



Introduction

- Youth with craniofacial conditions are at risk for psychosocial difficulties, including learning issues, anxiety, and social stigmatization (Kapp-Simon, 2017; Bous et al., 2020). Select studies have documented psychosocial challenges among parents and caregivers of affected youth (Habersaat et al., 2018).
- Use of a validated psychosocial risk screening instrument during team visits can improve risk identification and increase access to psychosocial consultation (Crerand et al., 2022).
- The Psychosocial Assessment Tool–Craniofacial Version ullet(PAT-CV) assesses risk across multiple domains (e.g., social support, craniofacial-specific problems, & resources) (Crerand et al., 2022).
 - Patients may fall into 3 risk categories—universal, targeted, or clinical.
- A QI project was initiated to pilot administration of the PAT-CV in a grant-funded, interdisciplinary craniofacial clinic.

Methods

- Parents of patients 18 and younger were administered the paper version of the PAT-CV.
- Changes were made to PAT–CV processes, including:
 - Administration in 100% of clinics (versus 50%)
 - Introduction of the PAT–CV at check-in with script
 - Implementation of phone calls between 3-6 months after completion by the clinic social worker to ensure access to recommended services and supports

Implementation of Psychosocial Risk Screening in an Interdisciplinary Craniofacial Clinic

Results



Number of families with completed PAT–CV by year



Percent of Patients per Risk Category by Year





Wolfson Children's Hospital

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- lacksquare
- - 52.4% male
- ranged from 25.2% to 30.2%.

Conclusion

- \bullet craniofacial populations.
- paced clinic setting.
- services.
- Next steps: \bullet

 - reduce staff burden.

* Full reference list available upon request

Data collection began in 2020; however, these data (n = 45) are excluded from comparison because of the onset of the COVID-19 pandemic.

Youth demographics (as reported by parents):

- 42.4% combined cleft lip and cleft palate

Between 2021 and 2023, percent of families that fell in the targeted or clinical risk categories

Percent of patients in the targeted and clinical risk categories (> 25%) highlights the importance of psychosocial risk assessment in the cleft and

Collaboration among multiple team members is required to administer screening measures in a fast-

Personalized follow-up via phone calls can potentially help reduce barriers to accessing care and increase likelihood of receiving psychosocial

> Determine success of connecting with families via follow-up phone calls (% families reached). > Examine rates of referral for mental health and neurodevelopmental assessment services (e.g., outpatient psychotherapy, psychiatric medication management, neuropsychological testing). > Electronic administration of the PAT-CV to help

Barriers to obtaining orthodontic care for patients with orofacial clefts: A Pediatric Health Information System (PHIS) Database Study

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Introduction

- Orofacial clefting (OFC) is a common birth defect, with healthcare costs 6-10 times higher than for unaffected children.
- Patients need long-term interdisciplinary care, at minimum including surgery, speech-language pathology, and orthodontics.
- Significant financial and non-financial barriers exist, especially for minorities and those with government-based insurance.
- The Affordable Care Act (ACA, 2010) aimed to expand Medicaid access, but state variations and optional expansion have limited their effectiveness. Currently, 40 states have chosen to expand.
- Separately, 32 states have mandates to provide care for atrisk children, with 18 states specifically targeting OFC and craniofacial disorders.
- Florida's non-Medicaid expansion status poses additional challenges despite state mandates guaranteeing OFC care.

Objectives

• To assesses barriers to OFC care in Florida and nationally, using the age of alveolar bone grafting (ABG) as a marker.

Materials and Methods

- Database: Pediatric Health Information System (PHIS), queried using ICD 9/10 codes for patients <18 years with cleft palate who underwent ABG.
- Data Range: 2010-2019 (pre-COVID-19 data).
- Variables:
- Age at ABG.
- -Medicaid expansion status and implementation date by state
- -Funding source: government, private, or other insurance.
- Statistical Analysis:
 - -Software: Microsoft Excel for descriptive analysis, SAS for statistical analysis.
 - -Tests Used: Mann-Whitney U, Kruskal Wallis, Dwass-Steel-Critchlow-Fligner test, and Bonferroni adjustment.
 - -Significance Level: p < 0.05.
 - -Multivariate Analysis: Linear regression with race and ethnicity as covariates.

all we do all for kids."

Results



Figure 2: Patient Demographics and ABG Age

	n (%)	Age of ABG (SD) (years)	p-value*	Group	Type of payment	n (%)	Age of ABG (SD	
Total Patients	1182 (100)	10.3 (±3.2)			In-State Medicaid (managed care)	223 (19)	10.5 (±3.1)	
Gender			0.243	ut	In-State Medicaid (other)	132 (11)	8.6 (±3.7)	
Male	670 (57)	10.3 (±3.0)		243 government	Out-of-State Medicaid (all)	41 (3)	10.1 (±2.4)	
Female	512 (43)	10.3 (±3.4)			Other Government	28 (2)	10.5 (±3.2)	
				Ŭ	Medicare	3 (0)	10.7 (±1.0)	
Race			0.070		CHIP	14 (1)	13.3 (±3.9)	
White	673 (57)	10.4 (±3.4)		क	Commercial PPO	267 (23)	9.6 (±2.8)	
Asian/Pacific Islander	228 (19)	9.8 (±2.7)		Private	Commercial Other	220 (19)	10.5 (±3.3)	
Other	166 (14)	10.3 (±2.7)			Commercial HMO	174 (15)	10.0 (±3.4)	
Black	60 (5)	10.8 (±3.8)			TRICARE	37 (3)	10.0 (±2.5)	
Unknown	41 (3)	10.8 (±3.1)			Unknown	21 (2)	11.3 (±3.4)	
American Indian	8 (1)	9.2 (±1.4)		Other	Other Payor	16 (1)	11.2 (±3.3)	
					Self Pay	5 (0)	10.8 (±2.1)	
Ethnicity			<.001		Charity	1 (0)	6.8 (±0.0)	
Not Hispanic or Latino	866 (73)	10.2 (±3.1)			olar Bone Graft; SD, Standard Deviation; CHIP, C		rance Program; PPO,	
Hispanic or Latino	196 (17)	10.8 (±3.2)		Preferred	provider organization; HMO, Health Maintenance	Organization		
Unknown	120 (10)	9.9 (±3.8)		Source of Payment				
Source of Payment			0.005		Nationally	Florida		
Government Funding	441 (37)	10.6 (±3.3)						
Private Funding	661 (56)	10.1 (±3.2)						
Other Funding	80 (7)	10.6 (±3.0)			7%			
Florida, Source of Payment	32 (100)	10.8 (±4.0)			5776	43%	57%	
Government Funding	16 (50)	11.6 (±4.5)			56%			
Private Funding	12 (38)	10.1 (±3.1)						

ABG, Alveolar Bone Grail, SD, Standard Deviatio *Kruskal Wallis tests used for statistical comparison. Dwass-Steel-Critchlow-Fligner used for Post hoc analysis



Figure 3: ABG Age by Medicaid Expansion Status and Source of Funding

Table 3: ABG Age by Sub-population

Total Patients

ABG in a non-expansion state

ABG in an expansion state

Pre-expansion*

Post-expansion*

Statistical Comparisons

A. Privately funded patients vs. B. Governme

- A. Expansion state vs. B. Non-expansion stat All insurances
- Government funded patients only
- Privately funded patients only

Medicaid Expansion State Sub-Analysis

- A. Privately funded patients vs B. Governm
- A. Pre vs. B. Post-expansion, all insurance
- A. Pre vs. B. Post-expansion, government
- A. Pre vs. B. Post-expansion, privately fund

xpansion and post-expansion sub-categories expressed at percentage of expansion state categor J tests used for univariate analysis Linear regression model with Race and Ethnicity as covariates used for multivariate analysis



Conclusion

- state interventions.
- access
- underestimated

JOHNS HOPKINS

MEDICINE

JOHNS HOPKINS

ALL CHILDREN'S HOSPITAL

Тс	otal	Gov	vernme	nt Funding	Private	Funding
n (%)	Age (SD)	n (%)		Age (SD)	n (%)	Age (SD)
1102 (100)		441 (1	100)		661 (100)	
283 (26)	10.7 (±3.1)	140 (3	32)	11.0 (±3.2)	143 (22)	10.3 (±3.0)
819 (74)	10.2 (±3.2)	301 (6	38)	10.4 (±3.3)	518 (78)	10.0 (±3.2)
354 (43)	9.9 (±3.4)	121	(40)	9.8 (±3.1)	233 (45)	9.8 (±3.6)
465 (57)	10.4 (±3.1)	180	(60)	10.8 (±3.4)	285 (55)	10.1 (±2.9)
		Age of AB	G (year	rs) Difference	p-'	value
	n	A vs	. В	(months)	univariate [†]	multivariate [‡]
ent funded patients	1102	10.1 vs	. 10.6	6.0	0.003	0.047
ate						
	1102	10.2 vs	. 10.7	6.0	0.003	0.023
	441	10.4 vs	. 11.0	7.2	0.045	0.109
	661	10.0 vs	. 10.3	3.6	0.078	0.188
ment funded patients	819	10.0 vs	• 10 /	4.8	0.036	0.004
es	819	9.9 vs			0.002	0.004
t funded patients only	301	9.8 vs			0.008	0.007
nded patients only	518	9.8 vs	. 10.1	3.6	0.025	0.153

• Patients with OFC face significant barriers to oral care despite federal and

• Medicaid expansion is linked to later ages of ABG; however, it is important to consider in the broader context of expanding healthcare

• Expanded coverage improves preventive care, timely diagnosis, and comprehensive OFC management and long-term benefits cannot be

• Florida shows that even with mandates, limited resources impede care delivery, reflecting common challenges in other states.

Publication Trends and Surgeon Perceptions: A Comprehensive Analysis of Gender **Disparities in Craniofacial Surgery**

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Introduction

- Women consistently face underrepresentation in various aspects of craniofacial surgery, including the number of surgeons, faculty positions, leadership roles, and conference representation.
- Academic productivity (often measured via publication) records), is a critical for career advancement and previous studies have shown female plastic surgeons consistently have fewer publications.
- Recent years have seen significant progress in DEI within medicine, surgery as a whole, and the plastic surgery, emphasizing the need for frequent and comprehensive updates in research to capture the changing landscape.
- While disparities exist, younger cohorts in craniofacial surgery show a more equitable distribution of leadership roles, signaling some advancements in DEI.

Objectives

- Examine the academic productivity, gauged by publication counts, of academic craniofacial surgeons in 2022. This will create a baseline to measure the impact of ongoing DEI initiatives.
- Directly assess craniofacial surgeons' perception concerning DEI, barriers obstructing progress, and suggestions for improvement.

Materials and Methods

- A dataset of 193 craniofacial surgeons and fellows in the United States and Canada were compiled using the websites of accredited plastic surgery training programs.
- Data collected for each surgeon included gender, total publications, first-author publications, senior-author publications in 2022, and fellow or program director status.
- A 19-question survey was distributed to craniomaxillofacial surgeons through ACPA's list service.
- The survey collected data on participants' demographics, practice details, leadership roles, and research funding sources.
- Optional free-response questions explored participants' perceptions of Diversity, Equity, and Inclusion in craniofacial surgery and their suggestions for improvement.

Results

Figure 1: Demographics and Publication Trends of Craniofacial Surgeons in 2022

	-			
	Total	Male (73%)	Female (27%)	p val
	n = 193	n = 140	n = 53	·
Craniofacial Surgeons n (colu	ımn %)			
Fellow	17 (9)	6 (35)	11 (65)	0.22
Attending	176 (91)	134 (76)	42 (24)	<0.0
Program director	29 (15)	26 (90)	3 (10)	<0.0
Total cohort	193 (100)	140 (73)	53 (27)	0.00
Program region <i>n (column %)</i>)			
Canada	9 (5)	8 (89)	1 (11)	0.50
Midwest	39 (20)	28 (72)	11 (28)	0.07
Northeast	46 (24)	32 (70)	14 (30)	0.01
South	58 (30)	41 (71)	17 (29)	0.06
West	41 (21)	31 (76)	10 (24)	0.02
Publications counts of Cranic	ofacial Surgeons in 2022 n	(column %)		
First author	52	27 (52)	25 (48)	0.78
Last author	513	433 (84)	80 (16)	<0.0
Total	1134	922 (81)	212 (19)	<0.0
Average publications per Cra	niofacial Surgeon in 2022	(SD)		
First author	0.27 (±0.88)	0.19 (±0.56)	0.47 (±1.41)	0.24
Last author	2.66 (±4.78)	3.09 (±5.27)	1.51 (±2.88)	0.00
Total	5.88 (±7.64)	6.59 (±8.42)	4.00 (±4.62)	0.04



Figure 2: Academic Experience, Leadership, and Research Among Survey Respondents

	Total	Male (65%)	Female (35%)	p value
Total Responses	n = 26	n =17	n = 9	
Mean years since fellowship (SD)	10.30 (±7.65)	10.47 (±7.35)	10.00 (8.65)	0.885
Leadership positions held <i>n (column %)</i>				
Department chair	4 (15)	3 (18)	1 (6)	1.000
Program or associate program director	14 (54)	9 (53)	5 (29)	1.000
National Society leadership	9 (35)	6 (35)	3 (18)	1.000
Regional Society leadership	5 (19)	3 (18)	2 (12)	1.000
Division Chief	3 (12)	2 (12)	1 (6)	1.000
Team Director	2 (8)	1 (6)	1 (6)	1.000
Prefer not to respond	2 (8)	2 (12)	O (O)	0.529
Total positions held	37	24	13	
Sources of research funding n (column %)			
Industry sponsorship	3 (12)	2 (12)	1 (11)	1.000
Institutional grants	4 (15)	3 (18)	1 (11)	1.000
NIH	1 (4)	O (O)	1 (11)	0.346
National societies	4 (15)	4 (24)	0 (0)	0.263
Other	2 (8)	2 (12)	0 (0)	0.529
Prefer not to respond/None	10 (38)	5 (29)	5 (56)	0.234
Total number of sources	14	11	3	



A Statistical Fragility Analysis of Bilateral Sagittal Split Osteotomies

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BACKGROUND

Randomized controlled trials (RCTs) evaluating bilateral sagittal split osteotomies (BSSO), a surgical procedure performed on the lower jaw to correct certain types of mandibular deformities, contain differing results regarding its efficacy. Fragility index (FI), reverse fragility index (rFI), and fragility quotient (FQ) values represent the statistical fragility of outcomes reported in RCTs evaluating BSSO.

METHODS

PubMed and MEDLINE were systematically searched for RCTs from January 1, 1998 to May 1, 2024 for outcome assessment of BSSO. Of 85 RCTs screened, 6 studies were included for analysis. We computed FI and rFI, denoting the quantity of outcome event reversals necessary to change the statistical significance for significant and non-significant outcomes, respectively. The FQ was determined by dividing the FI by the study sample size.

RESULTS

Author	Year	Journal	Total Sample Size
Leung et al.	2021	Int J Oral Maxillofac Surg	196
Baas et al.	2015	Int J Oral Maxillofac Surg	68
Ow et al.	2010	Int J Oral Maxillofac Surg	23
Kohnke et al.	2017	J Oral Maxillofac Surg	117
Baas et al.	2015	Int J Oral Maxillofac Surg	63
Baas et al.	2015	Int J Oral Maxillofac Surg	63

Table 2 – Fragility data based on trial and outcome characteristics					
	Number of Outcomes	Mean Fragility Index (SD)	Mean Fragility Quotient (SD)		
All RCT Outcomes	12	4.92 (2.61)	.050 (.033)		
Significant Outcomes (P<0.05)	3	6.67 (3.79)	.048 (.020)		
Nonsignificant Outcomes (P≥0.05)	9	4.33 (2.06)	.050 (.037)		
Fragility of Key Outcomes					

- IAN Deficit
- FI: 5, FQ: .026
- Wound Infection
- ∘ rFI: 1, FQ: .435
- Objective and Subjective Neurosensory
 Deficit
- Objective rFI: 4, FQ: .074
- $\circ~$ Subjective rFI: 7, FQ = .111

LIMITATIONS

- Fragility indices are only appropriate for dichotomous outcomes.
- There is no specific cutoff or lower limit of the fragility index to classify a study as "fragile" or "robust."



CONCLUSIONS

The efficacy of BSSO from RCTs is statistically fragile, particularly the outcome regarding wound infection. We recommend combined reporting of p-values with FI and FQ metrics to aid in interpreting clinical findings evaluating BSSO. Additionally, there should be a larger analysis, including a greater sample size of RCTs, to produce a more robust FI.

CLINICAL RELEVANCE

By determining how susceptible the results are to changes in a small number of events, clinicians can assess the reliability of the evidence supporting the efficacy and safety of BSSO procedures. This insight aids in better decision-making, risk assessment, and resource allocation, ensuring that clinical practices are grounded in solid, dependable evidence.

FCPA Characterizing Innovation in Cleft Palate and Craniofacial Surgery

FIGURES

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INTRODUCTION

The number of patent filings for surgery has increased tremendously over the past 50 years, yet trends within specific specialties remain poorly understood.

The pursuit of providing the best care for patients has driven cleft palate and craniofacial surgeons to build upon the foundations laid by their predecessors over the decades. Dr. Ralph Millard, renowned plastic surgeon, summarized this quest with the phrase, "Semper investigans, nunquam perficiens," meaning "Always searching, never fully attaining perfection"1

OBIECTIVES

This study aims to identify the areas and directionality of innovation within the field of cleft palate and craniofacial surgery (CPCS) through findings in intellectual property. The idea to profile innovation within this field came from Kwasnicki et al., a research article within plastic surgery.2

METHODS

A query of the LexisNexis TotalPatent One® database was performed to analyze patents filed worldwide regarding CPCS from 1974 to 2023. The Boolean keyword search "cleft palate' OR 'craniofacial surgery'"was employed.

Patents related to CPCS were defined as CPCS surgical methods, devices, implantables, introducers and sterilization equipment based on the Cooperative Patent Classification (CPC) code. Categorical data, including patent progress and demographic information, was obtained for each relevant CPC code.

Annual Cleft Palate and Craniofacial Patent Filings Filings Total Patent F $R^2 = 0.76$ the th 1980 1990 2000 2010 2020 Year

Figure 1. Annual Cleft Palate and Craniofacial Patent Filings





Figure 2. Patent Status Distribution

RESULTS

School of

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Medicine at

A total of 468 patents related to CPCS were filed over the period of study. Of the 468 patents filed, 183 (39.1%) were granted, 123 (26.3%) were ceased, and 162 (34.6%) are pending. The United States is the authority with the highest number of patent filings (160; 34.1%). The category with the most patent filings was CPC A61L: Methods or Apparatus for Material Sterilization (227; 48.5%). Nevertheless, the category with the most patents granted was CPC A61B: Diagnostic Surgical Devices (87; 47.5%).

- Actual Data

Legend

There was a general exponential (monotonic) growth in the + Fitted Model number of patent filings per year (Exponential $R_2 = 0.762$; rs = 0.910, p = 0.001), with 2022 having the greatest number of filings (46; 9.83%).

CONCLUSION

This study shows growth in patent filings and the diverse, evolving landscape of innovation within CPCS, corroborating the conclusions of recent literature.³ It also emphasizes the need for continued analysis on which patent codes are obtaining more or less filings to inform future advancements in the field.

LIMITATIONS

Although comprehensive, there remains the possibility that the LexisNexis TotalPatent® database does not contain all CPCS patents filed to all patent registries worldwide.

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GROW22q: A Referral Initiative to Expand a 22q Clinic

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Background / Description

- Patients with 22q11.2 deletion syndrome have a wide array of medical conditions and require yearly screening with close followup.¹⁻³
- Their care should be provided by an interdisciplinary team of subspecialists who are experienced with the condition.¹⁻³
- International care guidelines exist to help teams optimize care involving multiple organ systems. ¹⁻²

Aim

Increase referrals of patients with 22q11.2 deletion syndrome to our craniofacial clinic from < 1 per quarter to 2 per quarter starting January 1, 2023, and to sustain through December 31, 2024.

Design / Strategy

Our team sought to increase the number of 22q patients referred to our clinic through improving education and awareness in the community and improving accessibility to our clinic.

- Marketing
- Key stakeholder engagement

Run Chart



- Clinic coordination
- EMR tools



Measures

- Outcome Measures:
 - Number of patients referred to the clinic per quarter

Total patients scheduled since project initiation 15





Challenges/Barriers

- Low "n"
- Provider turnover
- Anticipate future access challenges
- Ensuring team is easily accessible and clear plan is in place for consults prior to education sessions
- With little improvement following our community engagement bundle, and following feedback received by providers, our team focused on our accessibility and education

Lessons Learned

- Our <u>Education and Accessibility Bundle</u> seemed to yield the biggest improvement in referrals
- Education sessions for physicians at division meetings (NICU, cardiology, genetics, etc.) reach a large amount of referring physicians
- Caring for children with 22q11.2 deletion syndrome in a multidisciplinary clinic ensures proper completion of screening per guidelines

Next Steps

- Pool data quarterly
- Continual conversations, education, marketing

- Process Measures:
 - Number of patients that received Best Practice Alert in EMR
 - Number of patients with new diagnosis of 22q11 Deletion Syndrome in EMR
- Balancing Measures:
 - Number of no-shows for referrals
 - Number of available timeslots in craniofacial clinic

Changes Made



Community Engagement Bundle

- "Patient Stories" shared with families and on social media
- Team attendance at "22q at the Zoo" event
- Social Media posts

Education and Accessibility Bundle

- Education of SLPs at OCPS
- NICU providers educated at division meeting
- "22q Team" created on hospital's secure messaging system

- Additional team clinic time beyond critical mass
- Improve academic presence at national/international meetings

References

McDonald-McGinn DM, Sullivan KE, Marino B et al. 22q11.2 deletion syndrome. Nat Rev Dis Primers. 2015 Nov 19;1:15071.

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Background

Cleft palate repair aims to reconstruct the levator veli palatini (LVP) muscle establish proper velopharyngeal and closure. Although morphology and function of the LVP have been studied, there is no consensus on the quantitative characterization of successful cleft palate repair procedures and further, to date, no studies have directly evaluated or measured the cleft morphology and defect immediately pre- and postpalatoplasty via intraoperative magnetic resonance imaging (MRI).

Methods

- study, we compare the • In this immediate pre- and post-palatoplasty anatomy of the LVP in six patients between the ages of 11 and 51 months via intraoperative MRI.
- Measurements of LVP length and were obtained using the thickness oblique plane coronal and measurements of velar length, velar thickness, and velopharyngeal gap were obtained using the sagittal plane.
- Paired sample T-test was used to assess statistical significance between for immediate pre- and post-palatoplasty.

Objective Analysis of Immediate Palatoplasty Results via Intraoperative MRI

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Results



Figure 1. Pre-operative (A) and post-operative (B) oblique coronal images of patient 1 displaying measurements of LVP length. Pre-operative (C) and post-operative (D) sagittal images of patient 1 displaying measurements of velum length and thickness



Figure 2. Pre-operative (dark grey) averages were compared to post-operative (white) averages with standard deviations for LVP length, LVP thickness, velum thickness, velum width, and pharyngeal depth (n=6). Statistical significance is labeled above the bar graph (* <0.05, ** <0.01, *** <0.001)

Results indicate that MRI quantification provides a valuable evaluation of the preoperative LVP anatomy. Early postsurgical results indicate no evidence of fistula formation or velopharyngeal insufficiency.

This study reports the first use of intraoperative MRI measurements in a cohort of patients to objectively assess cleft palate pre-operative anatomy and surgical repair results.



Results

• Mean LVP length increased from 47.8mm to 61.2mm (p=0.010)

• Mean LVP thickness at the central palate was reconstructed to 7.8mm (p<0.001).

• Mean velar length increased from 8.9mm to 26.3mm (p=0.002)

• Mean velar width increased from 5.8mm to 11.9mm (p<0.001)

• Mean velopharyngeal gap decreased from 6.9mm to 1.7mm (p=0.013)

Conclusions

College of Medicine UCF

The Interdisciplinary Care Team for Microtia and **Aural Atresia**



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INTRODUCTION

- Children with microtia and atresia require coordinated subspecialty care for optimal outcomes.
- Interdisciplinary care for patients with craniofacial differences, especially cleft palate, is the current standard.
- Children with microtia often receive isolated care.
- We aim to present our recommended comprehensive model to routinely address the needs of microtia and atresia patients through coordinated interdisciplinary care.
- We aim to enhance attendees' knowledge of the complex needs of patients with microtia, how these needs should be routinely addressed through coordinated interdisciplinary care.



Figure 3. Our NICEAR team

RESULTS

- Of n=76 patients with microtia evaluated at the interdisciplinary clinic, all were identified with conductive hearing loss secondary to atresia and were fitted with **bone conduction hearing devices**.
- **Microtia reconstruction surgeries** (n=6) were completed with both autologous (n=2) and allogeneic cartilage (n=4).
- Among patients with concerns for autism who were referred to behavioral health (n=11; 14.5%), a minority had a pre-existing diagnosis (n=5; 6.6%).
- As shown in Figure 2, n=10 referrals were provided for **genetics**. However, 9 patients with consults to genetics prior to coming to our clinic and another 10 consults placed by us are still waiting for evaluation. Only a small subset of patients identified as syndromic (n=6) and were already being followed by genetics prior to screening at the interdisciplinary clinic.
- A standardized screening tool for soft palate dysfunction has been implemented as part of our patients' evaluation with speech pathology (Figure 6). Hypernasality indicative of VPI was detected in n=3 out of 15 patients evaluated.

METHODS

We modeled an interdisciplinary clinic for patients with microtia after craniofacial centers nationwide.

Disciplines include plastic surgery, otolaryngology, audiology, speech pathology, genetics, developmental pediatrics, behavioral health, social work, ophthalmology, and 3D photography.

Team discussions facilitate review of microtia grade, surgical options, ear anatomy, and degree of hearing loss relative to language development.

The first 5 years focus on the **importance of hearing augmentation** as a crucial step for speech and language development. A CT scan is performed at age 5 to determine whether the patient is a candidate for hearing restoration surgery.

Reconstructive choice guides timing, type, and placement of hearing devices to optimize access to sound from an early age.

- Genetic counseling and social, educational events facilitate **dynamic** learning experiences.
- Social work, behavioral assessments, and psychological evaluations address patients' coping abilities, reinforcing holistic care.
- For many patients, this **integrated care** approach aids in **uncovering** coexisting conditions which may previously have been undiagnosed under isolated care.



Figure 4. Nemours Children's Hospital in Orlando, Florida.



Figure 6. Screening algorithm at NICEAR to evaluate for VPI. At the patient's initial visit, a certified bilingual speech-language pathologist (SLP) screens both English- and Spanish-speaking patients using the Cleft Audit Protocol for Speech-Augmented-Americleft Modification (CAPS-A-AM). This is a validated method for velopharyngeal evaluation that is often used in patients with cleft palate. When indicated, techniques are used to visualize and quantify soft palate dysfunction through measures of velopharyngeal closure and nasal air emission. Since VPI can impact the intelligibility, confidence, and quality of life (QOL) of our patients, we assess perception of speech and emotional impact on our patients and caregivers using the Intelligibility in Context Scale (ICS) and Velopharyngeal Insufficiency Effect on Life Outcomes (VELO) questionnaires.

CONCLUSIONS

A **holistic** approach that treats the patient rather than the disease is critical

RESULTS



INTERDISCIPLINARY CLINIC

Patients Evaluated at the **Interdisciplinary Clinic**



Figure 1. Distribution of unilateral versus bilateral microtia at our clinic. At the Nemours Interdisciplinary Center for Ear Reconstruction (NICEAR), we have evaluated n=76 patients with microtia since January 2023. The majority of patients (n=69) presented with unilateral microtia, of which n=18 have hemifacial microsomia (HFM).

Services and Referrals Provided to Patients Through the **Interdisciplinary Clinic**

Nasopharyngoscopy referrals

Figure 5. Interdisciplinary care is necessary for addressing our patient's complex needs. We implement this form of care at our NICEAR clinic, dedicated to providing a holistic approach in caring for our patients with microtia and atresia and providing support to the patients and families.

NICEAR Clinic Flow

- ~15-30 minutes each plastic surgery, with otolaryngology, audiology, speech pathology, and social work.
- If indicated, additional same-day referral may be provided for genetics, developmental pediatrics, **Initial Patient** behavioral health, ophthalmology, and/or 3D **Visit at NICEAR** photography.

- in patients diagnosed with microtia and atresia.
- Delivering care in the setting of an interdisciplinary team helps **reveal** trends, highlight new concerns, and select individualized reconstructive options, allowing for overall patient restoration.
- Incorporation of **social events** within our center allows for **kids to meet** other children living with the same condition and for their parents to form a support network.
- Our educational events allow parents to learn more about microtia and atresia as well as what our Ear Hub can offer.
- Interdisciplinary care of patients with microtia provides reassurance and education for the family, while standardizing precisely planned, coordinated intervention throughout the child's life.
- This allows for **focus on critical developmental periods** including speech and biaural hearing, allowing patients to achieve their full developmental potential and addressing needs beyond plastic surgery and otolaryngology care.





Figure 2. Services and referrals provided to patients through the NICEAR clinic. At each visit, patients see the core members of the interdisciplinary team—plastic surgery, otolaryngology, audiology, speech-language pathology, and social work. As these assessments occur during the visit, referrals may also be provided to other same-day services, and microtia reconstruction surgeries may be planned as a part of follow-up care when deemed appropriate following careful patient selection.



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Post-clinic meeting reviewing all cases. Review microtia grade, inner and middle ear anatomy, degree of hearing loss relative to language development.

- Discuss surgical and other therapeutic options considering timing, type of reconstruction, and placement of hearing devices to optimize access to sound.
- Individualized hearing enhancement and ear reconstruction options holistic guided by interdisciplinary approach.
- Additional support through speech therapy, audiology, social work, and/or behavioral health.

CONTACT US



THE INTERSECTIONALITY OF GLOBAL HEALTH, LEARNING ANALYTICS, AND MULTIDISCIPLINARY CLEFT CARE



Michelle Gross AdventHealth Sharing Smiles



- 1) Universal standards are difficult to achieve but we must continue to strive to make standards universally accessible and relevant.
- 2) Globalization means ever-changing iterative models that require adaptability and continued research to keep our finger on the pulse of what is happening in the field.
- 3) Outcomes can only be improved when they are measured.
- 4) Partnership is both inevitable and necessary for the sustainability of global cleft projects.

Key References

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