

STARCH INDUCED IMPLANT PERIAPICAL LESION:

A CASE REPORT

Rabah Nedir, DMD¹/ Mark Bischof, DMD¹/ Ollivier Pujol, PhD²/

Raymond Houriet, PhD³/ Jacky Samson, MD⁴/ Tommaso Lombardi MD, DMD, PD⁵

¹Assistant Professor, Department of Stomatology and Oral Surgery, School of Dental Medicine, University of Geneva, Switzerland; Clinical Director, Swiss Dental Clinics Group, Switzerland

²Researcher, Powder Technology Laboratory, Swiss Federal Institute of Technology, Lausanne, Switzerland

³Senior Researcher, Powder Technology Laboratory, Swiss Federal Institute of Technology, Lausanne, Switzerland

⁴Professor and Chairman, Department of Stomatology and Oral Surgery, School of Dental Medicine, University of Geneva, Switzerland

⁵Associate Professor, Laboratory of Oral Histopathology, Department of Stomatology and Oral Surgery, School of Dental Medicine, University of Geneva, Switzerland

Correspondence to:

Dr Rabah Nedir

Swiss Dental Clinics Group

CdR Clinique de Soins Dentaires SA

Rue du Collège 3, 1800 Vevey, Switzerland

Tel. +41-21-923 52 31. Fax : +41-21-922 22 98.

E-mail : rabah.nedir@swissdentalclinics.ch

STARCH INDUCED IMPLANT PERIAPICAL LESION:

A CASE REPORT

Abstract

This paper presents an unreported etiology of implant periapical lesion (IPL). The presence of an osteolytic area around the apex and around the middle portion of a stable Straumann hollow screw implant was found on periapical radiographs 3.5 years after implant placement. Case management involved curettage of the soft tissue surrounding the implant apex as well as resection of the non-osseointegrated portion of the implant. Histopathological examination revealed a connective fibrous tissue containing a dense chronic inflammatory infiltrate with a foreign body material. Polarized light microscopy and Fourier Transform Infrared (FTIR) micro-spectroscopy identified the foreign body material as starch particles. Etiology of this IPL was thus related to a foreign body reaction to starch particles. This exogenous contamination probably originated from starch coated gloves during the surgical procedure. This case report suggests that IPL may successfully be treated by debridement and implant resection instead of implant removal. Peri-implant apical soft tissue should be systematically submitted for histopathological examination.

Key words: dental implants, apical peri-implantitis, implant failure, radiolucency, granuloma, starch

Introduction

Implant periapical lesion (IPL) is a rare pathology. Single case reports¹⁻⁵ and two studies^{6,7} have been published. In 1995, Reiser & Nevins⁶ observed 10 IPLs out of 3800 examined implants. Recently, Quirynen et al.⁷ reported 10 IPLs out of 539 single implants. IPL, as a distinct entity, was first introduced by McAllister et al.⁸ Synonyms are “apical peri-implantitis” or “retrograde peri-implantitis”.⁹

IPL is usually suspected on clinical symptoms. The initial clinical manifestations of IPL are swelling and tenderness in the region of the affected implant, subsequently followed by a sinus tract.¹⁰ Radiographically, radiolucency is found at the apical portion of the implant, while the coronal portion is still supported by normal bone architecture in contact with a clinically stable implant. Differential diagnosis of IPL includes: pre-existing “necrotic bone” infection, infection from a neighbouring tooth or contamination during surgery. Pathogens (bacteria) are considered to be at the origin of IPL.

This paper reports on a case of IPL associated with a foreign body reaction to starch particles. To our best knowledge, this etiology has never been previously reported. The patient was successfully treated by implant apical resection and thorough curettage.

Case report

A 56-year-old Caucasian woman in good general health presented with a painful swelling in the area of the right cheek. Clinical examination revealed an abscess in the apical region of a 14 mm long Straumann hollow screw implant supporting a single crown in position 15 (ADA 4). A periapical radiograph showed the presence of an osteolytic area around both the apex and the middle portion of the implant (Fig 1).

Probing depth was less than 3 mm and the cervical peri-implant soft tissues showed no signs of inflammation. The implant had been placed 3.5 years ago. Before implant placement, the post-extraction healing period was 5 months. Tooth extraction was carried out due to a failing endodontic treatment with periapical radiological signs of pathosis.

Systemic antibiotics were prescribed during 12 days (amoxicillin Amoxi-Mepha[®], Mepha Pharma SA, Aesch, Switzerland; 750 mg, 3 times per day) as initial treatment. After one week, the inflammatory symptoms abated but a sinus tract with a mild purulent discharge developed in the following days. A mucoperiosteal exploratory flap was performed in order to determine the possible infection causes.

Flap reflection revealed a bone fenestration of the vestibular cortical plate with about 5 mm of residual cervical bone. An inflammatory tissue surrounded the apical portion of the implant. Upon debridement, yellowish exudate was present and an oroantral communication was observed. The perforated and hollow design of the implant apex did not allow a thorough curettage of the region. Thus, amputation of the non-osseointegrated implant extremity was decided, thereby eliminating the implant perforations. Resection was performed with a tapered fissure bur under copious irrigation. The resulting bone cavity communicated with the maxillary sinus but did not involve the adjacent teeth.

There were no post-operative complications and healing has been uneventful. Recurrence of the sinus tract was not observed over a two-year follow-up period, a nearly complete new bone formation was radiographically visible around the resected area (Fig 2). The implant is still in place, stable and functional.

Histopathological examination of the curetted tissue around the implant extremity revealed a connective fibrous tissue containing a dense chronic inflammatory

infiltrate consisting of lymphocytes, plasma cells and occasionally histiocytes (Figs 3 and 4). Numerous small round foreign bodies were observed and some of them were phagocytized by voluminous macrophages. These foreign particles were birefringent under polarized light and showed the typical aspect of a Maltese cross that might be attributed to starch.

To confirm the presence of starch particles in the inflammatory tissue, physico-chemical characterization was carried out by Fourier Transform Infrared (FTIR) micro-spectroscopy. The FTIR micro-spectroscope (Spectrum Spotlight 200 FTIR Microscope System, PerkinElmer, Schwerzenbach, Switzerland) was operated in reflection mode between 500 and 4000 cm^{-1} . This technique generates infrared signals of the whole inclusion area. It enables identification of differences in chemical compound and spatial distribution of sample features like individual contaminants.^{11,12} Tissue sample was embedded in paraffin wax and sections were deposited on a microscope glass slide coated with silver nanoparticles. The measurements were performed at room temperature.

A FTIR map of a 2.06 mm^2 cross section is showed in Fig 5a, built out of the total absorbance of the sample. It shows the specimen and the surrounding paraffin matrix. Inside the tissue, FTIR spectra were recorded at the three marked points 7, 8 and 9 (Fig 5a). They are displayed in Fig 5b. Spectrum 7 showed 2850-3000 cm^{-1} , 1450-1470 cm^{-1} and 1370-1380 cm^{-1} vibration bands. They are typical of alkane-like paraffin compound absorption. Spectrum 8 exhibited peaks at 1630 cm^{-1} and 1520 cm^{-1} , corresponding respectively to amide I (-C=O) absorption and amide II (-NH-) absorption bands of cellular tissue proteins. Compared with spectrum 8, spectrum 9 identified a new band at around 1030 cm^{-1} . This vibration, identified as C-O absorption band, is typical of a polysaccharide-like starch.

Fig 5c shows a FTIR map, obtained at high magnification, with a typical protein absorption band (1520 cm^{-1}) of a sample area. This mapping allowed discrimination between the curretted material of the IPL and the surrounding paraffin. Fig 5d presents a mapping of the spatial absorbance distribution of starch in this area. The starch distribution in the tissue is not homogeneous. Starch particles seem to agglomerate in clusters. Fig 5d shows isolated starch particles of about $15 \pm 5\ \mu\text{m}$ in diameter.

Discussion

The presence of a delimited endo-osseous radiolucency may evoke a retrograde apical pathosis. It can also be due to excessive drilling depth or it can be associated with an implant placed in a pre-existing intrabony scar lesion. These cases refer to "non-infected" retrograde peri-implant radiolucencies and they only require clinical and radiological follow-up.¹⁰

To date, little is known about the etio-pathogenesis of IPL. It appears to have a multifactorial origin.¹³⁻¹⁵ IPL is thought to be related to pre-existing bone infection¹⁶, residual root particles or foreign bodies⁶ introduced during surgery.⁴ Furthermore, overheating^{2,8}, excessive tightening, aseptic necrosis of the bone inside the hollow portion of the implant³, infected endodontic lesion from an adjacent tooth^{4,8,16,17} have all been proposed as etiologic factors. Nevertheless, these hypotheses are not supported by scientific data. Quirynen et al.⁷ observed that the incidence of IPL was significantly higher for implants with rough surfaces when compared with machined implants; however, the machined implant surface showed a higher failure rate than the rough implant surface.

In the field of oral and endodontic surgery, it is recommended in clinical practice to histologically analyze every endo-osseous or periapical lesion, in order to obtain a definitive diagnosis.¹⁸ Final diagnosis of the IPL should also fall under this recommendation. Histological examination should allow differentiation between inflammatory related and non inflammatory related pathologies (central giant cell granuloma, metastasis, etc.).^{19,20}

Histological examination was reported for five case reports. In those instances, final histopathological diagnoses were the following: acute localized osteomyelitis¹, aseptic bone necrosis²⁻⁴ and granulation tissue with acute inflammatory cells.⁵ This last diagnosis was obtained from biopsied tissues without implant removal whereas the four others were retrieved after implant removal.

IPL associated with a foreign body reaction due to starch particles has never been reported so far. Starch is added to rubber gloves as a lubricant, although many unwanted consequences are associated with its use like contact irritation and allergic reactions. Natural rubber latex proteins leak out of latex gloves and bind to surgical glove powder, this complex is responsible for latex allergy.²¹ For many years, gynaecological and visceral surgeons have already highlighted the risks of post-operative granuloma formation, due to glove powder contamination of the surgical wound.^{19,22-24} Starch granules are found in surgical wounds proportionally to the number of the surgical team members using powdered gloves.²⁵ In fact, different alternatives to powdered gloves do exist and certain medical institutions and centres have actually chosen to eliminate glove powder from their environment.²⁶

In the oral surgery literature, only a single case of foreign body reaction to starch particles was reported following extraction. It appeared as a firm submucosal nodule,

5 mm in diameter, in the buccal sulcus, which was excised 20 days after teeth extraction.²⁷

In dental implantology, the risk of starch contamination by medical gloves was evoked by Field²¹ and Belvedere and Lambert²⁸ as a source of complications. Although manufacturers recommend fixture insertion with a specific instrument, one cannot totally rule out some accidental contact. Moreover, most surgical instruments, like spirals drills or depth gauge, are manipulated with gloves.

Many foreign bodies are able to produce an inflammatory reaction and could explain some implant failures. They can be introduced at the implant site during dental procedures (endodontic and restorative obturation materials), extraction procedures or implant surgery.

In order to prevent the occurrence of the IPLs, it is important to thoroughly debride the infected socket following extraction. Implants have to preferentially be placed after bony socket maturation when lesions are no longer visible on radiographs.¹⁰ It also appears that foreign bodies may still dwell inside the bone and trigger IPL, even after thorough debridement, irrigation of the extraction sockets and sufficient healing time. Contamination of the implant site should be prevented. Implants should never come in contact with saliva, teeth, oral tissues or surgeon's gloves. Both careful implant site selection and surgical technique may further reduce the incidence of infected IPL.⁶

In the presence of an IPL, the choice of therapy is difficult. So far, there is no consensus report to which therapy should be favoured. To date, several treatment options are reported, ranging from the use of anti-infectious medication, lesion excision and debridement to implant removal. Different factors may influence treatment choice: implant stability, peri-implant probing depth, adjacent teeth status,

implant position and angulations. Moreover, the type and quality of the prosthetic rehabilitation should also be taken into account.

Anti-infectious therapy may be effective to treat the acute phase and adjunctive to the surgical treatment. This will rarely suffice to totally eliminate the underlying pathology.

Implant removal will usually be accompanied by considerable bone loss and the remaining bone along with the surrounding soft tissues will be altered. After several months of healing, an autologous bone and/or gingival graft may be necessary. The subsequent healing period as well as the difficulties inherent to this type of complex treatment may discourage the patient from pursuing implant therapy. Hence, a conservative surgical treatment, as described in this paper, may be proposed as a valid attemptable treatment option, as opposed to fixture removal. Furthermore, this will provide a histological diagnosis which may lead to complementary treatments. This treatment approach has the advantage of maintaining the implant in function.

Resection of the apical portion of the implant may be indicated when the geometry (implant type) does not allow proper debridement. Therefore, implant removal should be considered only as a last resort, when debridement has proven unsuccessful.

Any adjacent tooth which could be the source of infection should be considered for extraction if untreatable. It may therefore be possible to eliminate retrograde infection by debridement and hence maintain the implant if it clinically remains stable and if the infection is restricted to the endo-osseous region.^{8,9,29} Implant resection should facilitate complete mechanical debridement of the inflammatory granulation tissue.

Apex sectioning may be especially indicated in cases of perforated implant type.^{6,10,30} In addition to the conservative surgery, some authors have proposed the use of biomaterials.^{5,6,10,29} Literature data does not warrant their systematic use.

In the present case, only implant resection and debridement were performed. This treatment has been attempted for the following reasons: implant stability, presence of residual cervical bone in the coronal part, bone fenestration instead of dehiscence. Graft material has not been used because of the presence of cervical residual bone. After two years follow-up of the resective surgery, the radiograph shows a normal crestal bone level.

Conclusion

Exogenous contamination may provoke an IPL of an osseointegrated implant even several years after implant insertion. Although the occurrence of starch induced granuloma seems to be rare, the use of starch coated gloves should be avoided during surgical procedures. In order to identify the etiology of the pathosis, curetted tissues should be systematically histopathologically analyzed. The development of such a lesion should be considered as a complication rather than a failure. In this case, the IPL has been treated by implant debridement and resection, rather than by implant removal. This conservative approach has been effective and has maintained the implant in function.

Acknowledgement

Serge Szmukler-Moncler and Nathalie Nurdin are acknowledged for their efficient help during manuscript reviewing.

References

1. Piattelli A, Scarano A, Piattelli M. Abscess formation around the apex of a maxillary root form implant: clinical and microscopical aspects. A case report. *J Periodontol* 1995;66:899-903.
2. Piattelli A, Scarano A, Balleri P, Favero G. A. Clinical and histologic evaluation of an active "implant periapical lesion": a case report. *Int J Oral Maxillofac Implants* 1998;13:713-716.
3. Piattelli A, Scarano A, Piattelli M, Podda G. Implant periapical lesions: clinical, histologic, and histochemical aspects. A case report. *Int J Periodontics Restor Dent* 1998;18:181-187.
4. Scarano A, Di Domizio P, Petrone G, Lezzi G, Piattelli A. Implant periapical lesion: a clinical and histologic case report. *J Oral Implantology* 2000;26:109-113.
5. Chaffee NR, Lowden K., Tiffée JC, Cooper LF. Periapical abscess formation and resolution adjacent to dental implants: a clinical report. *J Prosthet Dent* 2001;85:109-112.
6. Reiser GM, Nevins M. The implant periapical lesion: etiology, prevention, and treatment. *Compend Contin Educ Dent* 1995;16:768-777.
7. Quirynen M, Vogels R, Alsaadi G, Naert I, Jacobs R, van Steenberghe D. Predisposing conditions for retrograde peri-implantitis, and treatment suggestions. *Clin Oral Implants Res* 2005;16:599-608.
8. McAllister BS, Masters D, Meffert RM. Treatment of implants demonstrating periapical radiolucencies. *Pract Periodontics Aesthet Dent* 1992;4:37-41.
9. Flanagan D. Apical (retrograde) peri-implantitis: a case report of an active lesion. *J Oral Implantol* 2002;28:92-96.
10. Jalbout ZN, Tarnow DP. The implant periapical lesion: four case reports and review of the literature. *Pract Proced Aesthet Dent* 2001;13:107-112.

11. Lamontagne J, Durrieu F, Planche JP, Mouillet V, Kister J. Direct and continuous methodological approach to study the ageing of fossil organic material by infrared microspectrometry imaging: application to polymer modified bitumen. *Anal Chim Acta* 2001;444:241-250.
12. Skrtic D, Antonucci JM, Eanes ED, Eidelman N. Dental composites based on hybrid and surface-modified amorphous calcium phosphates. *Biomater* 2004;25:1141-1150.
13. Esposito M, Hirsch JM, Lekholm U, Thomsen P. Biological factors contributing to failures of osseointegrated oral implants (I): success criteria and epidemiology. *Eur J Oral Sci* 1998;106:527-551.
14. Esposito M, Hirsch JM, Lekholm U, Thomsen P. Biological factors contributing to failures of osseointegrated oral implants (II): etiopathogenesis. *Eur J Oral Sci* 1998;106:721-764.
15. Oh TJ, Yoon J, Wang HL. Management of the implant periapical lesion: a case report. *Implant Dent* 2003;12:41-46.
16. Sussman HI. Periapical implant pathology. *J Oral Implantol* 1998;24:133-138.
17. Tseng CC, Chen YHM, Pang IC, Weber HP. Peri-implant pathology caused by periapical lesion of an adjacent natural tooth: a case report. *Int J Oral Maxillofac Implants* 2005;20:632-635.
18. Talacko AA, Alderd MJ, Abott PV, Smith ACH, Nerwich AH. Periapical Biopsy? *Oral Surg Oral Med Oral Pathol* 2000;89:532-533.
19. Selden HS. Central giant cell granuloma: a troublesome lesion. *J Endod* 2000;26:371-373.
20. Levi PA, Kim DM, Harsfield SL, Jacobson ER. Squamous cell carcinoma presenting as an endodontic-periodontic lesion. *J Periodontol* 2005;76:1798-1804.

21. Field EA. The use of powdered gloves in dental practice: a cause for concern? *J Dent* 1997;25:209-214.
22. Antopol W. Lycopodium granuloma: its clinical and pathological significance, together with a note on granuloma produced by talc. *Arch Pathol* 1933;16:326-331.
23. Neely J, Davies JD. Starch granulomatosis of the peritoneum. *Br Med J* 1971;3:625-629.
24. Steiner Z, Mogilner J, Bahar Y. Gastric outlet obstruction due to starch granuloma and peritonitis. *Eur J Pediatr Surg* 1996;6:364-366.
25. Hunt TK, Slavin JP, Goodsan WH. Starch powder contamination of surgical wounds. *Arch of Surg* 1994;129:825-828.
26. Cuming RG. Reducing the hazards of exposure to cornstarch glove powder. *AORN J* 2002;76:294-295.
27. Wilson DF, Garach V. Surgical glove starch granuloma. *Oral Surg Oral Med Oral Pathol* 1981;51:342-345.
28. Belvedere PC, Lambert DL. Negative effects of powdered latex gloves in clinical dentistry. *J Long Term Eff Med Implants* 1994;4:119-125.
29. Bretz WAG, Matuck AN, de Oliveira G, Moretti AJ, Bretz WA. Treatment of retrograde peri-implantitis: clinical report. *Implant Dent* 1997;6:287-290.
30. Shaffer MD, Juruaz DA, Haggerty PC. The effect of periradicular endodontic pathosis on the apical region of adjacent implants. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 1998;86:578-581.



Fig 1 Preoperative radiograph revealing the presence of an implant periapical lesion around a hollow screw perforated implant.

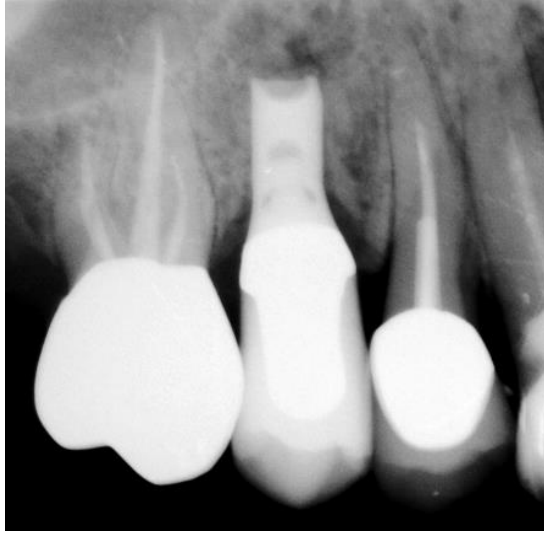


Fig 2 Radiograph taken two years after implant resection. Note the nearly complete new bone formation.

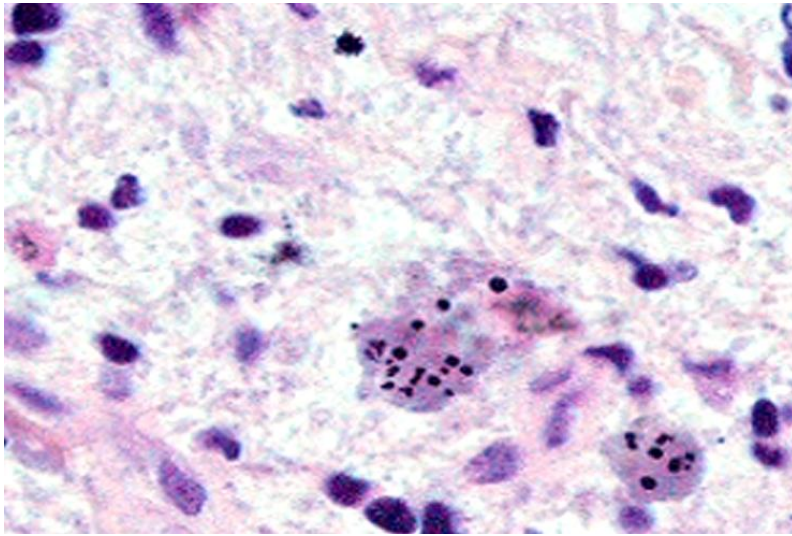


Fig 3 Histopathological examination of the tissue biopsy. Foreign particles were mainly phagocytized by macrophages.

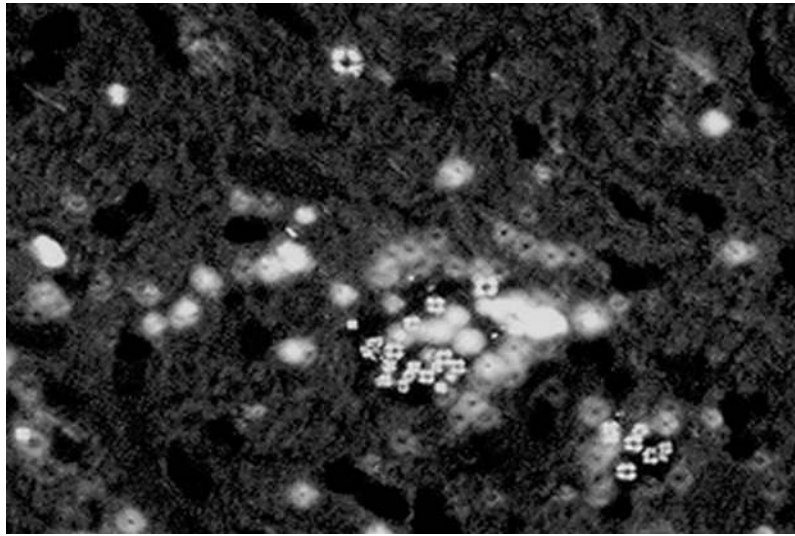


Fig 4 Histopathological examination under polarized light of the tissue biopsy. The foreign particles showed the typical Maltese cross aspect.

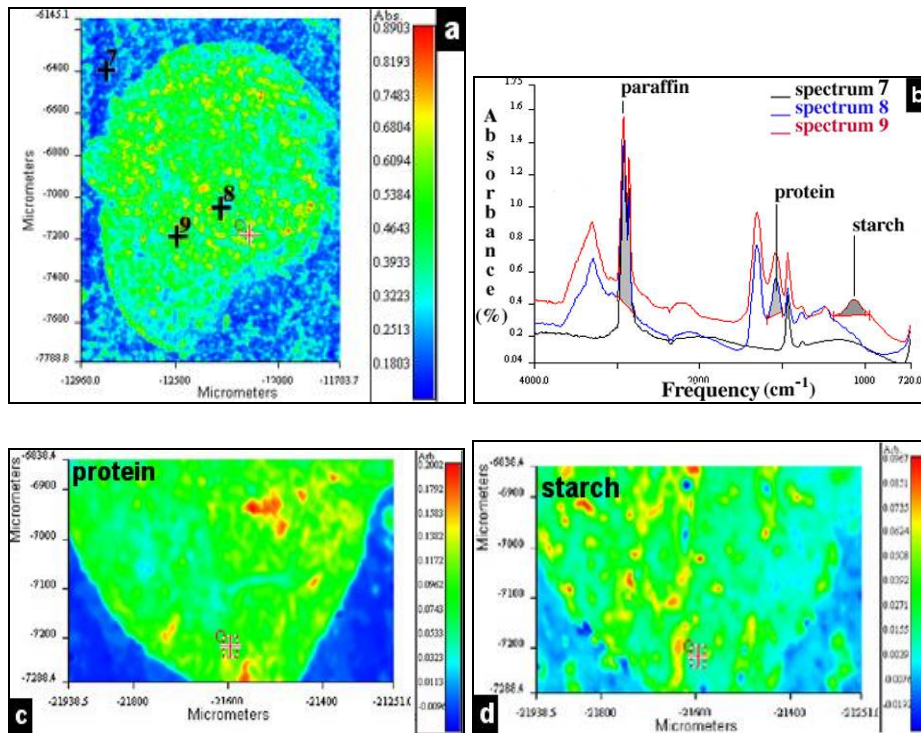


Fig 5 FTIR analysis of the biopsy.

- (a) FTIR map of the total absorbance of a granuloma cross section. The crosses marked 7, 8 and 9 correspond to three different points of analysis.
- (b) FTIR spectra recorded at the points marked 7, 8 and 9 in Fig 5a.
- (c) FTIR map obtained with a typical protein absorption band (1520 cm^{-1}).
- (d) FTIR map obtained with a starch absorption band (1030 cm^{-1}) of the same area.