



DOI: 10.6026/973206300212574



www.bioinformation.net **Volume 21(8)**

Review

Received August 1, 2025; Revised August 31, 2025; Accepted August 31, 2025, Published August 31, 2025

SJIF 2025 (Scientific Journal Impact Factor for 2025) = 8.478 2022 Impact Factor (2023 Clarivate Inc. release) is 1.9

Declaration on Publication Ethics:

The author's state that they adhere with COPE guidelines on publishing ethics as described elsewhere at https://publicationethics.org/. The authors also undertake that they are not associated with any other third party (governmental or non-governmental agencies) linking with any form of unethical issues connecting to this publication. The authors also declare that they are not withholding any information that is misleading to the publisher in regard to this article.

Declaration on official E-mail:

The corresponding author declares that lifetime official e-mail from their institution is not available for all authors

License statement

This is an Open Access article which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly credited. This is distributed under the terms of the Creative Commons Attribution License

Comments from readers:

Articles published in BIOINFORMATION are open for relevant post publication comments and criticisms, which will be published immediately linking to the original article without open access charges. Comments should be concise, coherent and critical in less than 1000 words.

Disclaimer

Bioinformation provides a platform for scholarly communication of data and information to create knowledge in the Biological/Biomedical domain after adequate peer/editorial reviews and editing entertaining revisions where required. The views and opinions expressed are those of the author(s) and do not reflect the views or opinions of Bioinformation and (or) its publisher Biomedical Informatics. Biomedical Informatics remains neutral and allows authors to specify their address and affiliation details including territory where required.

Edited by Ritik Kashwani E-mail: docritikkashwani@yahoo.com Phone: +91 8804878162

Citation: Beldar et al. Bioinformation 21(8): 2574-2580 (2025)

Success of pulpotomy with MTA in primary teeth: A systematic review and meta-analysis

Tejal Laxman Beldar*, Ashwin Muralidhar Jawdekar & Laresh Naresh Mistry

Department of Pediatric Dentistry and Preventive Dentistry, Bharati Vidyapeeth (Deemed to be) University Dental College and Hospital, Navi Mumbai, India; *Corresponding author

Affiliation URL:

https://www.bvuniversity.edu.in/dchmumbai/

Author contacts:

Tejal Laxman Beldar - E-mail: tejalbeldar@gmail.com

Ashwin Muralidhar Jawdekar - E-mail: jawdekar.ashwin@gmail.com

Laresh Naresh Mistry - E-mail: laresh.mistry@bharatividyapeeth.edu

Abstract:

The challenge of effectively managing primary teeth with irreversible pulpitis in pediatric dentistry is of interest. The clinical and radiographic success of different pulpotomy materials, including Mineral Trioxide Aggregate (MTA), in comparison to conventional materials like formocresol, ferric sulfate, calcium hydroxide, and Biodentine is reported. Therefore, it is of interest to evaluate the effectiveness of materials like MTA in pulpotomies for primary teeth with irreversible pulpitis. The meta-analysis revealed that MTA pulpotomies had a clinical success rate of 97.02% and a radiographic success rate of 94.21%, outperforming ferric sulfate, Biodentine and calcium hydroxide. CEM and Calcium Silicate Cements showed comparable success rates to MTA. Thus, MTA demonstrated superior clinical and radiographic outcomes for pulpotomy in primary teeth with irreversible pulpitis, showing statistically significant differences compared to other materials.

Keywords: Pulpotomy, mineral trioxide aggregate/ MTA, pulp therapy, formocresol, irreversible pulpitis, primary teeth

Background:

The conservative management of primary teeth with irreversible pulpitis presents a significant clinical challenge in pediatric dentistry [1]. Preserving the primary dentition is crucial for maintaining function, aesthetics, and proper alignment of the permanent teeth [2]. Pulpotomy, a widely accepted procedure for treating cariously exposed primary molars, is a critical dental procedure frequently employed to manage extensively decayed primary teeth and maintain their functionality until natural exfoliation [3]. The technique involves removing the coronal part of the dental pulp, followed by placing a medicament that preserves the vitality of the remaining radicular pulp [4]. For decades, numerous materials have been used in pulpotomy, each designed to achieve optimal clinical and radiographic outcomes [5]. Mineral trioxide aggregate (MTA) has emerged as a prominent material due to its excellent biocompatibility, practical sealing ability, and regenerative properties [6]. However, the quest for the ideal pulpotomy material is ongoing, with numerous studies comparing MTA to other traditional and contemporary materials [7]. Therefore, it is of interest to report the clinical and radiographic outcomes of pulpotomy in primary teeth using various materials compared to MTA.

Review:

This systematic review and meta-analysis investigates the success of MTA (calcium silicate-based cements) compared to other materials in pulpotomies of primary teeth with irreversible pulpitis. Using the PICOS framework, the Population (P) included primary teeth with irreversible pulpitis, Intervention (I) involved calcium silicate-based cements (e.g., MTA, CEM, Biodentine), and the Control (C) group used conventional materials (formocresol, ferric sulfate, calcium hydroxide). The Outcome (O) was clinical and radiographic success rates, and the Study design (S) focused on randomized controlled trials (RCTs). A thorough literature search was performed between March 1 and May 31, 2024, using PubMed, the Cochrane Library, Google Scholar, and Semantic Scholar. Only RCTs published in English or translated into English with full texts were included. Studies with a minimum 12-month follow-up were selected. The review adhered to PRISMA 2020 guidelines, and the detailed study selection process is shown in Table 1. The sources and methodology of this systematic review

and meta-analysis were designed to ensure accuracy and reliability. A comprehensive literature search was conducted by two researchers between March 1, 2024, and May 31, 2024, targeting English-language studies with full texts available. The primary databases searched were PubMed and the Cochrane Library, extended to Google Scholar and Semantic Scholar for comprehensive coverage. No date restrictions were applied, and only randomized controlled trials (RCTs) were included. The review followed PRISMA 2020 guidelines, ensuring a standardized approach. Eligibility criteria included RCTs involving children under 10 years with irreversible pulpitis in primary teeth and a minimum 12-month follow-up for clinical and radiographic success. The PRISMA 2020 flow diagram of the study selection process is shown in Figure 1.

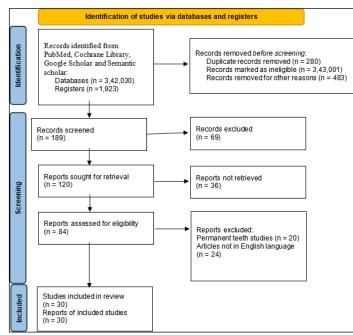


Figure 1: PRISMA 2020 flow diagram

The risk of bias (ROB) in the studies was assessed using the Cochrane Risk of Bias II tool, with independent assessments by both researchers and resolution of discrepancies through discussion or a third reviewer. The study is registered on PROSPERO (CRD42023468690, October 2023). Meta-analysis was conducted using forest plots to evaluate pooled clinical and radiographic success rates of pulpotomy materials. Heterogeneity was assessed with the I² statistic, and publication bias was examined through funnel plot analysis. Statistical analysis was performed with MedCalc software, ensuring a rigorous, unbiased comparison of MTA and other materials in primary teeth with irreversible pulpitis. This study follows PRISMA guidelines to provide reliable insights for pediatric dental practitioners. The clinical and radiographic successes of pulpotomy using MTA and other materials in primary teeth

with irreversible pulpitis for 30 studies (over 2500 participants) are reported in **Table 2**. Comparison of clinical and radiographic success of pulpotomy of calcium hydroxide, Biodentine, formocresol, ferric sulfate with MTA in primary teeth with irreversible pulpitis is reported in **Table 3**. The Risk of Bias summary was assessed using the Cochrane Risk of Bias II tool (**Figure 2**). The corresponding funnel plots for detecting publication bias are available in the additional information. Clinical and radiographic success of pulpotomy using MTA in primary teeth with irreversible pulpitis for 30 studies (over 2500 participants), the forest plot and funnel plot analyses are reported in **Figure 3 to Figure 6**.

Table 1: Data extraction

AUTHOR	SAMPLE SIZE	Comparison between	After 12 months follow-up				
total articles - 30)			CLINICAL SUCCESS	5 (%)	RADIOGRAPHIC SU	JCCESS (%)	
			Intervention	Control	Intervention	Control	
	40 s primary mandibular	MTA vs Bioceramic putty	Bioceramic putty	MTA	Bioceramic putty	MTA	
lnassar et al. [9]	molars in 40 healthy children		100%		100%		
	aged 6-8 years			95%		95	
Thorakian et al.	102 primary second molars in	CEM vs zinc oxide eugenol after	CEM	ES/ZOE	CEM	ES/ZOE	
10]	51 children aged between 4	electrosurgery (ES/ZOE)		E3/ ZOE		ES/ ZOE	
10]	and 6 years		100%	100%	97.90%	98.00%	
Haghgoo et al.	34 children aged 3-8 years	MTA, CH, or CEM	CH 96.7%	MTA 100%	CH 86.7%	MTA 100	
11]	54 children aged 5-6 years		CEM 100%	141171 100 /0	CEM 100%	141171 100	
Malekafzali <i>et al</i> .	80 teeth from Forty children	MTA and CEM	CEM	MTA	CEM	MTA	
12]	aged 4-8 years		100%	100%	100%	100%	
	44 mandibular primary molars	MTA vs Biodentine	MTA	Biodentine	MTA	Biodentir	
Çelik <i>et al</i> . [40]	in 44 children (24 boys, 20	(24 Month follow up)					
	girls) aged 5-9 years		100%	89.40%	100%	89.40%	
astor et al. [13]	90 primary Molars from	Biodentine and MTA	Biodentine	MTA	Biodentine	MTA	
astor et at. [15]	patients aged 4-9 years		100%	97.40%	94.40%	97.40%	
		calcium hydroxide mixtures and mineral	calcium hydroxide	MTA	calcium hydroxide	MTA	
	Forty-five primary mandibular	trioxide aggregate	mixtures		mixtures		
Bilva et al. [14]	molars		i)CH+saline33%	100%	i)CH+saline33%	100%	
			ii) CH+PEG		ii) CH+PEG		
			73%		73%		
	151 molars from 102 children	ProRoot MTA, OrthoMTA and RetroMTA	OrthoMTA 94.7%	ProRoot	OrthoMTA 94.7%	ProRoot	
Kang et al. [15]	of 3–10 years old	TOROUT WITH, OTHIOWITH AND REDUVITA	RetroMTA	MTA,	RetroMTA	MTA,	
	· ·		94.70%	100%,	94.70%	100%,	
Zhao et al. [16]	20 Children who had at least one pair of carious primary	iRoot BP Plus and mineral trioxide aggregate (MTA)	iRoot BP Plus	MTA	iRoot BP Plus	MTA	
znao ei ai. [10]	molars	aggregate (W171)	87%	96%	87%	96%	
		Biodentine™ and mineral trioxide					
Bani and Odabaş	primary molars from 32	aggregate (MTA)	Biodentine	MTA	Biodentine	MTA	
et al. [17]	children of 4- to 9-year-olds	aggregate (WIII)	96.80%	96.80%	93.60%	87.10%	
				ProRoot		07.1070	
		Mineral trioxide aggregate (ProRoot MTA),	ferric sulfate 100%	MTA	ferric sulfate 100%		
	32 healthy 5- to 7-year-old	ferric sulfate (15.5 % FS), formocresol (1:5					
Erdem et al. [18]	children with 128 carious	dilution of Buckley's FC) and zinc oxide	formocresol	100%	formocresol	ProRoot	
	primary molars	eugenol (ZOE)	1011110010001	10070	Tormocresor	MTA	
	primary monars	cugenor (202)	100%		100%		
			ZOE 92%		ZOE 92%		
	40 primary molars of 4- to 9-	MTA and CH	CH	MTA	CH	MTA	
.iu et al. [41]	vear-old children		64.70%	94.10%	64.70%	94.10%	
ernández et al.	total of 90 primary molars in	mineral trioxide aggregate and Biodentine	Biodentine	MTA	Biodentine	MTA	
19]	children aged 4–9 years	00.0	97%	92%	95%	97%	
•	ŭ ,	RetroMTA, OrthoMTA, and ferric sulfate	O-MTA 96.4%		O-MTA		
ilmaz et al. [20]	96 primary second molars from	, 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	R-MTA 92.8%	75% for FS	85.80%	50% for F	
[=2]	32 children aged 5 to 9 years				R-MTA 82.2%	20,2232	
		mineral trioxide aggregate and cold	11 .	3.655.4		3.6TF.4	
Rasteh et al. [21]	42 children aged 4-9 years old	ceramic	cold ceramic	MTA	cold ceramic	MTA	
. ,	5 ,		100%	100%	97%	100%	
T	90 bilateral primary molars	The Calculation					
lassanpour et al.	from 45 healthy	TheraCal and MTA	TheraCal	MTA	TheraCal	MTA	
2]	5- to 8-year-old children		99:4±3:8%	100%	97:2 ± 11:6%	98:8±7:7%	
	,	Biodentine; ProRoot White Mineral		ProRoot		ProRoot	
		Trioxide Aggregate (WMTA);	Biodentine	WMTA	Biodentine	WMTA	
lajashekharan et	Fifty-eight patients (82 teeth) in	Tempophore	95.24%	100%	94.40%	90.90%	
l. [23]	patients above 3 years of age	rempophore	Tempophore	10070	Tempophore	20.2078	
			95.65%		82.40%		
Randa et al. 2020	72- second primary molars in 4	Nanohydroxyapatite (NHA), Mineral					
24]	to 8 years old children	Trioxide Aggregate (MTA), Formocresol	NanoHA	MTA	NanoHA	MTA	
4 4]	to o years old children	moriue Aggregate (MTA), Formocresol					

Filt			(FC)				
Stake 1,25 52 children aged 3-6 years MTA and Biodentine 88.46%					87.50%		79.20%
State 1.25 52 children aged 3-6 years PRATTA PR						Formocresol 75 %	
Pocket P	E 1 1: / 1 [05]	F0 1:11 10 6	NETA ID: 1 c		00.460/	00.460/	00.468/
Cuven et al. [26] 26 healthy 5- to 7-year-old children	Eshghi et al. [25]	52 children aged 3–6 years		88.46%	88.46%	88.46%	88.46%
Minary M	Guven <i>et al.</i> [26]		[MTA-P], and Biodentine [BD]) and ferric		PR-MTA		PR-MTA
			· ·		93.10%		93.10%
Since Sinc							
Sakai et al. [27] Sakai et al. [27] molars of children aged 5-9 years old Moorollahian et al. [28] Say primary mandibular solution aged 5-9 years old Moorollahian et al. [28] Say primary malar in say solution ager gate and formocresol of age mineral trioxide aggregate and formocresol of aggregate of aggr							
Sakai et al. [27] molars of children aged 5-9 years old molars of children aged 5-9 years old Mra Mra Mra Mra Mra Mra Mra Mra							
Sakai et al. [27] molars of children aged 5-9 years old 100% 1					89.65%		89.65%
Molars of Children aged 5-9 years old 100% 100% 100% 100% 78.60%	Sakai et al. [27]	* *			MTA		MTA
Moretti et al. [29] 64 primary molars in 35 children between 5 and 9 years old Agamy et al. [32] 72 primary molars in 24 children 25 children (50 human primary molar teth) aged between 5 and 9 years Juneja et al. [34] 51 primary molars in 64 children aged 5-9 years old Sirohi et al. [35] 51 primary molar in children Sirohi et al. [35] 64 primary molar in children Formocresol mineral trioxide aggregate and formocresol aggregate		years old		100%	100%	100%	78.60%
Holan et al. [29] 64 primary molars in 35 children4 to 12 years 45 primary mandibular molars in 23 children between 5 and 9 years old 90 primary molars in 37 children aged 4-7 years 46 mineral trioxide aggregate and formocresol years old 90 primary molars in 37 children aged 4-7 years 46 mineral trioxide aggregate and formocresol 90% 80% 90% 90% 80% 90% 90% 80% 90% 90% 90% 90% 90% 90% 90% 90% 90% 9	- 100-0-0	molars of 46 children 5-7 years	mineral trioxide aggregate and formocresol	Formocresol	MTA	Formocresol	MTA
Moretti et al. [30]	Holan et al. [29]		mineral trioxide aggregate and formocresol	aggregate		aggregate	
Moretti et al. [30] in 23 children between 5 and 9 years old		· · · · · · · · · · · · · · · · · · ·		97%	83%	58%	52%
Olatosi et al. [31] 50 primary molars in 37 children aged 4-7 years mineral trioxide aggregate and formocresol FC 81% MTA 100% FC 81% MTA 96% Mineral Trioxide Aggregate and Formocresol White MTA Formocresol White MTA Agamy et al. [32] 72 primary molars in 24 children 90% 80% 90% 80% Carti et al. [33] 25 children (50 human primary molar teeth) aged between 5 and 9 years Juneja et al. [34] 51 primary molars of children aged 5-9 years old Sirohi et al. [35] Fifty primary molar in children aged 4 to 8 years Sirohi et al. [35] Fifty primary molar in children aged 4 to 8 years Mineral Trioxide Aggregate and Formocresol Biodentine MTA Mineral Trioxide Aggregate and Biodentine Biodentine MTA Biodentine MTA Biodentine MTA MTA Formocresol 73.3% Fifty primary molar in children aged 4 to 8 years Silicate Cement Silicate Cement MTA Biodentine FS Biodentine FS Biodentine FS	Moretti et al. [30]	in 23 children between 5 and 9		CH 36 %		CH 36 %	
Agamy et al. [32] 72 primary molars in 24 children Agamy et al. [32] 72 primary molars in 24 children Carti et al. [33] 25 children (50 human primary molar teeth) aged between 5 and 9 years Juneja et al. [34] 51 primary molars of children aged 5-9 years old Sirohi et al. [35] Fifty primary molar in children aged 4 to 8 years Mineral Trioxide Aggregate and Formocresol PC 81% MTA Formocresol White MTA Formocresol MTA Gray MTA Mineral Trioxide Aggregate and Biodentine MTA Biodentine MTA Biodentine MTA Biodentine MTA Biodentine MTA Biodentine MTA Biodentine MTA Formocresol 73.3% 100% 86.60% 100% Formocresol 73.3% Ferric Sulfate (FS) and Bioactive Tricalcium Silicate Cement Silicate Cement Fifty primary molar in children aged 4 to 8 years		J			MTA 100%		MTA 100%
Agamy et al. [32] 72 primary molars in 24 children	Olatosi et al. [31]		66 6	FC 81%	MTA 100%	FC 81%	MTA 96%
Agamy et al. [32] children Carti et al. [33] children Carti et al. [34] 25 children (50 human primary molar teeth) aged between 5 and 9 years Juneja et al. [34] 51 primary molars of children aged 5-9 years old Sirohi et al. [35] Fifty primary molar in children aged 4 to 8 years Sirohi et al. [35] Fifty primary molar in children aged 4 to 8 years Sirohi et al. [35] children (50 human primary molar in children aged 4 to 8 years said to 100%				Formocresol	White MTA	Formocresol	White MTA
Carti et al. [34] Juneja et al. [35] Sirohi et al. [35] Sirohi et al. [36] Zo children (50 human primary molar teeth) aged between 5 and 9 years Sirohi et al. [36] Sirohi et al. [37] Zo children (50 human primary molar in children aged between 5 and 9 years) Mineral Trioxide Aggregate and Biodentine Biodentine 96% 96% 96% 96% MTA Biodentine MTA Biodentine MTA Biodentine MTA Biodentine MTA Biodentine MTA Formocresol Fifty primary molars of children aged 4 to 8 years Silicate Cement Fifty primary molar in children aged 4 to 8 years	Agamy et al. [32]			90%	007-	90%	007-
Carti et al. [33] 25 children (50 human primary molar teeth) aged between 5 and 9 years Juneja et al. [34] 51 primary molars of children aged 5-9 years old Sirohi et al. [35] Fifty primary molar in children aged 4 to 8 years Sirohi et al. [35] 25 children (50 human primary molar in children aged 5-9 years old Mineral Trioxide Aggregate and Biodentine 96% 96% 60% 80% Biodentine, mineral trioxide aggregate and Biodentine 100% MTA Biodentine MTA Formocresol 73.3% 100% 86.60% 100% Formocresol 73.3 Ferric Sulfate (FS) and Bioactive Tricalcium Silicate Cement FS Biodentine FS Biodentine FS Biodentine		Cilidicii					
Carti et al. [33] molar teeth) aged between 5 and 9 years Biodentine Biodentine 96% 96% 60% 80% MTA Biodentine Formocresol Formocresol Formocresol Formocresol Formocresol Ferric Sulfate (FS) and Bioactive Tricalcium Silicate Cement FS Biodentine FS Biodentine FF Biodentine MTA Biodentine MTA Biodentine FF Formocresol Formocresol FF Formocresol FF Biodentine FF Biodentine FF Biodentine MTA Biodentine MTA Biodentine MTA Biodentine FF FF Formocresol FF FF FF FF FF FF FF Biodentine FF Biodentine MTA Biodentine MTA Biodentine MTA Biodentine MTA Biodentine MTA Biodentine FF FF FF FF FF FF FF FF FF					100%		100%
Juneja et al. [34] 51 primary molars of children aged 5-9 years old 51 primary molar in children Sirohi et al. [35] Fifty primary molar in children aged 4 to 8 years grant for the state of the state o	Carti et al.[33]	molar teeth) aged between 5					
Juneja et al. [34] 51 primary molars of children aged 5-9 years old formocresol formocreso		and 9 years		96%	96%	60%	80%
Sirohi et al. [34] aged 5-9 years old formocresol 73.3% 100% 86.60% 100% Formocresol 73.3 Ferric Sulfate (FS) and Bioactive Tricalcium Silicate Cement FS Biodentine FS Biodentine	Juneja <i>et al</i> . [34]		, 00 0	Biodentine 100%	MTA		
Formocresol 73.3 Sirohi et al. [35] Fifty primary molar in children aged 4 to 8 years aged 4 to 8 years aged 4 to 8 years aged 5 to 8 years aged 6 to 8 yea				formocresol 73.3%	100%		100%
Sirohi et al. [35] Fifty primary molar in children and 4 to 8 years and 4 years an		J ,					
96% 100% 84% 92%	Sirohi et al. [35]		` '	FS	Biodentine	FS	Biodentine
		ageu + 10 o years		96%	100%	84%	92%

 Table 2: Summary statistics: percentage clinical and radiographic success and 95% CI - all studies

Intervention	No of studies	Sample size	REM/ FEM	% Success	Confidence Interval		
CLINICAL							
MTA	33	2219	REM	97.02	95.46, 98.26		
Formocresol	8	542	REM	91.90	82.66, 97.84		
Ferric sulfate	4	303	REM	90.86	71.67, 99.72		
Biodentine	8	491	REM	95.82	92.16, 98.36		
Ca hydroxide	5	206	REM	76.37	50.95, 94.38		
Ca enriched mixture	3	216	FEM	99.67	97.71, 99.99		
Ca silicate	2	126	FEM	99.08	95.494, 99.96		
RADIOGRAPHIC							
MTA	33	2219	REM	94.21	91.20, 96.63		
Formocresol	8	542	REM	88.17	73.25, 97.51		
Ferric sulfate	4	303	REM	79.23	42.23, 99.31		
Biodentine	8	491	REM	88.47	81.40, 94.02		
Ca hydroxide	5	206	REM	73.35	50.09, 91.24		
Ca enriched mixture	3	216	FEM	99.67	97.71, 99.99		
Ca silicate	2	126	FEM	97.67	93.32, 99.52		

The values are percent success.

REM- Random effect model/ FEM- fixed effect model; CI- Confidence Interval

Table 3: Summary statistics: Odds ratios – various pulpotomy agents vs MTA (percentage clinical and radiographic success and 95% CI - all studies)

Intervention Vs. MTA	No of studies	Sample size	REM/ FEM	Odds ratio	Confidence Interval		
CLINICAL							
Ca hydroxide	5	206	FEM	0.046	0.02, 0.13		
Biodentine	8	491	REM	2.858	1.06, 7.67		
Formocresol	8	542	REM	0.304	0.06, 1.47		
Ferric sulfate	5	378	FEM	0.191	0.10, 0.36		
RADIOGRAPHIC							

Ca hydroxide	5	206	FEM	0.049	0.02, 0.13	
Biodentine	8	491	FEM	0.976	0.63, 1.51	
Formocresol	8	542	REM	0.779	0.22, 2.77	
Ferric sulfate	5	378	FEM	0.201	0.13, 0.32	

The values are Odds Ratios

REM- Random effect model/ FEM- fixed effect model; CI- Confidence Interval

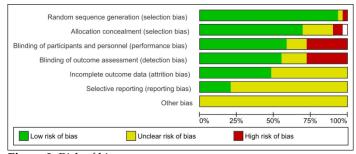


Figure 2: Risk of bias assessment summary.

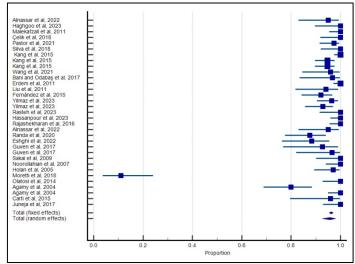


Figure 3: Forest plot - clinical success MTA

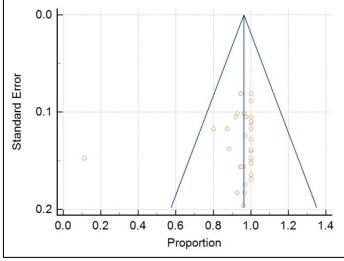


Figure 4: Funnel plot - clinical success MTA

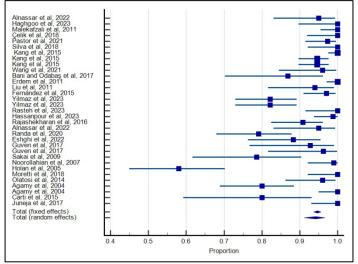


Figure 5: Forest plot - radiographic success MTA

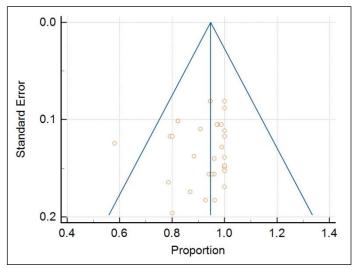


Figure 6: Funnel plot - radiographic success MTA

Discussion:

Irreversible pulpitis is when the dental pulp becomes inflamed and damaged to the point where it cannot heal independently. This condition is usually a result of deep decay, trauma, or repeated dental procedures that irritate the pulp [35]. Key characteristics of irreversible pulpitis include severe, intense, lingering pain, especially in response to hot or cold stimuli. The pain may also be spontaneous, without any external trigger. Severe inflammation leads to irreversible damage. As the condition progresses, the pulp may become necrotic, potentially leading to infection and an abscess at the root tip. Irreversible

pulpitis is believed to require more invasive treatments than pulpotomy because the pulp cannot recover independently [36]. The standard therapy for irreversible pulpitis is to remove the inflamed and damaged pulp to prevent further complications. In primary teeth, this is usually done through pulpotomy (removal of the pulp from the coronal portion) or pulpectomy (removal of the entire pulp), followed by filling the space with a suitable material [11]. If left untreated, irreversible pulpitis can lead to more severe dental issues, including abscess formation, bone loss around the tooth and potentially needing tooth extraction. The emergence of calcium silicate-based materials, particularly MTA and newer materials like Biodentine, has significantly transformed the practice of pulpotomy, especially in pediatric dentistry [37]. These materials have introduced a paradigm shift in the management of dental pulp therapy, primarily due to their superior biological properties (includes the inductive ability leading to dentin formation), clinical efficacy and longterm success rates [13]. These materials have revolutionized pulpotomy procedures by offering more biocompatible, effective and durable solutions for managing irreversible pulpitis in primary teeth. These materials have set a new standard in dental pulp therapy, leading to better patient outcomes and transforming the approach to pediatric dental care. Pulpotomy was used initially as a devitalisation procedure for inflamed pulp just for the pain to subside which is an obsolete concept now. We now prefer preservation/ regeneration approach to the mummification/ devitalization practice Devitalization, preservation and regeneration reflect the evolution of the procedure from a focus on simply managing symptoms to promoting long-term dental health and natural healing. These approaches offer more sustainable outcomes, especially in pediatric patients, by maintaining the function and health of the affected tooth until it can naturally exfoliate or continue to develop (as in the case of permanent teeth) . Although several randomised controlled trials have been available reporting success of these materials, some of these with recent evidence are available with sufficient follow-up. Pulpotomy treatment failures resulting in inflammation could be noticed over a period of 1 year and beyond; hence, our study assessed the success of pulpotomy with an inclusion criterion of minimum 1-year follow-up while assessing both individual and comparative performance of various materials. We found that calcium enriched mixture, calcium silicate, MTA and Biodentine cements to have the best clinical success followed by formocresol and ferric sulfate and was lowest for calcium hydroxide. Radiographically, a similar trend was observed. In general, both the clinical and radiographic success of these materials is comparable to that of reported studies for pulp therapies of primary teeth without irreversible pulpitis. Junior et al. [38] reported that the success rate of MTA was higher than that of formocresol, with a statistically significant difference. Formocresol pulpotomy success was not statistically different from ferric sulphate or electrosurgery. Tewari et al. [37] reported that pulpotomy medicaments, except calcium hydroxide, showed success rates of more than 80%, whereas most comparisons revealed no differences. MTA, however, was found

to be better than calcium hydroxide and formocresol. In comparison to MTA, calcium hydroxide, formocresol and ferric sulfate pulpotomies showed lower clinical and radiographic success. Biodentine exhibited superior clinical success however; radiographically the success was not significantly different. Junior et al. [38] reported that overall clinical and radiographic success rates Biodentine vs. MTA did not differ statistically in the 6-month follow-up. Coll et al. [39] reported that two calcium silicate cement pulpotomies success using mineral trioxide aggregate (MTA) and Biodentine were 94 percent and 90 percent, respectively. The current SRMA has a few limitations such as inclusion of fewer studies of direct comparison, Unavailability of trials with longer follow-up i.e. more than 2-3 years, variations in the identification of different calcium silicate materials. Despite such limitations, this study confirms the possibility of success of pulpotomy in primary teeth with irreversible pulpitis.

Conclusion:

Calcium silicate-based materials are superior to formocresol and ferric sulfate. Amongst calcium silicate-based materials, CEM and calcium silicate cement shows best outcomes followed by MTA and Biodentine. Hence, we conclude our findings; pulpotomy has potential for success over 90% in primary teeth with irreversible pulpitis using calcium silicate-based materials.

References:

- [1] Lin GSS *et al. Children (Basel)*. 2024 **11**:574. [DOI: 10.3390/children11050574]
- [2] Setty JV *et al. Int J Clin Pediatr Dent.* 2016 **9**:56. [DOI: 10.5005/jp-journals-10005-1334]
- [3] Farheen Chunara et al. African Journal of Biomedical Research, 2024 27:2254. [DOI: 10.53555/AJBR.v27i3S.2597]
- [4] Islam R et al. Jpn Dent Sci Rev. 2023 **59**:48. [DOI: 10.1016/j.jdsr.2023.02.002]
- [5] Pushpalatha C *et al. Front Bioeng Biotechnol.* 2022 **10**:941826. [PMID: 36017346]
- [6] Bossù M et al. J Clin Med. 2020 9:838. [PMID: 32204501]
- [7] Parisay I et al. Iran Endod J. 2015 **10**:6. [PMID: 25598803]
- [8] Khorakian F et al. European Archives of Paediatric Dentistry. 2014 **15**:223. [PMID: 24435546]
- [9] Alnassar I *et al. Clin Exp Dent Res.* 2023 **9**:276. [DOI: 10.1002/cre2.700]
- [10] Khorakian F *et al. Eur Arch Paediatr Dent.* 2014 **15**:223. [DOI: 10.1007/s40368-013-0102-z]
- [11] Haghgoo R et al. Journal of Evidence-Based Dental Practice. 2023 <u>23:</u>101920. [DOI: 10.1016/j.jebdp.2023.101920]
- [12] Malekafzali B et al. Eur J Paediatr Dent. 2011 12:189. [PMID: 22077689].
- [13] Vilella-Pastor S et al. Eur Arch Paediatr Dent. 2021 22:685. [DOI: 10.1007/s40368-021-00616-3]
- [14] Silva LLCE *et al. J Appl Oral Sci.* 2019 **27**:e20180030. [DOI: 10.1590/1678-7757-2018-0030]
- [15] Kang CM *et al. Oral Dis.* 2015 **21**:785. [DOI: 10.1111/odi.12348]

- [**16**] Zhao Y *et al. PeerJ.* 2024 **12**:e18453. [DOI: 10.7717/peerj.18453]
- [17] Bani M et al. Pediatr Dent. 2017 39:284. [PMID: 29122067]
- [18] Erdem AP et al. Pediatr Dent. 2011 33:165. [PMID: 21703067]
- [19] Cuadros-Fernández C *et al. Clin Oral Investig.* 2016 **20**:1639. [DOI: 10.1007/s00784-015-1656-4]
- [20] Yilmaz S *et al. Eur Oral Res.* 2023 **57**:144. [DOI: 10.26650/eor.2023950004]
- [21] Rasteh B et al. J Family Med Prim Care. 2023 12:3068. [DOI: 10.4103/jfmpc.jfmpc_412_23]
- [22] Hassanpour S *et al. Biomed Res Int.* 2023 **2023**:8735145. [PMID: 37124935].
- [23] Rajasekharan S *et al. Int Endod J.* 2017 **50**:215. [PMID: 26863893]
- [24] Abd Al Gawad RY *et al. Saudi Dent J.* 2021 33:560. [DOI: 10.1016/j.sdentj.2020.08.007]
- [25] Eshghi A *et al.* Int J Dent. 2022 **2022**:6963944. DOI: 10.1155/2022/6963944. [PMID: 35866144]
- [26] Guven Y et al. Biomed Res Int. 2017 2017:4059703. [DOI 10.1155/2017/4059703]
- [27] Sakai VT *et al. Br Dent J.* 2009 **207**:E5. [DOI: 10.1038/sj.bdj.2009.665]
- [28] Noorollahian H *et al. Br Dent J.* 2008 **204**:E20. [PMID: 18425074]

- [29] Holan G et al. Pediatr Dent. 2005 27:129. [PMID: 15926290]
- [30] Moretti AB et al. Int Endod J. 2008 41:547. [PMID: 18479381]
- [31] Olatosi OO *et al. Niger J Clin Pract.* 2015 **18**:292. [DOI: 10.4103/1119-3077.151071]
- [32] Agamy HA et al. Pediatr Dent. 2004 26:302. [PMID: 15344622].
- [33] Carti O *et al. Niger J Clin Pract*. 2017 **20**:1604. [DOI: 10.4103/1119-3077.196074]
- [34] Juneja P *et al. Eur Arch Paediatr Dent.* 2017 **18**:271. [DOI: 10.1007/s40368-017-0299-3]
- [35] Sirohi K *et al. Int J Clin Pediatr Dent.* 2017 **10**:147. [DOI: 10.5005/jp-journals-10005-1425]
- [36] Haghgoo R et al. Journal of Evidence-Based Dental Practice. 2023 23:101920. [DOI: 10.1016/j.jebdp.2023.101920]
- [37] Tewari N et al. Int J Paediatr Dent. 2022 32:828. [PMID: 35271753]
- [38] Junior ES et al. Clin Oral Investig. 2019 23:1967. [DOI: 10.1007/s00784-018-2616-6]
- [39] Coll JA et al. Pediatr Dent. 2023 45:474. [PMID: 38129755]
- [40] Çelik BN *et al. Clin Oral Investig.* 2019 **23**:661. [PMID: 29744721]
- [41] Liu H et al. Clinical oral investigations. 2016 **20**:1639. [DOI: 10.1007/s00784-015-1656-4]