutan Pathol, 2007. 34 Suppl 1: p. 37-40. [PMID: 17997737]
- 2. Ye, Y., et al., The clinical and trichoscopic features of syphilitic alopecia. J Dermatol Case Rep, 2014. 8(3): p. 78-80. [PMID: 25324910]
- 3. Pastuszczak, M. and A. Wojas-Pelc, Current standards for diagnosis and treatment of syphilis: selection of some practical issues, based on the European (IUSTI) and U.S. (CDC) guidelines. Postepy Dermatol Alergol, 2013. 30(4): p. 203-10. [PMID: 24278076]
- 4. Tipple, C. and G.P. Taylor, Syphilis testing, typing, and treatment follow-up: a new era for an old disease. Curr Opin Infect Dis, 2015. 28(1): p. 53-60. [PMID: 25485649]
- 5. Lee, J.H., et al., Comparison of an automated rapid plasma reagin (RPR) test with the conventional RPR card test in syphilis testing. BMJ Open, 2014. 4(12): p. e005664. [PMID: 25552608]
- 6. Liu, L.L., et al., Incidence and risk factors for the prozone phenomenon in serologic testing for syphilis in a large cohort. Clin Infect Dis, 2014. 59(3): p. 384-9. [PMID: 24803377]
- 7. Clement, M.E., N.L. Okeke, and C.B. Hicks, Treatment of syphilis: a systematic review. JAMA, 2014. 312(18): p. 1905-17. [PMID: 25387188]
- 8. Morales-Munera, C.E., P.A. Fuentes-Finkelstein, and M. Vall Mayans, Update on the Diagnosis and Treatment of Syphilis. Actas Dermosifiliogr, 2015. 106(1): p. 68-69. [PMID: 25245171]
- 9. Shields, M., et al., A longitudinal evaluation of Treponema pallidum PCR testing in early syphilis. BMC Infect Dis, 2012. 12: p. 353. [PMID: 23241398]
- 10. Wellinghausen, N. and H. Dietenberger, Evaluation of two automated chemiluminescence immunoassays, the LIAISON Treponema Screen and the ARCHITECT Syphilis TP, and the Treponema pallidum particle agglutination test for laboratory diagnosis of syphilis. Clin Chem Lab Med, 2011. 49(8): p. 1375-7. [PMID: 21619473]
- van Dommelen, L., et al., Confirmation of high specificity of an automated enzyme immunoassay test for serological diagnosis of syphilis: retrospective evaluation versus results after implementation. Sex Transm Dis, 2015. 42(3): p. 120-2. [PMID: 25668642]
- Castro, R., E.S. Prieto, and F. da Luz Martins Pereira, Nontreponemal tests in the diagnosis of neurosyphilis: an evaluation of the Venereal Disease Research Laboratory (VDRL) and the Rapid Plasma Reagin (RPR) tests. J Clin Lab Anal, 2008. 22(4): p. 257-61. [PMID: 18623120]