

### COMPANY UPDATE

#### **Rating: BUY**

Ticker: NRXP

Price: \$1.31

Target: \$45 (from \$44)

## NRx Pharmaceuticals, Inc.

Q3 EPS upside. Major opportunities in NRX-100 (IV ketamine) and NRX-101, with both NDAs expected soon. Raising P/T to \$45.

Q3 EPS upside: NRx recently (on November 14) reported its Q3 2024 (ending September) results. Net loss was \$1.6 million or EPS of (0.15) compared with our and consensus estimates of (0.66) - (0.73). There was no company guidance. NRx is a clinical stage drug development company so it generates no revenue.

**Operating expenses:** Operating expenses were \$3.0 million, down from Q2's \$7.1 million on much lower R&D and G&A costs.

No guidance: Management did not provide forward guidance.

Raising 2025 estimates: We are adjusting our 2024 EPS estimate to \$(1.96) from \$(2.84). We have raised our 2025 estimates for strong revenues and slight profitability.

**Focused on NRX-100 and NRX-101:** Its main drugs are NRX-100 (IV Ketamine) and NRX-101 (D-cylcoserine/Lurasidone) for the treatment of bipolar depression in patients with suicidality (the risk of suicide). The company plans to file NDA (New Drug Application) with the FDA for NRX-100 and NRX-101 by year end 2024.

NRX-101: NRX-101 is a dual-targeted therapy regimen consisting of an initial treatment with NRX-100 (intravenous ketamine) followed by 6-week treatment with NRX-101 (combined DCS and lurasidone). Ketamine is a generic drug and has been widely used for a long time as an antidepressant, although its effect does not last long (usually about a week). NRX-101 is designed to extend ketamine's proven antisuicidal and antidepressant benefits without its drawbacks.

Large market potential: There is no medicine approved to treat patients with bipolar depression suffering suicidal ideation. According to the NIH, an estimated 2.8% of the U.S. adult population had bipolar disorder in the past 12 months, and the lifetime prevalence is 4.4% of adults in the U.S. Lifetime suicide behavior occurs in 25% to 56% of people with bipolar depression.

**Positive clinical trials data:** In May, the company announced final positive clinical trials data in its Phase 2b/3 Clinical Trial of NRX-101 vs. Lurasidone for Treatment of Suicidal Bipolar Depression. The company believes that the findings when combined with the prior STABIL-B trial, demonstrate a basis for seeking accelerated drug approval of NRX-101 based on improved safety related to akathisia and suicidality in the setting of comparable antidepressant efficacy.

**NRX-100:** NRX-100 is Intravenous (IV) ketamine which has now become a standard of care for acute treatment of suicidal depression, in the absence of an FDA-labeled product. The company is developing NRX-100 (intravenous ketamine) as a labeled drug to treat acute depression and suicidality.

**HOPE Therapeutics spinoff:** The company plans to file a NDA for NRX-100 in 2024, with potential for revenue in 2025. The company plans to partially spin off HOPE Therapeutics (the first nationwide network of Interventional Psychiatry Clinics focused on suicidal depression and PTSD) to shareholders. HOPE Therapeutics is expected to be self-funded and profitable in its first year.

Signed LOI to acquire two precision psychiatry centers for HOPE Therapeutics: The company is in process to finalize the acquisitions for 2 psychiatry centers. It is also in negotiation with four additional psychiatry centers with the goal of assembling a network delivering \$25 million in ketamine-based precision psychiatric treatment for suicidal depression by year-end 2024.

**Balance sheet:** As of Q3, the company has \$2 million in cash and \$6 million in debt. In October (current Q4), NRx received \$5 million from its  $2^{nd}$  tranche financing.

**New CEO and CFO:** In October, Stephen Willard resigned as CEO and Jonathan Javitt, Chairman and co-founder and was previously CEO, was appointed Interim CEO while a search is on for a permanent CEO. In November, Michael Abrams was appointed as its permanent Chief Financial Officer succeeding Interim-CFO Richard Narido.

**Current valuation attractive:** We are maintaining our BUY rating, but raising our 12-month price target to \$45 from \$44 based on a NPV analysis. This represents significant upside from the current share price and we believe this valuation appropriately balances out the high risks with large upside opportunities.

#### **Company Description**

NRx Pharmaceuticals, based in Wilmington, DE, is a clinical stage biopharmaceutical company developing drugs to treat mental health disorders.

United States Healthcare

December 2, 2024

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#### Stock Data

Exchange:	NasdaqCM
52-week Range:	1.10 - 7.33
Shares Outstanding (million):	12
Market cap (\$million):	\$16
EV (\$million):	\$20
Debt (\$million):	\$6
Cash (\$million):	\$2
Avg. Daily Trading Vol. (\$million):	\$0.2
Float (million shares):	8
Short Interest (million shares):	0.7
Dividend, annual (yield):	\$0 (NA%)

#### Revenues (US\$ million)

	2024E (Cur.)	2024E (Old)	2025E (Cur.)	2025E (Old)
Q1 Mar	0A		5E	0E
Q2 Jun	0A		10E	0E
Q3 Sep	0A	0E	20E	0E
Q4 Dec	<u>0E</u>		<u>25E</u>	<u>0E</u>
Total	0E		60E	0E
EV/Revs	N/A		0.3x	

#### Earnings per Share (pro forma)

	2024E	2024E	2025E	2025E
	(Cur.)	(Old)	(Cur.)	(Old)
Q1 Mar	(0.74)A		(0.40)E	(0.60)E
Q2 Jun	(0.75)A		(0.22)E	(0.60)E
Q3 Sep	(0.15)A	(0.66)E	0.25E	(0.59)E
Q4 Dec	(0.39)E	(0.69)E	<u>0.37E</u>	(0.59)E
Total	(1.96)E	(2.84)E	0.02E	(2.39)E
P/E	N/A		66x	

<sup>\*</sup>Reflects a 1:10 reverse stock split in April 2024.

#### Important Disclosures

Ascendiant Capital Markets LLC seeks to do business with companies covered by its research team. Consequently, investors should be aware that the firm may have a conflict of interest that could affect the objectivity of this report. Investors should consider this report as only a single factor in making an investment decision.

For analyst certification and other important disclosures, refer to the Disclosure Section, located at the end of this report, beginning on page 20.



Exhibit 1: NRx Pharmaceuticals, Inc. Corporate Overview



# NRx Pharmaceuticals, Inc.

# Recent clinical developments enable FDA filing in 2024 with Potential for 2025 Commercial Revenue



NASDAQ: NRXP

- New Drug Application planned for NRX-100 (IV Ketamine) in Suicidal Depression
   Potential for \$100 million+ revenue in 2025 with peak in excess of \$1 billion
- Accelerated Approval application planned for NRX-101 in Bipolar Depression with Suicidality or Akathisia

Enabled by Prior Breakthrough Therapy designation for NRX-101

o Launch of HOPE Therapeutics with Expected 2024 Revenue; NRXP majority owner

Initial \$100 million mental health clinic rollup; independent funding via bond issuance; NRXP shareholders to receive share distribution

FUTURE YEARS

 Path to 2026 approval as first antibiotic to treat cUTI without risk of C. Difficile infection based on QIDP and Fast Track awards Requires Phase 2/3 Trial



**Exhibit 2: NRx's Investment Summary** 

# NRx Clinical Programs – 2025 Path to Profitability



# NRX-100 (IV Ketamine)

- Fast Track NDA in 2024
- Expected PDUFA 2025
- Approvable efficacy data from four studies in hand
- Manufacturing data has reached 12 month stability
- Six-month Ketamine tox data
- Alignment on pediatric plan with FDA
- \$2+ billion addressable market



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ipolar

## NRX-101 (Oral DCS/Lurasidone)

- Statistically-significant Phase 2 data: first antidepressant to decrease suicidality and/or akathisia
- NDA/Accelerated
  Approval for bipolar
  patients with akathisia in
  2H24
- Focused initial launch
- 7 million patient broad bipolar addressable market
- \$2 billion market potential



nterventional Psychiatry

# HOPE Therapeutics

- A network of Interventional Psychiatry Clinics focused on ketamine and other lifesaving interventions for Suicidal Depression and PTSD
- Targeting \$100 million revenue by year-end 2025
- Acquisition Financing in on is committed
- First acquisitions in 2024
- Expected in profit in 2025



Exhibit 3: NRX-101

# NRX-101: Oral medication with potential for 2025 NDA filing First oral antidepressant shown to reduce Suicidality & Akathisia

- Current efficacy and safety data support filing an NDA for Accelerated Approval in the narrow indication of patients with suicidal bipolar depression and akathisia
- Market potential for NRX-101 for suicidal bipolar depression and akathisia is well in excess of \$2 Billion
- 3 Narrow initial indication allows focused launch by NRx alone
- Additional phase 3 trial vs. placebo needed for the broad 7 million person bipolar market; planned to be financed by a partner or new investors
- NRX-101 Phase 3 investment is not part of use of current capital

# NRX-101™ For Suicidal Treatment-Resistant Bipolar Depression

First oral medicine in development for Suicidal Treatment-Resistant Bipolar Depression

- Non-addictive
- Non-neurotoxic
- Non-hallucinogenic

NRX-101 blocks the psychedelic effects of NMDA antagonists with evidence that the antidepressant and anti-suicidal properties can be preserved



Exhibit 4: NRx's Product Pipeline (as of July 2024)

# **Our Pipeline**

Leveraging our	Multi-Billion [	Dollar NMDA	Platform		
Product	Phase 1	Phase 2	Phase 3	NDA	Status
<u>Suicidal Depression</u>					
HTX-100 (IV Ketamine)	Suicidal Depressi	on			NDA Filing 2H24   PDUFA est. 2025
	*Collaboration Agree well-powered, acad	ement with Study Leade emic clinical trials	rship of two		
NRX-101™	Suicidal, Treatment- Suicidality	Resistant Bipolar Depre	ssion with Akathisia or		Filing NDA for Accelerated Approval 2H24; est PDUFA 2025
	Bipolar Depression				Phase 3 confirmatory trial, post approval
Chronic Pain					200 person, independent trial funded
D-Cycloserine (DCS)	Chronic Back Pair	1			by DOD Pending Data Readout
NRX-101™	Chronic Nocicept	ive Pain	$\supset$		IND Approved   Applied to NIH EPICNET & HEAL
PTSD			$\neg$		Nonclinical Evidence   Clinical
NRX-101™	PTSD		_>		Planning
Complicated UTI NRX-101™	Complicated UTI i	ncl. Pyelonephritis	$\supset$		In Vitro Data   QIDP and Fast Track granted



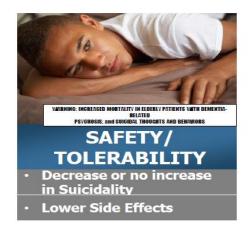
Exhibit 5: Targeting Suicidal Bipolar Depression Risks

### Why target Suicidal Bipolar Depression?

Suicide kills ~50,000 Americans annually\* - suicide is particularly high in bipolar disorder

#### **Selected Unmet Needs for New Antidepressants**





Source: Company reports.

**Exhibit 6: Bipolar Depression Suicide Market Opportunities** 

#### NRX-101 Market Opportunity in Bipolar Depression

Patients in clinics and outpatient being treated for Bipolar Depression with Suicidality

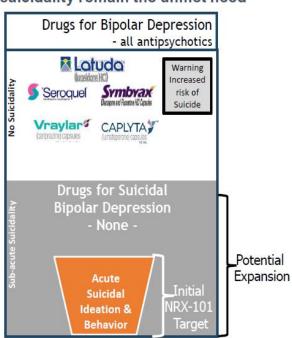




#### **Exhibit 7: Unmet Need for Bipolar Depression Suicidality**

Though numerous drugs have been approved for MDD and Bipolar Depression, faster, more robust response, and reduction of suicidality remain the unmet need





Source: Company reports.

#### **Exhibit 8: Science of Depression and Suicidality**

The Emerging Science of Depression and Suicidality

Depression and Suicidality – though overlapping is not the same

#### **Depression with Suicidality**

- Antidepressants (5HT2a / SSRIs) can increase suicidality - suicidality routinely an exclusion in depression studies
- NMDA antagonists (ketamine) can stabilize depression and suicidality
  - Suicidality improvement not strictly a function of improvements in depression
  - Ketamine can create hallucinations, may be highly addictive, requires supervised administration

#### Implications for Bipolar Depression with Suicidality

- Highest suicidality of depressive disorders ~ 50% attempt suicide
- Available drugs improve depression but can increase suicidality
- Drug abuse and overdose of great concern – addictive agents may require REMS

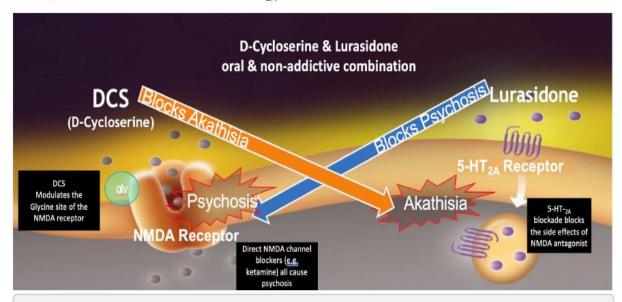
Development of Depression drugs has mostly avoided addressing Suicidality



#### **Exhibit 9: NRx Discovery**

### The NRx Discovery

Simultaneous Blockade of NMDA and 5-HT<sub>2A</sub>



### D-Cycloserine acts as an NMDA antagonist above certain dosages

Studies have shown that DCS + 5HT2a increases the antidepressant response and reduces suicidality

### **Understanding the NMDA Receptor**

The NMDA receptor is an ION Channel on the surface of Brain Cells

At high levels of NMDA activity (i.e. the channel is wide open) thoughts are slowed substantially, patients ruminate on negative, frequently suicidal thoughts. Brain cells stop making new connections to neighboring cells.

NMDA antagonists decrease symptoms of depression.

NMDA antagonists block the akathisia caused by SSRI antidepressants in nonclinical studies.

NMDA antagonists "rewire" the brain by stimulating new connections between brain cells.

#### NMDA RECEPTOR REGULATES SPEED OF THOUGHTS

TOO FAST and thoughts race uncontrollably (mania)
TOO SLOW and negative, self-destructive thoughts drive suicide





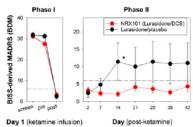
#### Exhibit 10: Phase 2 Study of D-Cycloserine in Depression / Suicidality Conclusions

Phase 2 Success: STABIL-B trial Showed Superiority of NRX-101 vs Lurasidone in Reducing Depression (primary endpoint) *after* 

**Ketamine Pre-treatment** 

Patients received one infusion of IV ketamine vs. placebo. Responders were randomized to NRX-101 vs lurasidone, a Standard of Care

- Mean 7.7 point benefit on MADRS (Primary Endpoint, P=.03) through day 42 vs. lurasidone.
- 40% relapse in control group, no relapse in NRX-101 group (P=.07)
- 1.5 point advantage vs SoC on Columbia Suicide Severity Rating Scale (C-SSRS) (P=.02)
- Decreased akathisia in the NRX-101 group on the BARS akathisia scale (P=.14)



	Effic	acy Measu	ıres: Repeat	ed Measu	res Mixed Mo	del LS Mea	n Difference	s
		Through	Day 28			Through	h Day 42	
	LOCF	No	LOCE	yes	LOCF	No	LOCF	yes
MADRS Depression	LS Mean A	p-value	LS Mean ∆	p-value	LS Mean Δ	p-value	LS Mean Δ	p-value
Score	-4.0	0.09	-7.7	0.03	-3.7	0.04	-7.7	0.04
Suicidality Rating	LS Mean A	p-value	LS Mean ∆	p-value	LS Mean ∆	p-value	LS Mean Δ	p-value
Scale C-SSRS	-0.5	NS	-1.3	0.04	-0.6	NS	-1.5	0.02
Clinical Global	LS Mean A	p-value	LS Mean ∆	p-value	LS Mean ∆	p-value	LS Mean Δ	p-value
Impression CGI-SS	-0.4	NS	-2.9	0.05	-0.6	NS	-2.9	0.02

Source: Company reports.

#### Exhibit 11: NRX-101 Phase 2b/3 Clinical Trial Program (SSIB & ASIB) Conclusions

NRX-101 demonstrates reduced Akathisia and Time to Suicidality Remission in Suicidal Bipolar Depression: *No Ketamine Pre-treatment* 

Phase 2b/3, randomized, double blind trial on NRX-101 vs Standard of Care (SoC) (lurasidone) in Suicidal Treatment Resistant Bipolar Depression (S-TRBD), n=74

- Similar (50% reduction) in depression vs. SoC
- Significant reduction in akathisia vs. SoC, p=0.03
- Time to Sustained Remission from Suicidality (C-SSRS ≤3)
   vs. SoC, p=0.05
  - We believe an antidepressant with Standard of Care level efficacy and a significant reduction in akathisia / suicidality vs SoC will become the new standard in bipolar depression
  - We believe NRX-101 can be that medication





#### Exhibit 12: NRX-101 Advantages and Objectives

# NRX-101 offers a differentiated profile for Suicidal Bipolar Depression with an FDA agreed upon path to NDA

Phase 3 with FDA Breakthrough Therapy designation

#### NMDA - A Validated Mechanism

- Depression & Suicidality
- · Esketamine, NRX-101 Phase 2, etc.

#### **Addresses High Unmet Need**

- · Treats depression and suicidality (bipolar space)
- · Oral, not addictive (not scheduled), avoids hallucinations
- Outpatient

#### **Composition of Matter Patent**

- NRx has a composition of matter patent for NRX-101 and an array of NMDA+5HT2A compounds,
- · Five patent families, 60+ applications, 30+ issued patents

#### FDA Agreed Upon Regulatory Path

 Special Protocol Agreement, Breakthrough Therapy designation: Treatment of Severe Bipolar Depression in Patients with ASIB after initial stabilization with ketamine or other effective therapy

#### Efficient Clinical Development Path to NDA

- · Seeking to replicate P2 study
- NRX-100 (144 pts.) NRX-101 (~80 pts.) pivotal study (severely depressed and acutely suicidal) to start 2H22
- Path to NDA filing in 2023

Path to ND/

Exploring expansion in earlier population

 NRX-101 Phase 2 trial (Bipolar Depression in sub-acute suicidality) initiated 2Q 2022

Source: Company reports.

#### **Exhibit 13: NRX-101 Market Opportunities**

#### Potential to Reach 75 Million Lives

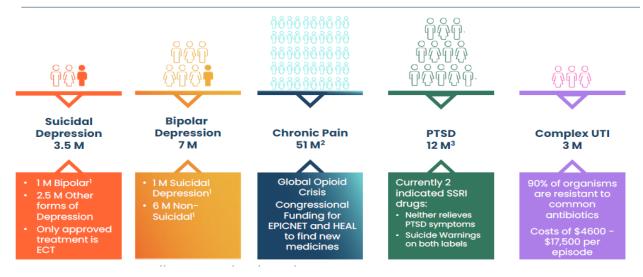
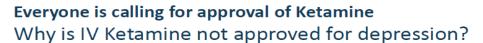




Exhibit 14: NRX-100 (IV Ketamine)

# NRX-100 (IV Ketamine) for Suicidal Depression

Aiming to be the first FDA-approved ketamine product to treat suicidal depression



- No Company has applied for FDA approval of IV Ketamine to treat depression
- No other Company has announced manufacture completion (i.e. FDA Module 3) of an IV Ketamine formulation targeted for the treatment of suicidal depression
- No other Company has announced patient-level efficacy data demonstrating the effectiveness of IV ketamine in the treatment of suicidal depression
- A No other Company has announced completion of FDA-required neurotoxicity data in support of an application to treat patients with IV ketamine
- No other Company has announced 12 month real-time stability data associated with a formulation of ketamine to treat suicidal depression

NRx Pharmaceuticals has achieved those milestones and expects to file an NDA under Fast Track designation this year



## NRx Has Toxicity and Manufacturing Data in Support of NDA

- NRx met with FDA on neurotoxicity protocol in 2016 Data were accepted
- 2023: NRx implemented formulation of preservative-free Ketamine
- 3 2023: First stability batch in BFS (no glass) with modern container closure
- 2024: Initial manufacturing completion of first preservative-free formulation
- Ketamine for anesthesia is on drug shortage and existing suppliers are under pressure to limit supply to approved uses

FDA has advised NRx that nonclinical requirements were met



#### **Exhibit 15: HOPE Therapeutics**

# HOPE Therapeutics: Why Spin Out a Separate Company? Expected 2025 profitability

1

NRx is a Biotechnology company focused on Research and Development **HOPE** is a Care Delivery company focused on Interventional Psychiatry and digital therapeutics

**2** 

Focus on TMS and Digital Therapeutics in addition to ketamine and oral meds

- 3
- Insurance-reimbursable: transforms a "cash and carry" market
- 4

Immediate sales of Ketamine under 503b pharmacy license by mid-2024 **Development** of HTX-100 (pH neutral Ketamine) – improved formulation

5

Financing independent of NRXP: equity and bond issuance

#### No FDA-Approved Medication today for Acute Suicidality

#### Only <u>FDA-approved</u> therapy is Electro-Convulsive Therapy (ECT)



#### IV Ketamine is used off-label But not <u>FDA-approved</u> Not reimbursed by Payers Inconsistent in quality



### Hope Therapeutics Near-Term Investor Highlights: Make Good Clinics Great

- Acquisition and management of ~30+ Ketamine clinics through 2025
  - Target \$100 million/yr run-rate with positive EBITDA
  - Starting with industry-leading mental health practices that set the standard and scale for future acquisitions; ongoing revenue and positive EBITDA
  - Make good clinics great: increase revenues via offerings and access, grow EBITDA via product mix, efficiencies and operational optimization
- Disciplined financing strategy
  - · Clinic acquisition:
    - · Market EBITDA multiples with performance based earn-outs
    - Cash via corporate bonds and/or other debt financing expected
    - IPO: planned as the company begins to scale & generate meaningful revenue
- Building shareholder value
  - Market value of clinics is largely based on EBITDA multiples
    - Growing total EBITDA, via increased integration of services, directly enhances shareholder value; debt financing avoids dilution



Exhibit 16: Q3 2024 Results and Recent Business Highlights (as of November 14, 2024)

# NRx Pharmaceuticals (NASDAQ:NRXP) Reports Third Quarter and Year to Date 2024 Financial Results and Provides Business Update

- On track to file New Drug Applications (NDAs) for NRX-100 (IV Ketamine) in treating suicidal ideation in depression, including bipolar depression and NRX-101 (Oral D-Cycloserine/Lurasidone) for Accelerated Approval in bipolar depression with suicidality or akathisia by year end 2024 with 2025 PDUFA date forecast.
- HOPE Therapeutics acquiring Interventional Psychiatry Clinics; key to developing a best-in-class network of care to prevent suicide, continues to expect first revenue by year-end 2024.
- 74% reduction in net operating losses compared to 3<sup>rd</sup> quarter 2023 with profitability forecast in 2025 from HOPE Therapeutics and from sales of medication.
- Management to host a conference call November 18, 2024 at 4:30 PM ET

WILMINGTON, Del., Nov. 14, 2024 / PRNewswire / -- NRx Pharmaceuticals, Inc. (Nasdaq: NRXP) ("NRx Pharmaceuticals", the "Company"), a clinical-stage biopharmaceutical company, today announced its financial results for the quarter and year to date ended September 30, 2024 and provided a business update.

#### **Key Clinical and Business Activities**

- NRX-100 (IV Ketamine) safety and efficacy data have been aggregated from clinical trials in more than 500 patients in collaboration with leading US and French Universities that demonstrate a highly significant reduction in suicidal ideation compared to placebo and active comparator. IV ketamine has been shown to be superior (in post-hoc analysis) to electroshock therapy in treating depression. Stability data has now been generated for three manufacturing lots required for NDA filing in suicidal depression, toxicology data are complete, and alignment with the FDA on its Pediatric Study Plan has been received. If approved, NRX-100 would be the world's first medicine to treat acute suicidality, a condition that kills 1 American every 11 minutes, according to the US Centers for Disease Control.
- NRX-101 NDA for accelerated approval in bipolar depression with suicidality or akathisia advancing, based
  on data from the Phase 2b/3 trial in suicidal patients with bipolar depression, which demonstrated
  depression efficacy comparable to standard of care and significant reduction of akathisia (P=0.025) and
  time to sustained remission from suicidality (P=.05). These data, along with data from our STABIL-B trial,
  support a possible best-in-class suicidal bipolar depression medication profile. This is a multi-billion-dollar
  market in the US.
- HOPE Therapeutics, the Company's wholly owned subsidiary, announced signing of two Letters of Intent
  (LOI) to acquire foundational Interventional Psychiatric Clinics, a key to developing a best-in-class network
  of clinics that offer a complete range of lifesaving therapies to patients with suicidal depression and PTSD.
  These are revenue generating, EBITDA positive acquisitions.
- Secured \$10.8 million in convertible-debt funding from an institutional investor; funds targeted to support FDA New Drug Applications for NRX-100 (ketamine) and NRX-101 and retirement of prior debt. Funding at a reduced in interest rate, conversion discount, and other financial terms compared to prior debt.
- Retired Streeterville debt and settlement of litigation at a substantial discount.



Exhibit 17: Positive Phase 2b/3 Clinical Trial (May 6, 2024)

# Safety Combined with Similar Efficacy in the Trial of NRX-101 Compared to Lurasidone in Suicidal Bipolar Depression



- Both drugs demonstrated > 50% response for treating depression. NRX-101
  demonstrated a mean 76% reduction in symptoms of akathisia compared to
  lurasidone that was sustained over 42 days (Effect Size .37; P=0.025), using prespecified
  analytic methodology memorialized in FDA Special Protocol Agreement. Levels of
  akathisia with NRX-101 were essentially zero at day 42
- This safety advantage was previously reported in the Company's published STABIL-B trial
- Akathisia is identified as a life-threatening side effect of nearly all antidepressants, reported in 10-15% of treated patients and is closely linked to suicide in FDA black box warning
- Akathisia was seen in 2% of participants treated with NRX-101 vs. 11% treated with lurasidone
- Company plans to seek accelerated approval of NRX-101 for use in patients with bipolar depression at risk of akathisia while continuing to broaden the indication to all patients with bipolar depression and perhaps schizophrenia
- Study will be presented at the American Society of Clinical Psychopharmacology
  (ASCP) meeting May 28-31, 2024 (Miami) together with study investigators,
  accompanied by a broadcast scientific presentation on akathisia and antidepressant
  safety, and investor Q&A

#### Commonly heard "But the recent trial did not meet its primary endpoint..."

- The trial did not demonstrate a superior antidepressant effect vs. a leading antidepressant (i.e. the declared primary efficacy endpoint)
- The trial <u>did</u> meet both primary (suicidality) and secondary (akathisia) pre-declared safety endpoints at a statistically-significant level and confirms the findings of two prior trials in this regard
- This provides a basis to file for time-limited (5 year) accelerated approval for patients who have no therapeutic alternative. During that period, the sponsor is required to provide confirmatory evidence of sustained benefit



#### Exhibit 18: Near Term Catalysts and Outlook (as of November 14, 2024)

#### **Key Upcoming 2024 Milestones**

- NRX-100 New Drug Application (NDA) filing for treatment of suicidal ideation in depression, including bipolar depression, based on data from four clinical trials in more than 500 participants demonstrating highly significant efficacy compared to placebo and active comparator, together with a 420-person noninferiority trial compared to electroshock therapy planned to be filed in 2024.
- Company plans to file a New Drug Application (NDA) for Accelerated Approval under Breakthrough Therapy
  Designation and Priority Review of NRX-101 (D-cycloserine+lurasidone) in treatment of bipolar depression
  in people akathisia or suicidality, based on the Phase 2b/3 and STABIL-B data in 2024. Akathisia is
  considered a lethal condition and there is no approved product for this indication.
- HOPE Therapeutics continuing to build its nationwide network of Interventional Psychiatry Clinics; the company is planned to be spun out as a separate entity to be owned by NRx, current NRx shareholders, and new investors.
- NRx continues to forecast first corporate revenues by year-end 2024 with profitability forecast in 2025.

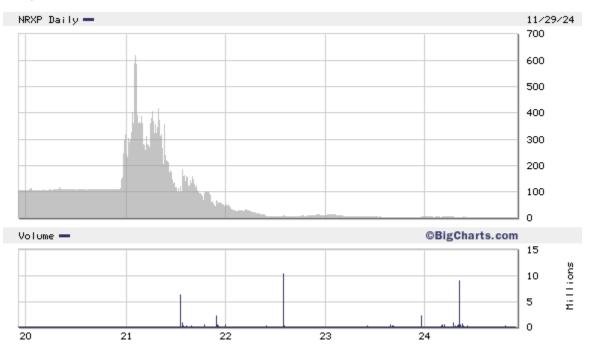


#### Exhibit 19: NRx Pharmaceuticals, Inc. Stock Price (5-Years)

SPAC (Big Rock Partners Acquisition Corp.) IPO - 11/20/17

SPAC Merger Announcement (with NeuroRx, Inc.) - 12/14/20

SPAC Merger Completion (to form NRx Pharmaceuticals, Inc.) - 5/25/21



<sup>\*</sup>Reflects a 1:10 reverse stock split in April 2024

Source: https://bigcharts.marketwatch.com/

Exhibit 20: 0	Consensus Expectations	s (as of November	14, 2024)

	Revenue (mils)	00055	, , ,	EPS	00055
	<u>2024E</u>	<u>2025E</u>		<u>2024E</u>	<u>2025E</u>
Q1 Mar	\$0A		Q1 Mar	\$(0.74)A	
Q2 Jun	\$0A		Q2 Jun	\$(0.75)A	
Q3 Sep	\$0E		Q3 Sep	\$(0.73)E	
Q4 Dec	\$0E		Q4 Dec	\$(0.78)E	
Total	\$0E	\$20E	Total	\$(3.04)E	\$(1.67)E

<sup>\*</sup>Quarterly estimates may not add to annual estimates due to variations in contributing estimates and rounding

Source: Company report, LSEG, and Ascendiant Capital Markets estimates

<sup>\*</sup>Reflects a 1:10 reverse stock split in April 2024



### **FINANCIAL MODEL**

NRx Pharmaceuticals. Inc.

NRx Pharmaceuticals	, Inc.																			
Income Statement (\$ mils)	Mar-22		Sep-22	Dec-22	2022			Sep-23		2023	Mar-24	Jun-24	Sep-24	Dec-24	2024		Jun-25			2025
Fiscal Year End: December 31	Q1A	Q2A	Q3A	Q4A	FY-A	Q1A	Q2A	Q3A	Q4A	FY-A	Q1A	Q2A	Q3A	Q4E	FY-E	Q1E	Q2E	Q3E	Q4E	FY-E
Total Revenue	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	5.0	10.0	20.0	25.0	60.0
Total Revenue	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	5.0	10.0	20.0	25.0	60.0
Cost of Revenues	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	2.5	5.0	7.5	10.0	25.0
Gross Profit	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	2.5	5.0	12.5	15.0	35.0
Research & development General and administrative	5.5 10.2	3.0 6.6	4.1 5.0	4.5 5.5	17.0 27.4	3.7 5.8	3.9 4.1	3.3 2.5	2.5 1.9	13.4 14.2	1.7 4.3	2.8 4.2	0.6 2.4	1.0 3.5	6.2 14.4	1.0 6.0	1.0 6.3	1.0 8.0	1.0 9.0	4.0 29.3
Restructuring and other	10.2	0.0	5.0	5.5	0.0	5.6	0.3	2.5	1.9	0.3	4.3	4.2	2.4	3.5	0.0	0.0	0.3	8.0	9.0	0.0
Total operating expenses	15.7	9.6	9.1	10.0	44.4	9.4	8.2	5.8	4.4	27.8	6.0	7.1	3.0	4.5	20.6	7.0	7.3	9.0	10.0	33.3
Operating income (loss)	(15.7)	(9.6)	(9.1)	(10.0)	(44.4)	(9.4)	(8.2)	(5.8)	(4.4)	(27.8)	(6.0)	(7.1)	(3.0)	(4.5)	(20.6)	(4.5)	(2.3)	3.5	5.0	1.7
Interest income (expense)	(0.0)	0.0	0.1	0.1	0.2	0.2	0.1	0.1	(0.0)	0.4	(0.2)	0.0	0.0	(0.2)	(0.4)	(0.3)	(0.3)	(0.3)	(0.3)	(1.4)
Other income (expense)	2.3	2.6	(0.0)	(0.5)	4.3	(1.8)	(0.7)	(0.3)	0.1	(2.7)	(0.3)	(0.9)	1.4	0.0	0.2	0.0	0.0	0.0	0.0	0.0
Income before income taxes	(13.4)	(7.0)	(9.1)	(10.3)	(39.8)	(11.0)	(8.7)	(6.1)	(4.3)	(30.2)	(6.5)	(7.9)	(1.6)	(4.7)	(20.8)	(4.8)	(2.6)	3.2	4.7	0.3
Income taxes	(40.0)	( <b>7</b> 0)	(0.4)	(40.0)	0.0	(44.0)	(0.7)	(0.4)	(4.0)	0.0	(0.5)	(7.0)	(4.0)	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Net income (loss)	(13.4)	(7.0)	(9.1)	(10.3)	(39.8)	(11.0)	(8.7)	(6.1)	(4.3)	(30.2)	(6.5)	(7.9)	(1.6)	(4.7)	(20.8)	(4.8)	(2.6)	3.2	4.7	0.3
Nonrecurring/noncash adjustme	ents				0.0					0.0					0.0					0.0
Net income (pro forma)	(13.4)	(7.0)	(9.1)	(10.3)	(39.8)	(11.0)	(8.7)	(6.1)	(4.3)	(30.2)	(6.5)	(7.9)	(1.6)	(4.7)	(20.8)	(4.8)	(2.6)	3.2	4.7	0.3
EBITDA																				
Shares, Basic	6.4	6.6	6.6	6.8	6.6	6.7	7.3	8.2	8.2	7.6	8.9	10.5	11.0	12.1	10.6	12.2	12.3	12.4	12.5	12.4
Shares, Diluted	6.4	6.6	6.6	6.8	6.6	6.7	7.3	8.2	8.2	7.6	8.9	10.5	11.0	12.1	10.6	12.2	12.3	12.4	12.5	12.4
EPS Basic (pro forma)	(\$2.11)	(\$1.06)	(\$1.37)	(\$1.53)	(\$6.05)	(\$1.64)	(\$1.19)	(\$0.74)	(\$0.53)	(\$3.98)	(\$0.74)	(\$0.75)	(\$0.15)	(\$0.39)	(\$1.96)	(\$0.40)	(\$0.22)	\$0.25	\$0.37	\$0.02
EPS Diluted (pro forma)	1							(\$0.74)					(\$0.15)				(\$0.22)		\$0.37	\$0.02
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Margins																				
Gross margin Research & development																				
General and administrative																				
Operating margin																				
Tax rate, GAAP																				
Net margin																				
Y/Y % change																				
Total Revenue																				
Gross margin	000/	-37%	2.40/	240/	160/	220/	240/	200/	420/	240/	E20/	200/	020/	640/	E 40/	420/	640/	6.40/	00/	250/
Research & development General and administrative	88% 387%	-37% -47%	-34% -64%	-31% -88%	-16% -63%	-33% -43%	31% -39%	-20% -50%	-43% -66%	-21% -48%	-52% -27%	-28% 4%		-61% 87%	-54% 1%		-64% 48%	64% 232%	0% 157%	-35% 103%
Operating income (loss)	-39%	-44%	-55%	-81%	-62%	-40%	-15%	-36%	-56%	-37%	-36%	-14%		2%			-67%	-216%	-211%	-108%
Net income (loss)	-47%	-97%	-56%	-78%	-89%	-18%	25%	-33%	-58%	-24%	-41%	-9%		9%	-31%		-66%	-294%	-199%	-101%
EPS Diluted (pro forma)	-70%	-98%	-66%	-81%	-92%	-23%	12%	-46%	-65%	-34%	-55%	-37%	-80%	-27%	-51%	-46%	-71%	-272%	-196%	-101%
						1														

Source: Company reports and Ascendiant Capital Markets estimates.

\*Reflects a 1-for-10 Reverse Stock Split in April 2024



NRx Pharmaceuticals, Inc.

NRx Pharmaceuticals, Inc.	May 20	lum 20	C-= 20	D 00	May 22	Lun 22	C 22	D 22	May 04	lum 24	Cam 24	Dec 24	May 25	lum 25	Cam 25	Dec 25
Balance Sheet (\$ mils) Fiscal Year End: December 31	Mar-22 Q1A	Jun-22 Q2A	Sep-22 Q3A	Q4A	Mar-23 Q1A	Q2A	Sep-23 Q3A	Q4A	Mar-24 Q1A	Jun-24 Q2A	Sep-24 Q3A	Dec-24 Q4E	Mar-25 Q1E	Jun-25 Q2E	Sep-25 Q3E	Dec-25 Q4E
Fiscal Teal Eliu. Decelliber 31	QIA	QZA	QSA	Q4A	QIA	QZA	QJA	Q4A	QIA	QZA	QJA	Q4E	Q IE	Q2E	Ų3E	Q4E
Assets																
Cash and cash equivalents	40.2	24.5	18.2	20.1	16.5	15.0	8.9	4.6	1.3	1.9	1.6	1.0	1.7	1.2	4.5	9.2
Short term investments												0.0	0.0	0.0	0.0	0.0
Account receivable																
Deferred income taxes												0.0	0.0	0.0	0.0	0.0
Prepaid expenses and other	3.4	7.9	6.6	5.7	5.3	4.8	4.2	2.3	2.0	3.0	2.5	2.5	2.5	2.5	2.5	2.5
Total current assets	43.6	32.4	24.8	25.8	21.8	19.8	13.1	6.9	3.3	4.9	4.1	3.5	4.1	3.6	6.9	11.7
Property and equipment, net												0.0	0.0	0.0	0.0	0.0
Intangibles, net												0.0	0.0	0.0	0.0	0.0
Deferred income tax												0.0	0.0	0.0	0.0	0.0
<u>Other</u>	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.4	0.4	0.4	0.4	0.4	0.0	0.0	0.0	0.0
Total assets	43.6	32.4	24.8	25.8	21.8	19.8	13.1	7.3	3.8	5.3	4.5	3.8	4.1	3.6	6.9	11.7
Liabilities and stockholders' equity																
Accounts payable	4.3	3.1	2.2	2.1	3.8	2.2	3.6	4.6	6.3	5.0	4.9	4.9	4.9	4.9	4.9	4.9
Accrued expenses	4.5	4.0	5.8	5.8	6.1	6.9	5.3	5.2	5.8	10.0	9.8	9.8	9.8	9.8	9.8	9.8
Deferred income tax												0.0	0.0	0.0	0.0	0.0
Warrant liabilities	0.1	0.0	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	1.9	1.9	1.9	1.9	1.9	1.9
Other	2.5					0.8	0.3				0.6	0.6	0.6	0.6	0.6	0.6
Short term debt	0.5			8.7	12.2	12.7	10.1	9.2	6.8	8.6	3.1	<u>7.0</u>	<u>7.0</u>	<u>7.0</u>	7.0	7.0
Total current liabilities	11.9	7.1	8.0	16.6	22.1	22.6	19.3	19.0	18.9	23.7	20.3	24.2	24.2	24.2	24.2	24.2
Deferred income taxes												0.0	0.0	0.0	0.0	0.0
Warrant liabilities												0.0	0.0	0.0	0.0	0.0
Other long term liabilities												0.0	5.0	7.0	7.0	7.0
Long term debt				<u>1.8</u>							3.0	3.0	<u>3.0</u>	3.0	3.0	3.0
Total other liabilities	0.0	0.0	0.0	1.8	0.0	0.0	0.0	0.0	0.0	0.0	3.0	3.0	8.0	10.0	10.0	10.0
Preferred stock							0.0	0.0				0.0	0.0	0.0	0.0	0.0
Common stock	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.0	0.0	0.0	0.2	0.3	0.4	0.6	0.7
Additional paid-in capital	228.3	229.0	229.5	230.4	233.6	239.9	242.5	241.3	244.6	249.2	250.4	250.4	250.4	250.4	250.4	250.4
Retained earnings	(196.7)	(203.7)	(212.8)	(223.1)	(234.0)	(242.8)	(248.8)	(253.1)	(259.7)	(267.6)	(269.2)	(273.9)	(278.8)	(281.4)	(278.3)	(273.6)
Other												0.0	0.0	0.0	0.0	0.0
Accumulated other comprehensive in					0.1	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)		0.0	0.0	0.0	0.0	0.0
Total stockholders' equity	31.7	25.4	16.8	7.4	(0.3)	(2.8)	(6.2)	(11.7)	(15.1)	(18.4)	(18.8)	(23.4)	(28.1)	(30.6)	(27.3)	(22.5)
Total stockholders' equity and liabil	43.6	32.4	24.8	25.8	21.8	19.8	13.1	7.3	3.8	5.3	4.5	3.8	4.1	3.6	6.9	11.7

Balance Sheet Drivers

	Mar-22	Jun-22	Sep-22	Dec-22	Mar-23	Jun-23	Sep-23	Dec-23	Mar-24	Jun-24	Sep-24	Dec-24	Mar-25	Jun-25	Sep-25	Dec-25
	Q1A	Q2A	Q3A	Q4A	Q1A	Q2A	Q3A	Q4A	Q1A	Q2A	Q3A	Q4E	Q1E	Q2E	Q3E	Q4E
Book & Cash Value (per share)																
Book Value per Share (diluted)	\$4.98	\$3.86	\$2.53	\$1.10	(\$0.04)	(\$0.38)	(\$0.76)	(\$1.43)	(\$1.70)	(\$1.75)	(\$1.72)	(\$1.93)	(\$2.30)	(\$2.49)	(\$2.20)	(\$1.80)
Cash per Share (diluted)	\$6.31	\$3.73	\$2.75	\$2.97	\$2.45	\$2.04	\$1.09	\$0.56	\$0.15	\$0.18	\$0.15	\$0.08	\$0.14	\$0.09	\$0.36	\$0.74
Net cash per Share (diluted)	\$6.23	\$3.73	\$2.75	\$1.41	\$0.64	\$0.31	(\$0.14)	(\$0.56)	(\$0.62)	(\$0.64)	(\$0.40)	(\$0.74)	(\$0.68)	(\$0.72)	(\$0.45)	(\$0.06)

Source: Company reports and Ascendiant Capital Markets estimates



NRx Pharmaceuticals, Inc.

NRx Pharmaceuticals, I					, .															
Cash Flow Statement (\$ mils)			Sep-22							2023	Mar-24	Jun-24		Dec-24	2024			Sep-25		
Fiscal Year End: December 31	Q1A	Q2A	Q3A	Q4A	FY-A	Q1A	Q2A	Q3A	Q4A	FY-A	Q1A	Q2A	Q3A	Q4E	FY-E	Q1E	Q2E	Q3E	Q4E	FY-E
Cash flow from operating activi																				
Net income	(13.4)	(7.0)	(9.1)	(10.2)	(39.8)	(11.0)	(8.7)	(6.1)	(4.3)	(30.2)	(6.5)	(7.9)	(1.6)	(4.7)	(20.8)	(4.8)	(2.6)	3.2	4.7	0.3
Depreciation	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Amortization					0.0					0.0					0.0					0.0
Debt related amortization exper		(0.0)	0.0	0.0	0.0					0.0			0.5		0.5					0.0
Stock comp	1.3	1.0	0.5	0.8	3.6	0.7	0.5	0.4	(1.2)	0.4	0.2	0.1	0.1	0.1	0.6	0.1	0.1	0.1	0.1	0.6
Deferred income taxes					0.0					0.0				0.0	0.0	0.0	0.0	0.0	0.0	0.0
Change in fair value of warrant		(0.1)	0.0	0.5	0.3	1.8	0.7	0.3	(0.1)	2.7	0.3	2.2	(3.7)		(1.2)					0.0
Change in fair value of earnout	(2.1)	(2.5)			(4.6)								1.3		1.3					
Writedowns and impairments					0.0					0.0			0.8		0.8					0.0
Other gains/losses					0.0		0.3			0.3			0.1		0.1					0.0
Other					0.0					0.0					0.0					0.0
Changes in operating assets and	liabilities:																			
Accounts receivable					0.0					0.0					0.0					0.0
Prepaid expenses & other curre	1.7	(4.5)	1.3	0.8	(0.6)	0.5	0.4	0.6	1.5	3.0	0.3	(0.9)	0.5		(0.1)					0.0
Income tax					0.0					0.0					0.0					0.0
Other assets					0.0					0.0				0.0	0.0	0.4	0.0	0.0	0.0	0.4
Accounts payable	0.6	(1.2)	(0.9)	(0.1)	(1.6)	1.7	(1.6)	1.5	1.0	2.7	2.1	2.1	0.7		4.9					0.0
Accrued expenses	1.6	(0.5)	1.8	(0.0)	2.9	0.3	0.6	(1.3)	(0.1)	(0.5)	(0.1)	0.9	(0.2)		0.6					0.0
Other liabilities					0.0					0.0		0.9	(1.0)	0.0	(0.1)	5.0	2.0	0.0	0.0	7.0
Net cash (used in) provided by	(10.4)	(14.8)	(6.3)	(8.3)	(39.8)	(6.1)	(7.8)	(4.6)	(3.2)	(21.7)	(3.7)	(2.6)	(2.3)	(4.6)	(13.1)	0.6	(0.5)	3.3	4.8	8.2
Cash flow from investing activi	ties																			
Purchases of property and equi	(0.0)	(0.0)	(0.0)	0.0	(0.0)	(0.0)	0.0	(0.0)	0.0	(0.0)				(0.0)	(0.0)	0.0	0.0	(0.0)	(0.0)	(0.0)
Purchases of short-term investr	nents				0.0					0.0					0.0					0.0
Acquisitions					0.0					0.0					0.0					0.0
Other					0.0					0.0					0.0					0.0
Net cash used in investing active	(0.0)	(0.0)	(0.0)	0.0	(0.0)	(0.0)	0.0	(0.0)	0.0	(0.0)	0.0	0.0	0.0	(0.0)	(0.0)	0.0	0.0	(0.0)	(0.0)	(0.0)
Cash flow from financing activi	ties																			
Issuance of debt				10.0	10.0		0.8		0.4	1.2			2.9	4.0	6.9	0.0	0.0	0.0	0.0	0.0
Repayment of debt		(0.5)			(0.5)		(0.1)	(2.7)	(0.3)	(3.1)	(2.2)	(0.0)	(3.2)		(5.3)					0.0
Issuance of stock	23.0	(0.3)	(0.0)	0.1	22.7	2.5	5.6	1.2	(1.2)	8.1	2.6	3.1	0.2	0.0	5.9	0.0	0.0	0.0	0.0	0.0
Proceeds from stock option exe	ercises			0.0	0.0					0.0			2.1		2.1					0.0
Other					0.0					0.0					0.0					0.0
Dividends and distributions					0.0					0.0					0.0					0.0
Cash provided by (used in) fina	23.0	(0.9)	(0.0)	10.1	32.2	2.5	6.3	(1.5)	(1.1)	6.2	0.4	3.1	2.1	4.0	9.5	0.0	0.0	0.0	0.0	0.0
		-	-												_					_
Effect of exchange rate on cash					0.0					0.0					0.0					0.0
Net increase (decrease) in cash	12.6	(15.7)	(6.3)	1.8	(7.6)	(3.5)	(1.5)	(6.1)	(4.3)	(15.5)	(3.3)	0.6	(0.3)	(0.6)	(3.6)	0.6	(0.5)	3.3	4.8	8.2
Beginning cash and equivalent	27.6	40.2	24.5	18.2	27.6	20.1	16.5	15.0	8.9	20.1	4.6	1.3	1.9	1.6	4.6	1.0	1.7	1.2	4.5	1.0
Ending cash and equivalents	40.2	24.5	18.2	20.1	20.1	16.5	15.0	8.9	4.6	4.6	1.3	1.9	1.6	1.0	1.0	1.7	1.2	4.5	9.2	9.2

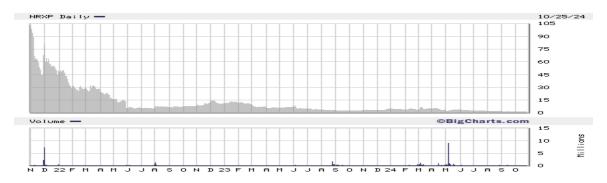
Source: Company reports and Ascendiant Capital Markets estimates



#### **ANALYST CERTIFICATION**

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### NRx Pharmaceuticals, Inc.



\*Reflects a 1:10 reverse stock split in April 2024

Source: https://bigcharts.marketwatch.com/

	Report Date		Price
Report	Date	Rating	Target
1	11/9/2022	В	40.00
2	11/18/2022	В	45.00
3	4/5/2023	В	47.50
4	5/23/2023	В	50.00
5	9/6/2023	В	52.50
6	12/22/2023	В	55.00
7	5/4/2024	В	50.00
8	6/6/2024	В	43.00
9	9/11/2024	В	44.00

Ascendiant Capital Markets, LLC has received compensation for advisory or investment banking services from the company
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Risks to attainment of our share price target include balance sheet/liquidity risks, failure of product candidates to demonstrate safety and efficacy in clinical trials, failure to gain regulatory approvals, ability to commercialize product, failure to obtain suitable reimbursement, competition, changing macroeconomic factors, investor sentiment for investing in biotech stocks, and changes in consumer or government priorities for healthcare.

#### **Ascendiant Capital Markets, LLC Rating System**

**BUY:** We expect the stock to provide a total return of 15% or more within a 12-month period.

**HOLD:** We expect the stock to provide a total return of negative 15% to positive 15% within a 12-month period.

SELL: We expect the stock to have a negative total return of more than 15% within a 12-month period.

Total return is defined as price appreciation plus dividend yield.

#### Ascendiant Capital Markets, LLC Distribution of Investment Ratings (as of October 11, 2024)

Investment	Banking	Services
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			Past 12 months	
Rating	Count	Percent	Count	Percent
Buy	58	98%	25	43%
Hold	0	0%	0	0%
Sell	1	2%	0	0%
Total	59	100%	25	42%



#### **Other Important Disclosures**

Our analysts use various valuation methodologies including discounted cash flow, price/earnings (P/E), enterprise value/EBITDAS, and P/E to growth rate, among others. Risks to our price targets include failure to achieve financial results, product risk, regulatory risk, general market conditions, and the risk of a change in economic conditions.

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