

BuyersStrike!

It Was Dark Over Westphalia

Christmas Week Quick Take: What Misleading BS is Cytodyn Spewing Today? (CYDY)

What nonsense is everyone's favorite reverse-merger pink sheet Coronacraper, **Cytodyn (CYDY)**, spewing in this holiday-shortened, and retail trader dominated week? Today, the 22nd of December, comes a [press release](https://www.cytodyn.com/investors/news-events/press-releases/detail/492/fda-resumes-eind-approval-for-severe-to-critical-covid-19) (<https://www.cytodyn.com/investors/news-events/press-releases/detail/492/fda-resumes-eind-approval-for-severe-to-critical-covid-19>) with a clearly misleading headline.

Notice the use of the word "Approval", which has a special meaning with regards to the FDA. INDs are never "approved", they are either rejected or accepted. EINDs (Emergency INDs) are never approved, they are "authorized". The use of "approval" in this context is **misleading**, as [loserlimab](https://buyersstrike.wordpress.com/2020/08/28/but-teh-science-be-so-gud-part-1-a-brief-history-of-loserlimab-cydy/) (<https://buyersstrike.wordpress.com/2020/08/28/but-teh-science-be-so-gud-part-1-a-brief-history-of-loserlimab-cydy/>) has not been approved for anything by the FDA.

FDA Resumes eIND Approval for Severe-to-Critical COVID-19 Patients Use of Vyrologix™ (Ieronlimab) Following Full Enrollment in CytoDyn's Phase 3 Trial

December 22, 2020 6:00am EST

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FDA's decision will enable CytoDyn to respond to ongoing requests for Ieronlimab until Phase 3 trial data is unblinded

(<https://buyersstrike.files.wordpress.com/2020/12/eind-approvalpr.png>)

You can read more about EINDs on the [FDA's web site](https://www.fda.gov/drugs/investigational-new-drug-ind-application/physicians-how-request-single-patient-expanded-access-compassionate-use) (<https://www.fda.gov/drugs/investigational-new-drug-ind-application/physicians-how-request-single-patient-expanded-access-compassionate-use>). What is important to note is that having an EIND accepted is neither noteworthy, nor newsworthy. It is a ho-hum, run-of-the-mill, everyday occurrence:

Upon receipt of the single patient expanded access application, FDA will review to make a decision. The FDA will either allow the treatment to proceed or place the IND on hold.

FDA allows over 99% of single patient expanded access requests to proceed. FDA may contact the physician to request more information or clarification in order to avoid placing the IND on hold.

(<https://buyersstrike.files.wordpress.com/2020/12/fda-eind-acceptance.jpg>)

What is newsworthy, however, is the Form 4 filed last night by **Cytodyn's Klown Krew CFO Mike Mulholland**, detailing the sale of millions of dollars worth of cheap stock. See the filing [here](https://www.cytodyn.com/investors/sec-filings/all-sec-filings/content/0001807094-20-000030/primary_doc.html?). (https://www.cytodyn.com/investors/sec-filings/all-sec-filings/content/0001807094-20-000030/primary_doc.html?)

Common Stock	Date	Code	Quantity	Type	Price
Common Stock	12/17/2020	M	32,000	A	\$ 0.39
Common Stock	12/17/2020	S(1)	32,000	D	\$ 4,352.3(2)
Common Stock	12/18/2020	M	155,500	A	\$ 0.39
Common Stock	12/18/2020	M	233,100	A	\$ 0.49
Common Stock	12/18/2020	M	98,402	A	\$ 0.57
Common Stock	12/18/2020	S	487,002	D	\$ 4,951.6(3)
Common Stock	12/21/2020	M	201,598	A	\$ 0.57
Common Stock	12/21/2020	M	300,000	A	\$ 0.80
Common Stock	12/21/2020	M	88,199	A	\$ 0.87
Common Stock	12/21/2020	S	585,797	D	\$ 5,582.4(4)

(<https://buyersstrike.files.wordpress.com/2020/12/mulholland4.png>)

Cytodyn is obviously hoping the press release, and curiously timed companion puff piece [article \(https://www.zerohedge.com/news/2020-12-22/fda-should-restore-doctors-rights-use-medicines-work\)](https://www.zerohedge.com/news/2020-12-22/fda-should-restore-doctors-rights-use-medicines-work) by a tout (who happens to be affiliated with a paid promoter called "Streetwise Reports"), draws attention far far away from it.

[Did you know Cytodyn is being sued by a group of former directors? What to know why? Start [here \(https://buyersstrike.wordpress.com/2020/12/18/update-update-whos-suing-cytodyn-now-and-who-is-admitting-the-plaintiffs-are-right-cydy/\)](https://buyersstrike.wordpress.com/2020/12/18/update-update-whos-suing-cytodyn-now-and-who-is-admitting-the-plaintiffs-are-right-cydy/) and [here \(https://buyersstrike.wordpress.com/2020/12/20/sunday-funday-what-else-doesnt-cytodyn-want-investors-to-learn-cydy/\)](https://buyersstrike.wordpress.com/2020/12/20/sunday-funday-what-else-doesnt-cytodyn-want-investors-to-learn-cydy/).]

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Posted in [Bad Directors](#), [Bio-Dreck](#), [Bucket Shops](#), [CoronaCrap](#), [Reverse Mergers](#) on [December 22, 2020](#) by [BuyersStrike!](#) [3 Comments](#)

3 comments

1. **Troy M** says:

[December 22, 2020 at 10:46 pm](#)

Lmao. The author of this is an idiot.

[Can you point out a single inaccuracy Troy? Didn't think so. – Editor]

[REPLY](#)

2. **jbperez808** says:

[December 23, 2020 at 9:11 am](#)

"THIS COMMENTARY IN NO WAY CONSTITUTES INVESTMENT ADVICE, AND SHOULD NEVER BE RELIED ON IN MAKING AN INVESTMENT DECISION, EVER."

'NUFF SAID 🤔🤔🤔

[Glad you are enjoying it. – Editor]

[REPLY](#)

3. Pingback: [Christmas Eve Quick Take: Who clearly has no clue what an Open Label Extension is? \(CYDY\) | BuyersStrike!](#)

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