

Optimizing Paediatric Clinical Trials:

Leveraging Milo AI for Efficient Recruitment and Data

Management

MILO Healthcare

Clinical trials are inefficient, costly, and lack real-time insight





30%

Clinical trial costs are high due to manual processes, slow patient recruitment and error-prone data entry.



Electronic clinical data capture powered with GenAl



Capabilities of the platform





What does Milo do?



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Technological evolution of MILO

			New capabilities required	New capabilities required	
70	Wall o	Wall 1	Wall 2	Wall 3	
70 60	Traditional Clinical trial	New capabilities required	1-Milo match: find patient immediately		
00	excel	SaaS Clinical Platform for	2-EHR to EDC solved 3- No connector to EHR is	⊳ milo	
50	Your smart way to compliance	1- eCRF	required		
40		2- ePRO 3- eConsent		MILO Medical Interface	
30		Milo	Computer use	Liaison	
0.		Studio		Officer	
20					
10					
\circ		Era 1 Scalling laws	Era 2 Taking Action	Era 3 Autonomy	
0	2020	2024	2025	2026	

Key differentiation current AI models









Features	Milo	Chatgbt 4o	Claude 3	Llama 3.3
MILO scribe	95%	66%	68%	54%
Milo match	85%	68%	71%	60%
MILO vision	75%	21%	25%	15%



Ongoing European clinical trials that demonstrate ROI in real life

Name of Hospitals	Country	Name of investigator	Trial Name	Comparative	N° of patients
CHU Montpellier	France	Prof. Luc Teot	Evidence I	Yes	100
Clinique Clémentville	France	Prof. Cristian Villanueva	Evidence II	Yes	400
CHU ROCHELLE	France	Dr Trijolet	Pilot	No	30
CHU Bordeaux	France	Dr Stéphanie BUI	Pilot	No	30
Alder Hey	UK	Dr Laura Thursfield	EVIDENCE III	Yes	100
Birmingham Children's Hospital	UK	Dr Priti Kenia	EVIDENCE III	Yes	100
Great Ormond Street Hospital	UK		EVIDENCE III	Yes	100
NA	Germany	Prof Axel Schulz	Pilot	No	100



Current product and current pricing

Human

Manually enters data from EMR and patient-doctor interaction into eCRF

Onsite monitoring for data error resolutions

Manually identify ptients using EMR

5000€ / patient

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Pushes the data into the EDC automatically

Flags inconsistencies

Milo is the support for data source and monitoring can be done remotely

Automatically pre-screen patients during consultations.

500€ / patient



Per 1000 patients

Introduction



Study Type: Prospective, open-label, randomized, multicentric, parallel-arm longitudinal study over 4 weeks.

Sponsor: Unither Pharma



Objectives:

•Primary Evaluate the performance and safety of hypertonic inhalation (3%, 6%, 7% sodium chloride) in infants, children, and adults with cystic fibrosis (CF) or non-CF bronchiectasis (NCFB).
•Secondary : Evaluate the time saved using EDC Milo thanks to automatic health data collection



Expected results





Primary: Change in lung clearance index (LCI) from baseline to week 4, measured via N2 Multiple Breath Washout.



Secondary: Time saved by the medical team with MILO VS without Milo:

- Patient recruitment
- Medical data collection
- eCRF completion



Methodology



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