Clinical Effectiveness of the Lateral Approach of the Maxillary Sinus Floor Augmentation using Different Gadgets with Simultaneous Implant Placement: A Systematic Review with Meta-Analysis

Ahmed Mortada Fikry^{1*}, Amr Zahran², Moamen Sheba² and Magdy Mostafa³

¹Assiut University, Egypt; mortadafikry@hotmail.com ²Faculty of Dentistry, Cairo University, Egypt; dramrzahran@gmail.com, moamen.sheba@dentistry.cu.edu.eg ³Faculty of Medicine, Cairo University, Egypt; imagdy@link.net

Abstract

Objectives: Help the clinician to choose between the newly introduced, specifically designed instruments for this procedure. **Methods/Statistical Analysis**: after thorough search of Cochrane CENTRAL and PubMed databases, the analysis was limited to clinical studies of the lateral approach sinus augmentation with simultaneous implant placement. **Statistical Analysis**: Comprehensive Meta Analysis version 2.2.048 software was used. **Results:** Eight articles were included in the review. Analysis of the formatted tables shows that the type of tool used in sinus lateral approach might affect the success of the procedure. **Findings**: There was a statistical significance difference of using the ultrasonic tips over the conventional use of rotary tips but it is not usually to be clinically significant; especially if a highly trained surgeon could perform the lateral osteotomy by any tool but the previous studies were highly supporting the ultrasonic tips over the rotary tips in reference to less traumatic surgery with minimal operative and postoperative complications. **Application/Improvements:** Randomized clinical trials are needed to compare the newly presented tools like DASK and SLA reamers to properly asses their clinical effectiveness. Studies of patient reported outcomes are needed to clearly evaluate the cost effectiveness of each tool.

Keywords: Augmentation, Implant, Lateral Approach, Maxillary Sinus, Simultaneous

1. Introduction

Damage of the membrane lining the maxillary sinus is the most common problem of the lateral approach sinus floor augmentation techniques which used as a corrective surgery to the upper posterior atrophic ridge with height problem that contradict implant placement at this site . Perforations were most likely accompanied with the use of traditional rotary tips and saws¹.

Recently different tools designed to minimize the possibility of sinus membrane tear .² Piezoelectric tips are ultrasonic auscultation tips that selectively cut the bone without damage of soft tissue and teeth^{3.4}.

Many reamers presented nowadays in the market which minimally invasive to the sinus membrane likes (Biomet 3i) but it was advised to be used with extreme caution⁵. LS and C reamers (SLA KIT, Neobiotech) are other types of drills of a conical trunk and specific penetration depth that selected for each case from the pre-operative cone beam volumetric tomography⁶. Moreover, it favors good accessibility to the surgical site; lessen the time of surgery when compared to the conventional rotary tips, also it considered of low cost when compared to the piezoelectric tips².

Altered type of reamers was presented that based on cutting the bone on the lateral surface of the maxillary

*Author for correspondence

sinus by thinning it gradually⁸. Also, there are new drills called Artificially Intelligent (AI- Water Lift System) that stops cutting the bone when they reach the membrane lining the sinus cavity by pressure sensitivity⁹.

Implant placement could be simultaneous with the sinus floor augmentation or in staged approach, the former provides the advantages of shortening the time of treatment and less surgical entries, but it requires minimal residual preoperative bone height of 5mm that guaranties implant primary stability¹⁰.

2. Material and Methods

2.1 Focus Question

Which is the best tool to be used in lateral sinus augmentation technique and to help clinicians choose between the newly introduced tools in the market? It established by PICO (Population, Intervention, Comparison Andoutcome).

Population: Patients with upper posterior edentulous area with insufficient bone height for implant placement.

Intervention: piezoelectric ultrasonic tips and reamers.

Comparison: Conventional rotary burrs.

Outcomes: implant survival and bone height changes

2.2 Type of Intervention

New tools which specifically designed for the lateral approach sinus floor augmentation technique.

2.3 Outcome Parameters

Outcomes to be assessed were membrane perforation, postoperative sinusitis, graft loss, implant survival rate, graft height reduction and postoperative bone height.

3. Study Type and Follow Up

Randomized clinical trials on patients (sample size = minimum 4 patients) need restoration of upper posterior edentulous area which complicated by sinus floor pneumatization; with residual bone height (at least 8 mm) requiring sinus floor augmentation via the lateral approach and simultaneous implant placement and at least follow up of 6 months after implant placement.

4. Exclusion Criteria

- In vitro studies.
- Case reports and case series studies.
- Implants survival rate not assigned in the paper or implant success not one of the outcomes.
- The technique of sinus augmentation not clearly described.

4.1 Search Strategy

A computerized systematic search strategy was conducted in Cochrane CENTRAL and PubMed searching for human studies with no language restrictions. The full text articles were obtained from reviews on sinus floor augmentation published to date. Additional publications were identified from the reference lists of the retrieved articles. No individual journal search was conducted but rather depending on reputable journals already indexed in the searched databases.

4.2 Search Combination

Search terms were ((((((sinus lift lateral) OR sinus lift lateral window) OR lateral sinus lift) OR external sinus lift) OR open sinus lift)) AND ((((simultaneous implant) OR implant placement) OR implants dental) OR implants)

4.3 Selection of Studies

Independent screening of all papers by the authors based on the inclusion criteria. The selected articles were then obtained in full text. Disagreements were resolved by discussion. PRISMA flow chart presented in Figure 1.

4.4 Data Extraction

The reviewers independently extracted the data using data extraction tables. Any disagreements were resolved by double-checking the original data and by discussion.

4.5 Quality Assessment

Retrospective studies were excluded because they are highly susceptible to recall bias so they considered of a low grade of evidence. Cochrane tool for risk of bias assessment was used to assess the quality of included clinical studies regarding sequence generation, allocation concealment, blinding, incomplete outcome data and selective outcome reporting.



Table 1.	List of excluded a	rticles
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Authors	Reasons of exclusion
9	Case report
10	Two stages implant placement (not simultaneous with sinus augmentation)
28	Crestal approach not lateral approach
29	Crestal approach – two stages implant placement
30	Staged implant placement
23	Two stages implant placement not simultaneous with sinus floor augmentation

5. Results

105 articles were identified by electronic searching. Eight original articles fulfilled the inclusion criteria as in (Figure 1).After reading the full text (Table 1): Three articles were excluded because implant placement were in two stages after augmenting the sinus, two articles were excluded also

Figure 1. PRISMA flow chart.

 Table 2. Study design and basic patient data and exclusion characters

The author	Design of study	No. patients	No. sinuses	Age (years)	Mean of age	Smokers	Systemic disease	Sinus pathology	Others
25	Randomized clinical trial	10	15	35-58		No	Yes	Yes	periodontitis
24	Retrospective	17	Not specified	Not specified	51	> 10 cigarettes per day	Yes	Yes	previous maxillary sinus surgery
14	Randomized clinical trial	12	13	20-50		Not specified	Yes	Yes	Limited mouth opening
11	Retrospective	33	Not specified	Not specified	55	Not specified	Yes	Not specified	Not specified
13	Retrospective	49	49			Yes	Yes	Yes	Not specified
26	Retrospective	73	81	29 to 78	53.79 +- 9.92	No	Yes	Yes	previous sinus surgery
27	Not specified	10	12	Not specified	Not specified	Not specified	Yes	Yes	Replacement of the bone window not possible
12	two-arm and split-mouth randomized controlled clinical trial	104 and 5	135	39 to 81	64.9	No	Yes	Yes	Pregnancy

Table 3. Char	acters of sur	gical procedures	S					
The authors	Residual	Simultaneous/	Membrane	The lateral	Tool used for	Healing	Grafting	Implants
	bone height	Staged implant placement	use	osteotomy characters	the lateral approach			
25	>5mm	Simultaneous	platelet-rich fibrin (PRF)	Not specified	ultrasonic tips	Loading after 6 months , 1 year postoperative ct	(MBCP) Biomatlante Inc., Vigneux de Bretagne, France	In-Kone TEKKA implants)
24	3.32 to 8.14mm	Simultaneous	No	bony window was used to cover the osteotomy site	a Piezo surgical tip or a number 4 diamond round bur	Follow up at 2 weeks, 4 weeks, and 4 months after the surgery for post surgical evaluation. Second stage surgery was scheduled at 5- to 6-month post sinus lift	calcium phosphate silicate (CPS) putty bone substitute (NovaBone Products, Alachua,fl)	(BioHorizons, Birmingham, AL)
14	>5mm	Simultaneous	collagen membrane	Window	3mm diameter round bur	Loading was 6 months after placement of implants.	Alloplastic bone graft (Bioactive glass putty)	Self-threaded, Tapering, Double thread, Acid etched and sand-blasted, selective Integrated surfaced Implants
3	5mm	Simultaneous	Not specified	trap-door, open-window	electric-motor drill	The prosthetic procedures of both 1 -stage or 2-stage systems were started at 7 to 8 months and with initial force loading at about 9 months after sinus lifting-combined implant surgery. follow up to 2 years	Not specified	-(ITI; Straumann, Waldenburg,Switzerland, and SwissPlus; Centerpulse Dental, Carlsbad, CA) -(Friate-2; FriadentGmbH,Mannheim, Germany)
٤١	>3mm	Not specified	Collagen	Window	hand and piezo -surgery burs	12 months of follow up after delivery of crowns.	(MinerOss, BioHorizons, Birmingham, allograft	(Nobel Biocare, Zurich- Flughafen, Switzerland)

Dental Inc.)	tem°, l Biocare AB, /eden)	aet 3i Inc., USA)			ate	eq	pa
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bone allografi (PUROS, Zimmer Dental Inc., Carlsbad, CA, USA)	Not specified	(Bio-Oss; Geistlich Pharmaceutical AG, Switzerland		rative Gra	No	cified No	ient No
lant	surgery ed is of	er		Postoper sinusitis	OLI	Not spec	One pati
delayed imp loading	Abutment connection s was perform after 6month healing.	Loading afte 6 months aft placement		1S Oration			: perforation, ed by bio- orbable nbrane.
o-surgery	llating saw sculap°, raun sungen AG, many)	specified		Sinu perf	eated no .30 nique ion, w	оц	t One) 3.7– seald). absc
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Not specifie	replaceable bone windo	Boyne & James (1980 and Tatum (1986)		ative bone he	ie of the new nm³, 0.70 ml proup and 3.4 sinus infiltrat the membrau .5 mm, Volui ied mean=2.9	14 mm (61.74 8.43 +- 2.08	n residual rid m (71.43 to m an average
ollagen ed only seal over rforations	ot specified	nrcine Ilagen ontrol oup)		Post-opera	The volum was 2.94 n sinus lift g ml in the s group Height of mean =13 bone form	Mean 13.3 Difference	Increase in ranged fro 6.1 mm (o
imultaneous C. or by staged us pproach to pe	Vot specified N	When residual \overline{Pc} some heightccvas ≥ 4 mm,(cmplantsgrwere placedinultaneouslyinultaneouslyinultaneously		Graft height reduction	1.7mm ³ Volume of bone graft material used, mean Correlation to new bone formed 0.079 mm ³	Not specified	0.3–0.7 mm 1 yeau after loading 0.2–0.3 mm 2 years after loading
<pre><6 mm s cc cc a a</pre>	Mean = 7 3.07 +- 1.35 mm	range, 4–10 T b mm v v w w w s s i n n n n	plications	Implant survival rate	Not specified	survival was 96.67% (29/30) during a mean follow- up of 15.74 months post loading.	100 %
56	27	12	Table 4. Com	The authors	25	24	14

11	100%	Not specified	Increases in lifted sinus bone height ranged from 3 mm to 9 mm with an average of 4.5 mm.	No	No	Not specified	Not specified
13	100%	Not specified	Residual sinus floor/initial bone height Mean control group particulate graft = 5.09 mm and in test group Dynablast =4.66 mm	2 patients , small perforations , covered by collagen	Not specified	Not specified	Not specified
26	Not specified	Not specified	11.32 (+- 2.48) mm	Mean perforation = 17.28%	Not specified	Not specified	Not specified
27	100%	Not specified	Mean (4–10) mm	Not specified	Not specified	Not specified	Not specified
12	95.3%	Not specified	Average of new bone 19 mm (-+ 6) covered by collagen 15 mm (-+5) uncovered by membrane	Not specified	Not specified	4	Not specified

because the augmentation were from the crest of the ridge and one article was excluded because it was a case report.

5.1 Included Studies

Eight articles were included in this review and presented in Tables 2-7. Most of the excluded patients in the included studies were the patients with history of sinusitis, immune system disorders, and uncontrolled systemic diseases; also most of the included studies excluded the smoker patients (Table 3). The surgical technique was by trap door technique or an access hole (Table 4). In the included studies, all patients underwent the lateral approach sinus floor augmentation with simultaneous implant placement.

5.2 Quality Assessment (Risk of Bias Assessment)

One randomized clinical trial was of low risk of bias, one randomized clinical trial was of unclear risk of bias because of unclear sequence generation, allocation concealment, blinding, incomplete outcome data and selective outcome reporting, one randomized clinical trial was of high risk of bias. (Table 5).

5.3 Statistical Analysis

Meta-analysis is the last step in systematic review formulation which aimed for analysis of the studies by Comprehensive Meta Analysis version 2.2.048 software, after measuring the heterogeneity to decide on the fixed or random effects approach by Cochran's Q, which provides P-value of the studies, but I² is more reliable in assessing inconsistency between studies, with values of 25%, 50% and 75% corresponding to low, moderate and high heterogeneity respectively¹¹.

Meta-analysis done for changes in bone height used the Standard Difference in Means (SDM) as the effect measure, while meta-analysis for survival rates used the rate as the effect measure.

The publication bias was checked by Funnel plot. Egger's test of the intercept was used to quantify the display of the funnel plot.

6. Results6.1 Survival Rate

Heterogeneity measures in Table 6 showed non-statistically significant Cochrane Q value (*P*-value = 1.000). I²

Author	Sequence generation	Allocation concealment	Blinding of participant, personnel and outcome assessor	Incomplete outcome data	Selective outcome reporting	Other sources of bias	Risk of bias
25	Unclear	Unclear	Unclear	Unclear	No	Unclear	Unclear
14	No	No	No	No	No	Unclear	high
12	Yes	Yes	Yes	Unclear	Unclear	No	Low

Table 5. Cochrane collaboration's tool for risk of bias assessment

value was 0.0% indicating no heterogeneity. So we can conclude that homogeneity hypothesis is not rejected and the fixed effects model will be used.



Figure 2. Forest plot for survival rate.

 Table 6. Heterogeneity measures of meta-analysis for survival rate

	Value	Df	P-value
Cochrane Q	0.1	2	0.942
I ²	0.0%		

*: Significant at $P \le 0.05$, df: degrees of freedom (n-1)

The fixed effects model (**Figure 2**) showed a survival rate (effect size) of 0.990 (99.0%) using conventional rotary tips and a survival rate of 1.000 (100.0%) using Ultrasonic tips. The overall survival rate is 0.993 (99.3%) and the effect size is statistically significant with *P*-value < 0.001.

The relative weight of the included studies, was the highest weight $(30.9\%)^{12}$ while showed the lowest weight $(7.9\%)^{13}$.

Funnel plot analysis for the included studies showed no publication bias. This was confirmed by Egger's regression intercept which showed non-statistically significant result (*P*-value = 0.548). As the results were not statistically significant, we concluded that there is no publication bias.

6.2 Changes in Bone Height

Heterogeneity measures shown in (Table 7) were statistically significant as Cochrane Q value (*P*-value < 0.001). I² value was 97.4% indicating high heterogeneity. So we can conclude that homogeneity hypothesis is rejected and the random effects model will be used.

The random effects model (Figure 4) resulted in standardized mean difference (effect size) of 0.376 using conventional rotary tips and 0.433 using ultrasonic tips. The overall change in bone height was in the direction of increase and the overall effect size was 0.410. The effect size was statistically significant with *P*-value < 0.001. So, we can conclude that there was a statistically significant change in bone height.

The relative weight of the included studies was the highest weight $(50.6\%)^{14}$ while showed the lowest weight $(32.7\%)^{15}$.

Funnel plot analysis for the included studies showed a publication bias. This was confirmed by Egger's regression intercept which showed a statistically significant result

То	ol used		The Author	•	Statistics for each study			ach study	
		Rate	Standard error	Varia	nce	Lower limit	Upper limit	Z-Value	p-Value
Rotary	27	1.000	0.289	0.083		0.434	1.566	3.464	0.001
Rotary	12	1.000	0.146	0.021		0.714	1.286	6.856	0.000
Rotary	25	1.000	0.213	0.045		0.582	1.418	4.690	0.000
Rotary	24	0.933	0.249	0.062		0.444	1.422	3.742	0.000
Rotary	14	1.000	0.277	0.077		0.456	1.544	3.606	0.000
Rotary Rotary		0.990	0.095	0.009		0.803	1.177	10.389	0.000
Ultrasoni	c 15	1.000	0.192	0.037		0.623	1.377	5.196	0.000
Ultrasoni	c 24	1.000	0.258	0.067		0.494	1.506	3.873	0.000
Ultrasoni	c	1.000	0.154	0.024		0.698	1.302	6.481	0.000
Overall		0.993	0.081	0.007		0.834	1.152	12.245	0.000

(*P*-value = 0.030). As the results were statistically significant, we concluded that there was a publication bias.



Figure 3. Funnel plot for survival rate.

Table 7. Heterogeneity measures of meta-analysis forchange in bone height (Group 2)

	Value	Df	P-value
Cochrane Q	154.9	4	< 0.001*
I^2	97.4%		

*: Significant at $P \le 0.05$, df: degrees of freedom (n-1)

7. Discussion

Many surgical tools designed recently aimed for minimizing the sinus trauma possibilities .¹⁶ Inthe contrast, some studies showed non-significant difference between piezo-



Figure 4. Forest plot for change in bone height.

electric tips and conventional rotary tips¹⁷, also some authors claimed that membrane perforation not correlate with implant survival¹⁸.



Figure 5. Funnel plot for change in bone height.

When comparing the piezoelectric tips with the conventional rotary tips, regarding the incidence of perforation was lowered from $7\%^{19}$ to $3.8\%^{20}$ while by conventional rotary tips was from 11%- $56\%^{21}$. Investigating the effect of the piezoelectric patch length on the required control voltage was the interest of recent engineering research²².

Regarding the operative time it was increased using piezoelectric tips and the rotary tips .other studies used the new reamers didn't measure the time, so more randomized clinical trials are needed which measure the time one of the outcomes. Post-operative sinusitis as an expected complication after the lateral approach sinus floor augmentation was not founded when ultrasonic tips used²³, it was founded in one patient by using 3mm round burr²⁴ and not specified in the rest of the included studies.

Regarding the implant survival rate which is the primary patient related outcome was 96.67% (29/30) during a mean follow-up of 15.74 months post loading, 95.3%, 100% when round burr used or electric motor drill or either use of hand or piezoelectric rotary tips or by using oscillating saw which means that survival implant rates not correlate to the tool used for the sinus augmentation and was not specified in the rest of studies²⁵

Graft loss was seen in four patients, not found when ultrasonic tips used, and was not specified in the other included studies²⁶ on the contrast the reduction of graft was 0.3–0.7 mm in height 1 year after loading and 0.2–0.3 mm 2 years after loading when 3mm round burr used but it was not clearly specified or calculated in the rest of the included studied²⁷.

8. Conclusions

The retrieved evidence provides a statistical significance difference of using the ultrasonic tips over the conven-

tional use of rotary tips but it is not usually to be clinically significant; especially if a highly trained surgeon could perform the lateral osteotomy by any tool also the perforation of the sinus could be managed easily especially if it of small size by covering it by collagen barrier or by suturing if it was of large size and accessible one, but the previous studies were highly supporting the ultrasonic tips over the rotary tips in reference to less traumatic surgery with minimal operative and postoperative complications. Clear reasons identified that should prompt the clinician to prefer ultrasonic tips or rotary tips or any other type of tools. Randomized clinical trials are needed to compare the newly presented tools like DASK and SLA reamers to properly asses their clinical effectiveness. Studies of patient reported outcomes are needed to clearly evaluate the cost effectiveness of each tool.

9. Conflict of Interest

The authors declare no conflict of interest. The authors don't have financial interests in the products or tools listed in the review.

10. Acknowledgment

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