

## Hyperion Therapeutics Inc

### COMPANY AND PIPELINE OVERVIEW REPORT

Coverage of the company and a summary of the drug pipeline portfolio.

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## GLOSSARY

### Number of Drugs in Active Development

Number of drugs associated with the company or subsidiary that are currently in active development, i.e. the development status for the drug(s) is one of the following: Discovery, Clinical, Phase I, Phase II, Phase III, Pre-registration, Registered, Launched, or Suspended.

### Number of Inactive Drugs

Number of drugs associated with the company or subsidiary that are currently classified as inactive, i.e. where the development status for the drug(s) is one of the following: No Development Reported, Discontinued, or Withdrawn.

### Number of Patents as Owner

Number of patents associated with the company where the company is listed as owner; i.e. the relationship type (or way the patent refers to the company) is: Patent Assignee/Owner, Patent owner (not assignee), Licensee for development and marketing, Licensee – marketing only (Distributor), Patent assignee of family member, Inferred assignee.

### Number of Patents as Third Party

Number of patents associated with the company where the company is listed as third party; i.e. the relationship type (or way the patent refers to the company) is: Patent assignee (not owner), Ex-Licensee for development and marketing, Ex-Licensee marketing only (Distributor), Customer of technology, Ex-Customer of technology, Patent opponent or infringer, Affiliate organization of inventor, Owner of underlying technology.

### Patents summary table

This table represents a summary of the core patent coverage for this company covering Therapeutic EP, US and WO patents since 1990 only.

### Number of Deals

A count of deals where the company or one of its subsidiaries is the primary company.

### Key Indications

Displays top ten key indications for the company and its subsidiaries based on frequency (indications occurring with high and identical frequency are always included, and this may result in more than ten Key Indications being listed). Includes both indications associated with patents where the company is patent owner and indications associated with drugs in active development. A drug is classified as 'active' if it features on a row (or rows) in the current development status table where the status is one of the following: Discovery, Clinical, Phase I, Phase II, Phase III, Pre-registration, Registered, Launched, or Suspended.

### Key Target-based Actions

Displays top ten key target-based actions for the company and its subsidiaries based on frequency (actions occurring with high and identical frequency are always included, and this may result in more than ten Key Target-based Actions being listed). Includes both target-based actions associated with patents where the company patent owner and target-based actions associated with drugs in active development. A drug is classified as 'active' if it features on a row (or rows) in the current development status table where the status is one of the following: Discovery, Clinical, Phase I, Phase II, Phase III, Pre-registration, Registered, Launched, or Suspended. A target-based action is one that is associated with a target.

### Key Technologies

Displays top ten key technologies for the company and its subsidiaries based on frequency (technologies occurring with high and identical frequency are always included, and this may result in more than ten Key Technologies being listed). Includes both key technologies associated with patents where the company relationship is patent owner and key technologies associated with drugs in active development. A drug is classified as 'active' if it features on a row (or rows) in the current development status table where the status is one of the following: Discovery, Clinical, Phase I, Phase II, Phase III, Pre-registration, Registered, Launched, or Suspended.

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# Hyperion Therapeutics Inc

## COMPANY OVERVIEW

|                                       |   |
|---------------------------------------|---|
| Company Name                          | Hyperion Therapeutics Inc   |
| Parent Company Name                   | Hyperion Therapeutics Inc   |
| Website                               | <a href="http://www.hyperiontx.com/">http://www.hyperiontx.com/</a> |
| Country                               | US  |
| Number of Drugs in Active Development | 2   |
| Number of Inactive Drugs              | 0   |
| Number of Patents as Owner            | 3   |
| Number of Patents as Third Party      | 0   |
| Number of Deals                       | 1   |
| Key Indications                       | Hepatic encephalopathy,Hyperammonemia,Uremia                        |
| Key Target-based Actions              |   |
| Key Technologies                      | Prodrug   |

## COMPANY PROFILE

### SUMMARY

Hyperion Therapeutics Inc focuses on gastrointestinal and hepatology programs and products that address unmet medical needs to improve patient care.

### FINANCIAL

In April 2012, Hyperion filed a registration statement with the SEC for the issuance of initial public offering of its common stock; at that time, the offering details were not disclosed; in July 2012, the company priced its initial public offering of 5 million shares of its common stock at \$10 per share. At that time, underwriters were granted 30-day option to purchase up to an additional 750,000 shares of common stock to cover any over-allotments. The offering was expected to close on July 31, 2012; later that month, the offering was completed, with underwriters exercising their overallotment option in full.

In June 2009, Hyperion raised \$60 million from a series C financing.

In September 2007, Hyperion raised \$4 million from a series B financing.

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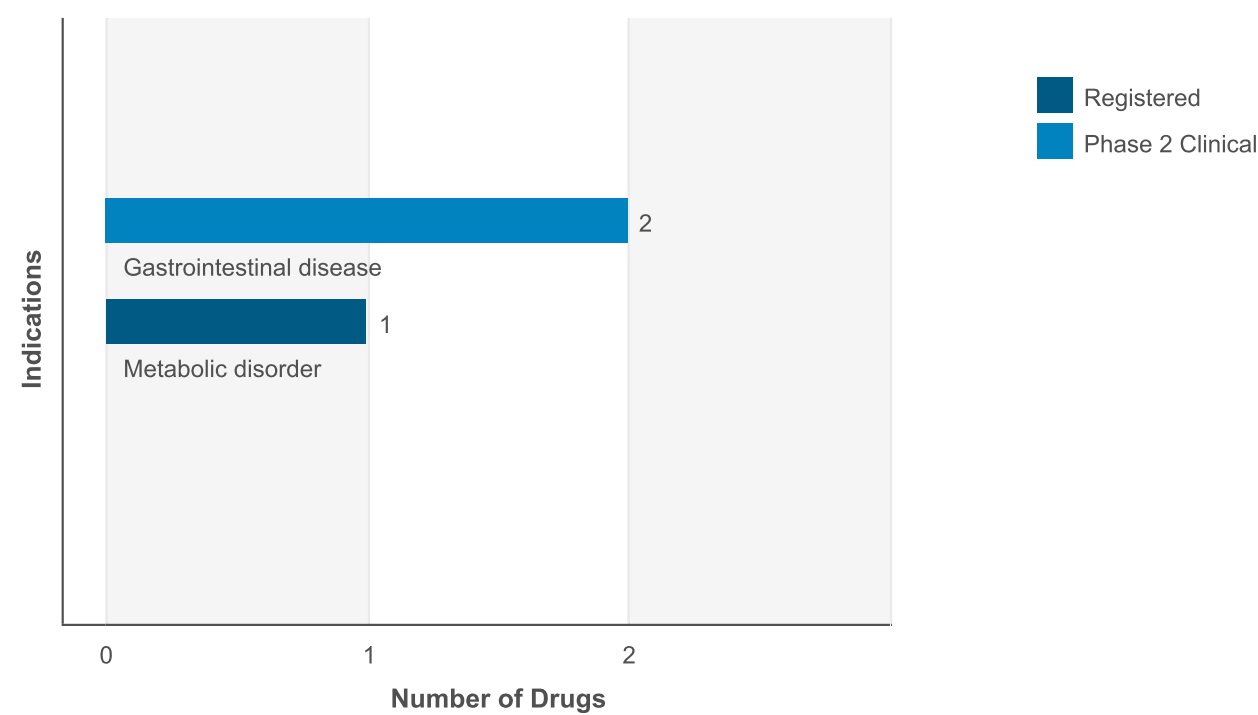


PRODUCT PORTFOLIO SUMMARY

DRUGS

Drugs by Indication

Active Drugs by Indication Chart



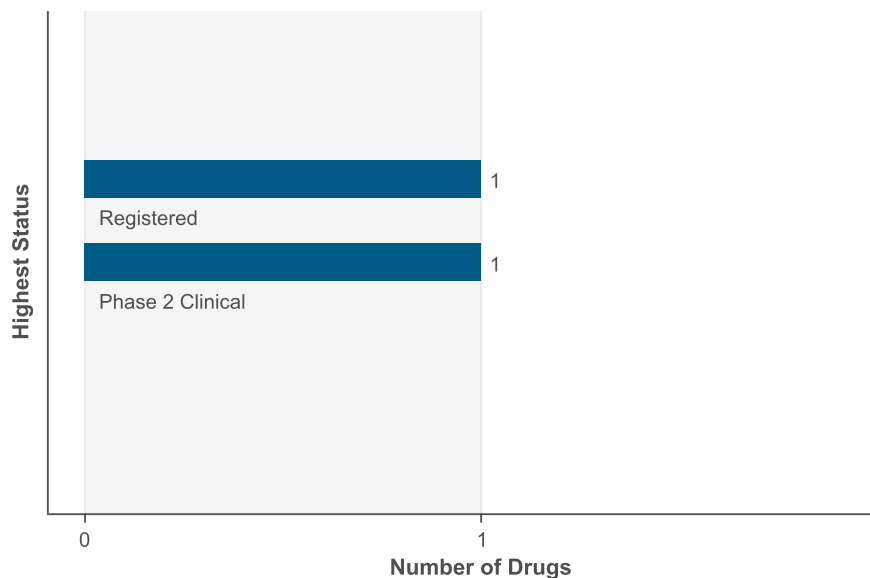
Drugs by Indication Table

| Indication               | Active | Inactive | Total |
|--------------------------|--------|----------|-------|
| Gastrointestinal disease | 2      | 0        | 2     |
| Metabolic disorder       | 1      | 0        | 1     |

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## Drugs by Highest Status

Active Drugs by Highest Status Chart



Drugs by Highest Status Table

| Development Status | Number of Drugs |
|--------------------|-----------------|
| Registered         | 1               |
| Phase 2 Clinical   | 1               |

## DEALS

| Deal Type                                    | Principal |          | Partner |          | Total |
|--|-----------|----------|---------|----------|-------|
|  | Active    | Inactive | Active  | Inactive |       |
| Drug - Development/Commercialization License | 0         | 0        | 1       | 0        | 1     |

## CLINICAL TRIALS

Trials by Condition Studied

| Condition Studied        | Ongoing | All |
|--------------------------|---------|-----|
| Metabolic disorder       | 1       | 9   |
| Gastrointestinal disease | 1       | 7   |
| Genitourinary disease    | 1       | 7   |
| Genetic disorder         | 0       | 1   |

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## Trials by Phase

| Phase   | Ongoing | All |
|---------|---------|-----|
| Phase 4 | 1       | 1   |
| Phase 3 | 0       | 3   |
| Phase 2 | 0       | 5   |
| Phase 1 | 0       | 4   |

### Phase Definitions

#### Phase 3 Clinical

Includes Phase 3, Phase 3b, Phase 3a, Phase 2/3 (where enrolment count is 300 or over)

#### Phase 2 Clinical

Includes Phase 2, Phase 2a, Phase 2b, Phase 1/2 (where enrolment count is 100 or over), Phase 2/3 (where enrolment count is under 300 or not specified)

#### Phase 1 Clinical

Includes Phase 1, Phase 1a, Phase 1, Phase 1/2 (where enrolment count is under 100 or not specified), Phase 0

## PATENTS \*

| Indication               | As Owner | As Third Party | Total |
|--------------------------|----------|----------------|-------|
| Gastrointestinal disease | 2        | 0              | 2     |
| Genitourinary disease    | 2        | 0              | 2     |
| Hematological disease    | 1        | 0              | 1     |
| Degeneration             | 1        | 0              | 1     |
| Neoplasm                 | 1        | 0              | 1     |
| Metabolic disorder       | 2        | 0              | 2     |
| Neurological disease     | 1        | 0              | 1     |

\* This table represents a summary of the core patent coverage for this company covering Therapeutic EP, US and WO patents since 1990 only.

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## PRODUCT PORTFOLIO DRUGS

**PLEASE NOTE:** Highest status refers to highest development of that drug for one of the active companies

### sodium phenylacetate + sodium benzoate (hyperammonemia), Ucyclid

|                             |   |
|-----------------------------|---|
| <b>Drug Name</b>            | sodium phenylacetate + sodium benzoate (hyperammonemia), Ucyclid      |
| <b>Key Synonyms</b>         | Ammonul   |
| <b>Originator Company</b>   | Ucyclid Pharma Inc  |
| <b>Active Companies</b>     | Hyperion Therapeutics Inc, Ucyclid Pharma Inc                         |
| <b>Inactive Companies</b>   |   |
| <b>Highest Status</b>       | Launched  |
| <b>Active Indications</b>   | Hepatic encephalopathy, Hyperammonemia                                |
| <b>Target-based Actions</b> |   |
| <b>Other Actions</b>        | Nitrogen metabolism modulator   |
| <b>Technologies</b>         | Drug combination, Intravenous formulation, Small molecule therapeutic |
| <b>Last Change Date</b>     | 18-Oct-2011   |

### glycerol phenylbutyrate

|                             |  |
|-----------------------------|--|
| <b>Drug Name</b>            | glycerol phenylbutyrate  |
| <b>Key Synonyms</b>         | Ravicti, glycerol phenylbutyrate   |
| <b>Originator Company</b>   | Ucyclid Pharma Inc   |
| <b>Active Companies</b>     | Hyperion Therapeutics Inc  |
| <b>Inactive Companies</b>   | Ucyclid Pharma Inc   |
| <b>Highest Status</b>       | Registered   |
| <b>Active Indications</b>   | Hyperammonemia, Hepatic encephalopathy   |
| <b>Target-based Actions</b> |  |
| <b>Other Actions</b>        | Nitrogen metabolism modulator  |
| <b>Technologies</b>         | Prodrug, Oral formulation, Oral liquid formulation, Small molecule therapeutic |
| <b>Last Change Date</b>     | 04-Feb-2013  |

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