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Introduction

- In 2014, 15.6 million cosmetic and 5.8 million reconstructive autologous fat grafting (AFG) procedures were performed in the United States.
- There is increasing evidence suggesting the safety and effectiveness of AFG.
- However, little is known about the efficiency of AFG.

Study Objective:

■ To conduct a literature review of the safety, effectiveness, and efficiency of AFG in all applications.

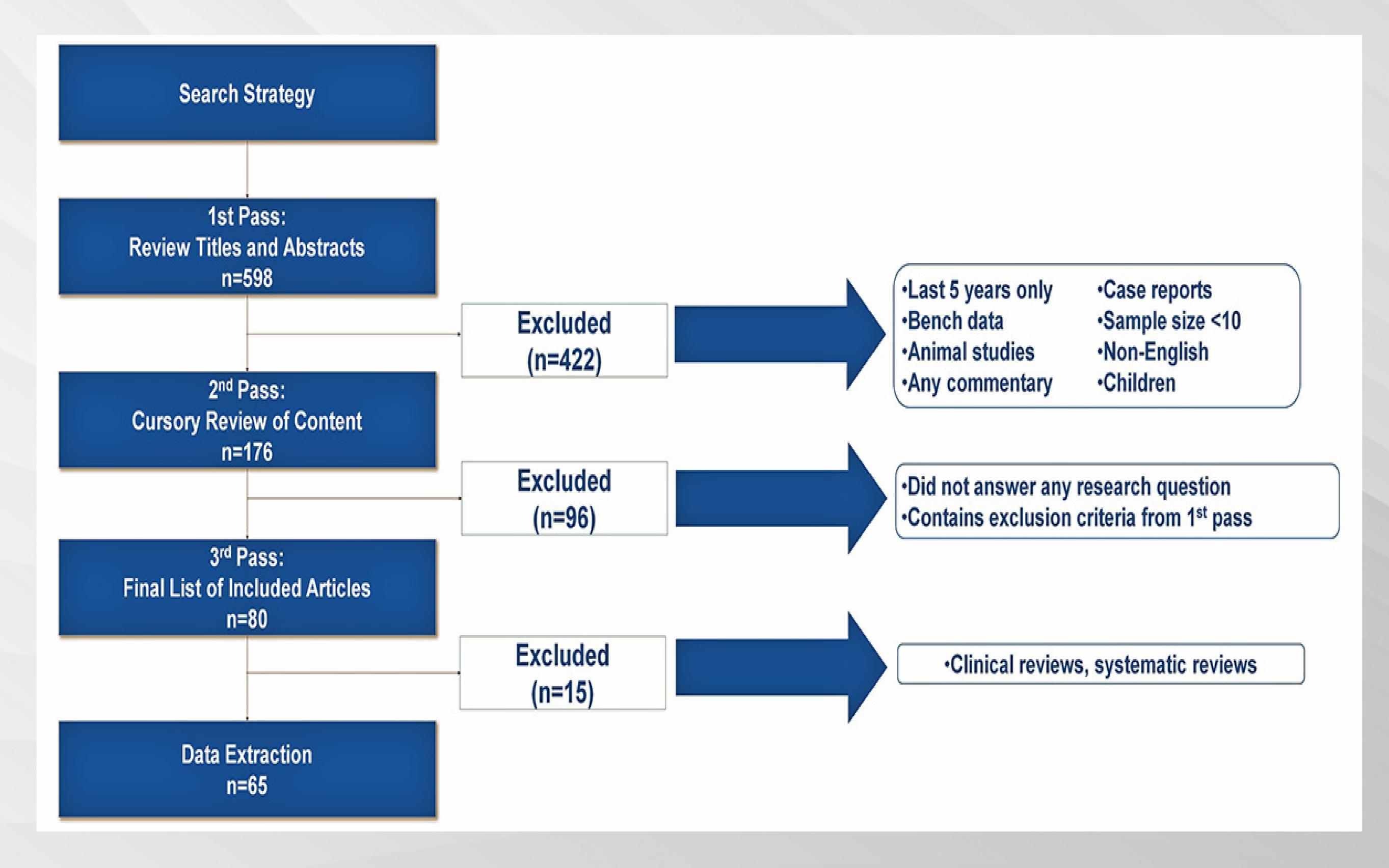
Methods

- Systematic literature review of fat grafting procedures
- Time Frame: April 1, 2010 and April 30, 2015
- Database: PubMed
- Variables collected: Study, patient, surgical characteristics, and safety, effectiveness and efficiency outcomes
- Data Output: Descriptive Statistics (Weighted Mean or Weighted Percentage), Univariate Analysis
- Two reviewers independently reviewed the articles and any differences were resolved by a third reviewer.

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Results



Studies Included:

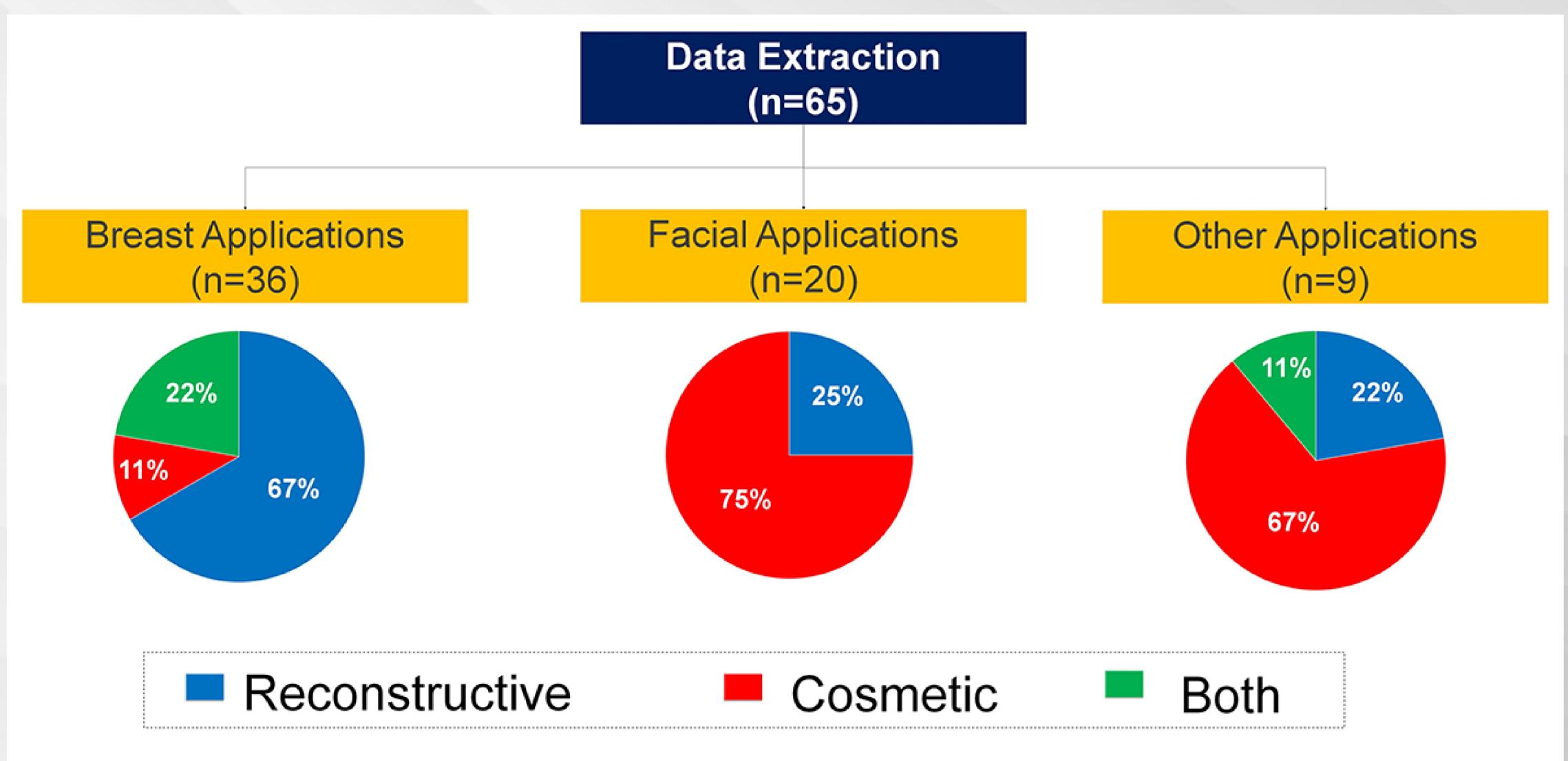
- Clinical trials
- Randomized Controlled trials
- Prospective/ Retrospective studies
- Epidemiology studies
- Burden and cost of illness

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Results (Cont'd)

Study characteristics



Variable	Breast	Facial	Other
Mean Follow-Up Time Months (Range)	25.4 (5-91)	15.1 (3-27)	52.2 (12-71.5)
Mean Study Size Subjects (Range)	121.1 (18-1000)	69.2 (12-500)	241.3 (12-789)

Patient characteristics

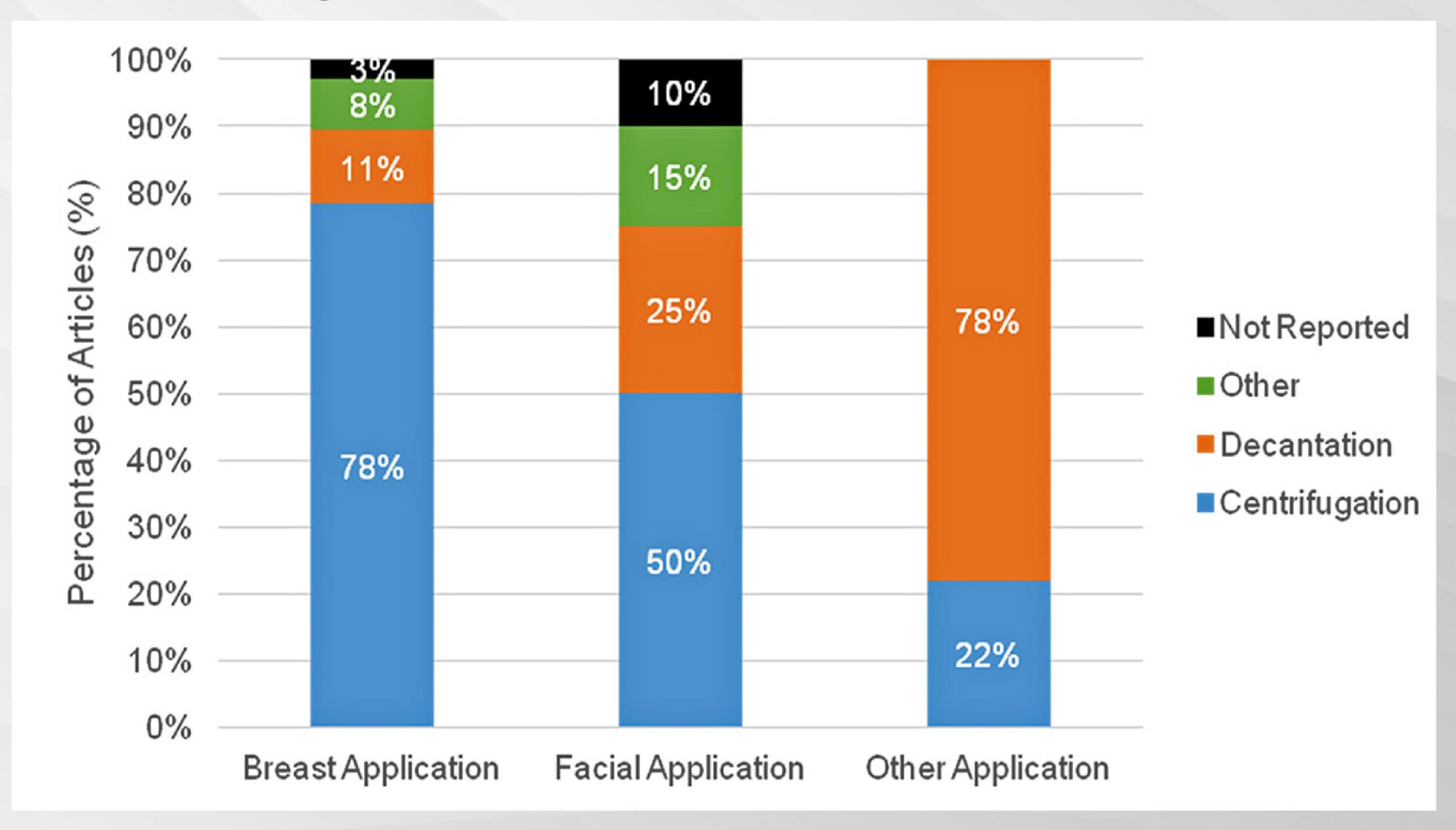
Facial application patient cohort had a higher mean age and
 BMI (body mass index) compared to breast or other applications.

Variable	Breast	Facial	Other
Age yrs, (Range)	44.5 (23-65)	50.2 (17-74)	31.9 (28-46)
BMI kg/m², Range)	22.6 (18.5-29.9)	23.9 (23.5-24.3)	21.5 (20.4-24.6)

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Results (Cont'd)

Fat processing technique



Safety & Effectiveness Outcomes

 Breast cancer recurrence rates post-reconstruction were about 3% using a weighted average (by sample size).

Variable	Breast	Facial	Other	
Reoperation	20%	3%	6.5%	
(Range)	(3-68%)	(0-12%)	(5-16%)	
Cyst Formation	5%	0%	Not	
(Range)	(0-25%)	(0%)	Reported	
Fat Necrosis	6%	1.1%	1.4%	
(Range)	(0-19%)	(0-2%)	(0-3%)	
Infection	1%	4.3%	0.4%	
(Range)	(0-4%)	(0-14%)	(0-1%)	
Retention	62%	63%	700/ *	
(Range)	(39-78%)	(54-85%)	70%*	
Patient Satisfaction (Range)	93% (68-100%)	89% (46-100%)	96% (91-100%)	

^{*} Range Not Available

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Results (Cont'd)

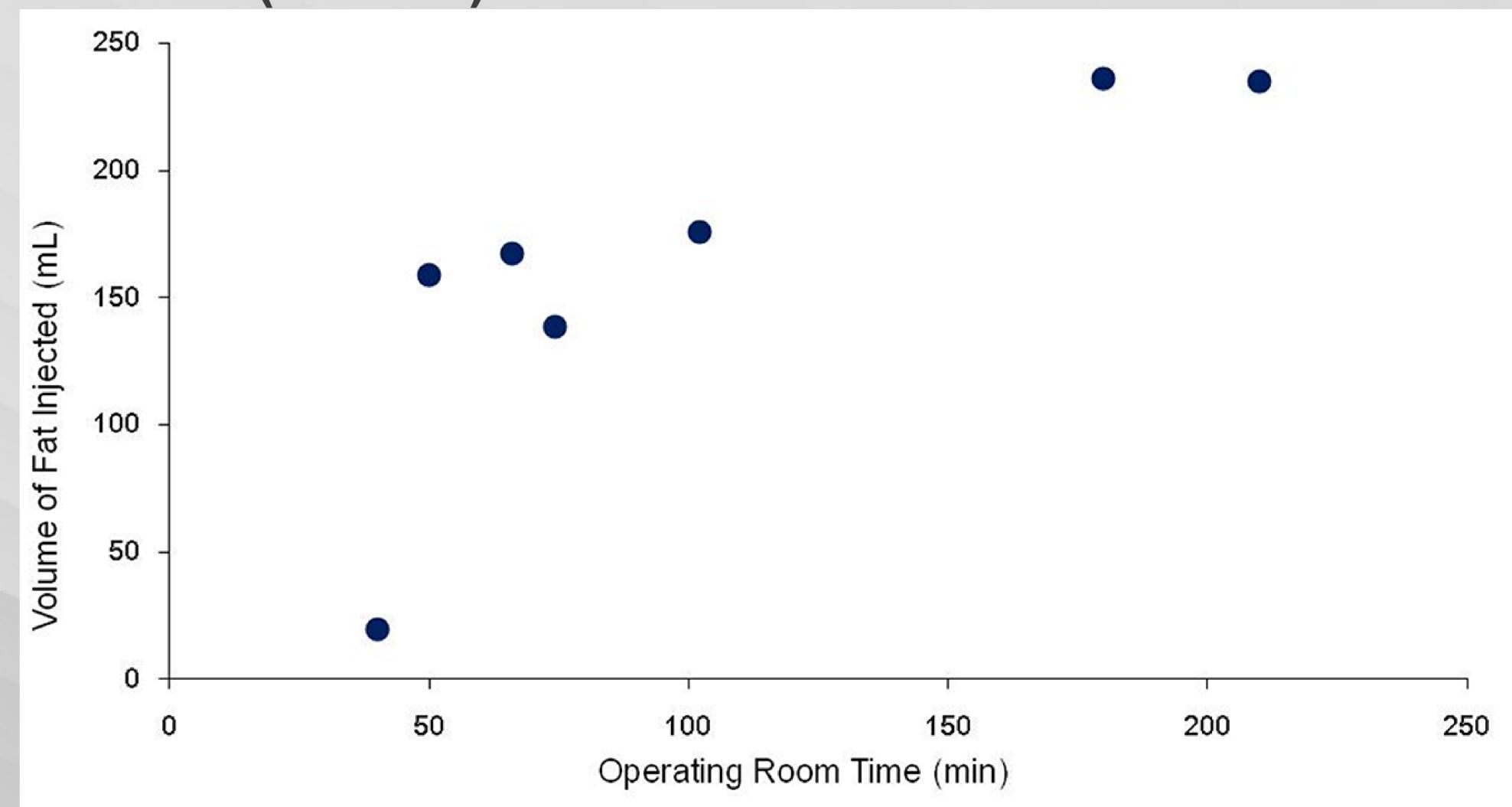
Efficiency outcomes

- For breast applications, there was an association between volume injected and operating room time.
- Data was not available for facial/other applications.

Variable	Breast	Facial	Other
Mean Harvest	558 ml	62 ml	1753.8 ml*
Volume (Range)	(120-1299ml)	(35-360ml)	
Mean Injection Volume (Range)	145 mL	14.5 ml	336.1 ml
	(20-606 ml)	(2.1-27.1 ml)	(18-692 ml)
Operating Room Time (Range)	125 minutes	Not	Not
	(40-210 minutes)	Reported	Reported

^{*} Range Not Available





Conclusions

 Safety/Effectiveness: The safety and effectiveness results were consistent and validate previous research published.

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Conclusions (Cont'd)

- Safety/Effectiveness: The safety and effectiveness results were consistent and validate previous research published.
- Efficiency: The efficiency data available, although limited, suggest that there is an opportunity to reduce OR time and resources.
- Limitation: There was high variability and lack of uniformity in reporting among these studies.
- Limited data suggests need for the ASPS effort to collect and standardize data by using the GRAFT registry.

References

Available upon request

Contact: PubsMgt@Acelity.com

Disclosures

- SLS: Consultant to LifeCell, an ACELITY Company,
 Allergan, Establishment Labs, and Novadaq and
 Endurance Labs
- CNC: Received funding to conduct research
- BKL: Employee of ACELITY
- MG: Received funding to conduct research
- MP, DM: Employees of LifeCell, an ACELITY
 Company