UNITED STATES SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): May 11, 2022

Hyperfine, Inc

(Exact name of Registrant as Specified in Its Charter)

Delaware (State or Other Jurisdiction of Incorporation) 001-39949 (Commission File Number)

Registrant's Telephone Number, Including Area Code: 866 796-6767

98-1569027 (IRS Employer Identification No.)

351 New Whitfield Street Guilford, Connecticut (Address of Principal Executive Offices)

06437 (Zip Code)

	N/A (Former Name or Former Address, if Changed Since Last Report)						
Che	Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:						
	Written communications pursuant to Rule 425 under the Securit	ies Act (17 CFR 230.425)					
	□ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)						
	Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))						
	Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))						
	Securities registered pursuant to Section 12(b) of the Act:						
	Trading Symbol(s) Class A common stock, \$0.0001 par value per share Trading Symbol(s) Name of each exchange on which registered HYPR The NASDAQ Stock Market LLC						

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§ 230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§ 240.12b-2 of this chapter).

Emerging growth company \boxtimes

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. \Box

Item 2.02 Results of Operations and Financial Condition.

On May 11, 2022, Hyperfine, Inc. (the "Company") issued a press release announcing its results for the first quarter and year ended March 31, 2022 and providing a business update. A copy of the press release is furnished as Exhibit 99.1 hereto.

The information in this Current Report on Form 8-K (including Exhibit 99.1) shall not be deemed to be "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such a filing.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit	Description
Number	
99.1	Press Release dated May 11, 2022
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

HYPERFINE, INC.

Date: May 11, 2022

/s/ Dave Scott
Dave Scott
President and Chief Executive Officer



Hyperfine Reports First Quarter 2022 Financial Results

GUILFORD, Connecticut, May 11, 2022 (GLOBE NEWSWIRE) – Hyperfine, Inc. (Nasdaq: HYPR), the creator of Swoop®, the world's first FDA-cleared portable MRI system TM , today reported financial results for the quarter ended March 31, 2022. Management will host a corresponding conference call to discuss the financial results and provide a business update today at 1:30 p.m. Pacific Time / 4:30 p.m. Eastern Time.

"We are pleased with our progress in the first quarter. We continued to build awareness of Swoop's tremendous value proposition and installed 11 commercial systems," said Dave Scott, President and CEO of Hyperfine. "We now have a commercial installed base* of 38 systems and a total, global installed base of 85 systems - and we look forward to continuing to execute our growth plan through 2022 and beyond."

First Quarter 2022 Financial Results

- Revenues for the first quarter of 2022 were \$1.509 million, compared to \$0.331 million in the first quarter of 2021.
- Gross margin for the first quarter of 2022 was \$0.084 million, compared to \$(0.277) million in the first quarter of 2021.
- · Research and development expenses for the first quarter of 2022 were \$8.334 million, compared to \$4.474 million in the first quarter of 2021.
- Sales, general, and administrative expenses for the first quarter of 2022 were \$15.521 million, compared to \$3.054 million in the first quarter of 2021.
- Net loss for the first quarter was \$23.775 million, equating to a net loss of \$0.34 per share, as compared to a net loss of \$7.794 million, or a net loss of \$4.86 per share, for the same period of the prior year.

Swoop Total Installed Base

		TOTAL INSTALLED UNITS					
	2020	2021				2022	
		Q1	Q2	Q3	Q4	Q1	Total
Commercial systems installations*	4	5	7	4	7	11	38
Grant fulfillment installations	_	2	2	4	10	2	20
	4	7	9	8	17	13	58
Research units	15	2	2	3	3	2	27
Total Installed Units	19	9	11	11	20	15	85

^{*} The Swoop total installed base consists of three components: Commercial system installations (which make up total revenue), grant fulfillment installations, and research unit installations. The Swoop total installed base (or total installed units) is the number of Swoop devices deployed to hospitals, other healthcare providers, and research institutions. We view the total installed base as a key metric of the growth of our business and is measured from period over period.

2022 Financial Guidance

- Management expects revenue for the full year 2022 to be \$10 to \$12 million.
- Management expects 50 to 60 commercial units installed in 2022.



Hyperfine will host a conference call at 1:30 p.m. PST / 4:30 p.m. ET today to discuss its first quarter 2022 financial results and provide a business update. The call may be accessed through an operator by calling (888) 708-1168 for domestic callers or (630) 652-5889 for international callers, using conference ID 4080148. A live and archived audio webcast will be available through the Investors page of Hyperfine's corporate website at https://investors.hyperfine.io/.

About Hyperfine

Hyperfine, Inc. is the groundbreaking medical device company that created Swoop®, the world's first FDA-cleared portable MRI system™. Hyperfine designed Swoop to enable rapid diagnoses and treatment for every patient regardless of income, resources, or location, pushing the boundaries of conventional imaging technology and expanding patient access to life-saving care. The Swoop Portable MR Imaging System™ produces high-quality images at a lower magnetic field strength, allowing clinicians to quickly scan, diagnose, and treat patients in various clinical settings. Swoop can be wheeled directly to the patient's bedside, plugged into a standard electrical wall outlet, and controlled by an iPad®. Designed as a complementary system to conventional MRIs at a fraction of the cost, Swoop captures images in minutes, providing critical decision-making capabilities in emergency departments, operating rooms outside the sterile field, and intensive care units, among others.

Investor Contact Marissa Bych Gilmartin Group LLC investors@hyperfine.io



Forward-Looking Statements

This press release includes "forward-looking statements" within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995. Hyperfine's actual results may differ from its expectations, estimates and projections and consequently, you should not rely on these forward-looking statements as predictions of future events. Words such as "expect," "estimate," "project," "budget," "forecast," "anticipate," "intend," "plan," "may," "will," "could," "should," "believes," "predicts," "potentia "continue," and similar expressions (or the negative versions of such words or expressions) are intended to identify such forward-looking statements. These forward-looking statements include, without limitation, expectations about Hyperfine's financial and operating results, the benefits of Hyperfine's products and services, and Hyperfine's future performance and its ability to implement its strategy. These forward-looking statements involve significant risks and uncertainties that could cause the actual results to differ materially from the expected results. Most of these factors are outside of Hyperfine's control and are difficult to predict. Factors that may cause such differences include, but are not limited to: the success, cost and timing of Hyperfine product development and commercialization activities, including the degree that Swoop is accepted and used by healthcare professionals; the impact of COVID-19 on Hyperfine's business; the inability to maintain the listing of Hyperfine's Class A common stock on the Nasdaq following the recently completed business combination; the inability to recognize the anticipated benefits of the business combination, which may be affected by, among other things, competition and Hyperfine's ability to grow and manage growth profitably and retain its key employees; changes in applicable laws or regulations; the inability of Hyperfine to raise financing in the future; the inability of Hyperfine to obtain and maintain regulatory clearance or approval for its products, and any related restrictions and limitations of any cleared or approved product; the inability of Hyperfine to identify, in-license or acquire additional technology; the inability of Hyperfine to maintain its existing or future license, manufacturing, supply and distribution agreements and to obtain adequate supply of its products; the inability of Hyperfine to compete with other companies currently marketing or engaged in the development of products and services that Hyperfine is currently marketing or developing; the size and growth potential of the markets for Hyperfine's products and services, and its ability to serve those markets, either alone or in partnership with others; the pricing of Hyperfine's products and services and reimbursement for medical procedures conducted using Hyperfine's products and services; Hyperfine's estimates regarding expenses, future revenue, capital requirements and needs for additional financing; Hyperfine's financial performance; and other risks and uncertainties indicated from time to time in Hyperfine's filings with the Securities and Exchange Commission, including those under "Risk Factors" therein. Hyperfine cautions readers that the foregoing list of factors is not exclusive and that readers should not place undue reliance upon any forward-looking statements, which speak only as of the date made. Hyperfine does not undertake or accept any obligation or undertaking to release publicly any updates or revisions to any forward-looking statements to reflect any change in its expectations or any change in events, conditions or circumstances on which any such statement is based.



HYPERFINE, INC. AND SUBSIDIARIES CONDENSED COMBINED AND CONSOLIDATED BALANCE SHEETS (in thousands, except share and per share amounts) (Unaudited)

	March 31, 2022		December 31, 2021	
ASSETS				
CURRENT ASSETS:				
Cash and cash equivalents	\$	161,580	\$	188,498
Restricted cash		1,983		2,662
Accounts receivable, net		1,944		553
Unbilled receivables		478		91
Inventory		4,538		4,310
Prepaid expenses and other current assets		3,205		1,357
Due from related parties		1		14
Total current assets	\$	173,729	\$	197,485
Property and equipment, net		3,877		3,753
Other long term assets		1,222		1,235
Total assets	\$	178,828	\$	202,473
LIABILITIES AND STOCKHOLDERS' EQUITY				
CURRENT LIABILITIES:				
Accounts payable	\$	1,745	\$	2,248
Deferred grant funding		1,983		2,662
Deferred revenue		973		730
Due to related parties		97		1,981
Accrued expenses and other current liabilities		6,616		8,115
Total current liabilities	\$	11,414	\$	15,736
Long term deferred revenue		851		510
Total liabilities	\$	12,265	\$	16,246
COMMITMENTS AND CONTINGENCIES				
STOCKHOLDERS' EQUITY				
Class A Common stock, \$.0001 par value; 600,000,000 shares authorized; 55,277,061 shares issued and outstanding at March 31, 2022 and December 31, 2021, respectively		5		5
Class B Common stock, \$.0001 par value; 27,000,000 shares authorized; 15,055,288 shares issued and outstanding at March 31, 2022 and December 31, 2021, respectively		2		2
Additional paid-in capital		326,651		322,540
Accumulated deficit		(160,095)		(136,320)
Total stockholders' equity	\$	166,563	\$	186,227
TOTAL LIABILITIES AND STOCKHOLDERS' EQUITY	\$	178,828	\$	202,473



HYPERFINE, INC. AND SUBSIDIARIES CONDENSED COMBINED AND CONSOLIDATED STATEMENT OF OPERATIONS AND COMPREHENSIVE LOSS (in thousands, except share and per share amounts) (Unaudited)

		Three months ended March 31,		
	-	2022		2021
Sales				
Device	\$	1,192	\$	169
Service		317		162
Total sales	\$	1,509	\$	331
Cost of sales				
Device	\$	1,037	\$	548
Service		388		60
Total cost of sales	\$	1,425	\$	608
Gross margin		84		(277)
Operating Expenses:				
Research and development	\$	8,334	\$	4,474
General and administrative		11,360		1,858
Sales and marketing		4,161		1,196
Total operating expenses		23,855		7,528
Loss from operations	\$	(23,771)	\$	(7,805)
Interest income	\$	1	\$	5
Other income (expense), net		(5)		6
Loss before provision for income taxes	\$	(23,775)	\$	(7,794)
Provision for income taxes				
Net loss and comprehensive loss	\$	(23,775)	\$	(7,794)
Net loss per common share attributable to common stockholders, basic and diluted	\$	(0.34)	\$	(4.86)
Weighted-average shares used to compute net loss per share attributable to common stockholders, basic and diluted		70,332,349		1,602,732



HYPERFINE, INC. AND SUBSIDIARIES CONDENSED COMBINED AND CONSOLIDATED STATEMENT OF CASH FLOWS (in thousands) (Unaudited)

	_	Three months ended March 31,		
		2022		2021
Cash flows from operating activities:				
Net loss	\$	(23,775)	\$	(7,794)
Adjustments to reconcile net loss to net cash used in operating activities:				
Depreciation and amortization		253		96
Stock-based compensation expense		4,111		267
Write-down of inventory		_		75
Payments received on net investment in lease		2		3
Changes in assets and liabilities:				
Accounts receivable		(1,391)		(467)
Unbilled receivables		(387)		(8)
Inventory		(228)		(672)
Prepaid expenses and other current assets		(1,848)		(326)
Due from related parties		13		882
Prepaid inventory		_		(16)
Other long term assets		11		(7)
Accounts payable		(565)		169
Deferred grant funding		(679)		_
Deferred revenue		584		600
Due to related parties		(1,884)		19
Accrued expenses and other current liabilities		(1,506)		(307)
Net cash used in operating activities	<u>\$</u>	(27,289)	\$	(7,486)
Cash flows from investing activities:				
Purchases of fixed assets		(308)		(170)
Net cash used in investing activities	\$	(308)	\$	(170)
Cash flows from financing activities:				
Proceeds from exercise of stock options		_		49
Proceeds from issuance of Series D convertible preferred stock		_		30,468
Stock issuance costs related to Series D convertible preferred stock		_		(7)
Investment from 4Bionics, LLC		_		700
Net cash provided by financing activities	\$		\$	31,210
Net (decrease) increase in cash and cash equivalents and restricted cash		(27,597)	<u> </u>	23,554
Cash, cash equivalents and restricted cash, beginning of period		191,160		64,286
Cash, cash equivalents and restricted cash, end of period	\$	163,563	\$	87,840
Reconciliation of cash, cash equivalents, and restricted cash reported in the statement of financial position		<u> </u>		
Cash and cash equivalents	\$	161,580	\$	86,230
Restricted cash	•	1,983	-	1,610
Total cash, cash equivalents and restricted cash	\$	163,563	\$	87,840
Supplemental disclosure of cash flow information:	<u>· </u>		_	
Cash received from exchange of research and development tax credits	\$		\$	324
Supplemental disclosure of noncash information:	5		Ψ	324
Noncash acquisition of fixed assets	\$	62	\$	58
Write-off of notes receivable	\$	90	\$	50
White-off of hotes received.	J	30	Ψ	_

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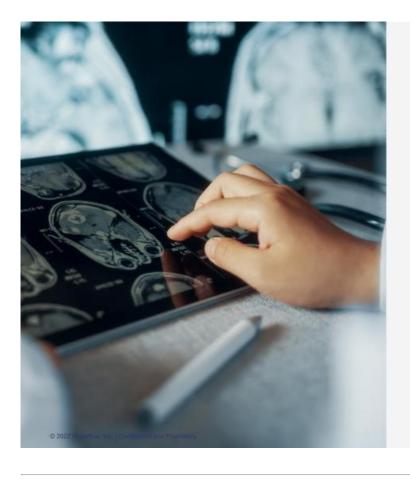
Defining the Future of Life-Saving Diagnostics at the Point of Care

Corporate Presentation | March 23, 2022

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Forward Looking Statements

This presentation includes forward-looking statements within the meaning of the federal securities laws, which are made pursuant to the Safe Harbor Provisions of the Private Securities Litigation Reform Act of 1995. Any statements contained in this call that relate to expectations or predictions of future events, results, or performance are forward-looking statements. All forward-looking statements, including, without limitation, those relating to our operating trends and future financial performance, the impact of COVID-19 or geo-political conflict such as the war in Ukraine, on our business and prospects for recovery, expense management, expectations for hiring, physician training and adoption, growth in our organization, market opportunity, commercial and international expansion, regulatory approvals, and product development are based upon our current estimates and various assumptions. These statements involve material risks and uncertainties that could cause actual results or events to materially differ from those anticipated or implied by these forward-looking statements. Accordingly, you should not place undue reliance on these statements. For a list and description of the risks and uncertainties associated with our business, please refer to the "Risk Factors" section of our S-1 filed with the Securities and Exchange Commission on January 24th, 2022.



Today, brain diagnostics are single point-in-time and delay the time from door to discharge.



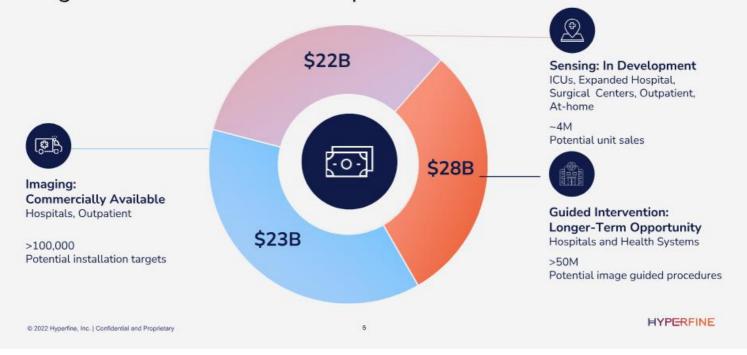
Our mission is to transform healthcare by creating access to life-saving diagnostics and actionable data at the patient bedside.

The Hyperfine Ecosystem

Democratizing Imaging, Sensing, and Guided Intervention to cover the care continuum



Imaging, Sensing, and Guided Intervention are Large Markets Poised for Disruption



We are Transforming Medical Imaging with $\mathsf{Swoop} \ensuremath{\mathbb{R}}$







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6

Swoop® is the Next Generation of MRI



Swoop® Brings MRI to the Patient



Acute Care Settings



Intensive Care Units and Operating Rooms



Global Health

- · Swoop is designed to enable rapid diagnoses and treatment for patients regardless of income, resources, or location
- · Produces high-quality images at low magnetic field strength, allowing clinicians to quickly scan, diagnose, and treat patients
- · Wheeled directly to a patient's bedside, plugged into a standard electrical wall outlet, and controlled by an iPad®

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8

Clinical & Workflow Benefits



Numerous **challenges** with conventional MRI:

High-cost limits accessibility	6
Complex site requirements and upgrades	
Scheduling delays lead	0
to longer length of stay	
Consumption of valuable	~\\$°
personnel resources	. A.
Risk of adverse events	- Th
during transportation	1875
Maintaining connection to	~-V-10./-
life support equipment	

Hyperfine Workflow Benefits



Traditional MRI workflow (25.8 hours)





Hyperfine workflow (90 mins, 94% reduction in total workflow time)



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Swoop Clinical Use Cases Today



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Intensive Care Unit

- Cerebral Edema

- Cerebrovascular Disease - Cranial Neuropathy

Follow-up Hematoma

- Tumor Pre- and Post-Op

Emergency Department
- Blurred Vision

- Traumatic Brain Injury

- Cranial Neuropathy - Dizziness - Headache - Numbness - Stroke

- Ataxia

- Stroke

- Tingling

- Vertigo

- Weakness

12

Clinical Validation of Hyperfine



Game changer is a good way to put it [...] being able to do the level of sophisticated imaging in an ICU that MRI can provide."

Dr. Fady Charbel, MD, FAANS, FACS





Hyperfine provides me with an opportunity to acquire the information, to interpret the information, and to make a decision based on the information that's in front of me."

Dr. Shahid Nimjee, MD, PhD, FAANS, FAHA



Northwell Health

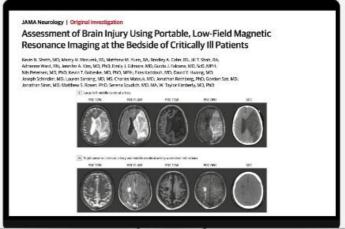


Portable MRI should be used to image any patients in ICUs in any [clinical] setting."

Dr. Michael Schulder, MD, FAANS

Over 40 conference presentations and publications discussing clinical benefits for:

Stroke | Hydrocephalus | Hematoma | Multiple scierosis | Tumor resection



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13

Use Case: Neuro-ICU

Patient Delays to Transfer in the ICU Creates Major Unnecessary Costs for Hospitals, is "Common and Costly"

Estimated \$300/hr for delays, >\$22,000/week for hospital (>\$1M/year) for large academic center

Imaging capabilities of MRI, CT and Ultrasound should be available 24/7/365 at all facilities.

In reality, patients can wait more than 24 hrs for MRI availability, resulting in cost for both the patient and the hospital, taking up an ICU bed.



If only there was a way to improve access to imaging...

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15

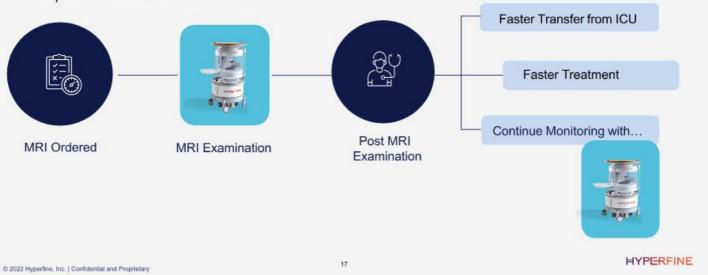
Current ICU Imaging Workflow with Conventional MRI

Traditional MRI workflow can lead to prolonged delays in patient care and higher resources consumption



Improved ICU Imaging Workflow with Swoop

Portable MRI workflow enables timely care for earlier discharge by bringing brain imaging to the patient's bedside



Word from the Clinician





Using the scanner in the ICU is an important use case. What Swoop can offer versus a conventional MRI is the **flexibility** and the usefulness of having it right there. It **favors time** in a situation when you need something acute.

Dr. Jennifer Moliterno-Gunel, Neurosurgeon Yale University School of Medicine

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8

Swoop's Potential Benefits in the ICU



Reduced time to diagnosis

Swoop workflow is significantly faster than conventional MRI



Reduced patient care interruption

Transport time (2-3 hours) interrupts patient care and impacts staffing for entire ICU¹



Reduced adverse events associated with patient transport

Adverse events occur in up to 46% of transported patients.





Reduced costs associated with length of stay

Shortening time to diagnosis, avoiding interruptions in care, and preventing adverse events



Optimized staffing in the ICU

Time consuming transport affects ICU staff: nurse, respiratory therapist, anesthesia, transport, and practitioner.



Reduced exposure to ionizing radiation

lonizing radiation from CT used for serial follow-up scans = risk to patient and staff

1. Hyperfine, Care Area - Acute Mental Status Change, Page 1a

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Use Case: Hydrocephalus

Pediatric Hydrocephalus Management is a Huge Problem

~400,000 hospital days, \$2B in hospital charges in the US



Children with hydrocephalus need life-long monitoring and use a disproportionate number of hospital days and resources.1,2



Children can receive 1-12 CTs5 each year, increasing their risk for radiation-associated malignancy6. Rapid MRI (T2 only) is preferred since it's radiation free but may not be available.



Any symptoms cause trips to hospital for a shunt check to ensure pressure on the brain remains normal. 50% of shunts fail in <2 years and 98% of shunts fail by year 10.2,3,4



Swoop helps overcome existing workflow barriers to enable safe and timely imaging at the point of care for an improved patient experience.

https://theins.org/focus/viewjournals/neurosurg-focus/37/5/article-pE5.xml

https://www.hydroassoc.org/osrebral-shunt-maifunctions/

https://www.gosh.nhs.uk/conditions-and-treatments/conditions-we-treat/ve-

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Personal Story from the Hydrocephalus Association



Received so many CT scans that we're waiting on a cancer diagnosis. **No radiation... Swoop is a parent's dream.**



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22

Hydrocephalus Workflow Improvement with Swoop®

Traditional workflow results in delayed diagnosis and potential radiation exposure



Hyperfine allows children to be imaged sooner, next to their loved ones, without radiation



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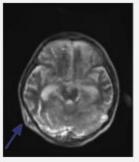
Hydrocephalus: Swoop's Potential from Early Cases

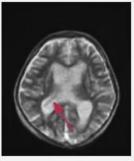


Hyperfine is an excellent addition to the neurosurgery clinic for screening of hydrocephalus patients. The convenience for the patient, reduced scan time, and cost of the machine make this a device that should be considered for any neurosurgery clinic.



When your child needs a hospital, everything matters.







5 y/o presents to Neurosurgery clinic with headache. Swoop® scan performed in the clinic demonstrates ventricular catheter (without artifact from valve) along with enlarged ventricles – child admitted to hospital for shunt revision immediately, saving radiation and delay.

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Use Case: Stroke

Hyperfine Provides Compelling Platform for Stroke Diagnosis

15 million people worldwide suffer a stroke annually

MRI scans are better at detecting ischemic stroke damage compared to CT scans



Stroke is the 2nd leading cause of death globally



MRI use for stroke has been limited due to lack of access to this expensive equipment and experienced neuroradiologists to interpret the results.



for stroke diagnosis at the patient's bedside, images can be shared securely with neuroradiologists around the world



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87% strokes

are ischemic strokes

Stroke Diagnosis Confirmed

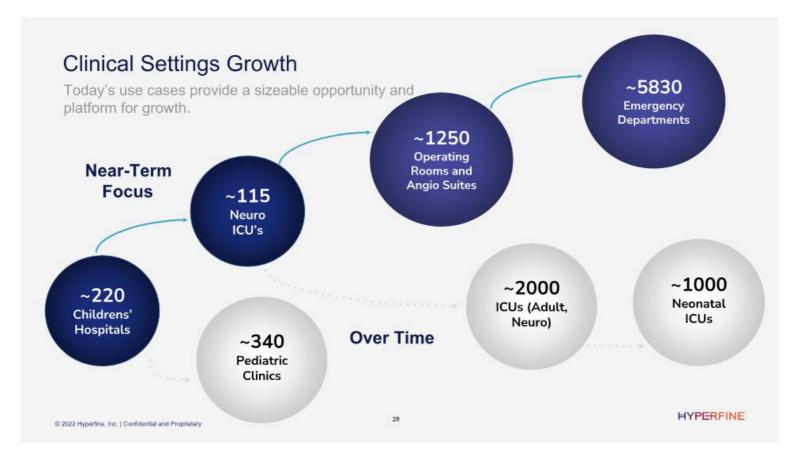
62-year-old male

Presented with new left sided weakness and tremor



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27



Pipeline Opportunities

Innovative R&D Engine Designed to Expand Product Roadmap



Developing a Non-Invasive Brain Vital Sensor

Breakthrough technology designed to unlock access to blood flow and pressure





Non-Invasive

Non-invasive use on every patient to enable broader access and earlier diagnosis



Continuous Trend Analysis

Designed for continuous sensing to build trends for data-backed treatment



Easy to use

Designed to be easy to use for immediate, precise care

*The first medical device is being developed to aid in the diagnosis and management of brain disorders through the development of novel acoustic sensing techniques and innovative algorithms for measuring key metrics of brain health.

Brain-Sensing Clinical Opportunities



Financial Profile

2021 Financial Results & Total Installed Units

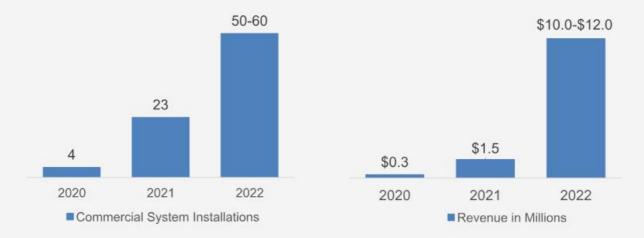
- · \$1.496 million in 2021 total revenue
- Total sales increased by \$1.2 million, or >400%, for the year ended December 31, 2021 compared to the year ended December 31, 2020.

	TOTAL INSTALLED UNITS					
	2020	2021				
		Q1	Q2	Q3	Q4	TOTAL
Commercial system installations*	4	5	7	4	7	27
Grant fulfillment installations		2	2	4	10	18
	4	7	9	8	17	45
Research units	15	2	2	3	3	25
Total Installed Units	19	9	11	11	20	70

Commercial system installations reflect device sales and subscription services through commercial agreements (commercial sales) or through research transfer agreements ("RTA") sales. Commercial sales are made to hospitals and other healthcare providers as direct sales of devices and software subscription services or through subscriptions for the use of the device and software. RTA sales represent the sale of Swoop units for research use purposes.

2022 Financial Guidance

- \$10.0-\$12.0 million in total revenue for full year 2022
- 50-60 commercial system installations in full year 2022



*2022 bar chart values reflect fiscal year financial guidance, not reported results

2021 & 2022: Major Accomplishments

- July 2021: Announced Definitive Agreement to be Listed on Nasdaq through a Business Combination with HealthCor Catalio Acquisition Corp.
- August 2021: Swoop® Demonstrates High Accuracy for Detection of Brain Hemorrhage in Study Published in Nature Communications
- September 2021: Announced Plans for Global Expansion Starting with Launches in the United Kingdom and Pakistan
- September 2021: Announced Receipt of Additional \$3.3 Million Grant from Bill & Melinda Gates Foundation to Improve
 Access to Neonatal and Pediatric Brain Imaging in Low-Resource Settings Globally
- November 2021: Received FDA Clearance for Deep Learning Portable MRI, Defining the Future of Life-Saving Diagnostics
- December 2021: Announced Expansion into Canadian Market with Medical Device License Issued by Health Canada
- December 2021: Closed Business Combination with HealthCor Catalio Acquisition Corp. and Liminal Sciences, Began Trading under the Ticker "HYPR" on the Nasdaq Global Market
- · January 2022: Placed Swoop system with Minnesota Medical Center to Grow its Advanced Imaging Systems Offering
- February 2022: Appointed Chip Truwit, M.D. as Senior Medical Director
- February 2022: Placed Swoop system with Queen's University Radiology to Improve Access to Care for Canadian Patients in Remote Northern Communities

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Named by Fierce Medtech as one of 2020's















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GLOBAL PORTABLE MAGNETIC RESONANCE IMAGING NEW PRODUCT INNOVATION AWARD

BEST 2020 PRACTICES

AWARD

Leadership Team

Management Team with Proven Track Record of Success



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Chief Executive Officer



Alok Gupta

Chief Financial Officer



Dr. Khan Siddiqui

Chief Strategy Officer & Chief Medical Officer



Tom Teisseyre

Chief Product Officer



Mark Hughes

Chief Operating Officer



Scott White

Chief Commercial Officer



Kyla Pavlina

Chief People Officer



Neela Paykel

General Counsel & Chief Compliance Officer

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38

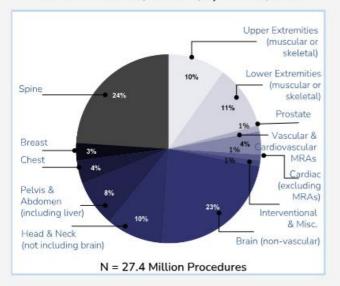
Thank You!



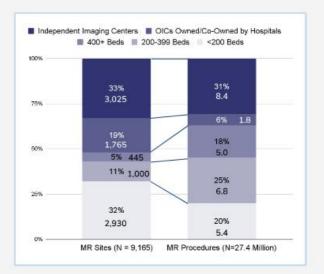
Appendix

Brain is the Largest MRI Market with Nearly 25% of MR Procedures

MR Procedure Mix, All Sites, by Percent, 2020



Distribution of MR Sites and Procedures, by Site Type, 2020



^{*}Source: 2020 IMV MR Benchmark Report © 2022 Hyperline, Inc. | Confidential and Proprietary