## Quoin Pharmaceuticals Provides Corporate Update and Announces Third Quarter 2023 Financial Results

November 8, 2023
Company reported positive clinical data for QRX003 from first six evaluable subjects in ongoing open-label Netherton Syndrome study
Positive benefits were observed across a number of clinical endpoints including pruritus, Investigator skin scoring system and patient global assessment

No safety concerns have been observed to date in either ongoing clinical study
Quoin expects its cash runway will extend through the end of 2024
ASHBURN, Va., Nov. 08, 2023 (GLOBE NEWSWIRE) -- Quoin Pharmaceuticals Ltd. (NASDAQ: QNRX) (the "Company" or "Quoin"), a clinical stage, specialty pharmaceutical company focused on rare and orphan diseases, today provides a business update and announces financial results for the quarter ended September 30, 2023.

Quoin CEO, Dr. Michael Myers, said, "While still early-stage, we are extremely excited by the positive clinical data generated to date across a number of endpoints from our ongoing open-label clinical study in Netherton Syndrome patients. We are particularly pleased with the pruritus results observed for five of the six subjects evaluated. Given the overall strong efficacy data and the exemplary safety profile demonstrated thus far, we are moving into an optimization phase for both of our Netherton Syndrome studies, which we believe could lead to an even more robust clinical data set."

## Corporate Highlights -

- On October $24^{\text {th }}$, Quoin announced positive clinical data from the first six evaluable patients in the company's open-label clinical trial in Netherton Syndrome patients.
- Five of the six subjects evaluated had negligible or absent pruritus, or itch, following treatment with QRX003, a significant improvement from prior to the study.
- All six subjects exhibited improvement in the Investigator assessed skin scoring system with three subjects showing improvement throughout the study and the other three at various points during the study.
- All six subjects expressed a favorable impression of QRX003 across multiple assessed metrics.
- No safety concerns have been reported to date for any subject in either of Quoin's studies.
- Company plans to further optimize its clinical program by making a number of protocol amendments including: eliminating the lower $2 \%$ dose in the double blinded study, changing the dosing frequency to twice-daily from once-daily and increasing the number of subjects in both studies.
- Company secured long term, exclusive supply of the only fully GMP grade active ingredient in QRX003.
- On September $6^{\text {th }}$, Quoin signed its ninth commercial agreement for QRX003, increasing the number of partnered countries to sixty-one.


## Financial Highlights

- Quoin had approximately $\$ 14.0$ million in cash, cash equivalents and marketable securities as of September 30, 2023.
- Net loss for the quarter ended September 30, 2023 was approximately $\$ 2.1$ million compared to approximately $\$ 2.3$ million for the quarter ended September 30, 2022, and net loss for the nine months ended September 30, 2023 was $\$ 7.2$ million compared to $\$ 7.2$ million for the nine months ended September 30, 2022.
- Investors are encouraged to read the Company's Quarterly Report on Form 10-Q when it is filed with the Securities and Exchange Commission (the "SEC"), which will contain additional details about Quoin's financial results as of and for the period ended September 30, 2023.

Quoin will host a conference call and webcast at 8:30am ET on Thursday, November 9, 2023. The call will include a discussion of third quarter 2023 financial results and a corporate update. The live call can be accessed by dialing 1-800-603-0527 (domestic) or 1-412-317-0688 (international). The
call webcast will be available at: https://event.choruscall.com/mediaframe/webcast.html?webcastid=sPhSaNGJ.

## About Quoin Pharmaceuticals Ltd.

Quoin Pharmaceuticals Ltd. is a clinical stage specialty pharmaceutical company focused on developing and commercializing therapeutic products that treat rare and orphan diseases. We are committed to addressing unmet medical needs for patients, their families, communities and care teams. Quoin's innovative pipeline comprises four products in development that collectively have the potential to target a broad number of rare and orphan indications, including Netherton Syndrome, Peeling Skin Syndrome, Palmoplantar Keratoderma, Scleroderma, Epidermolysis Bullosa and others. For more information, visit: www.quoinpharma.com or Linkedln for updates.

## Cautionary Note Regarding Forward Looking Statements

The Company cautions that statements in this press release that are not a description of historical facts are forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Forward-looking statements may be identified by the use of words referencing future events or circumstances such as "expect," "intend," "plan," "anticipate," "believe," and "will," among others. Because such statements are subject to risks and uncertainties, actual results may differ materially from those expressed or implied by such forward-looking statements. These forwardlooking statements are based upon the Company's current expectations and involve assumptions that may never materialize or may prove to be incorrect. Actual results and the timing of events could differ materially from those anticipated in such forward-looking statements as a result of various risks and uncertainties. More detailed information about the risks and uncertainties affecting the Company is contained under the heading "Risk Factors" included in the Company's Annual Report on Form 10-K for the year ended December 31, 2022 that the Company filed with the SEC. One should not place undue reliance on these forward-looking statements, which speak only as of the date on which they were made. The Company undertakes no obligation to update such statements to reflect events that occur or circumstances that exist after the date on which they were made, except as may be required by law.

## For further information:

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## -Tables Follow-

## QUOIN PHARMACEUTICALS LTD.

## Consolidated Balance Sheets

|  | $\begin{gathered} \text { September 30, } \\ 2023 \end{gathered}$ |  | $\begin{gathered} \text { December 31, } \\ 2022 \end{gathered}$ |  |
| :---: | :---: | :---: | :---: | :---: |
|  | (Unaudited) |  |  |  |
| ASSETS |  |  |  |  |
| Current assets: |  |  |  |  |
| Cash and cash equivalents | \$ | 3,163,426 | \$ | 2,860,628 |
| Investments |  | 10,818,051 |  | 9,992,900 |
| Prepaid expenses |  | 159,851 |  | 516,584 |
| Total current assets |  | 14,141,328 |  | 13,370,112 |
| Prepaid expenses - long term |  | 300,000 |  | 383,390 |
| Intangible assets, net |  | 626,529 |  | 704,561 |
| Total assets | \$ | 15,067,857 | \$ | 14,458,063 |
| LIABILITIES AND SHAREHOLDERS' EQUITY |  |  |  |  |
| Current liabilities: |  |  |  |  |
| Accounts payable | \$ | 239,978 | \$ | 605,600 |
| Accrued expenses |  | 2,594,199 |  | 1,175,705 |
| Accrued interest and financing expense |  | 1,146,251 |  | 1,146,251 |
| Due to officers - short term |  | 600,000 |  | 600,000 |
| Total current liabilities |  | 4,580,428 |  | 3,527,556 |
| Due to officers - long term |  | 3,073,733 |  | 3,523,733 |
| Total liabilities | \$ | 7,654,161 | \$ | 7,051,289 |

Commitments and contingencies

Shareholders' equity:
Ordinary shares, no par value per share, $8,333,334$ ordinary shares
authorized - 987,220 ( 987,220 ADS's) ordinary shares issued and outstanding at

Treasury stock, 45 ordinary shares
Additional paid in capital
Accumulated deficit
Total shareholders' equity

Total liabilities and shareholders' equity

## QUOIN PHARMACEUTICALS LTD

## Statements of Operations (Unaudited)

|  | Nine months ended September 30, |  |  |  | Three months ended September 30, |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 2023 |  | 2022 |  | 2023 |  | 2022 |  |
| Operating expenses |  |  |  |  |  |  |  |  |
| General and administrative | \$ | 4,685,241 | \$ | 5,112,002 | \$ | 1,366,464 | \$ | 1,582,059 |
| Research and development |  | 2,475,596 |  | 2,059,769 |  | 758,759 |  | 745,506 |
| Total operating expenses |  | 7,160,837 |  | 7,171,771 |  | 2,125,223 |  | 2,327,565 |
| Other (income) and expenses |  |  |  |  |  |  |  |  |
| Forgiveness of accounts payable |  | - |  | $(416,000)$ |  | - |  | - |
| Warrant liability (income) expense |  | - |  | $(77,237)$ |  | - |  | - |
| Unrealized loss (gain) |  | 11,926 |  | 3,053 |  | $(2,119)$ |  | 3,053 |
| Interest income |  | $(536,068)$ |  | $(15,132)$ |  | $(196,425)$ |  | $(15,132)$ |
| Interest and financing expense |  | - |  | 714,081 |  | - |  | 714,081 |
| Total other (income) expense |  | $(524,142)$ |  | 208,765 |  | $(198,544)$ |  | 702,002 |
| Net loss | \$ | $(6,636,695)$ | \$ | $(7,380,536)$ | \$ | $(1,926,679)$ | \$ | $(3,029,567)$ |
| Deemed dividend on warrant modification |  | - |  | $(65,266)$ |  | - |  | $(65,266)$ |
| Net loss attributable to shareholders | \$ | $(6,636,695)$ | \$ | $(7,445,802)$ | \$ | $(1,926,679)$ | \$ | $(3,094,833)$ |

Loss per ADS
Loss per ADS

## Basic

Fully-diluted

Weighted average number of ADS's outstanding Basic
Fully-diluted
(7.61) $\$ \quad(55.79) \quad \$ \quad$ (1.95) \$
(7.61) $\$ \quad$ (55.79) \$ (1.95) \$

| 871,835 | 133,450 | 987,220 | 274,317 |
| :--- | :--- | :--- | :--- |
| 871,835 | 133,450 | 987,220 | 274,317 |

