UNITED STATES SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549

FORM 6-K

Report of Foreign Private Issuer Pursuant to Rule 13a-16 or 15d-16 of the Securities Exchange Act of 1934

For the Month of Feb 2021

Commission File Number 001-35948

Kamada Ltd.

(Translation of registrant's name into English)

2 Holzman Street Science Park, P.O. Box 4081 Rehovot 7670402 Israel

(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F.
Form 20-F ⊠ Form 40-F □
Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):
Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7):
Indicate by check mark whether the registrant by furnishing the information contained in this Form is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.
Yes □ No ⊠
If "Yes" is marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b): 82
This Form 6-K is being incorporated by reference into the Registrant's Form S-8 Registration Statements, File Nos. 333-192720, 333-207933 333-215983, 333-222891 and 333-233267

99.1 Kamada Reports Fourth Quarter and Fiscal Year 2020 Financial Results, Recent Achievements and Corporate Development Activities

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.

Date: February 10, 2021 KAMADA LTD.

By: /s/ Yifat Philip

Yifat Philip Vice President General Counsel and

Corporate Secretary

EXHIBIT INDEX

EXHIBIT NO. DESCRIPTION

99.1 Kamada Reports Fourth Quarter and Fiscal Year 2020 Financial Results, Recent Achievements and Corporate Development Activities

Kamada Reports Fourth Quarter and Fiscal Year 2020 Financial Results, Recent Achievements and Corporate Development Activities

- Full-Year 2020 Revenues were \$133.2 Million, In-Line with the Company's Guidance; Adjusted EBITDA for 2020 was \$25.1 Million
- Kamada Enters U.S. Plasma Collection Market Through Acquisition of an FDA-Approved Facility, Furthering its Strategic Goal of Becoming a Fully Integrated Specialty Plasma Company Focused on Growing its Hyperimmune IgG Portfolio
- Israeli Ministry of Health (IMOH) Initiated Treatment of Hospitalized COVID-19 Patients with Kamada's Plasma-Derived COVID-19
 Immunoglobulin (IgG) Investigational Product Primarily as Part of a Multi-Center Clinical Study Led by the IMOH; Current Planned Supply to
 IMOH is Sufficient to Treat an Estimated 500 Patients
- Kamada Continues to Expand Existing Business Lines as Evidenced by the Recent Additions to its Israeli Distribution Biosimilar Products
 Portfolio; Also Exploring Business Development Opportunities and Intends to Leverage IgG Platform Technology as Response to Future
 Pandemic Situations

REHOVOT, Israel – February 10, 2020 -- Kamada Ltd. (NASDAQ: KMDA; TASE: KMDA.TA), a plasma-derived biopharmaceutical company, today announced financial results for the three and 12 months ended December 31, 2020.

"We are pleased to have overcome meaningful operational challenges caused by the global COVID-19 pandemic and met our key financial targets for 2020," said Amir London, Kamada's Chief Executive Officer. "For full-year 2020, we recorded total revenues of \$133.2 million, in-line with our guidance of \$132 million to \$137 million, and a 5% increase compared to 2019 revenues. These results are indicative of the fundamental strengths of our business."

"The acquisition of the Blood and Plasma Research (B&PR) collection center and establishment of our wholly-owned subsidiary, Kamada Plasma LLC, represents our entry into the U.S. plasma collection market, furthers our strategic goal of becoming a fully integrated specialty plasma company, and is expected to improve our IgG competitive profile in multiple markets. We intend to significantly expand our hyperimmune plasma collection capacity by investing in this center and leveraging its FDA license to open additional centers in the U.S.," continued Mr. London.

"We continue to advance the rapid development and manufacturing of our plasma-derived COVID-19 IgG investigational product, which is being supplied to the Israeli Ministry of Health (IMOH) for the treatment of an estimated 500 hospitalized patients, through an agreement that is expected to generate approximately \$3.4 million in revenue to Kamada. We continue to ramp up production of the product using plasma collected by our partner Kedrion Biopharma, in anticipation of a potential expansion of the IMOH supply agreement and possible demand from additional international markets," added Mr. London.

"In addition, we continue to develop our pipeline, primarily focusing on the pivotal Phase 3 InnovAATe clinical trial of Inhaled AAT for the treatment of Alpha-1 Antitrypsin Deficiency and on exploring new strategic business development opportunities that will utilize and expand our core plasma-derived development, manufacturing, and commercialization expertise. Moreover, we intend to leverage our IgG platform technology as a strategic business line, with the ability to respond to future potential pandemic situations," concluded Mr. London.

As previously reported, the transition of Glassia manufacturing to Takeda and the continued uncertainty in the operating environment created by the ongoing global COVID-19 pandemic are expected to result in reduced revenues and profitability in 2021. At the same time, Kamada continues to focus on expanding its existing growth drivers, which include:

- Continued market share growth for KedRAB in the U.S.;
- Expanding the sales of Glassia and the Company's IgG portfolio in ex-U.S. markets, including registration and launch of the products in new territories;
- Generating royalties from Glassia, projected to be in the range of \$10 million to \$20 million per year commencing in 2022;
- Launching nine biosimilar products in the Israeli-based Distribution segment between 2022 and 2025, pending regulatory approval, with estimated maximum sales in the range of \$25 million to \$35 million; and
- Leveraging the Company's plasma-derived products manufacturing facility and expertise to offer contract manufacturing services of specialty IgG products, including an FDA-approved and commercialized IgG product that is expected to add between \$8 million to \$10 million in annual revenues, beginning in 2023.

Financial Highlights for the Three Months Ended December 31, 2020

- Total revenues were \$31.5 million in the fourth quarter of 2020, a 2% decrease from the \$32.1 million recorded in the fourth quarter of 2019.
- Gross profit was \$10.2 million in the fourth quarter of 2020, compared to \$12.1 million reported in the fourth quarter of 2019.
- Net income was \$1.6 million, or \$0.04 per share, in the fourth quarter of 2020, as compared to net income of \$5.4 million, or \$0.13 per share, in the fourth quarter of 2019.
- Adjusted EBITDA, as detailed in the tables below, was \$4.0 million in the fourth quarter of 2020, as compared to \$6.8 million in the fourth quarter of 2019.
- Cash provided by operating activities was \$12.7 million in the fourth quarter of 2020, as compared to cash provided by operating activities of \$8.6 million in the fourth quarter of 2019.

Financial Highlights for the Year Ended December 31, 2020

- Total revenues were \$133.2 million in the year ended December 31, 2020, a 5% increase from the \$127.2 million recorded in the year ended December 31, 2019.
- Gross profit was \$47.6 million in the year ended December 31, 2020, compared to \$49.7 million in the same period of 2019.
- Proprietary Product segment gross margins in the year ended December 31, 2020 were 43%, down three percentage points from the year ended December 31, 2019, and in-line with the Company's expectations of an annual decrease of three to five percentage points, primarily attributable to a change in product sales mix and reduced plant utilization.
- Operating expenses, including Research and Development, Sales & Marketing, General and Administrative, and Other Expenses, totaled \$28.3 million in the year ended December 31, 2020, as compared to \$27.0 million in the year ended December 31, 2019.
- The 4% increase in Research and Development expenses for full-year 2020 as compared to 2019 was below Kamada's guidance of a 13%-15% year-over-year increase. This is primarily due to the delay in patient enrollment in the Company's InnovAATe clinical trial due to the impact of the COVID-19 pandemic.
- Net income was \$17.1 million, or \$0.38 (fully diluted) per share, in the year ended December 31, 2020, as compared to net income of \$22.3 million, or \$0.55 per share, in the year ended December 31, 2019.
- Adjusted EBITDA, as detailed in the tables below, was \$25.1 million in the year ended December 31, 2020, as compared to \$28.5 million in the
 year ended December 31, 2019.
- Cash provided by operating activities was \$19.1 million in the year ended December 31, 2020, as compared to cash provided by operating activities of \$27.6 million in the year ended December 31, 2019.

Balance Sheet Highlights

As of December 31, 2020, the Company had cash, cash equivalents, and short-term investments of \$109.3 million, as compared to \$73.9 million on December 31, 2019. This increase was attributable to the issuance and sale of \$25 million of equity to FIMI Opportunity Fund, the leading private equity investor in Israel, as well as positive operational cash flow.

Recent Corporate Highlights

- Acquired privately held B&PR's plasma collection center in Beaumont, TX, which specializes in the collection of hyper-immune plasma used by Kamada in the manufacture of its anti-D products.
- Began supplying its COVID-19 plasma-derived IgG product to the IMOH for the treatment of COVID-19 patients in Israel; agreement expected
 to generate approximately \$3.4 million in revenue to Kamada.
- Entered into agreements with two undisclosed international pharmaceutical companies to commercialize three biosimilar product candidates in Israel. Subject to approval by the European Medicines Agency and subsequently by the IMOH, the three products are expected to be launched in Israel between 2022 and 2024 and Kamada estimates the potential collective maximum sales generated by the sale of these three products, achievable following regulatory approval and within several years of launch, to be in the range of \$5 million to \$7 million annually.
- Added to the NASDAQ Biotechnology Index® (Nasdaq: NBI) designed to track the performance of a set of either biotechnology or pharmaceutical companies listed on The NASDAQ Stock Market®.

Conference Call

Kamada management will host an investment community conference call on Wednesday, February 10, 2020, at 8:00am Eastern Time to discuss these results and answer questions. Shareholders and other interested parties may participate in the conference call by dialing 877-407-0792 (from within the U.S.), 1-809-409-247 (from Israel), or 201-689-8263 (International) and entering the conference identification number: 13715277. The call will also be webcast live on the Internet on the Company's website at http://public.viavid.com/index.php?id=143096.

About Kamada

Kamada Ltd. ("the Company") is a commercial stage plasma-derived biopharmaceutical company focused on orphan indications, with an existing marketed product portfolio and a late-stage product pipeline. The Company uses its proprietary platform technology and know-how for the extraction and purification of proteins from human plasma to produce Alpha-1 Antitrypsin (AAT) in a highly-purified, liquid form, as well as other plasmaderived immune globulins. The Company's flagship product is GLASSIA®, the first liquid, ready-to-use, intravenous plasma-derived AAT product approved by the U.S. FDA. The Company markets GLASSIA in the U.S. through a strategic partnership with Takeda Pharmaceuticals Company Limited and in other countries through local distributors. Pursuant to an agreement with Takeda, the Company will continue to produce Glassia for Takeda through 2021 and Takeda will initiate its own production of Glassia for the U.S. market in 2021, at which point Takeda will commence payment of royalties to the Company. The Company's second leading product is KamRab®, a rabies immune globulin (Human) for post-exposure prophylaxis against rabies infection. KamRab is FDA approved and is being marketed in the U.S. under the brand name KEDRAB® through a strategic partnership with Kedrion S.p.A. In addition to Glassia and KEDRAB, the Company has a product line of four other plasma-derived pharmaceutical products administered by injection or infusion, that are marketed through distributors in more than 15 countries, including Israel, Russia, Brazil, India and other countries in Latin America and Asia. The Company has late-stage products in development, including an inhaled formulation of AAT for the treatment of AAT deficiency. In addition, the Company's intravenous AAT is in development for other indications, such as GvHD and prevention of lung transplant rejection, and during 2020, the Company initiated the development of a plasma derived hyperimmune immunoglobulin (IgG) product as a potential treatment for coronavirus disease (COVID-19). The Company leverages its expertise and presence in the plasma-derived protein therapeutics market by distributing more than 20 complementary products in Israel that are manufactured by third parties. FIMI Opportunity Fund, the leading private equity investor in Israel, is the Company's lead shareholder, beneficially owning approximately 21% of the outstanding ordinary shares.

Cautionary Note Regarding Forward-Looking Statements

This release includes forward-looking statements within the meaning of Section 21E of the U.S. Securities Exchange Act of 1934, as amended, and the safe harbor provisions of the U.S. Private Securities Litigation Reform Act of 1995. Forward-looking statements are statements that are not historical facts, including statements regarding: 1) 2020 financial results being indicative of the fundamental strengths of Kamada's business, 2) benefits of the B&PR acquisition, including furthering Kamada's strategic goal of becoming a fully integrated specialty plasma company, the acquisition's ability to improve Kamada's IgG competitiveness in different markets and ability to expand Kamada's hyperimmune plasma collection capacity by investing in the acquired center and leveraging its FDA license to open additional centers in the U.S., 3) plasma derived COVID-19 IgG investigational product supplied to the IMOH is estimated to be used to treat 500 hospitalized patient, through an agreement expected to generate approximately \$3.4M in revenue to Kamada, 4) raping up of manufacturing of Kamada's plasma derived COVID-19 immunoglobulin investigational product in anticipation of a potential expansion of the IMOH supply agreement and possible demand from additional international markets, 5) optimism about the pipeline, which is primarily focused on the pivotal Phase 3 InnovAATe clinical trial of Inhaled AAT for the treatment of Alpha-1 Antitrypsin Deficiency and exploring new strategic business development opportunities that will utilize and expand Kamada's core plasma-derived development, manufacturing, and commercialization expertise, 6) Kamada's intention to leverage its IgG platform technology as a strategic business line, with the ability to respond to future potential pandemic situations, 7) expectation to reduced revenues and profitability in 2021 due to the recently reported transition of Glassia manufacturing to Takeda and the continued uncertainty in the operating environment created by the ongoing global COVID-19 pandemic, 8) expectation for a continued increase in KedRAB market share in the U.S., 9) Expansion of sales of Glassia and IgG portfolio in ex-U.S. markets, including the registration and launch of the products in new territories, 10) Glassia's royalties estimated at a range of \$10 Million to \$20 Million per year commencing 2022, 11) the nine biosimilar products portfolio in the Israeli-based Distribution segment launching between 2022 and 2025 and estimation for maximum sales in the range of \$25 Million to \$35 Million, and 12) the leveraging of the Company's plasma-derived products manufacturing facility and expertise to offer contract manufacturing services of specialty IgG products, including those of an FDA-approved and commercialized IgG product expected to add between \$8 Million to \$10 Million in annual revenues starting in 2023. Forwardlooking statements are based on Kamada's current knowledge and its present beliefs and expectations regarding possible future events and are subject to risks, uncertainties and assumptions. Actual results and the timing of events could differ materially from those anticipated in these forward-looking statements as a result of several factors including, but not limited to, the continued evolvement of the COVID-19 pandemic, its scope, effect and duration, availability of sufficient raw materials required to maintain manufacturing plans, the effects of the COVID-19 pandemic and related government mandates on the availability of adequate levels of work-force required to maintain manufacturing plans, disruption to the supply chain due to COVID-19 pandemic, continuation of inbound and outbound international delivery routes, ability to offset significant revenue loss associated with GLASSIA manufacturing transitioning to Takeda, continued demand for Kamada's products, including GLASSIA and KEDRAB, in the U.S. market and its Distribution segment related products in Israel, financial conditions of the Company's customer, suppliers and services providers. ability to reap the benefits of the B&PR acquisition, ability to obtain regulatory approval for clinical trials of the plasma-derived hyperimmune IgG product for COVID-19, ability to obtain regulatory approval for the nine biosimilar products portfolio, the fruition of the contract manufacturing services of the FDA-approved and commercialized specialty IgG product, the ability to continue enrollment of the pivotal Phase 3 InnovAATe clinical trial, unexpected results of clinical studies and on-going compassionate-use treatments, Kamada's ability to manage operating expenses, additional competition in the markets that Kamada competes, regulatory delays, prevailing market conditions and the impact of general economic, industry or political conditions in the U.S., Israel or otherwise. The forward-looking statements made herein speak only as of the date of this announcement and Kamada undertakes no obligation to update publicly such forward-looking statements to reflect subsequent events or circumstances, except as otherwise required by law.

CONTACTS:

Chaime Orlev Chief Financial Officer IR@kamada.com

Bob Yedid LifeSci Advisors, LLC 646-597-6989 Bob@LifeSciAdvisors.com

CONSOLIDATED BALANCE SHEETS

Assets 2020 2019 U.S. Dollars in thousands Current Assets Cash and cash equivalents \$70,197 \$42,662 Short-term investments 39,069 31,245 Trade receivables, net 22,108 23,210 Other accounts receivables 4,524 3,272 Inventories 42,016 43,173 Total Current Assets 177,914 143,562 Non-Current Assets 25,679 24,550 Right-of-use-assets 3,440 4,022 Other long term assets 1,573 352 Contract asset 2,059 - Deferred taxes 2,059 - Total Non-Current Assets 32,751 30,235 Total Assets \$210,665 \$173,797
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Lighilities
Liquinties
Current Liabilities
Current maturities of bank loans \$ 238 \$ 489
Current maturities of lease liabilities 1,072 1,020
Trade payables 16,110 24,830
Other accounts payables 7,547 5,811
Deferred revenues - 589
Total Current Liabilities 24,967 32,739
Non-Current Liabilities
Bank loans 36 257
Lease liabilities 3,593 3,981
Deferred revenues 2,025 232
Employee benefit liabilities, net 1,406 1,269
Total Non-Current Liabilities 7,060 5,739
<u>Shareholder's Equity</u>
Ordinary shares 11,706 10,425
Additional paid in capital net 209,760 180,819
Capital reserve due to translation to presentation currency (3,490)
Capital reserve from hedges 357 8
Capital reserve from financial assets measured at fair value through other comprehensive Income - 145
Capital reserve from share-based payments 4,558 8,844
Capital reserve from employee benefits (320) (359)
Accumulated deficit (43,933) (61,073)
Total Shareholder's Equity 178,638 135,319
Total Liabilities and Shareholder's Equity \$\frac{\$10,665}{\$}\$\$ \frac{\$173,797}{\$}\$

CONSOLIDATED STATEMENTS OF COMPREHENSIVE INCOME

		For the year ended December 31,			Th		s period ended lber 31,	
		2020		2019		2020		2019
	U	J.S. Dollars	in the	ousands, oth				
Revenues from proprietary products	\$	100,916	\$	97,696	\$	23,283	\$	25,175
Revenues from distribution		32,330	-	29,491	*	8,259	*	6,896
Total revenues	_	133,246		127,187		31,542		32,071
Cost of revenues from proprietary products		57,750		52,425		13,933		14,013
Cost of revenues from distribution		27,944		25,025		7,444		5,969
Total cost of revenues		85,694		77,450		21,377		19,982
Gross profit		47,552		49,737		10,165		12,089
Research and development expenses		13,609		13,059		3,274		3,329
Selling and marketing expenses		4.518		4,370		1.221		929
General and administrative expenses		10,139		9,194		3,006		2,343
Other expenses and (incomes)		49		330		15		3
Operating income		19,237		22,784		2,649		5,485
Financial income		1,027		1,146		162		259
Income (expense) in respect of securities measured at fair value, net		1,027		(5)		102		(2)
Income (expense) in respect of currency exchange differences and derivatives		102		(3)		_		(2)
instruments, net		(1,535)		(651)		(839)		(148)
Financial expenses		(266)		(293)		(62)		(76)
Income before taxes		18,565		22,981		1,910		5,518
Taxes on income		1,425		730		281		156
Net Income	\$	17,140	\$	22,251		1,629	\$	5,362
Other Comprehensive Income (loss):								
Amounts that will be or that have been reclassified to profit								
or loss when specific conditions are met:								
Gain from securities measured at fair value through other comprehensive		(100)		1.42				11
income		(188)		143 92		360		11
Gain (loss) on cash flow hedges Net amounts transferred to the statement of profit or loss for cash flow		876		92		300		(7)
hedges		(528)		(23)		(255)		(3)
Items that will not be reclassified to profit or loss in subsequent periods:		(326)		(23)		(233)		(3)
Remeasurement gain (loss) from defined benefit plan		64		(388)		64		(388)
Tax effect		19		(11)		(10)		22
Total comprehensive income	\$	17,383	\$	22,064	\$	1,788	\$	4,997
Esminar and have established to esmit a later of the Community								
Earnings per share attributable to equity holders of the Company:	C	0.20	Ф	0.55	Ф	0.04	Ф	0.12
Basic income per share	\$	0.39	\$	0.55	\$	0.04	\$	0.13
Diluted income per share	\$	0.38	\$	0.55	\$	0.04	\$	0.13

CONSOLIDATED STATEMENTS OF CASH FLOWS

	For the year ended December 31,					ee months Decem		
		2020	20	19	20	020	2	2019
			U.S	in thous	sands			
Cash Flows from Operating Activities								
Net income	\$	17,140	\$	22,251	\$	1,629	\$	5,362
Adjustments to reconcile net income to net cash provided by (used in)								
operating activities:								
Adjustments to the profit or loss items:								
Depreciation and impairment		4,897		4,519		1,265		1,140
Financial income, net		672		(197)		739		(33)
Cost of share-based payment		977		1,163		124		176
Taxes on income		1,425		730		281		156
Gain from sale of property and equipment		(7)		(2)		-		-
Change in employee benefit liabilities, net		201		94		208		(3)
		8,165		6,307		2,617		1,436
Changes in asset and liability items:								
Decrease (increase) in trade receivables, net		1,332		5,117		6,872		709
Increase in other accounts receivables		115		(214)		(857)		(1,418)
Increase in inventories		1,157		(13,857)		602		(9,142)
Decrease (increase) in Contract asset and deferred expenses		(3.085)		399		(621)		66
Increase (decrease) in trade payables		(9,560)		6,259		928		10,844
Increase (decrease) in other accounts payables		1,736		863		1,310		484
Decrease in deferred revenues		1,204		(283)		14		(62)
		(7,101)		(1,716)		8,248		1,481
Cash received (paid) during the period for:		(1,5202)		(1,110)		<u> </u>		3,102
Interest paid		(209)		(243)		(51)		(61)
Interest received		1,211		1,106		320		552
Taxes paid		(101)		(134)		(14)		(109)
		901		729		255		382
Net cash provided by operating activities	\$	19,105	\$	27,571	\$	12,749	\$	8,661

CONSOLIDATED STATEMENTS OF CASH FLOWS

	For the year ended December 31,					nree months Decem		
	2020			2019		2020		2019
	U.S. Dollars in					ousands		
Cash Flows from Investing Activities								
Proceeds of investment in short term investments, net	\$	(7,646)	\$	1,727	\$	8,000	\$	7,887
Purchase of property and equipment and intangible assets		(5,488)		(2,300)		(2,116)		(812)
Proceeds from sale of property and equipment		7		9		<u>-</u>		_
Net cash provided by (used in) investing activities		(13,127)		(564)		5,884		7,075
Cash Flows from Financing Activities								
Proceeds from exercise of share base payments		65		16		4		4
Repayment of lease liabilities		(1,103)		(1,070)		(288)		(276)
Repayment of long-term loans		(492)		(476)		(119)		(123)
Proceeds from issuance of ordinary shares, net		24,894		<u> </u>				
Net cash used in financing activities		23,364		(1,530)		(403)		(395)
Exchange differences on balances of cash and cash equivalent		(1,807)		(908)		(520)		(128)
Increase in cash and cash equivalents		27,535		24,569		17,710		15,213
Cash and cash equivalents at the beginning of the period		42,662		18,093		52,487		27,449
Cash and cash equivalents at the end of the period	\$	70,197	\$	42,662	\$	70,197	\$	42,662
Significant non-cash transactions								
Purchase of property and equipment through leases	\$	539	\$	5,035	\$	-	\$	51
Purchase of property and equipment	\$	722	\$	992	\$	722	\$	992

ADJUSTED EBITDA

	 For the year ended December 31,				ree months Decem		period ended ber 31,		
	2020		2019		2019 2020		2020		2019
		J	J.S. Dollars	in thousands					
Net income (loss)	\$ 17,140	\$	22,251	\$	1,629	\$	5,362		
Taxes on income	1,425		730		281		156		
Financial expense (income), net	672		(197)		739		(33)		
Depreciation and amortization expense	4,897		4,519		1,265		1,140		
Non-cash share-based compensation expenses	 977		1,163		124		176		
Adjusted EBITDA	\$ 25,111	\$	28,466	\$	4,038	\$	6,801		

ADJUSTED NET INCOME

		For the year ended December 31,					period ended ber 31,		
	2020		2019		2020			2019	
			Ţ	J.S. Dollars	in tho	usands			
Net income (loss)	\$	17,140	\$	22,251	\$	1,629	\$	5,362	
Share-based compensation charges		977		1,163		124		176	
Adjusted net income	\$	18,117	\$	23,414	\$	1,753	\$	5,538	