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Page 4/24

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RESEARCH

TRIANGLE PARK –

Kriya Therapeutics, which has operations in the Triangle and Silicon Valley, is even more flush with cash than a year ago, adding a second round of financing totalling \$100

...

**Gene therapy startup
in Triangle lands \$100**

Page 5/24

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Gene Therapy

million in new funding

New research results show that a new form of gene editing is efficient in correcting a mutation in patient cells with the monogenic disease

Alpha-1 antitrypsin deficiency (AATD), a common inherited ...

New form of gene editing efficient for treatment of AATD,

Page 6/24

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liver and lung disease

"Prime Editing can work in probably more than 90% of the known disease-causing mutations," said Keith Gottesdiener, CEO of Prime Medicine, which launched on Tuesday after receiving \$315 million in ...

Broad Institute

spinout launches with

Page 7/24

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**\$315M for new gene-
editing tech**

Scientists reported striking, early-stage results from a small trial of a gene therapy for AADC deficiency, an ultra-rare genetic disorder.

**Landmark gene
therapy trial points to
a wider window to
alter course of rare**

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disease

Jul 05, 2021 (The
Expresswire) -- "Final
Report will add the

analysis of the impact of
COVID-19 on this
industry." Global "Gene
Therapy Market" 2021

...

**Gene Therapy Market
Size, Global Industry
Trends, Share,
Growth, Emerging**

Page 9/24

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Gene Therapy

Technologies,

Opportunity and

Forecast 2021 to 2026

Facility to Support

Allogeneic CAR T,

Gene Therapy, and

Stem Cell-Derived

Therapies Facility Able

to Support Clinical Trial

and Early Commercial

Launches of Multiple

Programs State-of-the-

Art ...

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**Sana Biotechnology
Enters Long-Term
Lease to Establish
Manufacturing
Facility to Enable Its
Gene and Cell
Therapy Product
Candidates**

In a new study on mice,
Johns Hopkins
Medicine researchers
report that using MRI
scans to measure blood
volume in the brain can

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serve as a noninvasive way to potentially track the progress of gene ...

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Gene therapy in early stages may slow down Huntington's disease

Treatments from Bluebird bio and CRISPR Therapeutics have shown considerable promise and could soon be nearing regulatory

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review. A full pipeline
is growing behind them.

Technology

Essment In Its

**Gene therapy for
sickle cell disease:**

**progress and
competition**

In a new study on mice,
Johns Hopkins

Medicine researchers
report that using MRI
scans to measure blood
volume in the brain can
serve as a noninvasive

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way to potentially track
the progress of gene ...

**Gene Editing Therapy
in Early Stages of
Huntington's Disease
May Slow Down**

**Symptom Progression,
Mouse Study Shows**

With the goal of training
and attracting top talent,
Columbus-based gene
therapy company Forge
Biologics has launched

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the Forge Forward gene
therapy workforce
development program in
partnership with ...

**Forge Biologics,
CWRU launch gene
therapy workforce
development
partnership**

Hot Gene Editing

Stocks To Watch In The
Stock Market Today.

Finding the best stocks

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to invest in 2021 is all about understanding the stock market trajectory. With the healthcare s ...

Societal

What Are The Best Stocks To Invest In? 5 Gene Editing Stocks To Watch Now

Forge Biologics, a gene therapy-focused contract development and manufacturing organization (CDMO),

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has launched the Forge
Forward gene therapy
workforce development
program in partnership
with the ...

Context

**Forge Biologics and
Case Western Reserve
University Announce
Gene Therapy
Workforce
Development
Partnership**

Codexis enzymes have

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Applications in the
sustainable
manufacturing of
pharmaceuticals, food,
and industrial products;
the creation of the next
generation of life
science tools; and as
biologic ...

**Codexis Announces
Completion of
CodeEvolver® License
Technology Transfer**

Page 18/24

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Gene Therapy

with Global

Pharmaceutical

Leader

Gene editing could revolutionize the way we treat diseases. Along the way, it could also create substantial opportunity for related stocks.

**The Top 3 Gene
Editing Stocks to Own
for the Next Decade**

Page 19/24

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In a new study on mice, Johns Hopkins Medicine researchers report that using MRI scans to measure blood volume in the brain can serve as a noninvasive way to potentially track the progress of gene ...

Study: Gene editing therapy for early-stage Huntington's disease may have long-lasting

Page 20/24

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benefits

BioMarin

Pharmaceutical Inc.

BMRN announced that the European Medicines Agency (EMA) has granted accelerated assessment for the review of its investigational gene therapy valoctocogene ...

BioMarin (BMRN)

Page 21/24

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Hemophilia Gene Therapy Gets EMA's Speedy Review

ASC Therapeutics, a privately-held biopharmaceutical company pioneering the development of transformative in-vivo gene replacement, gene editing and allogeneic cell therapies for hematologic and other

...

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**ASC Therapeutics
Receives IND**

**Clearance From the
U.S. Food and Drug**

**Administration for
ASC618 Second-**

**Generation Gene
Therapy for**

Hemophilia A

The researchers say that
by identifying and
treating the mutation
known to cause

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Huntington's disease
with this type of gene
therapy, before a patient
starts ... a reliable
biomarker that can track

Context

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