Oral carcinoma cuniculatum, an unacquainted variant of oral squamous cell carcinoma: A systematic review

Amina Fouad Farag1,*, Dalia Ali Abou-Alnour2, Noha Saleh Abu-Taleb3

1Oral Pathology Department, Faculty of Dentistry, October 6 University, Egypt
2Oral Radiology Department, Faculty of Dentistry, Modern University for Technology and Information, Egypt
3Oral and Maxillofacial Radiology Department, Faculty of Dentistry, Cairo University, Egypt

ABSTRACT

Purpose: Oral carcinoma cuniculatum is a rare well-differentiated variant of oral squamous cell carcinoma. The purpose was to systematically review its unique features to differentiate it from other variants as verrucous carcinoma, papillary squamous cell carcinoma and well-differentiated squamous cell carcinoma.

Materials and Methods: A systematic review was performed using MEDLINE, Dentistry and Oral Sciences Source and PubMed databases and any existing articles related to the research subject missed in the search strategy to screen ones reporting cases occurring exclusively in the oral cavity in English literature. Variables analyzed included clinical, etiologic, imaging, histopathological features, treatment, follow-up and survival rates.

Results: From 229 hits, 17 articles with 43 cases were included in the systematic review. Clinically it showed a female predilection with pain and/or ulceration of a relatively long duration and exudation being the most common symptoms. Histologically, it showed more endophytic features comprising well-differentiated squamous epithelium with absent or minimal cytological atypia and multiple keratin filled crypts or cuniculus. Inflammatory stromal reaction and discharging abscesses were reported in most of the cases. Bone destruction was predominant in most imaging features. Complete surgical resection with a safety margin was the treatment of choice in most of the cases with few recorded recurrence cases.

Conclusion: Apprehensive knowledge of oral carcinoma cuniculatum unique features is essential to avoid its misdiagnosis and provide proper treatment especially for recurrent cases. (Imaging Sci Dent 2018; 48: 233-44)

KEY WORDS: Mouth; Carcinoma, Squamous Cell; Carcinoma, Verrucous

Introduction

Oral carcinoma cuniculatum (CC) is a rare, unacquainted variant of oral squamous cell carcinoma (OSCC).1 It is similar to cutaneous CC, which was first described by Aird et al. in 1954, in its clinicopathological findings and biological behavior.2

Apart from the 3 cases of cutaneous CC described by Aird et al.2 on the plantar aspect of the foot, this type of tumor has been reported at other sites, including the face, esophagus, abdomen, palm, leg, penis, and cervix.3-7

In 1977, Flieger and Owiński8 were the first to report CC in the oral cavity, but they failed to establish the relationship of this tumor to ordinary OSCC because of its unique clinical features, including locally aggressive behavior with both endophytic and exophytic growth patterns. In addition, it has a unique histological architecture that comprises well-differentiated epithelial cells with no or rare cytological atypia, unlike OSCC.9

Oral CC is a locally aggressive tumor formed of multiple, branching, keratin-filled crypts (rabbit burrows or cuniculi) lined by well-differentiated hyperplastic stratified squamous epithelium with minimal cellular atypia, but evident local bone invasion and rare metastasis.10

The World Health Organization (WHO) 2005 classi-
Oral carcinoma cuniculatum, an unacquainted variant of oral squamous cell carcinoma: A systematic review

Fication of tumors included oral CC as a new variant of OSCC characterized by the presence of keratin-filled branching crypts and keratin cores, resulting in the tumor having a cuniculatum architecture, similar to rabbit burrows.\textsuperscript{11}

Although it is a variant of OSCC, the diagnosis of oral

| Author            | No | Age/ Gender | Site                  | Clinical appearance/signs and symptoms                                                                 | Duration (months) | Possible etiologic or predisposing factors                                      |
|-------------------|----|-------------|-----------------------|--------------------------------------------------------------------------------------------------------|-------------------|--------------------------------------------------------------------------------|
| Allon et al.\textsuperscript{25} | 1  | 56/M        | Ant Max gingiva       | Red, exophytic, nodular soft lesion with focal white patches, mobile teeth                          | 6                 | Smoking                                                                      |
| Heasman et al.\textsuperscript{26} | 1  | 44/F        | Mand gingiva          | Swelling, pain, soreness, endophytic ulceration, white patches, area of exposed bone on the right alveolar ridge | N/A               | Smoking and alcohol consumption                                               |
| Raguse et al.\textsuperscript{27} | 1  | 81/F        | Ant Mand region (2 lesions; 1 right and 1 left) | Pain                                                                                                  | N/A               | None                                                                          |
| Kruse and Grätz\textsuperscript{28} | 1  | 74/F        | Ant Max gingiva       | Ulceration                                                                                           | 3                 | None                                                                          |
| Hutton et al.\textsuperscript{29}  | 1  | 7/F         | Ant Max gingiva       | Swelling, ulceration, white verrucous areas                                                          | N/A               | None                                                                          |
| Pons et al.\textsuperscript{10}   | 3  | 65.7/ 3M    | Ant mandible          | Dental pain, Carcinoma cuniculatum developed on an Mand cyst, dental pain                            | 3-8               | None                                                                          |
| Sepulveda et al.\textsuperscript{30} | 1  | 49/M        | Left Max region       | Swelling, pain, bone weakness of palate, pus, inflammation and purulent fistulae                   | N/A               | N/A                                                                          |
| Sun et al.\textsuperscript{9}      | 15 | 67/7M, 8F  | Tongue: 8 Vestibule: 1 | Pain: 12, ulceration: 8, induration: 7, swelling: 6, white patches: 5, bleeding: 4, exudation: 1, leukoplakia: 3 patients, leukoplakia and lichen planus: 1, nodular or keratotic patch co-existed in some cases (10 cases were exophytic, 5 cases were endophytic) | 1-24 (mean 8.5)  | Smoking or Alcohol: 7                                                         |
| Suzuki et al.\textsuperscript{11}  | 1  | 68/M        | Mand gingiva          | Pain, exophytic verrucous leukoplakia, exudation                                                       | Over 19           | Smoking and alcohol consumption                                               |
| Thavaraj et al.\textsuperscript{32} | 1  | 61/M        | Tongue                | Swelling without ulceration, restriction of tongue mobility, dysphagia, difficulty with articulation | 12                | N/A                                                                          |
| Fonseca et al.\textsuperscript{33} | 2  | 62/F        | Ant Mand gingiva      | Pain, swelling, exudation, reddish lesion with numerous whitish areas and small nodules            | 2                 | None                                                                          |
| Goh et al.\textsuperscript{3}     | 1  | 62/M        | Tongue                | Ulceration                                                                                           | 72                | Smoking                                                                      |
| Padilla and Murrah\textsuperscript{13} | 10 | 71.9/ 3M, 7F | Mand gingiva: 7 Max gingiva: 2 Vestibule: 1         | Erythroleukoplakia: 1, erythroplakia with induration: 1, red and white lesion: 3, swelling: 1, gingival enlargement: 1, ulceration: 1, exudation: 1, lichen planus: 1, some cases showed cobblestone-like or pebbly exophytic surface | 3, 11 in two cases N/A in other cases | Smoking: 1 Smokeless tobacco: 1                                               |
| Shakil et al.\textsuperscript{34}  | 1  | 63/F        | Buccal mucosa         | Reddish white ulceroproliferative cauliflower-like growth                                           | 8                 | N/A                                                                          |
| Shapiro et al.\textsuperscript{35} | 1  | 71/F        | Mand gingiva          | White and red patches, pain, history of leukoplakia                                                  | Over 108          | N/A                                                                          |
| Shay et al.\textsuperscript{36}    | 1  | 58/M        | Mand gingiva, parotid gland, deep parapharyngeal muscles | Swelling, induration, exudation                                                               | 36                | None                                                                          |
| Datar et al.\textsuperscript{1}   | 1  | 58/F        | Mand gingiva          | Ulceration, white and red growth, pebbly surface                                                    | 3                 | None                                                                          |

Ant: anterior, M: male, F: female, Max: maxillary, Mand: mandibular, N/A: not available, Post: posterior, *: mean age of collected data, †: mean duration of collected data
CC has remained very difficult and challenging because of clinicians’ lack of awareness and familiarity with the tumor, which in turn has resulted in under-reporting of oral CC and its suggested low incidence. Correlation of histological findings with clinical and radiographic features is essential for the diagnosis of oral CC and to avoid confusing it with other tumors, mainly verrucous carcinoma (VC), well-differentiated SCC, papillary squamous cell carcinoma (PSCC), and other tumors included in the histological differential diagnosis of oral CC.9-13

The aim of the current study was to conduct a systematic review of the studies describing oral CC by pooling patients from multiple studies, with the goal of illustrating and further distinguishing the clinical features, etiologic agents, predisposing factors, imaging features, histopathological findings, treatment, and follow-up and survival rates of this distinctive rare tumor.

Materials and Methods

A systematic review of the published cases of oral CC in the literature was performed according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) statement, an evidence-based set of items for presenting these types of studies.14

Search strategy

A broad literature searches in Medline (full text), the Dentistry and Oral Sciences Source via EbscoHost research databases, and PubMed was performed. Relevant studies published from 1954, the first mention of CC in the literature, to January 2018 were identified. These electronic databases were searched using both MeSH terms and free text words. A search strategy was developed that combined the terms (carcinoma cuniculatum) AND (palatal OR palate OR gingiva OR tongue OR oral OR maxilla OR maxillofacial OR mandible OR mandibular OR ramus OR alveolar OR jaw). In the advanced search, all fields were included in the PubMed search and all texts in Dentistry and Oral Sciences Source databases and in Medline (for full texts). Three independent researchers, including 1 oral pathologist and 2 oral radiologists, examined the retrieved hits and discarded duplicates. The references of relevant articles were screened for papers missed in the initial search.

Selection criteria

The inclusion criteria were: articles published in the English language, with human subjects, that described oral CC in the oral or maxillofacial region. All titles and abstracts were analyzed. Full-text articles were obtained only for abstracts found to satisfy the inclusion criteria. Articles published in languages other than English, those regarding non-human subjects, and studies on any variant of OSCC other than oral CC were excluded from the review.

Data extraction

Individual and aggregated data describing the clinical features, etiologic agents, predisposing factors, radiographic features, histopathological and immunohistochemical features, treatment, and follow-up and survival rates of patients with oral CC were collected. Other variables were also gathered, including author names, publication year, and number of cases. The quality and internal validity of the studies were independently assessed by the authors according to the CARE (CasEReport) Statement and Checklist Tool (Fig. 1). Any disagreement among the authors was resolved by discussion.

Results

Search results

The search strategy retrieved 229 hits; 27 were relevant for our study, of which 10 studies were excluded because they were not written in the English language or they did not contain any oral cases. This resulted in a total of 17 articles that were included in the systematic review, which presented 43 cases (Fig. 2).

Clinical features

The clinical features of the included cases are presented in Table 1. The number of included cases was 43 (19 males and 24 females) with a mean age of 62.7 years (range, 7-81 years). The most common sites were the mandibular gingiva in 19 cases (44.2%), followed by the tongue in 10 cases (23.2%), the maxillary gingiva in 6 cases (13.9%), the mandible in 4 cases (9.3%), the vestibule in 2 cases (4.6%), the maxillary region in 1 case (2.3%), and the buccal mucosa in 1 case (2.3%).

The clinical presentations of the tumors varied, and patients presented with various symptoms as single entities or co-existing with each other. The most common symptom was pain in 21 cases (48.8%), followed by ulceration in 16 cases (37.2%), swelling in 13 cases (30.2%), induration in 9 cases (20.9%), a red and white lesion in 8 cases (18.6%), white patches/areas in 7 cases (16.3%), exuda-
tion in 7 cases (16.3%), leukoplakia in 5 cases (11.6%), bleeding in 4 cases (9.3%), erythroleukoplakia in 2 cases (4.6%), lichen planus in 1 case (2.3%), leukoplakia and lichen planus in 1 case (2.3%), soreness in 1 case (2.3%), bone exposure in 1 case (2.3%), teeth mobility in 1 case (2.3%), restriction of tongue mobility with dysphagia and
difficulty with articulation in 1 case (2.3%), and gingival enlargement in 1 case (2.3%). Roughly one-fourth of the cases showed a nodular, pebbly, cobblestone, verrucous, or cauliflower-like surface.

The lesion duration was recorded in most cases, with a mean of 19.5 months. However, the duration of the lesion was not recorded in 13 cases.

Thirteen cases (30.2%) were reported to be in tobacco smokers, alcoholics, or both, while 1 case (2.3%) reported to be in a smokeless tobacco user and in 26 cases (60.5%), the patient did not consume any alcohol or smoked. The corresponding information was missing in the remaining cases.

In addition, the size of the lesion was reported in only 7 of the 43 cases; it ranged from 1 to 6 cm.

**Imaging findings**

The imaging findings of the tumors are listed in Table 2. In 25 of the 43 cases, no imaging modality was used. Computed tomography (CT) was the modality of choice in 5 cases involving the bone, where it showed osteolytic lesions. The tumors were occasionally well-defined, but in 14 cases the lesions were almost ill-defined and caused bone destruction (32%). Erosion of the underlying bone occurred in tumors located within soft tissue. Magnetic resonance imaging (MRI) was the modality of choice in 3 cases involving the tongue and the gingiva to detect the actual extent of the lesion. MRI showed tumorous lesions of heterogeneous intensity affecting most of the tongue, but sparing most of the extrinsic muscles, and also showed inflammation in the gingiva. Both CT and MRI were used together in 2 other cases. Panoramic radiographs were used in 8 cases as screening tools through which the abnormality was detected. Panoramic findings included an ill-defined irregular osteolytic radiolucent lesion in 6 cases (13.9%), erosion of the superficial cortical bone in 1 case (2.3%), and a ‘moth-eaten’ osteolytic lesion in 1 case (2.3%).

**Preoperative diagnosis**

The preoperative diagnoses are listed in Table 2. They ranged from inflammatory conditions such as granuloma, abscess, or osteomyelitis in 5 cases to reactive epithelium with hyperkeratosis or pseudoepithelomatous hyperplasia in 1 case and malignancy, including VC and OSCC, in 5 cases. The diagnosis changed from an inflammatory condition to malignancy in 3 cases. In 3 cases, the preoperative diagnosis was a keratocyst, and it was a mandibular cyst in 1 case. No preoperative diagnoses were reported in 28 cases.

**Histopathological features**

The histopathological features of the tumors are listed in Table 3. Histopathological features consistent with oral CC were reported in all 43 cases (100%). These comprised well-differentiated squamous epithelium that extended deep into the connective tissue with multiple, branching, keratin-filled crypts (rabbit burrows or cuniculi), with absent or mild cytological atypia that was usually limited to the basal and parabasal layers. In 39 cases (90.7%), an inflammatory stromal reaction mainly consisting of neutrophils was observed, along with discharging abscesses reported in 28 cases (65.1%).

The margins were free from tumor in 35 cases (81.4%) and showed infiltration in 2 cases (4.6%); however, no data were reported from the remaining 6 cases (13.9%). Cases that were reported to involve infiltration, invasion, destruction, and penetration of the surrounding tissue and cases with deep projections or down-growths with an invasive front were considered to be locally invasive. All cases showed local invasion, which did not extend any further in 39 cases (90.7%), while the remaining 4 cases showed metastasis: 3 to the regional lymph nodes and 1 to the lungs.

**Immunohistochemical findings or special staining**

The immunohistochemical findings are listed in Table 3. Neither special nor immunohistochemical staining was used in 37 cases (86%). Positive Gomori methenamine silver (GMS) and periodic Acid-Schiff (PAS) staining indicated the presence of fungal organisms resembling Candida both within the crypts and the epithelium in 1 case.
| Author                  | Modality                        | Appearance                                                                 | Bone destruction | Preoperative Diagnosis/Initial Diagnosis                                      |
|-------------------------|---------------------------------|-----------------------------------------------------------------------------|------------------|--------------------------------------------------------------------------------|
| Allon et al.            | Panoramic radiography          | Unilocular radiolucent lesion with irregular borders in the Ant Max from the right lateral incisor to the left canine; partial resorption of the lamina dura of the right lateral incisor and both central incisors. | Yes              | N/A                                                                            |
| Heasman et al.          | CT (bone window)               | Large osteolytic lesion with poorly defined margins and irregular trabeculation in the Ant Max where the Max sinuses on either side were not involved. | Yes              | Osteomyelitis                                                                 |
| Raguse et al.           | Panoramic radiography          | Mott-eaten appearance of the right alveolar ridge                           | Yes              | Osteomyelitis, well-differentiated OSCC                                         |
| Kruse and Grätz         | MRI                             | Low-intensity mass causing destruction of the bone of the upper Ant jaw with infiltration into the Max sinus. | Yes              | OSCC                                                                           |
| Hutton et al.           | CT                              | Swelling buccal to the upper right and left permanent central incisors in addition to lymphadenopathy around the left carotid artery and jugular vein. | Yes              | Abscess, well-differentiated OSCC                                               |
| Pons et al.             | Case 1: panoramic radiography  | Osteolytic lesion in the area of the second and third Mand molars           | None             | Inflammatory granuloma or abscess                                              |
|                         | Case 2: CT (bone window)       | Well-defined lesion with marginal cortical osteolysis                       | Yes              | Ant Mand cyst                                                                 |
|                         | Case 3: CT (bone window)       | Well-defined osteolytic lesion of the left Mand ramus, Max, zygomatic, and petrous bones, as well as the infra temporal fossa. | Yes              | Keratocyst                                                                    |
| Sepulveda et al.        | CT                              | Great expansive destructive and heterodense process of the left half-face with current extension into the skull base, left Max region, part of the Ant zygomatic bone and lower lateral wall of the left orbital cavity and sphenoid of the same side. | Yes              | Max osteomyelitis                                                             |
| Sun et al.              | N/A                             | N/A                                                                         | N/A              | Osteomyelitis with leukoplakia                                                  |
| Suzuki et al.           | CT (soft tissue window)         | Osteolytic lesion of the left Mand with sequestration of the alveolar bone. | Yes              | Osteomyelitis with leukoplakia                                                  |
| Thavaraj et al.         | MRI                             | Extensive mass with hyperintense margin replacing almost all of the tongue, sparing the extrinsic muscles apart from the superior genioglossus. | None             | N/A                                                                            |
| Fonseca et al.          | Case 1: panoramic radiography  | Initial: ill-defined osteolytic area on the right Mand extending from the incisors to the ascending ramus. Follow-up: invasion of the left side of the Mand by an ill-defined osteolytic lesion extending up to the molar region, causing extensive bone destruction. | Yes              | Infected orthokeratinized keratocyst                                           |
| Goh et al.              | MRI                             | Extensive destruction of the left Max with ill-defined borders              | None             | Osteomyelitis/abscess                                                          |
| Padilla and Murrań       | Panoramic radiography in 1 case | Erosion of the superficial cortical bone                                     | None             | Pseudoepithelomatous hyperplasia                                               |
| Shakil et al.           | N/A                             | Hyper intensity tumorous lesion within the left posterolateral aspect of the tongue | None             | N/A                                                                            |
| Shapiro et al.          | CT (soft tissue and bone window) | Initial: Destructive lesion in the Mand body. Follow-up: thick-walked, enhancing fluid collection anteroinferior to the Mand extending into the submandibular space with erosion and sclerosis of the anterior cortex of the Mand and multiple mildly enlarged level I nodes. | Yes              | Multiple misdiagnoses: dental abscess, osteomyelitis, verrucous hyperplasia and well-differentiated OSCC |
| Shuy et al.             | CT and MRI                      | Very large, destructive heterogeneous mass centered on the Mand obliterating the masticator space, involving the parotid gland and extending to the deep parapharyngeal muscles. Extensive surrounding inflammation and phlegmon, abutting the internal carotid artery just below the skull base and obliterating the infra temporal fossa. | Yes              | Superinfected odontogenic keratocyst                                           |
| Datar et al.            | Panoramic radiography          | Irregular radiolucent lesion involving the alveolar bone around the Mand second molar | Yes              | Well-differentiated OSCC                                                       |

Mand: mandibular, Max: maxillary, Ant: anterior, CT: computed tomography, MRI: magnetic resonance imaging, N/A: not available, OSCC: oral squamous cell carcinoma, VC: verrucous carcinoma
In 1 case, there was a high rate of positive p53 immu-
noexpression (indicative of mutations in the TP53 gene,
where tumors with more than 10% positively stained p53
cells are highly suggestive of malignant features), which
was found in more than 90% of basal and suprabasal
cells, whereas it was minimally expressed with no muta-
tions in another case and showed negative results for ex-
pression in 2 other cases. High positive Ki-67 expression
(a protein that increases with cell division preparation; the
more positive cells for Ki-67, the more quickly a tumor is
dividing, indicating the uncontrolled cell proliferation as-
associated with malignant cells) in the basal and suprabasal
layers was reported in 1 case, while low levels of expres-
sion were reported in 4 cases. In addition, findings for

| Author          | Well-differentiated squamous epithelium with multiple, branching, keratin-filled crypts (rabbit burrows or cuniculus) | Cytological atypia | Discharging abscesses / Stromal reaction | Margins | Immunohistochemical findings or special stains | Growth pattern |
|-----------------|-------------------------------------------------------------------------------------------------------------|-------------------|------------------------------------------|---------|-----------------------------------------------|---------------|
| Allon et al.25  | Yes                                                                                                          | Mild              | Yes / N/A                                | N/A     | Negative p53                                  | Exophytic     |
|                 |                                                                                                              |                   |                                         |         | Negative HPV                                  |               |
| Heasman et al.26| Yes                                                                                                          | Absent to mild    | Yes / Intense                           | N/A     | N/A                                           | N/A           |
| Raguse et al.27 | Yes                                                                                                          | Mild              | Yes / Prominent                         | Free    | N/A                                           | N/A           |
| Kruse and Grätz28| Yes                                                                                                          | Mild              | N/A                                     | N/A     | Positive Ki-67 in basal and suprabasal layers, Negative p53 | Both exophytic and endophytic |
|                 |                                                                                                              |                   |                                         |         | Negative HPV                                  |               |
| Hutton et al.29  | Yes                                                                                                          | N/A               | N/A                                      | Free    | N/A                                           | Endophytic    |
| Pons et al.10    | Yes                                                                                                          | Absent to mild    | Yes / N/A                                | Free: 2 | N/A                                           | Both exophytic and endophytic |
|                 |                                                                                                              |                   |                                         | Infiltrated: 1 |         |               |               |
| Sepulveda et al.26| Yes                                                                                                          | N/A               | Yes / Yes                               | N/A     | N/A                                           | Endophytic    |
| Sun et al.5      | Yes                                                                                                          | Absent            | Yes / Moderate                           | Free    | N/A                                           | N/A           |
| Suzuki et al.31  | Yes                                                                                                          | Mild              | N/A / Prominent                         | Free    | N/A                                           | N/A           |
| Thavaraj et al.32| Yes                                                                                                          | Mild              | Yes / Moderate                           | N/A     | Positive Ki-67 in basal & suprabasal layers (low, <15% of cells), Positive p53 in >90% of basal and suprabasal cells Negative HPV | Endophytic    |
| Fonseca et al.33 | Yes                                                                                                          | Mild              | Yes / Prominent                         | Free    | Positive Ki-67 in basal and suprabasal layers (low, <5% of cells) | Exophytic    |
| Goh et al.3      | Yes                                                                                                          | Absent to mild    | Yes / Intense                           | Free    | Positive GMS and PAS for fungal organisms, pseudohyphae resembling candida both within the crypts and the epithelium Negative p16 Positive weak to moderate p53 expression, but with no mutations | Endophytic    |
| Padilla and Murrah13 | Yes                                                                                                      | Absent to mild     | N/A / Prominent                         | Free    | N/A                                           | Both exophytic and endophytic |
| Shakil et al.34  | Yes                                                                                                          | N/A               | N/A / Intense                           | N/A     | N/A                                           | Both exophytic and endophytic |
| Shapiro et al.35 | Yes                                                                                                          | Mild              | Yes / Prominent                         | Infiltrated | N/A   | Endophytic                                   |               |
| Shay et al.36    | Yes                                                                                                          | Mild              | Yes / Prominent                         | Free    | N/A                                           | Endophytic    |
| Datar et al.1    | Yes                                                                                                          | Mild              | None                                     | Free    | N/A                                           | Endophytic    |

GMS: Gomori methenamine silver, HPV: human papilloma virus, N/A: not available, PAS: periodic acid-schiff
Oral carcinoma cuniculatum, an unacquainted variant of oral squamous cell carcinoma: A systematic review

*p16 expression (a protein that acts as a tumor suppressor by decelerating the cell progression from the G1 phase to the S phase) were negative in the single tested case. Furthermore, human papillomavirus (HPV) was tested in 3 cases, with negative results.

Growth pattern

Both exophytic and endophytic growth patterns were histologically reported in 15 cases (34.9%), while an endophytic pattern was reported in 7 cases (16.3%), an exophytic pattern was reported in 2 cases (4.6%), and records were missing in the remaining cases.

Treatment and follow-up

Results are shown in Table 4. Complete surgical resection with a safety margin was the treatment of choice in 42 cases (97.7%), with a subtotal maxillectomy or mandibulectomy performed in cases with bone invasion, in addition to neck dissection in cases with enlarged lymph nodes. Only one case was treated by chemotherapy and radiotherapy.

Following the surgical removal of the tumor, 37 patients (86%) continued to attend follow-up appointments. The follow-up duration ranged from 4 months to 14 years, with a mean of 31.1 months. Of the 37 patients, 29 (67.4%) were reported to be disease-free and did not show recurrence or mortality. However, 4 cases (9.3%) showed local recurrence. In the cases of recurrence, the treatment was second surgical removal in 1 case, second surgical removal with concurrent chemotherapy in 1 case, concurrent chemoradiotherapy in 1 case, and chemotherapy in 1 case.

Table 4. Treatment, follow-up, and status at follow-up of cases included based on the search strategy

| Author | Treatment of choice | Follow-up (months) | Status at follow-up (Number of patients) |
|--------|---------------------|--------------------|-----------------------------------------|
| Allon et al. | Surgical removal: anterior maxillectomy | 20 | FOD: 1 REC: 0 |
| Heasman et al. | Surgical removal: subtotal mandibulectomy and right supramylohyoid neck dissection repaired using a radial forearm flap | N/A | N/A |
| Raguse et al. | Surgical removal: Mandibular resection and neck dissection (tumour enucleation with free margins) | 24 | FOD: 1 REC: 0 |
| Kruse and Grätz | Surgical removal: anterior maxillectomy with bilateral selective neck dissection | 24 | FOD: 1 REC: 0 |
| Hutton et al. | Surgical removal: anterior maxillectomy | 24 | N/A | N/A |
| Pons et al. | Surgical removal: bloc resection associated with modified radical neck dissection followed by reconstruction with bone and skin grafts | 9.3 | FOD: 2 REC: 0 Mortality: 1DUC |
| Sepulveda et al. | Chemotherapy and radiotherapy | N/A | Tumor persistence |
| Sun et al. | Surgical removal: wide surgical excision for cases with no bone invasion while those with bone invasion, a subtotal maxillectomy or mandibulectomy with safety margin. Upon recurrence in 3 cases, 1 patient underwent repeated surgical resection, the second concurrent chemoradiotherapy, and the third chemotherapy only | 45 | FOD: 12 REC: 3 with partial SCC transformation of one case of them and Mortality: 2 (DOD) |
| Suzuki et al. | Surgical removal: (hemi-mandibulectomy with unilateral supraomohyoid neck dissection followed by abdominal rectus myocutaneous free flap reconstruction). Upon recurrence, second Surgery added by chemotherapy | 14 | FOD: 1 REC: 1 then Mortality due to aspiration pneumonia (DOD) |
| Thavaraj et al. | Surgical removal: total glossectomy with bilateral selective neck dissection with safety margin of 5 mm | 24 | FOD: 1 REC: 0 |
| Fonseca et al. | Surgical removal: wide surgical excision | 45 | FOD: 2 REC: 0 |
| Goh et al. | Surgical removal: left hemiglossectomy and an ipsilateral selective neck dissection with safety margin | 14 | FOD: 1 REC: 0 |
| Padilla and Murrah | Surgical removal: total surgical excision of the soft tissue and bloc resection with a 1-cm safety margin | 3 cases: N/A 7 cases: 33.86 | FOD: 7 REC: 0 |
| Shakil et al. | Surgical removal | N/A | N/A |
| Shapiro et al. | Surgical removal / radiotherapy (composite mandibular resection with a 1-cm safety margin, bilateral selective neck dissection, and fibula free flap reconstruction) | N/A | N/A |
| Shay et al. | Surgical removal: composite mandibulectomy, radical parotidectomy, left modified radical neck dissection, and left fibular free flap for mandibular reconstruction) | 5 | FOD: 1 REC: 0 |
| Datar et al. | Surgical removal: (hemimandibulectomy with ipsilateral type II modified neck dissection) | 24 | FOD: 1 REC: 0 |

DOD: dead of disease, DUC: dead of unknown cause, FOD: free of disease, REC: recurrence
diotherapy in 1 case, and chemotherapy in 1 case.

**Differential diagnosis**

The criteria for the differential diagnosis between oral CC and other similar lesions were collected from the included articles (Table 5). A reliable knowledge of these criteria could provide the basis for a proper differential diagnosis between oral CC and other similar lesions.

**Discussion**

Oral CC is a rare variant of OSCC that was added to the WHO classification in 2005, however, most oral clinicians are unfamiliar with this entity and often misdiagnose it, leading to fatal outcomes due to faulty treatment of the affected patients.

The current systematic review is the first to systematically analyze the existing oral CC cases described in the English-language literature in an effort to illustrate and further characterize the etiologic agents, predisposing factors, clinical and imaging features, histopathological findings, treatment, and follow-up and survival rates. The goal of this review was to enable oral clinicians to diagnose this rare entity promptly and to differentiate it from other tumors with similar features, facilitating its adequate curative treatment.

A total of 55 reported oral CC cases were found in the literature between 1954 and January 2018, of which only 43 cases were included for analysis in the current systematic review, as the remaining 12 cases were de-

### Table 5. Differential diagnosis between oral carcinoma cuniculatum, verrucous carcinoma, and papillary squamous cell carcinoma

|                      | Oral carcinoma cuniculatum | Verrucous carcinoma | Papillary squamous cell carcinoma |
|----------------------|----------------------------|---------------------|----------------------------------|
| **Etiologic agents** | Not identified             | Snuff and tobacco chewing, human papillomavirus | Alcohol, smoking, human papillomavirus |
| **Clinical appearance** | Broad, red or white, blunt mildly papillary/cobblestone like architecture | Broad white warty lesion showing prominent papillary, verrucous surface projections (cauliflower-like) with downward pushing border | A soft, friable, polypoid, exophytic, papillary broad-based tumor frequently arises from a thin stalk |
| **Growth pattern** | Chiefly endophytic but sometimes show both endophytic and exophytic growth(locally destructive, infiltrative tumor) | Exclusively exophytic limited to the lamina propria (laterally growing tumor) | Exophytic or papillary growth (occurs in situ or as an invasive tumor) |
| **Radiological features** | Ill-defined osteolytic lesion showing invasion into the underlying tissues and bone destruction | Superficial bone erosion | Ill-defined osteolytic lesion showing invasion into the underlying tissues and bone destruction |
| **Histological feature** | Complex branching interconnecting network of multiple keratin filled crypts and cores (canaliculi) infiltrating adjacent tissues | A verrucous/frond-like hyperkeratotic surface with vertical church-spire like keratinization and parakeratin plugging | Characteristic thick papillary, finger-like, cauliflower-like growth of neoplastic cells surrounding narrow fibrovascular cores with minimal keratosis and single or multiple nests of tumor cells invading the underlying stroma. Cell nests and keratin pearls but no micro-abscesses |
| **Neoplastic cells** | Well differentiated | Well differentiated | Immature basaloid cells or more pleomorphic cells |
| **Cytological atypia** | Absent or minimal cellular atypia and mitoses | Absent or minimal cellular atypia and mitoses | Prominent many cytological atypias and dysplastic features |
| **Keratin** | + | +++ | ++ |
| **Other features** | Presence of neutrophil micro-abscesses | Broad pushing infiltrative front but with no invasion into underlying tissues | Frequent necrosis and hemorrhage |
| **Nodal metastasis** | Rare | Never | Frequent |
| **Immunohistochemical findings** | Less expression of p53 and Ki-67 | Strong expression of p53 and Ki-67 | Strongest expression for p53 and Ki-67 |
| **Therapy of choice** | Surgery | Surgery | Surgery/Radiotherapy/Chemotherapy |
| **Prognosis** | Better than papillary squamous cell carcinoma, not good when compared to verrucous carcinoma | Better than oral carcinoma cuniculatum and papillary squamous cell carcinoma | Worst prognosis |
Oral carcinoma cuniculatum, an unacquainted variant of oral squamous cell carcinoma: A systematic review

Although Hutton et al.²⁹ reported the presence of oral CC in a 7-year-old patient, the present search showed a predominance in the sixth and seventh decades of life, with a mean age of 62.7 years. This finding is consistent with those of Sun et al.,⁹ Pons et al.,¹⁰ and Padilla and Murrah.¹³

In the current review, oral CC was found to occur with a slight female predominance: 24 females (55.8%) and 19 males (44.2%). This matches the research of Sun et al.⁹ on 15 cases, Padilla and Murrah¹³ on 10 cases, and Fonseca et al.³³ on 2 cases. However, a slight male predominance of oral CC was reported by Pons et al.,¹⁰ who studied 3 cases, all of whom were males. Since oral CC is a variant of OSCC, and OSCC has a male predominance, this female predominance is surprising.¹¹

The current analysis revealed that the mandibular gingiva was the most common site for oral CC, as it was reported in more than half of the cases. This finding is in agreement with those of Pons et al.,¹⁰ Padilla and Murrah,¹³ and Fonseca et al.³³ However, Sun et al.⁹ reported the tongue to be the most common site.

The most common clinical presentation was pain,³,⁹,¹⁰,²⁶,²⁷,³⁰,³¹,³³,³⁵ followed by ulceration,¹,³,⁹,¹³,²⁶,²⁸,³⁰,³²,³³,³⁶ swelling,⁹,¹³,₂⁶,₂⁹,₃⁰,₃₂,₃³,₃₆ and induration.⁹,₁³,₃₆ This differs from the clinical presentation of OSCC, which may vary according to the affected intraoral subsite and for which pain is not the most common feature. This tendency is also distinct from VC, which usually presents as a well-demarcated, thin white keratotic plaque which quickly thickens and develops papillary (blunted tips) or verruciform (pointed tips) surface projections and from PSCC, for which hoarseness and airway obstruction are the most common presenting symptoms, as the larynx and the hypopharynx are among the most common sites of involvement.¹¹

The size of oral CC ranged from 1 to 6 cm; however, this range was derived from only 7 cases,³,¹³,₂⁵,₂⁶,₂₈,₃₄,₃₆ which is insufficient for a conclusive estimation of the average size of these tumors.

The duration of the lesion was recorded in 90.7% of the cases,¹,³,⁹,₁₀,₁₃,₂₅,₂₆,₂₈,₃₁-₃₆ with a mean of 19.5 months. This relatively long duration most likely indicates the slowly growing clinical course of oral CC,⁹,₁₀,₃₃,₃₆ which matches VC; however, this finding is in contrast to the behavior of OSCC, which shows a rapidly growing course.¹¹

In the current analysis, 13 patients (30.2%) were tobacco smokers, alcoholics, or both,³,⁹,₁₀,₁₃,₂⁵,₂₆,₃₁-₃₆ and 1 patient (2.3%) was a smokeless tobacco user.¹³ Smoking and alcohol consumption are known to be predisposing factors for oral malignancy,¹¹ however, a clear etiology for oral CC has yet to be established.

Premalignant lesions/conditions as leukoplakia, erythroleukoplakia, and lichen planus had been found in 10 cases (23.2%), suggesting that they underwent a malignant transformation to oral CC, which matches with what has been reported by several authors.¹,⁹,₁₀,₂₅,₂₆,₂₉,₃₁,₃₃

Preoperative imaging was reported for 18 cases (41.7%); some studies used panoramic radiographs,¹,₁₀,₁₃,₂₅,₂₇-₃₃ while others used CT,¹₀,₂₅,₂₉-₃₁,₃₆ MRI,³,₈,₃₁,₃₂ or both.³⁵ Most of the lesions were osteolytic and almost ill-defined, with cortical destruction and erosion of the underlying bone in tumors located in soft tissue. Ill-defined borders of a lesion indicate an inflammatory condition or malignancy; therefore, a correlation with the clinical features of the lesion is mandatory. Moreover, erosion of the underlying bone indicates the infiltrative nature of the lesion, as found in malignant conditions.

It is worth mentioning that the preoperative diagnoses were incorrect in all cases. To a large extent, this occurred due to clinicians’ lack of awareness about this entity. This also may have occurred due to the improper biopsy depth as biopsies were taken only from the superficial part of the lesion in some cases.

Regarding the histopathological features, all the included cases exhibited the same features specified by the WHO classification¹¹ and the diagnostic system purposed by Chen et al.,³⁹ wherein the presence of a defined set of histological features is required for the accurate diagnosis of oral CC and its differentiation from other similar lesions, such as VC and PSCC. In all cases, oral CC had a distinctive histopathologic appearance that revealed multiple branching keratin-filled crypts resembling rabbit burrows or cuniculi (thus the name cuniculatum), lined by well differentiated squamous epithelium with absent or minimal cytological atypia and normal or little mitosis. Moreover, micro-abscesses with discharge of yellowish secretion through the crypts and a prominent stromal reaction, mainly consisting of neutrophils, were reported in most cases.³,⁹,₁₀,₁₃,₂₅,₂₇-₃₀,₃₆

Oral CC has a locally aggressive invasive nature due to its deep epithelial projections or down-growths that extend into the connective tissue. In the current analysis, all the cases showed local invasion, with evidence of bone erosion and destruction noted radiographically, and 4 cases showed metastasis: 3 to the regional lymph nodes and 1 to the lungs.⁹ In our opinion, the occurrence of regional lymph node and/or distant metastasis should be taken into
consideration, as the early diagnosis and knowledge of appropriate treatments will prevent further spread of the tumor.

The presence of mutated forms of p53 was found to be associated with malignant features, yet, studies on its association with oral CC are lacking, since only 4 studies reported findings on p53 expression. These included case reports on 4 patients, with discordant results, as 2 studies showed positive results for p53 expression, while the other 2 showed negative results. Similar conflicting reports on Ki-67 expression, which is associated with cell proliferation in malignancy, were found: 1 study reported its increased expression, while other studies reported its consistently low expression. Regarding p16, which is a tumor suppressor, it was tested in only 1 study, which reported negative results for its expression. Extensive further research on the immunohistochemistry of oral CC is required.

Although a correlation between HPV and cutaneous CC has been observed, such a correlation was not established in oral cases, where no record of HPV presence was reported, except in 3 of the included cases, all of which showed negative results. As a result, the role of HPV in oral CC pathogenesis remains unclear.

The growth pattern of the lesion is reflected in its clinical presentation; both exophytic and endophytic growth patterns were recorded in 15 cases (34.9%), which most commonly presented as ulcerations, red and white lesions, and pain of a relatively long duration, in contrast to the prominent exophytic growth of VC.

Regarding the treatment of oral CC, complete surgical resection with a safety margin was the treatment of choice. Subtotal maxillectomy or mandibulectomy and/or neck dissection was performed in patients with extensive tumors because of the tendency of oral CC for local invasion. This treatment yielded effective results and an excellent (disease-free) prognosis in 29 of the 37 patients who continued to attend follow-up appointments for a duration ranging from 4 months to 14 years following surgery.

However, 4 patients (9.3%) experienced local recurrence. Three cases of tumor recurrence were reported by Sun et al.; 1 patient underwent repeated surgical resection following the initial recurrence, but a second recurrence occurred in the following year, together with partial malignant transformation to OSCC (hybrid tumor of oral CC and conventional OSCC), and the patient died of uncontrolled cancer with cachexia 6 months later. The second patient received concurrent chemoradiotherapy after recurrence, but the tumor was not controlled and metastasis developed to the deep cervical lymph nodes and the lungs, ultimately leading to death. The third received chemotherapy after recurrence, but the tumor had increased in size and treatment was not effective. Suzuki et al. reported a case in which multiple recurrences occurred after multiple surgical excisions, and finally the patient was treated unsuccessfully with chemotherapy before his death from aspiration pneumonia due to the involvement of the parapharyngeal space. The aforementioned findings encompass 3 cases of mortality: 1 case died of lung metastasis, while the other 2 died due to the increased tumor size. Therefore, mortality from oral CC is thought to primarily result from locally destructive growth, rather than metastasis.

Furthermore, such findings underscore the necessity of the cautious use of chemotherapy and/or radiotherapy, especially in the treatment of recurrent cases that did not respond to these types of treatment. This is supported by the assumption that radiotherapy may provoke anaplastic transformation. It is worth noting that recurrent oral CC demands careful attention due to its more resistant and aggressive behavior than its primary form.

In conclusion, oral CC is a rare, well-differentiated variant of OSCC with unique clinical and histopathological features. It has a good prognosis following appropriate surgical treatment with a relatively low recurrence rate. However, it shows resistant behavior in cases of recurrence. Proper knowledge of the diagnostic criteria of this lesion, together with a thorough correlation of the histological findings with the clinical and imaging features, is essential for its diagnosis and differentiation from other similar lesions. Due to its locally aggressive and invasive nature, it is advisable to perform multiple deep biopsies of the suspected lesions in addition to a thorough sampling of these biopsies to avoid underdiagnosis.

References

1. Datar UV, Kale A, Mane D. Oral carcinoma cuniculatum: a new entity in the clinicopathological spectrum of oral squamous cell carcinoma. J Clin Diagn Res 2017; 11: ZD37-9.
2. Aird I, Johnson HD, Lennox B, Stansfeld AG. Epithelioma cuniculatum a variety of squamous carcinoma peculiar to the foot. Br J Surg 1954; 42: 245-50.
3. Goh GH, Venkateswaran K, Leow PC, Loh KS, Thamboo TP, Petersson F. Carcinoma cuniculatum of the esophagus and tongue: report of two cases, including TP53 mutational analysis. Head Neck Pathol 2014; 8: 261-8.
4. Lawrence-Brown MM, Gollow IJ, Lam TP, Frost FA. Carcinoma cuniculatum of the abdominal wall. Med J Aust 1984;
Oral carcinoma cuniculatum, an unacquainted variant of oral squamous cell carcinoma: A systematic review

140: 668-9.
5. Barreto JE, Velazquez EF, Ayala E, Torres J, Cubilla AL. Carcinoma cuniculatum: a distinctive variant of penile squamous cell carcinoma: report of 7 cases. Am J Surg Pathol 2007; 31: 71-5.
6. Smyth M, Jaaback K, Tabrizi SN, Garland SM, Yin H, Scurry J. Carcinoma cuniculatum of the cervix. Pathology 2014; 46: 353-5.
7. Kotwal M, Poflee S, Bobhate S. Carcinoma cuniculatum at various anatomical sites. Indian J Dermatol 2005; 50: 216-20.
8. Flieger S, Owiński T. Epithelioma cuniculatum an unusual form of mouth and jaw neoplasm. Czas Stomatol 1977; 30: 395-401.
9. Sun Y, Kuyama K, Burkhardt A, Yamamoto H. Clinico pathological evaluation of carcinoma cuniculatum: a variant of oral squamous cell carcinoma. J Oral Pathol Med 2012; 41: 303-8.
10. Pons Y, Kerrary S, Cox A, Guerre A, Bertolus C, Gruffaz F, et al. Mandibular cuniculatum carcinoma: apropos of 3 cases and literature review. Head Neck 2012; 34: 291-5.
11. Barnes L, Eveson J, Reichart P, Sidransky D. World Health Organization classification of tumours. Pathology and genetics of head and neck tumours. Lyon: WHO IARC Press; 2005. p. 163-75.
12. Devaney KO, Ferlito A, Rinaldo A, El-Naggar AK, Barnes L. Verrucous carcinoma (carcinoma cuniculatum) of the head and neck: what do we know now that we did not know a decade ago? Eur Arch Otorhinolaryngol 2011; 268: 477-80.
13. Padilla RJ, Murrah VA. Carcinoma cuniculatum of the oral mucosa: a potentially underdiagnosed entity in the absence of clinical correlation. Oral Surg Oral Med Oral Pathol Oral Radiol 2014; 118: 684-93.
14. Moher D, Liberati A, Tetzlaff J, Altman DG, PRISMA Group. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. Ann Intern Med 2009; 151: 264-9.
15. Gagnier JJ, Kienle G, Altman DG, Neumann RA, Bodemer W, Radl-wimmer B, Aberer E, et al. DNA dot-blot hybridization implicate human papillomavirus type 11-DNA in epithelioma cuniculatum. J Med Virol 1989; 29: 33-7.
16. Burkhardt A. Verrucous carcinoma and carcinoma cuniculatum - forms of squamous cell carcinoma? Hautarzt 1986; 37: 373-83.
17. Kahn JL, Blez P, Gasser B, Weill-Bousson M, Vetter JM, Champy M. Carcinoma cuniculatum. Apropos of 4 cases with orofacial involvement. Rev Stomatol Chir Maxillofac 1991; 92: 27-33.
18. Delahaye JF, Janser JC, Rodier JF, Auge B. Cuniculatum carcinoma. 6 cases and review of the literature. J Chir (Paris) 1994; 131: 73-8.
19. Huault M, Laroche J, Levy J, Laxenaire A, Roucayrol AM, Scheffer P. Epithelioma cuniculatum. Apropos of a case in the anterior gingiva with involvement of the mandibular symphysis bone and reconstruction using a fibular osteocutaneous flap and integrated implants. Rev Stomatol Chir Maxillofac 1998; 99: 143-8.
20. Gassler N, Helmké B, Schweigert HG, Hassfeld S, Otto HF, Flechtenmacher C. Carcinoma cuniculatum of the oral cavity. A contribution to the differential diagnosis of potentially malignant papillary lesions of mouth mucosa. Pathologe 2002; 23: 313-7.
21. Neill S, Calnan C, Rahim G. Carcinoma cuniculatum. Clin Exp Dermatol 1984; 9: 309-11.
22. Lozzi GP, Peris K. Carcinoma cuniculatum. CMAJ 2007; 177: 249-51.
23. Prasad SG, Kaur K, Gupta S. Carcinoma cuniculatum: a review. Indian J Dent Sci 2012; 4: 87-9.
24. Kubik MJ, Rhatigan RM. Carcinoma cuniculatum: not a verrucous carcinoma. J Cutan Pathol 2012; 39: 1083-7.
25. Allon D, Kaplan I, Manor R, Calderon S. Carcinoma cuniculatum of the jaw: a rare variant of oral carcinoma. Oral Surg Oral Med Oral Pathol Oral Radiol Endod 2002; 94: 601-8.
26. Heasman P, Smith D, Martin I, Soames J. Carcinoma cuniculatum presenting as a gingival lesion. Perio 2005; 2: 199-203.
27. Raguse JD, Menneking H, Scholmenn JH, Bier J. Manifestation of carcinoma cuniculatum in the mandible. Oral Oncol Extra 2006; 42: 173-5.
28. Kruse AL, Graetz KW. Carcinoma cuniculatum: a rare entity in the oral cavity. J Craniolaf Surg 2009; 20: 1270-2.
29. Hutton A, McKaig S, Bardsley P, Monaghan A, Parmar S. Oral carcinoma cuniculatum in a young child. J Clin Pediatr Dent 2010; 35: 89-94.
30. Sepolveda I, Spencer L, Piatin E, Bravo F. Maxillary cuniculatum carcinoma: a case report and literature review. Int J Odontostomatol 2012; 6: 281-4.
31. Suzuki J, Hashimoto S, Watanabe K, Takahashi K, Usabuchi H, Suzuki H. Carcinoma cuniculatum mimicking leukoplakia of the mandibular gingiva. Auris Nasus Larynx 2012; 39: 321-5.
32. Thavaraj S, Cobb A, Kalavrezos N, Beale T, Walker DM, Jay A. Carcinoma cuniculatum arising in the tongue. Head Neck 2012; 6: 130-4.
33. Fonseca FP, Pontes HA, Pontes FS, de Carvalho PL, Sena-Filho M, Jorge J, et al. Oral carcinoma cuniculatum: two cases illustrative of a diagnostic challenge. Oral Surg Oral Med Oral Pathol Oral Radiol 2013; 116: 457-63.
34. Shakil M, Mohtesham J, Jode M, Prabhu V. Carcinoma cuniculatum of the oral cavity- a rare entity. J Adv Med Dent Sci 2014; 2: 124-6.
35. Shapiro MC, Wong B, O’Brien MJ, Salama A. Mandibular destruction secondary to invasion by carcinoma cuniculatum. J Oral Maxillofac Surg 2015; 73: 2343-51.
36. Shay S, Choy W, Christensen RE, St John MA. Extensive carcinoma cuniculatum of the mandible. Am J Otolaryngol 2015; 36: 446-50.
37. Knobler RM, Schneider S, Neumann RA, Bodemer W, Radl-wimmer B, Aberer E, et al. DNA dot-blot hybridization implicates human papillomavirus type 11-DNA in epithelioma cuniculatum. J Med Virol 1989; 29: 33-7.
38. Wastiaux H, Dreno B. Recurrent 144...