

Comparison of osteoclastogenesis and local invasiveness of ameloblastoma and keratocystic odontogenic tumor

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ABSTRACT

Objectives: The aim of this study was to compare the expression of receptor-activated nuclear factor kappa B (RANK) with its ligand (RANKL) and matrix metalloproteinase-2 (MMP2) in solid/multicystic ameloblastomas (ABs) and keratocystic odontogenic tumors (KOTs). **Materials and Methods:** The expression of MMP2, RANK, and RANKL molecules was evaluated in 13 ABs and 14 KOTs by immunohistochemistry. The expressions were calculated in the odontogenic epithelial cells as well as the stromal cells. **Results:** Odontogenic epithelia of AB expressed MMP2, RANK, and RANKL significantly higher than that of KOTs ($P < 0.05$). The expression of MMP2, RANK, and RANKL was highest in plexiform subtype (79.9%, 81.08%, and 65.1%, respectively). KOTs without daughter epithelia nests expressed both MMP2 and RANK the least (56.06% and 47.5%, respectively). Stromal cells, on the other hand, expressed similar MMP2 pattern in odontogenic epithelia of both AB and KOT. RANKL was expressed weaker in the stromal cells of both lesions. **Conclusion:** Invasive biological and osteolytic behaviors of both lesions were evaluated in this study. It was found to be more in AB than keratocystic odontogenic. A significant expression of MMP2, RANK, and RANKL in both KOTs associated with microcyst and plexiform type AB as well.

Key words: Keratocystic odontogenic tumor, matrix metalloproteinase-2, odontogenic keratocysts, receptor-activated nuclear factor kappa B, receptor-activated nuclear factor kappa B ligand, solid/multicystic ameloblastoma

INTRODUCTION

Keratocystic odontogenic tumors (KOTs) and ameloblastoma (AB) are characterized by slow growth with locally invasive behavior and high risk of recurrence.^[1] Both show infiltration into cancellous bone and destructive growth. That is why in 2005, odontogenic keratocyst is renamed as “KOTs” by the World Health Organization.^[2] The receptor-activated nuclear kappa B (RANK) is a central activator of necrosis factor-kappa B, which controls the transcription of DNA, and is the signaling receptor for RANK

ligand (RANKL). RANKL binds to RANK on the surface of preosteoclasts and stimulates the development and activation of osteoclasts.^[3] Many authors suggested that odontogenic epithelial cells are an important source of RANK.^[4,5] Sandra *et al.*^[6] suggested that AB could induce osteoclastogenesis by secreting RANKL and tumor necrosis factor α (TNF- α). da Silva *et al.*^[5] in their study on AB and odontogenic keratocyst (OKC) found that >90% of odontogenic epithelium of both

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lesions expressed RANK. They suggested that RANK originating from odontogenic epithelium enhances bone resorption due to its effect on tumor growth, cell proliferation, or delaying apoptosis. MMP2 is a gelatinase of the matrix metalloproteinase (MMP) family. It degrades many types of collagens such as native type IV, V, and X collagen and denatured fibrillar type I, II, and III collagens.^[7] It has been found that many members of MMP family take part in cancer invasion and metastasis.^[8] Interest in AB and KOTs has risen due to their aggressive clinical behavior, variety of morphological patterns, and high recurrence rates. Since these tumors generally exhibit localized bone destruction and tooth resorption, the differential expressions of MMP2, RANK, and RANKL may be expected to find in these lesions. The aim of the present study is to examine and evaluate the expression of MMP2, RANK, and RANKL in both lesions, in an attempt for better understanding of biological behavior associated with these odontogenic tumors from the points of invasiveness and osteoclastogenesis.

MATERIALS AND METHODS

Tissue samples

The study was approved by the Research and Ethics Committee of College of Dentistry at University of Baghdad (REC. 25-2014). The paraffin blocks were retrieved from the archives of the Oral Diagnosis Department, College of Dentistry, University of Baghdad. The diagnosis of each case was confirmed by two oral pathologists through the examination of hematoxylin and eosin-stained sections. Data, regarding name, age, sex, and site of the lesion, were collected from the routine histopathology files of the same cases. Recurrent AB and KOTs were excluded from the study.

The expression of MMP2, RANK, and RANKL was evaluated in the tissue specimens of 14 KOT and 13 solid/multicystic AB (SMA). The SMA was subdivided into eight follicular and five plexiform patterns, while KOTs were categorized into those associated with daughter epithelial nests (KOT with + dentigerous cyst [DC]) group ($n = 5$) and those without (KOT with - DC) group ($n = 9$). All tests were carried out on 4 μ m sections from formalin-fixed paraffin-embedded blocks of SMA and KOT. The expressions were calculated in the odontogenic epithelial cells as well as the stromal cells.

Immunohistochemical staining

The paraffin blocks were cut serially into approximately 5 μ m thick sections on charged slides. Using Dako

Autostainer, the slides were deparaffinized with xylene for 30 min, washed with absolute 100% alcohol for 15 min, and then washed with 95% alcohol and distilled water. Dako REAL™ EnVision™, peroxidase/diaminobenzidine (DAB), and Mouse IG detection system were used for MMP2, RANK, and RANKL. For antigen retrieval, 20-min heat-induced antigen retrieving was performed with RANK and RANKL, while in MMP2, enzyme block solution was used only (chemical retrieving). The retrieving solution used for RANK and RANKL was citrate buffered (pH = 9 under 80°C for RANKL and pH = 6 under 85°C for RANK) and then washed in phosphate buffered saline for 5 min. To block endogenous peroxidase activity, slides were incubated with 3% hydrogen peroxide, washed in distilled water and then soaked in phosphate buffer saline for 5 min.

MMP2 antibody at a dilution of 1:50 (Abcam No: Ab3158, Monoclonal Mouse Anti-Human MMP2 Antigen, CA-4001/7CA19E3C, Abcam, USA), RANK antibody at a dilution of 1:50 (Abcam No: Ab12008, Monoclonal Mouse Anti-Human MMP2 Antigen, 9A725, Abcam, USA), and RANKL at a dilution of 1:75 (Abcam No: Ab 45039, Monoclonal Mouse Anti-Human RANKL Antigen, 12A668, Abcam, USA) were used as primary antibodies. Slides were incubated overnight at 4°C with MMP2, RANK, and RANKL. Positive control slides were obtained according to the antibodies manufacturer's data sheet. For MMP2 monoclonal antibodies, tissue blocks of breast carcinoma were used.^[7] For RANK, the bone tissue located at the periphery of the lesions of SMA and KOT was used, while that for RANKL, tissue blocks of peripheral giant cell granuloma were used.^[5] The negative control slides were prepared by substituting the primary specific antibodies with nonimmune serum. All the above control samples were fixed and processed in a way similar to that of test samples. The secondary antibody Dako REAL™ EnVision™/HRP, Rabbit/Mouse IgG was reacted for 30 min, followed by DAB-chromogen solution for 30 min being used to visualize the reaction. Finally, the sections were counterstained with Mayer's hematoxylin, dehydrated, coverslipped, and evaluated by light microscopy.

Signal specificity was demonstrated by the absence of immunostaining in the negative control slides and its presence in the recommended positive controls. Odontogenic epithelia and stromal cells in both SMA and KOT with clear brown membranous/cytoplasmic staining were considered to be positive for MMP2, RANK, and RANKL immunostaining within a violet-blue tissue section background of

hematoxylin staining. All the slides were assessed blindly by two pathologists without prior knowledge of the corresponding clinicopathological data, and the average of the two readings was obtained.

The specimens were examined and counted at ×200 magnification using integration graticule with an Olympus C ×30 microscope. In the epithelia, positive cells were counted in five contiguous and consecutive microscopic high-power fields. In the stroma, positive endothelial and fusiform cells (fibroblasts) were quantified in five contiguous and consecutive areas adjacent to neoplastic odontogenic epithelium;^[5,9] each field of the integration graticule had an area of 0.0061 mm².^[9] For accuracy, the counting rechecked at × 400 for each field. Results were obtained and the mean of the percentage of positive cells was determined for each representative field.

Statistical analyses were done using SPSS version 21 (Chicago., IL, USA). The statistical significance of differences in RANK, RANKL, and MMP 2 reactivity was analyzed in both lesions by the Mann–Whitney U-test and *P* < 0.05 was considered to indicate statistical significance.

RESULTS

The demographic data of the studied sample are shown in Table 1. The mean age of patients from which the AB specimens were obtained ranged between 10 and 45 years (mean = 30.6 years). For KOT cases, the age range was between 15 and 45 years (mean = 31.9 years). All cases of AB occurred in the mandible, and for KOT, the mandible was more frequently involved than maxilla (64.3% and 35.7%, respectively) [Table 1].

MMP2, RANK, and RANKL immunostaining was detected in the odontogenic epithelial cells as well as in the stromal cells.

Matrix metalloproteinase-2 immunostaining

Strong expression of MMP2 was seen in ABs and KOTs, with higher value in SMA tumor cells than keratocyst lining epithelia (73.29% and 63.87%, respectively). However, the difference was not statistically significant (*P* = 0.09) [Table 2]. MMP2 expression was higher in epithelial cells of plexiform AB (79.9%), followed by KOT associated with daughter epithelia nests (76.37%) and then follicular AB (68.57%), and the least was in KOT not associated with daughter epithelial nests (56.06%) as shown in Figure 1 and Table 3. MMP2 expression in stromal cells was almost the same in SMA and KOT (48.66% and 50%, respectively), with no significant differences between the histological subgroups [Table 2].

Receptor-activated nuclear factor kappa B immunohistochemical staining

Odontogenic epithelial cells of AB strongly expressed RANK, which is significantly higher than that of KOT (67.55 ± 16.83 versus 51.8 ± 17.15) as seen in Table 4 and Figure 2, whereas RANK expression in stromal cells did not show any significant differences between lesions.

Plexiform AB highly expressed RANKL in their odontogenic epithelium in 81.08%, followed by follicular type (62%), KOT associated with daughter epithelia nest (58.7%), and then KOT with no daughter epithelia nest group (47.5%). A statistically significant difference was seen only when plexiform pattern

Table 1: Clinical parameters (age, sex, and site) for ameloblastoma and keratocystic odontogenic tumors cases

	n	Sex		Range	Age		Site	
		Male (%)	Female (%)		Mean±SD	SE	Mandible (%)	Maxilla (%)
AB	13	7 (53.8)	6 (46.2)	10-45	30.6±9.16	2.64	13 (100)	0
KOT	14	6 (42.9)	8 (57.1)	15-45	31.9±8.94	2.38	9 (64.2)	5 (35.7)

AB: Ameloblastoma, KOT: Keratocystic odontogenic tumor, SD: Standard deviation, SE: Standard error

Table 2: The comparison between matrix metalloproteinase-2 expression of the odontogenic epithelia and stromal cells of solid/multisystem ameloblastoma and keratocystic odontogenic tumor

Tissue group	Cells	Total number	Positive MMP2	Negative MMP2	Mean±SD	SE	P
SMA	Odontogenic epithelium	13	12	1	73.29±10.92	3.16	0.09
KOT		14	13	1	63.87±16.36	4.53	
SMA	Stromal cells	13	12	1	48.66±12.03	3.47	0.7
KOT		14	13	1	50.00±14.69	4.07	

MMP2: Matrix metalloproteinase-2, SMA: Solid/multicystic ameloblastoma, KOT: Keratocystic odontogenic tumor, SD: Standard deviation, SE: Standard error

Table 3: Comparison between matrix metalloproteinase-2 area percentage in the odontogenic epithelia and stromal cells of follicular ameloblastoma, plexiform ameloblastoma, keratocystic odontogenic tumor (negative dentigerous cysts), and keratocystic odontogenic tumor (positive dentigerous cysts)

Tissue group	Cells	Total number	Positive MMP2	Negative MMP2	Mean±SD	SE	P
Follicular AB	Odontogenic epithelia	8	7	1	68.57±8.52	3.22	0.06
Plexiform AB		5	5	0	79.9±11.29	5.05	
KOT (negative DC)		9	8	1	56.06±9.56	3.38	0.01*
KOT (positive DC)		5	5	0	76.37±18.65	8.07	
Follicular AB	Stromal cells	8	7	1	52.57±4.98	1.88	0.16
Plexiform AB		5	5	0	43.2±17.23	7.70	
KOT (negative DC)		9	8	1	46.87±13.15	4.65	0.3
KOT (positive DC)		5	5	0	55.0±17.16	7.67	

*Significant at $P \leq 0.05$. AB: Ameloblastoma, KOT: Keratocystic odontogenic tumor, SD: Standard deviation, SE: Standard error, MMP2: Matrix metalloproteinase-2, DC: Dentigerous cysts

Table 4: Comparison between receptor-activated nuclear factor kappa B area percentage of the odontogenic epithelia of solid/multisystem ameloblastoma and keratocystic odontogenic tumor

Tissue group	Total number	Cells	Positive RANK	Negative RANK	Mean±SD	SE	P
SMA	13	Odontogenic epithelia	12	2	67.55±16.83	5.07	0.02*
KOT	14		13	1	51.8±17.15	4.75	
SMA	13	Stromal cells	10	3	45.22±18.74	5.92	0.59
KOT	14		13	1	41.34±18.49	5.13	

*Significant at $P \leq 0.05$. RANK: Receptor-activated nuclear factor kappa B, SMA: Solid/multicystic ameloblastoma, KOT: Keratocystic odontogenic tumor, SD: Standard deviation, SE: Standard error

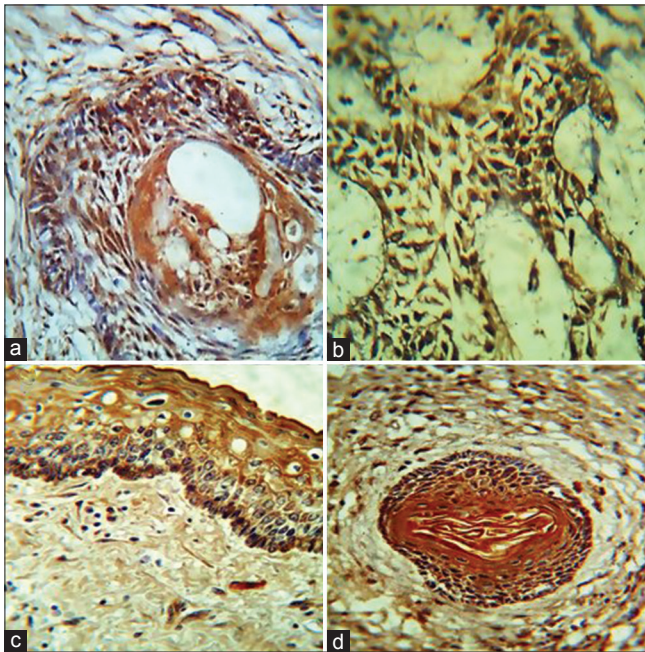


Figure 1: (a-d) Matrix metalloproteinase-2 expression in odontogenic epithelium, follicular ameloblastoma, plexiform ameloblastoma, keratocystic odontogenic tumor (-dentigerous cysts), keratocystic odontogenic tumor (+dentigerous cysts)

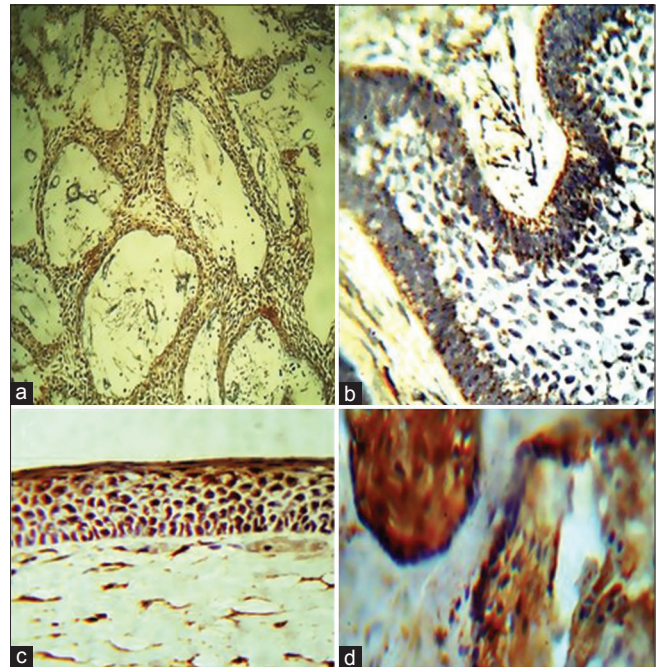


Figure 2: (a-d) Receptor-activated nuclear factor kappa B expression in odontogenic epithelium, follicular ameloblastoma, plexiform ameloblastoma, keratocystic odontogenic tumor (-dentigerous cysts), keratocystic odontogenic tumor (+dentigerous cysts)

compared with both subtypes of KOT ($P = 0.03$ and $P = 0.009$, respectively) [Table 5]. In the stroma, the expression was almost similar with a bit elevation of SMA (45.22% and 41.34%) [Table 4].

Receptor-activated nuclear factor kappa B ligand immunostaining

In this study, RANKL was detected in the membrane and cytoplasm and in the extracellular of AB and KOT cells as seen in Figure 3.

Table 5: The comparison between receptor-activated nuclear factor kappa B area percentage of the odontogenic epithelia and stromal cells of follicular ameloblastoma, plexiform ameloblastoma, keratocystic odontogenic tumor (positive dentigerous cysts), and keratocystic odontogenic tumor (negative dentigerous cysts)

Tissue group	Total number	Cells	Positive RANK	Negative RANK	Mean±SD	SE	P
Plexiform AB	5	Odontogenic epithelia	3	2	81.08±3.74	2.16	0.03*
Follicular AB	8		8	0	62.48±17.11	6.05	
KOT (negative DC)	9		8	1	47.5±16.87	5.96	0.25
KOT (positive DC)	5		5	0	58.7±16.97	7.59	
Plexiform AB	5	Stromal cells	3	2	38.33±2.88	1.66	0.009**
Follicular AB	8		8	0	48.17±2.14	8.36	
KOT (negative DC)	9		8	1	48.2±14.53	6.49	0.25
KOT (positive DC)	5		5	0	37.06±20.28	7.17	

*Significant at $P \leq 0.05$, **Significant at $P \leq 0.01$. AB: Ameloblastoma, KOT: Keratocystic odontogenic tumor, SD: Standard deviation, SE: Standard error, RANK: Receptor-activated nuclear factor kappa B, DC: Dentigerous cysts

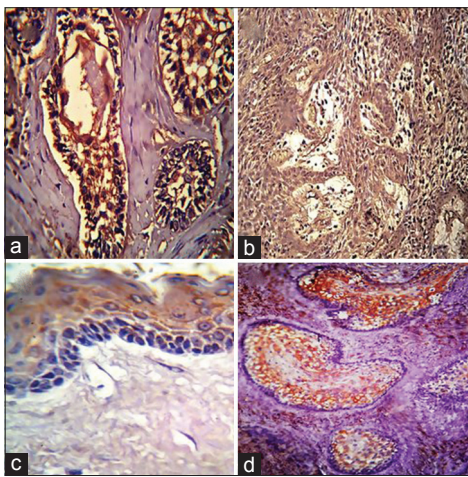


Figure 3: (a-d) Receptor-activated nuclear factor kappa B ligand expression in odontogenic epithelium, follicular ameloblastoma, plexiform ameloblastoma, keratocystic odontogenic tumor (-dentigerous cysts), keratocystic odontogenic tumor (+dentigerous cysts)

Epithelial cells of AB expressed more RANKL than that of KOT without reaching statistical significance ($P = 0.06$). However, stromal cells expressed RANKL more in AB than KOT with almost significant differences [Table 6].

Regarding histological subtypes, epithelia of plexiform type expressed more RANKL (65.1%) than follicular type (53.53%). Epithelial and stromal cells of KOT, on the other hand, showed almost same expression in both subtypes without observable statistical significant value [Table 7].

DISCUSSION

Assessment of matrix metalloproteinase-2 expression in ameloblastoma and keratocystic odontogenic tumor
MMP2 plays a major role in extracellular matrix remodeling because of its ability to initiate and continue

Table 6: Comparison between receptor activator of nuclear factor kappa-B ligand area percentage in the odontogenic epithelia of solid/multicystic ameloblastoma and keratocystic odontogenic tumor

Tissue group	Total number	Positive RANKL	Negative RANKL	Mean±SD	SE	P
SMA	13	12	1	57.93±20.6	5.94	0.06
OKC	14	13	1	43.78±17.43	4.83	
SMA	13	11	2	34.68±20.42	6.15	0.09
OKC	14	12	2	23.96±10.03	2.89	

OKC: Odontogenic keratocyst, SMA: Solid/multicystic ameloblastoma, RANKL: Receptor activator of nuclear factor kappa-B ligand, SD: Standard deviation, SE: Standard error

degradation of fibrillar collagen, especially type IV collagen which is the major component of basement membrane.^[10] Although numerous researches have verified that MMP2 plays an important role in tumor invasion,^[11] yet there are limited researches about its behavior in AB and KOT. The present study as well as the others showed that the high expression and activity of MMP2 is related to the aggressive behavior in both AB^[12-15] and KOT,^[16,17] with an overall higher level in the first than in the latter.^[12,14,16] There was a strong expression of MMP2 in the stroma of SMA and KOT, which is in accordance with other studies,^[18,19] although one study observed the opposite.^[15] The presence of MMPs in the tumor stroma could be attributed to tumor induction since neoplastic cells express the protein extracellular inducers (EMMPRIN/CD147) on their surface, and this acts as a potent inducer for the production of MMPs by stromal fibroblasts and endothelial cells.^[20]

Furthermore, the high expression of MMP2 in the odontogenic epithelia of plexiform type and KOT with daughter cysts (KOT with + DC) could suggest more aggressive behavior of plexiform type of AB. KOTs have a tendency to form satellite cysts through

Table 7: The comparison between receptor activator of nuclear factor kappa-B ligand area percentage in the odontogenic epithelia and stromal cells of follicular ameloblastoma, plexiform ameloblastoma, keratocystic odontogenic tumor (positive dentigerous cysts), and keratocystic odontogenic tumor (negative dentigerous cysts)

Tissue group	Total number	Cells	Positive RANKL	Negative RANKL	Mean±SD	SE	P
Plexiform AB	5	Odontogenic epithelia	5	-	65.1±23.75	10.62	0.34
Follicular AB	8		7	1	53.53±18.65	7.05	
KOT (with negative DC)	9		8	1	43.16±19.57	6.92	0.87
KOT (with positive DC)	5		5	0	44.78±15.47	6.92	
Plexiform AB	5	Stromal cells	5	0	38.44±25.72	11.5	0.56
Follicular AB	8		6	2	31.55±16.70	6.82	
KOT (with negative DC)	9		7	2	26.9±11.0	4.15	0.23
KOT (with positive DC)	5		5	2	19.86±7.68	3.43	

AB: Ameloblastoma, KOT: Keratocystic odontogenic tumor, SD: Standard deviation, SE: Standard error, RANKL: Receptor activator of nuclear factor kappa-B ligand, DC: Dentigerous cysts

detachment of epithelial cell lining from the connective tissue wall. This suggests the important role of MMP2 in this detachment. Wahlgren *et al.*^[17] mentioned that MMP2 can induce epithelial migration by fragmenting the basement membrane material laminin-5 gamma-2 chain, which induced migration, tendency to detach from the connective tissue capsule. This molecular behavior has been thought to be responsible for the frequent recurrence of KOT after surgical enucleation.

Assessment of receptor-activated nuclear factor kappa B expression in ameloblastoma and keratocystic odontogenic tumor

RANK, a member of the tumor necrosis factor receptor superfamily, is the signaling receptor for RANKL. RANKL binds to RANK on the surface of preosteoclasts and stimulates the development and activation of osteoclasts.

RANK originating from the epithelium has an effect on bone resorption, cell proliferation, or delaying apoptosis and hence the tumor growth. The present study could indicate greater bone resorption activity in SMA than KOT. The RANK expression in odontogenic epithelia of SMA is significantly higher than in KOT. Tekkesin *et al.*^[9] observed the opposite findings. However, da Silva *et al.*^[5] observed almost the same findings of the present study.

RANK in stromal cells other than osteoclasts and their precursors may have influential effects in several functions, such as immune cell regulation and inhibition of apoptosis of osteoclasts.^[6,21] da Silva *et al.*^[5] and Tekkesin *et al.*^[9] showed different levels of positive stroma cells in AB and OKC; the present study observed similar stromal RANK expression in both SMA and KOT with heavy staining at the periphery of AB and KOT.

Expression of RANK, in both the epithelial and connective tissue cells of KOTs and AB, indicates that RANK plays a role in local bone resorption of both lesions.

Assessment of receptor-activated nuclear factor kappa B ligand expression in ameloblastoma and keratocystic odontogenic tumor

RANKL is a membrane-bound protein found on osteoblastic and activated T-cells. RANKL binds to RANK on the surface of preosteoclasts and stimulates the development and activation of osteoclasts. Osteoprotegerin (OPG) is a soluble decoy receptor for RANKL that inhibits the pro-osteoclastogenic interaction between RANK and RANKL, thereby inhibiting bone resorption.^[22] The RANKL, RANK, and OPG system have been shown to be abnormally regulated in several malignant osteolytic pathologies, including neoplastic and nonneoplastic odontogenic lesions. The enhanced RANKL expression or decreased OPG levels play an important role in tumor-associated bone destruction.^[23] OPG and RANKL have also been detected in periodontal ligament cells and their expression are considered to play a role in osteoclastogenesis and bone resorption in periodontal diseases.^[24]

RANKL expression in the present study confirmed the findings of previous studies where tumor cells and stroma could act as a source of this osteoclastogenic factor. This indicates that RANKL-producing tumor cells of AB and cystic cells of KOT are crucial for osteoclastogenic behavior. The results of the present study also showed that both factors RANK and RANKL had higher ratio when comparing SMA to KOT. This would account to more aggressive nature and higher rate of recurrence in AB than KOT.^[5,6,9,25] This fact could be also true when

comparing plexiform pattern with other histological subtypes of SMA and KOT.^[26] There is no known link in the pathway of synthesis, mechanism of action, overexpression of MMP2, and the RANK and RANKL in both SMA and KOT.^[27] MMP2 role of action is mainly degradation of extracellular matrix, while RANKL and RANK are involved in mechanism of how the tumor expands in the bone, through stimulation proliferation and activation of osteoclast and hence bone resorption.^[28] Bone resorption mediated by activation of RANK and RANKL signalling pathway and matrix degradation by activation of MMP2 pathway in AB and KOT will identify the aggressive biological behavior of both lesions. The fact that RANKL is required for osteoclast development suggests that agents that inhibit its activity may be therapeutic. This has provided the rationale for the development of targeted molecular therapy with the ability to modulate RANK-induced osteoclastogenesis.

CONCLUSION

The high expression of MMP2, RANK, and RANKL is related to the biological behavior of AB and KOT and may indicate close behavior of KOT to AB, which reinforce the possibility of its recent classification as an odontogenic tumor. In addition, tumor epithelial cells of plexiform AB and epithelial nests or satellite microcyst-associated KOT exhibited distinct stronger expression of these markers compared with follicular AB and KOT without microcyst or epithelial nests, respectively. This could reveal a more aggressive behavior of these types of lesions.

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Conflicts of interest

There are no conflicts of interest.

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