UNITED STATES SECURITI

SECURITIES	AND EXCHANGE (Washington, D.C. 20549	COMMISSION
	FORM 8-K	
of :	CURRENT REPORT Pursuant to Section 13 or 15(d the Securities Exchange Act of	
Date of Repo	rt (Date of earliest event reported): A	august 10, 2020
	Morphic Holding, Inc. Name of Registrant as Specified in its	
Delaware (State or other jurisdiction of incorporation or organization)	001-38940 (Commission File Number)	47-3878772 (I.R.S. Employer Identification No.)
35 Gatehouse Drive, A2 Waltham, Massachusetts (Address of principal executive of	fices)	02451 (Zip Code)
-	lephone number, including area code Not Applicable e or Former Address, if Changed Sin	
Check the appropriate box below if the Form 8-K of the following provisions:	filing is intended to simultaneously sati	sfy the filing obligation of the registrant under any
☐ Written communications pursuant to Rule 425	under the Securities Act (17 CFR 230.	425)
\square Soliciting material pursuant to Rule 14a-12 un	der the Exchange Act (17 CFR 240.14a	:-12)
☐ Pre-commencement communications pursuant	to Rule 14d-2(b) under the Exchange A	Act (17 CFR 240.14d-2(b))
☐ Pre-commencement communications pursuant	to Rule 13e-4(c) under the Exchange A	act (17 CFR 240.13e-4(c))
Securities registered pursuant to Section 12(b)) of the Act:	
Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock, \$0.0001 par value per shar		Nasdaq Global Market
Indicate by check mark whether the registrant is a of this chapter) or Rule 12b-2 of the Securities Ex		

X

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 August 10, 2020, Morphic Holding, Inc. issued a press release announcing its financial results for the quarter ended June 30, 2020. A copy of the press release is attached as Exhibit 99.1 to this report.

The information in this Item 2.02, including Exhibit 99.1 to this report, 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 or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended (the "Securities Act"). The information contained in this Item 2.02 and in the accompanying Exhibit 99.1 shall not be incorporated by reference into any other filing under the Exchange Act or under the Securities Act, except as shall be expressly set forth by specific reference in such filing.

Item 9.01 Financial Statements and Exhibits.

Exh

Exhibit Number	Description							
99.1	Press release issued by Morphic Holding, Inc. regarding its financial results for the quarter ended June 30, 2020, dated August 10, 2020.							

SIGNATURE

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.

MORPHIC HOLDING, INC.

Date: August 10, 2020 By: /s/ Marc Schegerin

Marc Schegerin

Chief Financial Officer and Chief Operating Officer



Morphic Announces Corporate Highlights and Second Quarter 2020 Financial Results

Company's first IND accepted by FDA for MORF-057 in IBD phase 1 study expected to begin in third-quarter 2020 Positive data presented at DDW 2020 further support MORF-057 preclinical profile

WALTHAM, Mass. – August 10, 2020 – Morphic Therapeutic (NASDAQ: MORF), a biopharmaceutical company developing a new generation of oral integrin therapies for the treatment of serious chronic diseases, today reported corporate highlights and second quarter 2020 financial results.

"Morphic has made tremendous strides with our lead program, MORF-057, an oral small molecule inhibitor of $\alpha_4\beta_7$ in development to treat inflammatory bowel disease. I'm so proud of the efforts of our team that has taken this program from the bench in Dr. Tim Springer's Lab to the clinic in just a few short years," commented Praveen Tipirneni, M.D., president and chief executive officer of Morphic Therapeutic. "We recently presented a substantive preclinical data set at Digestive Disease Week that support MORF-057's profile as a candidate for the treatment of IBD through the proven mechanism of $\alpha_4\beta_7$ inhibition with many potential advantages, including oral administration. While we are just beginning our clinical journey, I congratulate the Morphic team for the achievement of these important milestones on our mission to deliver a new generation of oral integrin therapeutics."

Peter Linde, M.D., Morphic's chief medical officer, added "We expect to initiate a robust clinical development program for MORF-057 imminently, beginning with a phase 1 clinical trial designed to evaluate the safety, pharmacokinetic and pharmacodynamic profile of this compound with single and multiple ascending dose cohorts. The phase 1 trial will also generate important receptor occupancy data that we believe could provide early clinical proof of concept and dose selection guidance for use in future studies within the MORF-057 program in IBD patients."

Morphic is developing MORF-057 as a selective, oral small molecule inhibitor of the $\alpha_4\beta_7$ integrin, a target for the treatment of inflammatory bowel disease (IBD) with an initial focus on moderate-to-severe ulcerative colitis (UC). The mechanism of $\alpha_4\beta_7$ inhibition to treat IBD has been clinically validated by the success of the approved infused antibody therapy, vedolizumab.

Second quarter and recent corporate highlights:

- Announced acceptance of the Investigational New Drug (IND) filing for MORF-057 by the U.S. Food and Drug Administration (FDA), with a phase 1 clinical trial in healthy subjects expected to begin in the third quarter of 2020
 - O The phase 1 trial is a randomized, double-blind placebo-controlled single and multiple ascending dose study in healthy adults evaluating endpoints including safety, tolerability, and pharmacokinetic and pharmacodynamic measures, as well as a concurrent food effect study
 - O The full data set from the MORF-057 phase 1 trial is expected to be presented by mid-2021
- Presented new positive preclinical data supporting MORF-057 in models of increasing complexity at the virtual Digestive Disease Week 2020
 - O These data demonstrated MORF-057's inhibition of $\alpha_4\beta_7$ -expressing lymphocyte migration to the gut, a fundamental contributor to IBD, acting through the same mechanism of action as the approved antibody therapeutic, vedolizumab
 - O Further, oral administration of a MORF-057 analog in non-human primate studies demonstrated saturation of the $\alpha_4\beta_7$ receptor throughout the entire 7-day dosing period receptor saturation at the dosages assessed in preclinical studies provide further evidence that MORF-057 is an extremely potent inhibitor of $\alpha_4\beta_7$ and may produce clinical efficacy
- Continued progress in our collaboration with AbbVie on the development of integrin inhibitors, including $\alpha_v \beta_6$ and other integrin targets
- Appointed Marc Schegerin, M.D., M.B.A., as chief operating officer and chief financial officer; Dr. Schegerin was previously chief financial officer of ArQule until its acquisition by Merck. His prior roles include senior positions in finance, business development and healthcare investment banking

COVID-19 Preparedness

Morphic is not currently aware of any significant delay to its timelines due to the COVID-19 pandemic. In light of the evolving circumstances, Morphic will continue to assess any potential impact of the COVID-19 pandemic in dialogue with regulators, partners and vendors.



Financial Results for Second Quarter 2020

Net loss for the quarter ended June 30, 2020 was \$15.9 million or \$0.52 per share compared to a net loss of \$9.4 million or \$4.73 per share for the same quarter last year.

- Revenue was \$7.7 million for the quarter ended June 30, 2020 compared to \$5.6 million for the same quarter last year. The increase was due to higher level of research & development efforts in collaboration agreements signed with AbbVie in October 2018 and Janssen in February 2019
- Research and development expenses were \$19.9 million for the quarter ended June 30, 2020, compared to \$13.9 million in the same quarter last year. The \$6.0 million increase year-over-year reflects development and manufacturing costs associated with lead wholly owned product candidate, MORF-057; research costs associated with preclinical studies related to our partnered product candidate MORF-720 with AbbVie; as well as increased personnel-related costs to support continued progress with the company's pipeline
- General and administrative expenses were \$4.2 million for the quarter ended June 30, 2020, compared to \$2.1 million in the same quarter last year. The \$2.1 million increase year-over-year was primarily attributable to increased headcount and higher professional fees to operate as a public company along with consulting fees associated with ongoing business development activities

As of June 30, 2020, Morphic had cash, cash equivalents, and marketable securities of \$202.5 million, compared to \$237.0 million at the end of 2019. Morphic believes its cash, cash equivalents, and marketable securities balance as of June 30, 2020 will be sufficient to fund operating expenses and capital expenditure requirements at least through the end of 2022.

Financial Tables Follow



Morphic Holding Inc. Condensed Consolidated Statements of Operations

(unaudited)

(in thousands, except share and per share data)

	Three Months Ended June 30			Six Months Ended June 30								
2020			2019		_	2020				2019		
Collaboration revenue	\$	7,693	\$		5,567	\$	5	13,287	\$		11,635	
Operating expenses:												
Research and development		19,918			13,907			38,878			24,278	
General and administrative		4,195			2,077			8,618			3,908	
Total operating expenses		24,113			15,984			47,496			28,186	
Loss from operations		(16,420)			(10,417)			(34,209)			(16,551)	
Interest income, net		407			1,119			1,293			2,182	
Total other income		407			1,119			1,293			2,182	
Loss before benefit from (provision for) income taxes Benefit from (provision		(16,013)			(9,298)			(32,916)	_		(14,369)	
for) income taxes		155			(135)			312	_		(264)	
Net loss	\$	(15,858)	= =	\$	(9,433)	4	5	(32,604)	_	\$	(14,633)	
Net loss per share, basic and diluted Weighted-average common shares	\$	(0.52)		\$	(4.73)	9	\$	1.08		\$	(7.54)	
outstanding, basic and diluted		30,360,851			1,992,410		;	30,274,713	_		1,940,923	



Morphic Holding Inc. Condensed Consolidated Balance Sheets

(unaudited) (in thousands)

	June 30, 2020		D	December 31, 2019			
Assets							
Cash, cash equivalents and marketable securities	\$	202,538	\$	237,016			
Other current assets		6,072		6,557			
Total current assets		208,610		243,573			
Other assets		3,389		3,862			
Total assets	\$	211,999	\$	247,435			
Liabilities and Stockholders' Equity							
Current liabilities	\$	45,284	\$	35,350			
Long-term liabilities		51,798		71,167			
Total liabilities		97,082	<u> </u>	106,517			
Total stockholders' equity		114,917		140,918			
Total liabilities and stockholders' equity	\$	211,999	\$	247,435			

About Morphic Therapeutic

Morphic Therapeutic is a biopharmaceutical company developing a new generation of oral integrin therapies for the treatment of serious chronic diseases, including autoimmune, cardiovascular and metabolic diseases, fibrosis and cancer. In collaboration with AbbVie, Janssen and Schrödinger, Morphic is advancing its pipeline and discovery activities using its proprietary MInT technology platform which leverages the Company's unique understanding of integrin structure and biology. For more information, visit www.morphictx.com.

Cautionary Note Regarding Forward-Looking Statements

This press release contains "forward-looking" statements within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995, including, but not limited to: Morphic's plan to develop and commercialize oral small-molecule integrin therapeutics and Morphic's expectations about timing and ability to commence or complete clinical studies and to obtain regulatory approvals for MORF-057, MORF-720, and other candidates in development, the ability of MORF-057 to treat inflammatory bowel disease, the potential impact of the COVID-19 pandemic and the sufficiency of our cash, cash equivalents and investments to fund our operations. Statements including words such as "believe," "plan," "continue," "expect," "will be," "develop," "signal," "potential," or "ongoing" and statements in the future tense are forward-looking statements. These forward-looking statements involve risks and uncertainties, as well as assumptions, which, if they do not fully materialize or prove incorrect, could cause our results to differ materially from those expressed or implied by such forward-looking statements. Forward-looking statements are subject to risks and uncertainties that may cause Morphic's actual activities or results to differ significantly from those expressed in any forward-looking statement, including risks and uncertainties related to Morphic's ability to develop, obtain regulatory approval for and commercialize MORF-057, MORF-720, and other product candidates, the timing and results of preclinical studies and clinical trials, the potential impact of the COVID-19 pandemic, Morphic's ability to protect intellectual property; and other risks set forth in our filings with the Securities and Exchange Commission. These forward-looking statements speak only as of the date hereof and Morphic specifically disclaims any obligation to update these forward-looking statements or reasons why actual results might differ, whether as a result of new information, future events or otherwise, except as required by law.



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