

Kintor Pharmaceutical (9939 HK)

Multiple out-license collaborations for Proxalutamide

Kintor reported attributable net loss of RMB325.8mn in 1H21 vs RMB195.4mn in 1H20, while adjusted net loss widened from RMB163.7mn in 1H20 to RMB299.9mn in 1H21. R&D expenses increased 90% YoY to RMB282.2mn in 1H21.

- Proxalutamide has clear MoA for treatment of COVID-19.** Proxalutamide has a dual mechanism of action, which inhibits the androgen receptor competitively and decreases the expression of AR, effectively lowering the expression of the proteins ACE2 and TMPRSS2, which the coronavirus uses to invade host cells. In addition, Proxalutamide also promotes the clearance of pathogens and decreases inflammation by activating the Nrf2 pathway, which activates several antioxidative genes and proteins and reduces the intensity of the cytokine response, which is of clinical benefit to the most seriously ill COVID-19 patients. The drug also has potential to treat coronavirus variants. In vitro studies in the P3 laboratory have demonstrated that Proxalutamide can effectively inhibit infections caused by Alpha and Delta variants. The outcome of genome sequencing on COVID-19 inpatients in Brazil has shown that Proxalutamide has effectively treated inpatients infected by Gamma variant.
- Proxalutamide has achieved multiple commercial collaborations in overseas regions.** In Jul 2021, Kintor entered into an agreement with Fosun Pharma on the commercialization of Proxalutamide for COVID-19 indication in India and 28 African countries. In Aug 2021, Kintor entered into a licensing agreement with Etana Biotech in relation to the commercialization of Proxalutamide for COVID-19 treatment in Indonesia. Commercial rights out-licenses reflect peers' approbation on Proxalutamide for COVID-19. Proxalutamide is currently undergoing two registered phase 3 MRCT for the treatment of COVID-19 outpatients with mild to moderate symptoms, and one registered phase 3 MRCT for COVID-19 in-patients, in regions including the US, South America, EU and Asia. In 1H21, Kintor obtained its first emergency use authorization (EUA) from Paraguay's MSPBS.
- Pyrilutamide to release PoC data soon.** Kintor has completed the phase 2 trial of Pylrutamide with 120 enrolled androgenetic alopecia (AGA) patients in China. Preliminary data is expected to be released in Sep 2021, which could be the first PoC data. On 11 Jul, the US FDA has greenlighted Pylrutamide's phase 2 clinical trial for treating AGA. The population of patients with AGA reached over 133.7mn in China and 83.1mn in the US in 2019, respectively. The large prevalence and lack of effective treatment indicates huge unmet medical need for AGA.
- Maintain BUY.** We maintain TP unchanged at HK\$98.07 based on 10-year DCF model (WACC: 10.0%, terminal growth rate: 3.0%). **Risks:** Clinical trial failure in proxalutamide for COVID-19; Delay in pipelines.

Earnings Summary

(YE 31 Dec)	FY19A	FY20A	FY21E	FY22E	FY23E
Revenue (RMB mn)	0	0	0	10,132	8,351
Attributable net profit (loss) (RMB mn)	(233)	(508)	(450)	5,899	5,030
R&D expenses	(214)	(329)	(400)	(600)	(400)
EPS (RMB)	N/A	(1.64)	(1.16)	15.22	12.98
Consensus EPS (RMB)	N/A	N/A	(1.57)	12.66	6.09
ROE (%)	(63)	(34)	(22)	75	39
ROA (%)	(42)	(27)	(19)	67	37
Net gearing (%)	Net cash	Net cash	Net cash	Net cash	Net cash
Current ratio (x)	1.5	8.4	11.0	11.0	21.5

Source: Company data, Bloomberg, CMBIS estimates

BUY (Maintain)

Target Price **HK\$98.07**
 (Previous TP **HK\$98.07**)
 Up/Downside **+36.21%**
 Current Price **HK\$72.00**

China Healthcare Sector

Sam Hu, PhD
 (852) 3900 0882
 samhu@cmbi.com.hk

Jill Wu, CFA
 (852) 3900 0842
 jillwu@cmbi.com.hk

Jonathan Zhao
 (852) 6359 1614
 jonathanzhao@cmbi.com.hk

Mkt. Cap. (HK\$ mn)	27,906
Avg. 3mths t/o (HK\$ mn)	184.39
52W High/Low (HK\$)	86.60/7.20
Total Issued Shares (mn)	388

Source: Bloomberg

Shareholding Structure

Management	34.03%
Pre-IPO & corner stone investors	34.07%
Free float	31.90%

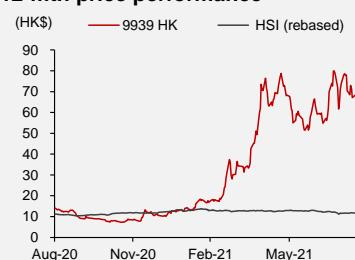
Source: HKEx, Bloomberg

Share performance

	Absolute	Relative
1-mth	12.0%	12.3%
3-mth	5.4%	20.8%
6-mth	326.5%	386.5%

Source: Bloomberg

12-mth price performance



Source: Bloomberg

Auditor: PWC
Web-site: www.kintor.com.cn

Related report:

- Encouraging real-world data of proxalutamide in COVID-19 treatment in Paraguay – 26- Jul 2021
- Proxalutamide may become an effective treatment for COVID-19 – 4 May 2021
- Fast clinical progress for Proxalutamide and other core assets – 29 Mar 2021

Valuation

We use DCF method to value the Company and we derive TP of HK\$98.07 based on 10-year risk-adjusted DCF model (WACC: 10.0%, terminal growth rate: 3.0%).

Figure 1: Risk-adjusted DCF valuation

DCF Valuation (in Rmb mn)	2021E	2022E	2023E	2024E	2025E	2026E	2027E	2028E	2029E	2030E
EBIT	(465)	6,883	5,786	3,854	2,870	2,596	2,499	2,606	2,757	2,996
Tax rate	0%	15%	15%	15%	15%	15%	15%	15%	15%	15%
EBIT*(1-tax rate)	(465)	5,851	4,918	3,276	2,439	2,207	2,124	2,215	2,344	2,547
+ D&A	12	16	20	24	28	31	34	37	39	42
- Change in working capital	0	(661)	(185)	(143)	90	34	18	(21)	(54)	(73)
- Capex	(90)	(80)	(80)	(80)	(80)	(80)	(80)	(80)	(80)	(80)
FCFF	(543)	5,126	4,673	3,077	2,476	2,192	2,096	2,151	2,249	2,436
Terminal value										35,894
FCF + Terminal value	(543)	5,126	4,673	3,077	2,476	2,192	2,096	2,151	2,249	38,330
Present value of enterprise	29,961									
Net Debt	(1,588)									
Minorities	0									
Equity value (RMB mn)	31,549									
Equity value (HK\$ mn)	38,011									
Equity value (US\$ mn)	4,905									
Target price (HK\$)	98.07									
Terminal growth rate	3.0%									
WACC	10.0%									
Cost of Equity	3.0%									
Cost of Debt	10.0%									
Equity Beta	12.5%									
Risk Free Rate	5.0%									
Market Risk Premium	0.9									
Target Debt to Asset ratio	3.0%									
Effective Corporate Tax Rate	10.5%									

Source: CMBIS estimates

Figure 2: Sensitivity analysis (HK\$)

		WACC				
		9.0%	9.5%	10.0%	10.5%	11.0%
Terminal growth rate	4.0%	123.94	114.03	105.75	98.71	92.66
	3.5%	117.57	108.94	101.61	95.31	89.83
	3.0%	112.27	104.63	98.07	92.36	87.35
	2.5%	107.78	100.94	95.00	89.78	85.17
	2.0%	103.94	97.75	92.32	87.51	83.23

Source: Company data, CMBIS estimates

Figure 3: Key pipeline drugs of Kintor (as of 27 Aug 2021)

Drug Candidate	Target / Mechanism	Indication	Country/Region	Pre-Clinical	IND Filing (Filed)(Accepted)	Phase I	Phase II	Phase III	NDA
Proxalutamide (GT0918)	Second generation AR antagonist	COVID-19 (Outpatients)	US & Intl		Completed first patient enrolment on Apr 24, 2021				
		COVID-19 (Inpatients)	US & Intl		FDA greenlighted to conduct on May 17, 2021				
		COVID-19 (Outpatients)	Brazil & Intl		IND was approved on Jun 11, 2021				
		mCRPC	China		Expected to submit NDA in 2021				
		Combination therapy with Abiraterone for mCRPC	China		Expected to complete patients enrolment in 2021				
		mCRPC	US		Expected to complete phase II in 2021				
		Metastatic breast cancer	China						
Pyrilutamide (KX-826)	AR antagonist (for external use)	Androgenetic alopecia	China		Completed patients enrolment in Dec 2020				
		Androgenetic alopecia	US		FDA greenlighted to conduct on Jul 7, 2021				
		Acne vulgaris	China		Completed FPI on Apr 16, 2021				
		Acne vulgaris	US						
ALK-1 (GT90001)	Angiogenesis inhibitor	Combination therapy with a PD-1 for metastatic HCC (2L)	Taiwan		Interim data was released at ASCO GI in Jan 2021				
		Combination therapy with a PD-1 for metastatic HCC (2L)	US & Intl		IND was approved on Feb 18, 2021				
		HCC (1 st -line combination therapy)	China		Preparing for IND				
		Combination therapy with KN046 (PD-1/CTLA-4) for HCC, GC, GEJ adenocarcinoma, UC, ESCC	Taiwan						
Detorsertib (GT0486)	mTOR kinase inhibitor	Metastatic solid tumours	China						
GT1708F	Hedgehog/SMO inhibitor	Leukaemia	China						
		Basal-cell carcinoma	US						
GT20029	AR-PROTAC compound	AGA and acne vulgaris	China		First batch of patients were dosed on Jul 28, 2021				
		AGA and acne vulgaris	US		IND clearance was granted on Jul 8, 2021				
GT90008	PD-L1 / TGF-β dual targeting antibody	Multiple types of solid tumours	China		IND was accepted on Aug 16, 2021				
Pre-clinical	Other AR-PROTAC compounds	Multiple indications							
	c-Myc inhibitor	Blood cancer							
	ALK-1/VEGF bispecific antibody	Solid tumours							

■ Trials initiated by Kintor ■ Trials initiated by Kintor and partners

HCC = hepatocellular carcinoma, GC = gastric carcinoma, GEJ = gastroesophageal junction, UC = urothelial carcinoma, ESCC = esophageal squamous cell carcinoma.

Source: Company data, CMBIS; Notes: mCRPC = metastatic castration resistant prostate cancer, MRCT = Multi Regional Clinical Trial, HCC = hepatocellular carcinoma, BCC = basal cell carcinoma, PROTAC = proteolysis targeting chimera, ESCC = Esophageal squamous cell carcinoma, * Subject to regulators' approval

Figure 4: CMBIS estimates revision

RMB mn	New			Old			Diff (%)		
	FY21E	FY22E	FY23E	FY21E	FY22E	FY23E	FY21E	FY22E	FY23E
Revenue	0	10,132	8,351	0	10,132	8,351	N/A	0.0%	0.0%
Gross Profit	0	8,106	6,764	0	8,106	6,764	N/A	0.0%	0.0%
Operating Profit	(441)	6,949	5,927	(449)	6,934	5,912	N/A	0.2%	0.2%
Net profit	(450)	5,899	5,030	(457)	5,887	5,018	N/A	0.2%	0.3%
EPS (RMB)	(1.16)	15.22	12.98	(1.18)	15.19	12.95	N/A	0.2%	0.3%
Gross Margin	N/A	80.00%	81.00%	N/A	80.00%	81.00%	N/A	+0.00 ppt	+0.00 ppt
Operating Margin	N/A	68.58%	70.97%	N/A	68.44%	70.79%	N/A	+0.14ppt	+0.18 ppt
Net Margin	N/A	58.22%	60.23%	N/A	58.10%	60.08%	N/A	+0.12 ppt	+0.15 ppt

Source: Company data, CMBIS estimates

Figure 5: CMBIS estimates vs consensus

RMB mn	CMBIS			Consensus			Diff (%)		
	FY21E	FY22E	FY23E	FY21E	FY22E	FY23E	FY21E	FY22E	FY23E
Revenue	0	10,132	8,351	40	8,553	4,653	N/A	18%	79%
Gross Profit	0	8,106	6,764	34	7,634	4,048	N/A	6%	67%
Operating Profit	(441)	6,949	5,927	(634)	2,972	1,871	N/A	134%	217%
Net profit	(450)	5,899	5,030	(613)	5,339	2,322	N/A	10%	117%
EPS (RMB)	(1.16)	15.22	12.98	(1.57)	12.66	6.09	N/A	20%	113%
Gross Margin	N/A	80.00%	81.00%	85.00%	89.25%	87.00%	N/A	-9.25 ppt	-6.00 ppt
Operating Margin	N/A	68.58%	70.97%	-1605.06%	34.75%	40.21%	N/A	+33.83 ppt	+30.76 ppt
Net Margin	N/A	58.22%	60.23%	-1550.63%	62.42%	49.90%	N/A	-4.20 ppt	+10.33 ppt

Source: Company data, CMBIS estimates

Financial Statements

Income statement

YE 31 Dec (RMB mn)	FY19A	FY20A	FY21E	FY22E	FY23E
Revenue	0	0	0	10,132	8,351
Proxalutamide China sales - risk adjusted	0	0	0	29	183
Proxalutamide US sales - risk adjusted	0	0	0	10,058	8,038
Pyrilutamide China sales - risk adjusted	0	0	0	46	123
Pyrilutamide US sales - risk adjusted	0	0	0	0	6
ALK-1 China sales - risk adjusted	0	0	0	0	0
Others	0	0	0	0	0
Cost of sales	0	0	0	(2,026)	(1,587)
Gross profit	0	0	0	8,106	6,764
Other income	19	25	39	81	155
Selling & distribution expenses	(33)	(77)	(80)	(608)	(501)
R&D expenses	(214)	(329)	(400)	(600)	(400)
Administrative expenses	(0)	(9)	0	(30)	(92)
Other expenses	(1)	(116)	0	0	0
Operating profit (loss)	(229)	(505)	(441)	6,949	5,927
Finance costs	(4)	(3)	(9)	(9)	(9)
Pre-tax profit (loss)	(233)	(508)	(450)	6,940	5,918
Income tax	0	(0)	0	(1,041)	(888)
Minority interests	0	0	0	0	0
Attributable net profit (loss)	(233)	(508)	(450)	5,899	5,030

Cash flow summary

YE 31 Dec (RMB mn)	FY19A	FY20A	FY21E	FY22E	FY23E
Profit before tax	(233)	(508)	(450)	6,940	5,918
Depreciation and amortization, etc.	5	7	12	16	20
Change in working capital	0	(13)	0	(661)	(185)
Others	(0)	134	(1)	(1,042)	(888)
Net income tax paid	0	(0)	0	(1,041)	(888)
Operating cash flow	(228)	(381)	(439)	5,254	4,864
Purchase of PP&E	(67)	(69)	(90)	(80)	(80)
Purchase of land use right	0	0	0	0	0
Purchases of financial assets at FV through profit or loss	0	(253)	0	0	0
Purchases of financial assets measured at amortized cost	(55)	0	0	0	0
Others	115	(118)	0	0	0
Investing cash flow	(7)	(440)	(90)	(80)	(80)
Proceeds from borrowings	59	239	0	0	0
Repayments of borrowings	(65)	(79)	0	0	0
Capital contribution from equity holders	348	1,653	950	0	0
Others	(46)	(32)	0	0	0
Financing cash flow	296	1,780	950	0	0
FX changes	(3)	(91)	0	0	0
Net change in cash	61	960	421	5,174	4,784
Cash at the beginning year	138	196	1,066	1,487	6,661
Cash at the end	196	1,065	1,487	6,661	11,445

Balance sheet

YE 31 Dec (RMB mn)	FY19A	FY20A	FY21E	FY22E	FY23E
Non-current assets	333	431	510	574	634
PP&E	98	175	255	320	382
Intangible assets	179	210	210	209	209
Right-of-use assets	14	12	11	10	9
Other non-current assets	41	34	34	34	34
Current assets	221	1,421	1,840	8,261	13,086
Inventories	0	0	0	167	174
Trade receivables	0	0	0	833	915
Other receivables and prepayments	25	32	30	278	229
Financial assets at FV through P&L	0	0	0	0	0
Cash and cash equivalents	196	1,066	1,487	6,661	11,445
Restricted cash	0	0	0	0	0
Non-current liabilities	41	174	174	174	174
Borrowings	0	135	135	135	135
Lease liabilities	2	0	0	0	0
Deferred income tax liabilities	39	39	39	39	39
Current liabilities	143	169	168	754	610
Trade and other payables	80	81	80	666	522
Borrowings	59	84	84	84	84
Lease liabilities	3	3	3	3	3
Deferred income	1	0	0	0	0
Amounts due to related parties	0	1	1	1	1
Total net assets	370	1,508	2,008	7,907	12,937
Minority interest	0	0	0	0	0
Shareholders' equity	370	1,508	2,008	7,907	12,937

Key ratios

YE 31 Dec	FY19A	FY20A	FY21E	FY22E	FY23E
Sales mix (%)					
Proxalutamide China sales adjusted	0	0	0	0	2
Proxalutamide US sales	0	0	0	99	96
Pyrilutamide China sales - adjusted	0	0	0	0	1
Pyrilutamide US sales	0	0	0	0	0
ALK-1 China sales -	0	0	0	0	0
Others	0	0	0	0	0
Total	100	100	100	100	100
Profit & loss ratios (%)					
Gross margin	N/A	N/A	80	80	81
EBITDA margin	N/A	N/A	N/A	68	70
Pre-tax margin	N/A	N/A	N/A	68	71
Net margin	N/A	N/A	N/A	58	60
Effective tax rate	0	0	0	15	15
Balance sheet ratios					
Current ratio (x)	2	8	11	11	21
Net debt to equity (%)	Net cash	Net cash	Net cash	Net cash	Net cash
Returns (%)					
ROE	-63	-34	-22	75	39
ROA	-42	-27	-19	67	37
Per share value					
EPS (RMB)	N/A	(1.64)	(1.16)	15.22	12.98
DPS (RMB)	N/A	0.00	0.00	0.00	0.00
BVP (RMB)	N/A	4.87	5.18	20.40	33.38

Source: Company data, CMBIS estimates

Disclosures & Disclaimers

Analyst Certification

The research analyst who is primary responsible for the content of this research report, in whole or in part, certifies that with respect to the securities or issuer that the analyst covered in this report: (1) all of the views expressed accurately reflect his or her personal views about the subject securities or issuer; and (2) no part of his or her compensation was, is, or will be, directly or indirectly, related to the specific views expressed by that analyst in this report.

Besides, the analyst confirms that neither the analyst nor his/her associates (as defined in the code of conduct issued by The Hong Kong Securities and Futures Commission) (1) have dealt in or traded in the stock(s) covered in this research report within 30 calendar days prior to the date of issue of this report; (2) will deal in or trade in the stock(s) covered in this research report 3 business days after the date of issue of this report; (3) serve as an officer of any of the Hong Kong listed companies covered in this report; and (4) have any financial interests in the Hong Kong listed companies covered in this report.

CMBIS Ratings

BUY : Stock with potential return of over 15% over next 12 months
HOLD : Stock with potential return of +15% to -10% over next 12 months
SELL : Stock with potential loss of over 10% over next 12 months
NOT RATED : Stock is not rated by CMBIS

OUTPERFORM : Industry expected to outperform the relevant broad market benchmark over next 12 months
MARKET-PERFORM : Industry expected to perform in-line with the relevant broad market benchmark over next 12 months
UNDERPERFORM : Industry expected to underperform the relevant broad market benchmark over next 12 months

CMB International Securities Limited

Address: 45/F, Champion Tower, 3 Garden Road, Hong Kong, Tel: (852) 3900 0888 Fax: (852) 3900 0800

CMB International Securities Limited ("CMBIS") is a wholly owned subsidiary of CMB International Capital Corporation Limited (a wholly owned subsidiary of China Merchants Bank)

Important Disclosures

There are risks involved in transacting in any securities. The information contained in this report may not be suitable for the purposes of all investors. CMBIS does not provide individually tailored investment advice. This report has been prepared without regard to the individual investment objectives, financial position or special requirements. Past performance has no indication of future performance, and actual events may differ materially from that which is contained in the report. The value of, and returns from, any investments are uncertain and are not guaranteed and may fluctuate as a result of their dependence on the performance of underlying assets or other variable market factors. CMBIS recommends that investors should independently evaluate particular investments and strategies, and encourages investors to consult with a professional financial advisor in order to make their own investment decisions.

This report or any information contained herein, have been prepared by the CMBIS, solely for the purpose of supplying information to the clients of CMBIS or its affiliate(s) to whom it is distributed. This report is not and should not be construed as an offer or solicitation to buy or sell any security or any interest in securities or enter into any transaction. Neither CMBIS nor any of its affiliates, shareholders, agents, consultants, directors, officers or employees shall be liable for any loss, damage or expense whatsoever, whether direct or consequential, incurred in relying on the information contained in this report. Anyone making use of the information contained in this report does so entirely at their own risk.

The information and contents contained in this report are based on the analyses and interpretations of information believed to be publicly available and reliable. CMBIS has exerted every effort in its capacity to ensure, but not to guarantee, their accuracy, completeness, timeliness or correctness. CMBIS provides the information, advices and forecasts on an "AS IS" basis. The information and contents are subject to change without notice. CMBIS may issue other publications having information and/ or conclusions different from this report. These publications reflect different assumption, point-of-view and analytical methods when compiling. CMBIS may make investment decisions or take proprietary positions that are inconsistent with the recommendations or views in this report.

CMBIS may have a position, make markets or act as principal or engage in transactions in securities of companies referred to in this report for itself and/or on behalf of its clients from time to time. Investors should assume that CMBIS does or seeks to have investment banking or other business relationships with the companies in this report. As a result, recipients should be aware that CMBIS may have a conflict of interest that could affect the objectivity of this report and CMBIS will not assume any responsibility in respect thereof. This report is for the use of intended recipients only and this publication, may not be reproduced, reprinted, sold, redistributed or published in whole or in part for any purpose without prior written consent of CMBIS.

Additional information on recommended securities is available upon request.

For recipients of this document in the United Kingdom

This report has been provided only to persons (I) falling within Article 19(5) of the Financial Services and Markets Act 2000 (Financial Promotion) Order 2005 (as amended from time to time) ("The Order") or (II) are persons falling within Article 49(2) (a) to (d) ("High Net Worth Companies, Unincorporated Associations, etc.") of the Order, and may not be provided to any other person without the prior written consent of CMBIS.

For recipients of this document in the United States

CMBIS is not a registered broker-dealer in the United States. As a result, CMBIS is not subject to U.S. rules regarding the preparation of research reports and the independence of research analysts. The research analyst who is primary responsible for the content of this research report is not registered or qualified as a research analyst with the Financial Industry Regulatory Authority ("FINRA"). The analyst is not subject to applicable restrictions under FINRA Rules intended to ensure that the analyst is not affected by potential conflicts of interest that could bear upon the reliability of the research report. This report is intended for distribution in the United States solely to "major US institutional investors", as defined in Rule 15a-6 under the US, Securities Exchange Act of 1934, as amended, and may not be furnished to any other person in the United States. Each major US institutional investor that receives a copy of this report by its acceptance hereof represents and agrees that it shall not distribute or provide this report to any other person. Any U.S. recipient of this report wishing to effect any transaction to buy or sell securities based on the information provided in this report should do so only through a U.S.-registered broker-dealer.

For recipients of this document in Singapore

This report is distributed in Singapore by CMBI (Singapore) Pte. Limited (CMBISG) (Company Regn. No. 201731928D), an Exempt Financial Adviser as defined in the Financial Advisers Act (Cap. 110) of Singapore and regulated by the Monetary Authority of Singapore. CMBISG may distribute reports produced by its respective foreign entities, affiliates or other foreign research houses pursuant to an arrangement under Regulation 32C of the Financial Advisers Regulations. Where the report is distributed in Singapore to a person who is not an Accredited Investor, Expert Investor or an Institutional Investor, as defined in the Securities and Futures Act (Cap. 289) of Singapore, CMBISG accepts legal responsibility for the contents of the report to such persons only to the extent required by law. Singapore recipients should contact CMBISG at +65 6350 4400 for matters arising from, or in connection with the report.