UNITED STATES

SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934 Date of Report (Date of earliest event reported) November 14, 2000 Aastrom Biosciences, Inc. (Exact name of registrant as specified in charter) Michigan 0-22025 94-3096597 (State or other jurisdiction (Commission (IRS Employer of incorporation) File Number) Identification N Identification No.) of incorporation) File Number) 24 Frank Lloyd Wright Drive, P.O. Box 376, Ann Arbor Michigan (Address of principal executive offices) (Zip Code) Registrant's telephone number, including area code (734) 930-5555 Not Applicable (Former name or former address, if changed since last report)

Item 7. Financial Statements and Exhibits.

Exhibit

No. Description

99 Slides used in presentations

Item 9. Regulation FD Disclosure.

Attached hereto as Exhibit 99, which is incorporated herein by reference, is a copy of certain slides used by the Company in making presentations to analysts and shareholders. This information is not "filed" pursuant to the Securities Exchange Act and is not incorporated by reference into any Securities Act registration statements. Additionally, the submission of this report on Form 8-K is not an admission as to the materiality of any information in this report that is required to be disclosed solely by Regulation FD.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Aastrom Biosciences, Inc.

Date: November 14, 2000 By: /s/ Todd E. Simpson

Vice President Finance & Administration

Vice President, Finance & Administration and Chief Financial Officer (Principal Financial and Accounting Officer)

EXHIBIT INDEX

Exhibit

No. Description

99 Slides used in presentations

[SLIDE 1]

Aastrom Biosciences, Inc.

Bringing Therapeutic Cells Into Medical Practice

November, 2000

Artwork: Technician working with machine. Illustration of body cells.

[SLIDE 2]

SAFE HARBOR

This presentation contains forward-looking statements, including without limitation statements concerning product development objectives, Aastrom's business model for its products, clinical trial timing and anticipated results, potential revenues, margins and markets for Aastrom's products and products under development, and potential advantages and applications of the AastromReplicell System, which involve certain risks and uncertainties. Actual results may differ significantly from the expectations contained in the forward-looking statements.

Among the factors that may result in differences are the results obtained from clinical trials and development activities, regulatory approval requirements, competitive developments and the availability of resources.

These and other significant factors are discussed in greater detail in Aastrom's Annual Report of Form 10-K and other filings with the Securities and Exchange Commission.

[SLIDE 3]

Aastrom Biosciences, Inc. Company Overview

- . Develops and commercializes proprietary clinical products and processes used in medical practice for the ex vivo production of human cells
- . Patient-specific cells would then be used therapeutically to enable treatments for multiple diseases including:

cancer leukemia genetic blood diseases osteoporosis viral infections

[SLIDE 4]

Aastrom Biosciences, Inc. Highlights

- . Pioneering products line to commercialize and standardize therapeutic cell production
- . Flagship stem cell products in final phases of clinical development in US
- . CE Mark approvals received for stem cell and cord blood products in Europe and pilot launch initiated
- . Broad patent position for ex vivo production of stem and other cells, and for clinical systems to enable these processes
- . Business model contemplates a two-tiered revenue stream from sales of consumable therapy kits and of instrumentation platform

[SLIDE 5]

Aastrom Biosciences, Inc. Company at a Glance (10/00)

. Founded: 1991

Location: Ann Arbor, MIEmployees: Approximately 35Public listing: ASTM (Nasdaq; 1997)

Common Shares: 33.8 million

. Market Cap: Approximately \$65 million (10/31/00)

(52 week range: \$7 million to \$235 million)

[SLIDE 6]

Cell Therapy The Need for Products

- . Expanding use of human cells to treat disease, or to replace diseased or damaged tissues
- . Patient-specific cells need to be produced outside of the body (ex vivo), from small starting cell samples
- . Increasing market need for a product that enables therapeutic cell production to move into standard medical practice

[SLIDE 7]

Cell Therapy Commercialization Pathway

Research Lab

Artwork: Body cells on one side of brick wall. Patient on other side of brick wall.

- . Bone Marrow
- . Cord Blood Cells
- . Stem Cells
- . T-cells
- . Chondrocytes
- . Neuronal Cells
- . Dendritic Cells

[SLIDE 8]

Cell Therapy Commercialization Pathway

Research Lab AastromReplicell System

Artwork: Body cells passing through open door in brick wall to patient.

- . Bone Marrow
- . Cord Blood Cells
- . Stem Cells
- . T-cells
- . Chondrocytes
- . Neuronal Cells
- . Dendritic Cells

[SLIDE 9]

AastromReplicell System Meeting Market Needs

- . A turn-key platform medical product line in development that provides a standard commercialization pathway for high quality therapeutic cell production
- . Automated production of cells with superior biologic function compared to standard culture approaches in bags/flasks
- Business model provides pharmaceutical-type revenue stream through sales of single-use therapy-specific kits that operate on a fixed instrument platform

[SLIDE 10]

AastromReplicell System Platform Automated Cell Production

Artwork: Illustration of system platform. Two illustrations of technician using system.

[SLIDE 11]

AastromReplicell System
Therapy Kit Single-Use Components

Cell Cassette

Artwork: Illustration of cell cassette

- . sterile fluid pathway
- . used for all cell types

Application Key

Artwork: Illustration of application key

- . therapeutic cell specific program software
- . attaches to cell cassette
- . programs the instruments

Liquid Products

Artwork: Illustration of liquid products

- . therapeutic cell specific growth medium
- . added to cell cassette with cell inoculum

[SLIDE 12]

AastromReplicell System Therapy Kits

- . Single-use product to produce a specialized mixture of cells to treat a specific medical indication
- . Provides an automated, fixed-process cell production under GMP compliance
- . Provides a way for Aastrom to get paid for each patient cell therapy procedure
- Enables indication-specific product development capability for regulatory approval, sales and marketing strategies, and pricing

[SLIDE 13]

AastromReplicell System Therapy Kits - Anticipated Use A New Generation of Therapies

Hospital or Service Provider Purchases Therapy Kits from Aastrom

CB-I Therapy Kit

SC-I Therapy Kit

OC-I Therapy Kit

DC-I Therapy Kit

TC-I Therapy Kit

Physician Prescribes Therapy

Lab/Service Selects Therapy Kit

Cell Prescription Filled

Patient Infused with Therapeutic Cells

[SLIDE 14]

AastromReplicell System
Product Oveview - Launch + 5 years

Graph of Revenue Distribution

Graph of Contribution to Overall Margin

[SLIDE 15]

AastromReplicell System Therapy Kit Status

Therapy Kit	Target 	Europe	US
SC-I (stem cells)	Solid Cancers	Approved	Phase III
CB-I (cord blood)	Leukemia	Approved	Phase III
OC-I (bone cells)	Osteoporosis		Phase I/II
CB-II	Leukemia		Pre-clinical
DC-I (dendritic cells)	Cancers		Pre-clinical
T-Cell #1	Viruses		Pre-clinical

[SLIDE 16]

Cord Blood Market

Stem Cell Transplants (US/Europe)

- . 60,000+ annual cases of leukemia
- Curative treatment potential with donor stem cell transplant35,000+ could be candidates for cord blood derived SCT, if there was a large enough cell dose
- Potential market size expansion with use in genetic diseases such as sickle cell anemia

[SLIDE 17]

CB-I Therapy Kit Development Status

- . Aastrom multi-center trials have shown:
 - production of desired cord blood doses

excellent 100 and 400 day survival rates in pediatric leukemia patients enablement of disease-free blood and immune system recovery in adult leukemia patients

(published: April, 2000 Bone Marrow Transplantation)

- US Phase III Trial planned in FY 2001
- Approved for sale in Europe

[SLIDE 18] Bone Disease Markets Osteoporosis

- . Estimated 10 million with disease in US
- . Drug treatments for osteoporosis generally deter further degeneration, but do not repair/form new bone
- . Bone progenitor cells can be used to restore and build bone tissue
- . Severe osteoporosis candidates projected for cell therapy: approximately 350.000
- . Potential cell therapy market: \$1.5 billion

[SLIDE 19] OC-I Therapy Kit Development Plan

- . Aastrom pre-clinical studies demonstrating bone progenitor cell expansion completed
- . Transplant (compassionate use) of genetic bone disease patient with OC-I Therapy Kit produced cells, proved successful in restoring bone density
- . Phase I/II feasibility trial in patients with severe osteoporosis approved by FDA (initiated 10/00)

[SLIDE 20] AastromReplicell System Target Markets (US/Europe)

Therapy Kit	Disease	Estimated Annual Patient Candidates	Potential Market
SC-I	Solid Cancers	15,000	\$110 M
CB-I	Leukemia	35,000	\$250 M
CB-II	Leukemia+	45,000	\$335 M
OC-I	Osteoporosis	350,000	\$1500 M
DC-1	Cancers	150,000	\$1200 M

[SLIDE 21]

Immunotherapy Markets Dendritic Cells and T-Cells

- . Ex vivo produced dendritic cells or T-Cells can trigger immune responses against specific targets
- . Potential treatment for all types of cancer and leukemia, and for certain viral infections
- Ex vivo produced/modified dendritic cells shown effective in multicenter trial to eradicate large bulky human tumors (Published: Nature Medicine, March 2000)
- . Rapidly emerging major market opportunity

[SLIDE 22]

DC-I Therapy Kit Development Status

- . Successful clinical approaches completed using limited manual research lab culture procedures non-automated process restricts move to multi-centers and regulatory approval
- . Aastrom has completed clinical scale human dendritic cell production improved dendritic cell production and biological function observed
- . First dendritic cell product (DC-I Therapy Kit) under development and expected into US and European clinical research markets in FY 2001

[SLIDE 23]

Aastrom Biosciences, Inc. 12 Month Milestones (projected)

- . Formal European launch of the SC-I and CB-I Therapy Kits
- . Initiation of US Phase III trial of the CB-I Therapy Kit
- . Initiation of the Phase I/II trial of the OC-I Therapy Kit for severe osteoporosis $\,$
- . Completion of the DC-I Therapy Kit and entered into US clinical research market and European medical market
- . Completion of additional financing (strategic and/or equity)

[SLIDE 24]

Aastrom Biosciences, Inc. Growing in All Ways

- Pioneering product line for standardized therapeutic cell production Strong and diverse patent position
- Lead stem cell therapy products now approved for sale in Europe
- Positioned to enter new osteoporosis and cancer immunotherapy markets
- In forefront of apparent blockbuster dendritic cell therapy market
- New strategic partnership potential

[SLIDE 25]

Aastrom Biosciences, Inc. Bringing Therapeutic Cells Into Medical Practice November, 2000

Artwork: Technician working with machine. Illustration of body cells.