WEDBUSH QUICK NOTE

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May 4, 2010

Ironwood Pharmaceuticals (IRWD – OUTPERFORM): Digestive Disease Week Linaclotide Chronic Constipation Phase III Data Oral Presentation

- Additional secondary endpoint data was presented for the 2 linaclotide Phase III studies in the chronic
 constipation (CC) setting yesterday at the DDW meeting in New Orleans. The results are consistent with the
 data that was previously presented and there was nothing in the results that should be cause for concern.
- The street is focused on the differences between existing approved treatments (osmotics, lubiprostone/Amitiza, motility agents) and linaclotide, and in particular for Amitiza, the concern is that this approved agent is not doing well commercially.
- Key details emerged from discussions with Dr. Rao and other thought leaders at the reception for the lack
 of Amitiza's market penetration. The Physicians stated that linaclotide's impact on pain and bloating will
 differentiate it from Amitiza; nausea is an adverse event that is thought to limit the market uptake of Amitiza and
 the absence of dose flexibility impacts this treatment's utility (i.e. too low a dose, no efficacy, too high, nausea).
 Furthermore, the perception of linaclotide's safety could lead to see physicians readily prescribing this therapy in
 the front line setting.
- We expect an update on the timing of the IBS-C Phase III data when the company reports Q1 financial results (date not set). One IBS-C Phase III study completed recruitment on February 19, the other on March 5, and we believe that patients in the second study are now in the open label extension study. Data timing will be dependent upon whether the company wants the open label phase (to week 26) of the data in hand before releasing primary endpoint top-line (to week 12).
- Reiterate our OUTPERFORM rating with a \$24/share price target. We estimate that US linaclotide sales will
 peak at approximately \$2.4 billion in 2019 in the CC and IBS-C settings. We arrive at our valuation by discounting
 back the product of the net present value of losses and profits through 2015 plus 18X 2016 linaclotide royalties
 and US revenues (25% discount rate, estimated current diluted share count of 119.8 million).
- Risks to the attainment of our price target include potential negative data from the IBS-C Phase III studies, regulatory risk associated with the NDA expected to be filed with the FDA and failure to achieve meaningful sales penetration of linaclotide in the IBS-C and/or CC settings.

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I, Gregory Wade, certify that the views expressed in this report accurately reflect my personal opinion and that I have not and will not, directly or indirectly, receive compensation or other payments in connection with my specific recommendations or views contained in this report.

IMPORTANT DISCLOSURES

Disclosure information regarding historical ratings and price targets is available at http://www.wedbush.com/ResearchDisclosure/DisclosureQ110.pdf

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OUTPERFORM – Expect the total return of the stock to outperform relative to the median total return of the analyst's (or the analyst's team) coverage universe over the next 6-12 months.

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The Investment Ratings are based on the expected performance of a stock (based on anticipated total return to price target) relative to the other stocks in the analyst's coverage universe (or the analyst's team coverage).*

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WS is acting as financial advisor to Dynavax (DVAX).

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Greg Wade, the analyst providing research coverage of Adolor (ADLR), BioMimetic (BMTI), Cubist (CBST), CombinatoRx (CRXX), OncoGenex Pharmaceuticals (OGXI) and ZymoGenetics (ZGEN), maintains long positions in the common stocks.

Jeremiah Shepard, the associate providing research coverage of Ironwood Pharmaceuticals (IRWD) and Xenoport (XNPT) maintains a long position in the common stocks.

* WS changed its rating system from (Strong Buy/Buy/Hold/Sell) to (Outperform/ Neutral/Underperform) on July 14, 2009.

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